

## Product Information

### D-Cycloserine

Catalog Numbers:  
**C6880, C3909, C7670**

Storage Temperature -20°C

CAS #: 68-41-7  
Synonyms: D-4-amino-3-isoxazolidone, D-oxamycin,  
Seromycin, K300, NJ-21

#### Product Description

Appearance: White powder  
Molecular Formula: C<sub>3</sub>H<sub>6</sub>N<sub>2</sub>O<sub>2</sub>  
Molecular Weight: 102.09  
E<sup>1%</sup> = 402 (226 nm)  
[α]<sub>D</sub><sup>23</sup> = +115° (c=1.0%, water)<sup>1</sup>

D-Cycloserine, a structural analog of D-alanine, is a broad spectrum antibiotic produced by certain strains of *Streptomyces orchidaceus* or *S. garphalus*.<sup>1-5</sup> D-cycloserine (at 100-200 µg/ml) inhibits the synthesis of bacterial cell walls (involving peptidoglycan synthesis) by preventing formation of D-alanine from L-alanine and hence the formation of peptide bonds involving D-alanine.<sup>4</sup> D-cycloserine has antibiotic activity *in vitro* against growth phase Gram-negative bacteria including *Escherichia coli* (working concentration of approx. 200 µg/ml)<sup>4</sup>, strains of *Staphylococcus aureus*, *Nocardia* species and *Chlamydia*,<sup>3</sup> and some mycobacteria including *Mycobacterium tuberculosis*. The minimum inhibitory concentrations (MIC) *in vitro* for *M. tuberculosis* range from 5-20 µg/ml. Studies *in vitro* show no suppression of growth in cultures made in media containing D-alanine which appears to block the antibacterial action of D-cycloserine.<sup>3</sup>

D-cycloserine is an excitatory amino acid and partial agonist at the glycine binding site of the N-methyl-D-aspartate (NMDA) receptor.<sup>6-8</sup> At low doses it is a cognitive enhancer that improves learning and memory in several experimental models of disease and cognitive deficit.<sup>6,7,9-14</sup> At high doses, D-cycloserine is an anti-convulsant.<sup>15,16</sup> Intermediate doses potentiate the anti-convulsant action of phenytoin but block its long-term memory impairment.<sup>16</sup>

The HPLC determination of D-cycloserine in plasma and urine<sup>17</sup> and a colorimetric method for determination of cycloserine in biological fluids<sup>18</sup> have been reported. UV, IR, NMR and mass spectra and pharmacokinetics of D-cycloserine have been reported.<sup>2</sup>

#### Reagents

These products are supplied as powders.

**C7670** is convenience packaged for use in molecular biology; it is pre-weighed in quantities to give typical working concentrations when the entire package is added to 1 L of agar preparations (for 50 plates of 20 ml per plate). Furthermore, C 7670 is γ-irradiated for sterility and septum-capped for ease in injecting sterile diluent. C 7670 is also USP tested for potency following γ-irradiation to assure full biological activity.

#### Preparation Instructions

D-cycloserine is soluble in deionized water up to 100 mg/ml. A solution of 50 mg/ml cycloserine in water is clear and colorless or very faintly yellow. D-cycloserine is also soluble at 1 in 50 parts of 96% ethanol, but practically insoluble in chloroform and ether. It is also slightly soluble in methanol or propylene glycol. Stock solutions (e.g. 10 mg/ml) of D-cycloserine may also be prepared immediately before use in 0.1 M sodium phosphate buffer, pH 8.0.

#### Storage/Stability

D-Cycloserine powder is stable for at least four years when stored unopened and desiccated at -20 °C. It is generally recommended to prepare solutions immediately before use because neutral or acidic solutions are unstable.<sup>4</sup> However, aqueous solutions buffered to pH 10 with sodium carbonate may be stored for up to one week if stored at 2-8 °C. In addition, aqueous solutions of D-cycloserine have been stored in aliquots at -20 °C and thawed just prior to use.<sup>12</sup>

#### Precautions and Disclaimer

This product is for R&D use only, not for drug, household, or other uses. Please consult the Material Safety Data Sheet for information regarding hazards and safe handling practices.

#### References

1. Stammer, C.H., et al., *J. Am. Chem. Soc.*, **77**, 2346 (1955).

2. El-Obeid, H.A. and Al-Badr, A.A, Analytical Profile of D-Cycloserine in *Analytical Profiles of Drug Substances*, **Vol. 18**, p. 567 (Academic Press, New York, 1989).
3. Goodman and Gilman's The Pharmacological Basis of Therapeutics, 9th Ed., Hardman, J.G. et al., (Eds.), p. 1164 (McGraw-Hill, New York, NY, 1995).
4. Raleigh, E.A., et al., Selected Topics from Classical Bacterial Genetics in *Short Protocols in Molecular Biology*, 4th Ed., Ausubel, F.M. et al., (Eds.), Unit 1.4, p. 1-9 (John Wiley & Sons, Inc., New York, 1999).
5. Kuehl, F.A., Jr. et al., *J. Am. Chem. Soc.*, **77**, 2344 (1955).
6. Sirvio, J., et al., *Neurosci. Lett.*, **146**, 215 (1992).
7. Ohno, M. and Watanabe, S., *Eur. J. Pharmacol.*, **318**, 267 (1996).
8. Watson, G.B., et al., *Brain Res.*, **510**, 158-160 (1990).
9. Flood, J.F., et al., *Eur. J. Pharmacol.*, **221**, 249 (1992).
10. Schuster, G.M. and Schmidt, W.J., *Eur. J. Pharmacol.*, **224**, 97 (1992).
11. Temple, M.D. and Hamm, R.J., *Brain Res.*, **741**, 246 (1996).
12. Nakazato, E., et al., *Life Sci.*, **67**, 1139-1147 (2000).
13. Schneider, J.S., et al., *Brain Res.*, 860, 190-194 (2000).
14. Pussinen, R., and Sirvio, J., *J. Psychopharmacol.*, **13**, 171-179 (1999).
15. Chessell, I.P., et al., *Brain Res.*, **565**, 345 (1991).
16. Wlaz, P., et al., *Epilepsia*, **37**, 610-617 (1996).
17. Musson, D.G. et al., *J. Chromatog.*, **414**, 121 (1987).
18. Jones, L.R., *Anal. Chem.*, **28**, 39 (1956).

RC,MM,PHC 05/13-1