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# **ProductInformation**

Thioredoxin from *E. coli*, recombinant, overexpressed in *E. coli* 

Product Number **T 0910** Storage Temperature –20 °C

CAS<sup>#</sup> 52500-60-4 GenBank Accession Number M26133

## **Product Description**

Thioredoxin from *E. coli* has a molecular weight of approximately 11.7 kDa (SDS-PAGE). The active site of thioredoxin contains two vicinal cysteine residues having the amino acid sequence Cys-Gly-Pro-Cys. In the reduced form two sulfhydryl groups are present and when oxidized they form a disulfide bridge. The thioredoxin system includes thioredoxin, which is reduced by thioredoxin reductase with NADPH that serves as the hydrogen donor.

Thioredoxin is implicated in ribonucleotide reduction, <sup>1</sup> in assimilatory sulfate reduction, <sup>2</sup> and in a regulatory scheme involving oxidation and reduction of protein sulfhydryl groups. <sup>3</sup> It has been shown to be required for filamentous phage assembly in vivo <sup>4</sup> and it catalyzes refolding of various proteins. <sup>5</sup>

The product is supplied as an essentially salt-free, lyophilized powder,

Purity: minimum 90% (SDS-PAGE)

Specific Activity: minimum 3 units/mg protein

Unit definition: One unit will cause a  $\Delta A_{650}$  of 1.0 in 1 minute at pH 7.5 at 25 °C in the insulin reduction assay. Thioredoxin activity is assessed in an insulin reduction assay, based on the formation of reduced insulin, which precipitates in the presence of a fixed amount of dithiothreitol and suitable amounts of thioredoxin. Precipitation of reduced insulin is monitored by an increase in absorbance at 650 nm.

#### **Precautions and Disclaimer**

This product is for laboratory research use only. Please consult the Material Safety Data Sheet for information regarding hazards and safe handling practices.

## **Preparation Instructions**

The product is soluble in water (1 mg/ml), yielding a clear, colorless solution.

#### Storage/Stability

It is recommended to store the product at –20 °C. Thioredoxin from *E. coli* was found to be stable after 16 hours at 50 °C in both the lyophilized and reconstituted forms. It was also stable after 96 hours at 37 °C.

## References

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