

User Manual

Version 2.0

Product name: Inorganic Pyrophosphatase, yeast

Cat #: PI-100, PI-200, PI-OEM

Description:

The Pyrophosphatase, Inorganic (PPase) catalyzes the hydrolysis of inorganic pyrophosphate to two orthophosphates. The enzyme requires a divalent metal cation, with Mg²⁺ conferring the highest activity.

Application:

- High yield synthesis of RNA by in vitro transcription^(1,2)
- DNA polymerization reactions: preventing accumulation of pyrophosphate^(3, 4)
- Removal of contaminant PPi in reagents used for SNP genotyping by methods based on the detection of pyrophosphate $^{(5)}$

Source: E. coli cells with a cloned ppa gene from Sacharomyces cerevisiae.

Unit Definition:

One unit is the amount of enzyme that will generate 1 μ mol of phosphate per minute from inorganic pyrophosphate under standard reaction conditions (a 10 minute reaction at 25°C in 100 mM Tris-HCl, pH 7.2, 2 mM MgCl₂ and 2 mM PPi in a reaction volume of 0.5 ml).

Storage Conditions:

20 mM Tris-HCl 100 mM KCl 1 mM Dithiothreitol 0.1 mM EDTA 50% Glycerol pH 8.0 @ 25°C

Recommended Storage Condition: -20°C

References:

- 1. Cooperman, B.S., The mechanism of action of yeast inorganic pyrophosphatase, Meth. Enzymol., 87, 526-548, 1982.
- 2. Cunningham, P.R. and Ofengand, J., Use of inorganic pyrophostase to improve the yield of in vitro transcription reactions catalyzed by T7 RNA polymerase, Biotechniques, 9, 713-714, 1990.
- 3. Tabor, S., Richardson, C.C., DNA sequence analysis with a modified bacteriophage T7 DNA polymerase. Effect of pyrophosphorolysis and metal ions, J. Biol. Chem., 265, 8322-8328, 1990.
- 4. Dean, B.F., et al., Rapid amplification of plasmid and phage DNA using phi29 DNA polymerase and multiply primed Rolling Circle amplification, Genome Res., 11, 1095-1099, 2001.
- 5. Zhou, G.H., et al., Quantitative detection of single nucleotide polymorphisms for a pooled sample by a bioluminometric assay coupled with modified primer extension reactions (BAMPER), Nucleic Acids Res., 29, E93,2001.