

## PRODUCT INFORMATION SHEET

# Thermostable Inorganic Pyrophosphatase

### PRODUCT SPECIFICATIONS

**Product Name:** Recombinant Thermostable Inorganic Pyrophosphatase, RUO Grade

**Product number/code:** PyroPh-SAC-Pyrophosphatase enzyme, His purified, 274-002

**Product Description:** Thermostable Inorganic Pyrophosphatase is prepared using Crisbio®, a novel expression system that utilizes a recombinant baculovirus carrying a *Sulfolobus acidocaldarius* gene to inoculate *Trichoplusia ni* chrysalises

**Source of Protein:** Engineered Baculovirus carrying the Thermostable Pyrophosphatase gene from *S. acidocaldarius* inoculated in *Trichoplusia ni*

**Unit Availability:** 2000 U and 1000 U

**Unit Definition:** One unit (U) will release 1.0 µmol of inorganic orthophosphate per minute at pH 9.0 at 75°C

**Shelf Life:** 2 years from manufacturing date. Shelf life currently under assessment with stability studies

**Storage Temp:** -20°C

**Storage Buffer:** 10 mM Tris-HCl pH 7.5, 50 mM NaCl, 1 mM DTT, 0.1 mM EDTA and 50% Glycerol

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#### Limitations of Use:

**For research and development use only.** Not suitable for human or animal therapeutic or clinical diagnostic use. SDS sheets relevant to this product are available upon request.

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D-PS-PyroPh-SAC\_v04

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Assay	SDS-PAGE Purity	SDS-PAGE Western Blot Identity <sup>1</sup>	Enzyme Activity	DNase Activity	Residual gDNA
Specification	≥ 95%	Specific Band Detection	≥ 2000 U/ml	Not detected <sup>2</sup>	≤ 100 pg/mg

<sup>1</sup> performed during manufacture process

<sup>2</sup> < LOD of the assay (< RFU of 0.02 U control, in 50 enzyme units tested)

### QUALITY CONTROL ANALYSIS

**Enzyme Activity** is determined by Absorption Spectrophotometry:

The phosphate released after pyrophosphate hydrolysis by pyrophosphatase is measured by means of a color reaction. Dilutions of the enzyme in 50mM Tris-HCl pH 9.0, 10mM MgCl<sub>2</sub> and 10mM sodium pyrophosphate are assayed. After a 10 min incubation at 75°C, the 2-orthophosphate product reacts with ammonium molybdate to form phosphomolybdic acid. The phosphomolybdic acid is reduced by ferrous sulphate giving a blue color to the reaction. This color can be measured at 660 nm. A phosphate standard curve is used to quantify the product formed.

**Protein Purity** is evaluated by SDS-PAGE of concentrated and diluted enzyme solutions followed by Coomassie stain. Purity is assessed by comparing the aggregate mass of non-specific bands in the concentrated sample to the mass of the protein of interest band in the diluted sample.

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**DNase Contamination** is detected by Fluorometric Assay:

Use of a fluorogenic substrate that emits fluorescence after cleavage by the presence of DNases.

**Residual (genomic) DNA Contamination** is evaluated by quantitative PCR:

Amplification of specific genomic sequence of *T ni*.

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