

Antarctic Phosphatase from E. coli, Recombinant

Cat. No. NATE-1399

Lot. No. (See product label)

Introduction

Description Antarctic Phosphatase catalyzes the removal of 5 ' phosphate from DNA and RNA.

Since phosphatase-treated fragments lack the 5´ phosphoryl termini required by ligases, they cannot self-ligate. This property can be used to decrease the vector

background in cloning strategies.

Applications Removing 5´ phosphates from DNA, RNA, rNTPs and dNTPs Preparation of

templates for 5´ end labeling Prevention of recircularization of cloning vectors Removal of dNTPs and pyrophosphate from PCR reactions Dephosphorylation of

proteins

Synonyms Antarctic Phosphatase

Product Information

Species E. coli

Source E. coli

Form 10 mM Tris-HCl (pH 7.4), 1 mM MgCl2, 0.01 mM ZnCl2, 1 mM DTT and 50% glycerol.

Molecular Weight Apparent: 35 kDa Theoretical: 69 kDa

Purity >95% estimated by SDS-PAGE

Concentration 5,000 units/ml

Unit Definition One unit is defined as the amount of enzyme that will dephosphorylate 1 μg of

pUC19 vector DNA cut with HindIII (5 $^{\prime}$ protruding ends), HincII (blunts ends) or PstI (5 $^{\prime}$ recessed ends) in 30 minutes at 37 $^{\circ}$ C. Dephosphorylation is defined as > 95 $^{\circ}$

1/1

inhibition of recirculation in a self-ligation reaction and is measured by

transformation into E. coli.

Storage and Shipping Information

Storage at -20°C

Tel: 1-631-562-8517 1-516-512-3133 **Email:** info@creative-enzymes.com