Casein Kinase II (CK2)





P6010S

R\{

500.000 U/ml Lot: 0151210 10.000 units RECOMBINANT Store at -70°C Exp: 10/13

Description: Casein Kinase II (CK2) is a constitutively active serine/threonine protein kinase composed of two 44 kDa catalytic α -subunits and two 26 kDa regulatory β -subunits in an $\alpha_{\alpha}\beta_{\alpha}$ configuration to form stable heterotetramers. CK2 holoenzyme undergoes autophosphorylation at two serine residues (S2/S3) of its β-subunit. Recently it has been shown that CK2 α -subunits undergo intermolecular tyrosine-autophosphorylation at Y182, which may represent a specific regulatory

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mechanism. Also, CK2 is able to phosphorylate. under special circumstances, tyrosyl residues in proteins. CK2 is implicated in a variety of cellular functions (1,2).

Recognition Determinants: The CK2 substrate specificity is invariably determined by multiple acidic residues located at positions between -2 and +5 relative to the target amino acid (mostly Ser and rarely Thr). The general recognition motif for phsophorylation by CK2 is **\$**XXE/D, although SXE/D and S/D, and variations of these sequences are also phosphorylated. Polyanionic compounds, like heparin, inhibit CK2 activity with a K, of 1.4 nm (4,5).

Source: Isolated from a strain of E. coli expressing both α and β CK2 subunits derived from a human glioblastoma cDNA library (kindly provided by Dr. D. Marshak) (3).

Supplied in: 350 mM NaCl, 20 mM Tris-HCl (pH 7.5 @ 25°C), 1 mM Na₂EDTA, 2 mM DTT and 0.1% Triton X-100.

Reagents Supplied with Enzyme: 10X CK2 Reaction Buffer

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(pH 7.5 @ 25°C), 1 mM Na₂EDTA, 2 mM DTT and 0.1% Triton X-100.

42 kDa.

Reaction Conditions: 1X CK2 Reaction Buffer. supplement with 200 µM ATP and gamma-labeled ATP to a final specific activity of 100-500 µCi/ µmol. (CK2 will also accept GTP as a phosphory) donor in place of ATP). Incubate at 30°C.

1X CK2 Reaction Buffer:

20 mM Tris-HCI 50 mM KCI 10 mM MgCl₂ pH 7.5 @ 25°C

Note that optimal incubation times and enzyme concentrations must be determined empirically for each particular substrate.

Unit Definition: One unit is defined as the amount of CK2 required to catalyze the transfer of 1 pmol of phosphate to CK2 Peptide Substrate, RRRADDSDDDDD (100 µM, NEB #P6012), in 1 minute at 30°C in a total reaction volume of 25 ul (4.5).

Specific Activity: ~ 859.000 units/mg.

Molecular Weight: α -subunit (45 kDa), β -subunit (25 kDa). The apparent molecular weight of the α-subunit estimated by SDS-PAGE is about 42 kDa.

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Specific Activity: ~ 859,000 units/mg.

Molecular Weight: α -subunit (45 kDa), β -subunit (25 kDa). The apparent molecular weight of the α -subunit estimated by SDS-PAGE is about

Quality Assurance: CK2 contains no detectable protease or phosphatase activities.

Quality Control Assays

Protease Activity: After incubation of 5,000 units of Casein Kinase II (CK2) with a standard mixture of proteins for 2 hours at 30°C, no proteolytic activity could be detected by SDS-PAGE analysis.

Phosphatase Activity: After incubation of 5,000 units of Casein Kinase II (CK2) with 50 mM p-nitrophenyl phosphate for 2 hours at 30°C, no phosphatase activity could be detected by spectrophotometric analysis.

References:

- 1. Donella-Deana, A. et al. (2001) Biochem J. 357. 563-567.
- 2. Marin, O. et al. (1999) J. Biol. Chem. 274, 29260-29265.
- 3. Chester, N. and Marshak, D.R. (1993) Anal. Biochem. 209, 284-290.
- 4. Marin, O. et al. (1994) BBRC, 198, 898-905.
- 5. Sarno, S. et al. (1996) J. Biol. Chem. 271. 10595-10601.

CERTIFICATE OF ANALYSIS

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References:

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expressing both α and β CK2 subunits derived from a human glioblastoma cDNA library (kindly provided by Dr. D. Marshak) (3).

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