

Product Information

CAMK2 α , active, GST-tagged, human PRECISIO® Kinase recombinant, expressed in *Sf9* cells

Catalog Number **C9869**
Storage Temperature $-70\text{ }^{\circ}\text{C}$

Synonyms: CAMKA, KIAA0968

Product Description

CAMK2 α is a serine/threonine protein kinase that is a member of the Ca²⁺/calmodulin-dependent protein kinase family. CAMK2 α is abundant in the brain as a major constituent of the postsynaptic density and is required for hippocampal long-term potentiation (LTP) and spatial learning.¹ In addition to its Ca²⁺/calmodulin-dependent activity, CAMK2 α can undergo autophosphorylation, resulting in Ca²⁺/calmodulin-independent activity. The protein level of CAMK2 α fluctuates during neuronal activity in cultured rat pup hippocampal neurons. The levels of CAMK2 α increased with heightened neuronal activity.²

This recombinant product was expressed by baculovirus in *Sf9* insect cells using an N-terminal GST-tag. The gene accession number is NM 171825. It is supplied in 50 mM Tris-HCl, pH 7.5, with 150 mM NaCl, 0.25 mM DTT, 0.1 mM EGTA, 0.1 mM EDTA, 0.1 mM PMSF, and 25% glycerol.

Molecular mass: ~74 kDa

Precautions and Disclaimer

This product is for R&D use only, not for drug, household, or other uses. Please consult the Material Safety Data Sheet for information regarding hazards and safe handling practices.

Storage/Stability

The product ships on dry ice and storage at $-70\text{ }^{\circ}\text{C}$ is recommended. After opening, aliquot into smaller quantities and store at $-70\text{ }^{\circ}\text{C}$. Avoid repeated handling and multiple freeze/thaw cycles.

Figure 1.
SDS-PAGE Gel of Typical Lot:
 $\geq 70\%$ (SDS-PAGE, densitometry)

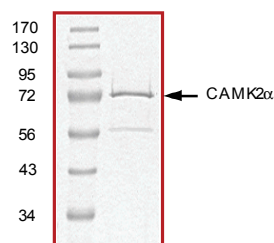
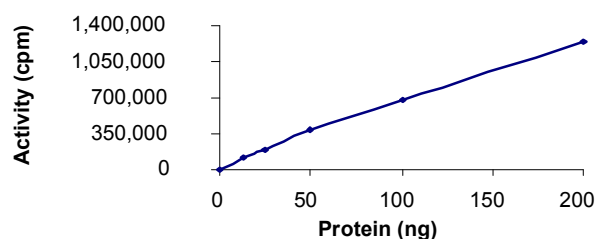


Figure 2.
Specific Activity of Typical Lot:
218–294 nmole/min/mg



Procedure

Preparation Instructions

Kinase Assay Buffer – 25 mM MOPS, pH 7.2, 12.5 mM glycerol 2-phosphate, 25 mM MgCl₂, 5 mM EGTA, and 2 mM EDTA. Just prior to use, add DTT to a final concentration of 0.25 mM.

Kinase Dilution Buffer – Dilute the Kinase Assay Buffer 5-fold with a 50 ng/μl BSA solution.

Kinase Solution – Dilute the active CAMK2 α (0.1 $\mu\text{g}/\mu\text{l}$) with Kinase Dilution Buffer to the desired concentration.

Note: The specific activity plot may be used as a guideline (see Figure 2). It is recommended the researcher perform a serial dilution of active CAMK2 α kinase for optimal results

10 mM ATP Stock Solution – Dissolve 55 mg of ATP in 10 ml of Kinase Assay Buffer. Store in 200 μl aliquots at $-20\text{ }^{\circ}\text{C}$.

γ - ^{32}P -ATP Assay Cocktail (250 μM) – Combine 5.75 ml of Kinase Assay Buffer, 150 μl of 10 mM ATP Stock Solution, 100 μl of γ - ^{32}P -ATP (1 mCi/100 μl). Store in 1 ml aliquots at $-20\text{ }^{\circ}\text{C}$.

Substrate Solution – Dissolve the Autocamtide 2 synthetic peptide substrate (KKALRRQETVDAL-amide) in water at a final concentration of 1 mg/ml.

1% phosphoric acid solution – Dilute 10 ml of concentrated phosphoric acid to a final volume of 1 L with water.

Kinase Assay

This assay involves the use of the ^{32}P radioisotope. All institutional guidelines regarding the use of radioisotopes should be followed.

1. Thaw the active CAMK2 α , Kinase Assay Buffer, Substrate Solution, and Kinase Dilution Buffer on ice. The γ - ^{32}P -ATP Assay Cocktail may be thawed at room temperature.
2. In a pre-cooled microcentrifuge tube, add the following solutions to a volume of 20 μl :
 - 10 μl of Kinase Solution
 - 7.5 μl of Substrate Solution
 - 2.5 μl of 5 mM CaCl_2 solution containing 0.75 μg Calmodulin
3. Set up a blank control as outlined in step 2, substituting 7.5 μl of cold water ($4\text{ }^{\circ}\text{C}$) for the Substrate Solution.
4. Initiate each reaction with the addition of 5 μl of the γ - ^{32}P -ATP Assay Cocktail, bringing the final reaction volume to 25 μl . Incubate the mixture in a water bath at $30\text{ }^{\circ}\text{C}$ for 15 minutes.
5. After the 15 minute incubation, stop the reaction by spotting 20 μl of the reaction mixture onto an individually pre-cut strip of phosphocellulose P81 paper.

6. Air dry the pre-cut P81 strip and sequentially wash in the 1% phosphoric acid solution with constant gentle stirring. It is recommended the strips be washed a total of 3 times of ~ 10 minutes each.
7. Set up a radioactive control to measure the total γ - ^{32}P -ATP counts introduced into the reaction. Spot 5 μl of the γ - ^{32}P -ATP Assay Cocktail on a pre-cut P81 strip. Dry the sample for 2 minutes and read the counts. Do not wash this sample.
8. Count the radioactivity on the P81 paper in the presence of scintillation fluid in a scintillation counter.
9. Determine the corrected cpm by subtracting the blank control value (see step 3) from each sample and calculate the kinase specific activity

Calculations:

1. Specific Radioactivity (SR) of ATP (cpm/nmole)

$$\text{SR} = \frac{\text{cpm of } 5\ \mu\text{l of } \gamma\text{-}^{32}\text{P}\text{-ATP Assay Cocktail}}{\text{nmole of ATP}}$$

cpm – value from control (step 7)

nmole – 1.25 nmole (5 μl of 250 μM ATP Assay Cocktail)

2. Specific Kinase Activity (SA) (nmole/min/mg)

$$\text{nmole/min/mg} = \frac{\Delta\text{cpm} \times (25/20)}{\text{SR} \times \text{E} \times \text{T}}$$

SR = specific radioactivity of the ATP (cpm/nmole ATP)

Δcpm = cpm of the sample – cpm of the blank (step 3)

25 = total reaction volume

20 = spot volume

T = reaction time (minutes)

E = amount of enzyme (mg)

References

1. Silva, A.J. et al., Impaired spatial learning in alpha-calcium calmodulin kinase II mutant mice. *Science*, **257**, 206-211 (1992).
2. Thiagarajan, T.C. et al., Alpha- and beta-CaMKII: inverse regulation by neuronal activity and opposing effects on synaptic strength. *Neuron*, **36**, 1103-1114 (2002).

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