

Effect of metal ions on the activity of casein kinase II from *Xenopus laevis*

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CaSein kinase II purified from the nuclei of *Xenopus laevis* oocytes as well as the recombinant α and β subunits of the *X. laevis* CKII, produced in *E. coli* from the cloned cDNA genes, were tested with different divalent metal ions. The enzyme from both sources was active with either Mg^{2+} , Mn^{2+} , or Co^{2+} . Optimal concentrations were 7-10 mM for Mg^{2+} , 0.5-0.7 mM for Mn^{2+} and 1-2 mM for Co^{2+} . In the presence of Mn^{2+} or Co^{2+} the enzyme used GTP more efficiently than ATP as a phosphate donor while the reverse was true in the presence of Mg^{2+} . The apparent K_m values for both nucleotide triphosphates were greatly decreased in the presence of Mn^{2+} as compared with Mg^{2+} . Addition of Zn^{2+} (above 150 μM) to an assay containing the optimal Mg^{2+} ion concentration caused strong inhibition of both holoenzyme and β subunit. Inhibition of the holoenzyme by 400 μM Ni^{2+} could be reversed by high concentrations of Mg^{2+} but no reversal of this inhibition was observed with the α subunit. © 1993.