

# An inactive mutant of the $\beta$ subunit of protein kinase CK2 that traps the regulatory CK2 $\alpha$ subunit

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Protein kinase CK2 (casein kinase 2) is a ubiquitous Ser/Thr protein kinase involved in cell proliferation. Mutation of the  $\beta$  subunit of the *Xenopus laevis* CK2 to change aspartic acid 156 to alanine (CK2 $\beta$ A156) resulted in an inactive enzyme. The CK2 $\beta$ A156 mutant, however, binds the regulatory subunit as measured by retention of  $\alpha$  on a nickel chelating column mediated by (His)<sub>6</sub>-tagged CK2 $\beta$ A156. Addition of CK2 $\beta$ A156 also caused  $\alpha$  to shift sedimentation in a sucrose gradient from a ( $\alpha$ )<sub>2</sub> dimer (52 kDa) to an ( $\alpha$ )<sub>2</sub>( $\beta$ )<sub>2</sub> tetramer (130000 kDa). CK2 $\beta$ A156 can trap the  $\alpha$  subunit in an inactive complex reducing the stimulation of casein phosphorylation caused by addition of  $\alpha$  to wild-type  $\beta$ . This competitive effect depends on the ratio of  $\alpha$ / $\beta$ A156 and on the amount of  $\alpha$  available. Since  $\alpha$  inhibits the phosphorylation of calmodulin by CK2 $\beta$ , the addition of CK2 $\beta$ A156, in this case, increases calmodulin phosphorylation by the  $\alpha$  and  $\beta$  combination. These results suggest that CK2 $\beta$ A156 may be a useful dominant