

Fig. 1. Alignment of the plant CK2 α catalytic subunits with the human CK2 α and CK2 α ' sequences. The amino acid sequence of the three maize CK2 α subunits, Zm a-1 (CAA43659), Zm a-2 (CAA72290), Zm a-3 (AAG36872) and the two Arabidopsis CK2 α subunits At a-1 (Q08467) and At a-2 (Q08466) has been aligned with human CK2 α ', H alfa' (AAA51548) and CK2 α , H alfa (AAA355503). Invariant residues are indicated by asterisks and by shaded boxes, and dash indicates a gap introduced to maximize alignment. Characteristics domains of CK2 α catalytic subunits, ATP binding site and the basic stretch (NLS) are underlined.

activity [23], it confers stability to the enzyme [24] and it provides specificity for the interaction with substrates and inhibitors [25]. Whereas in most organisms only one gene for CK2 β has been described, two genes (CKB1 and CKB2) have been identified in Saccharomyces cerevisiae [26]. At least two genes exist in Drosophila melanogaster [27, 28], plus the β -like protein Stellate [22]. In contrast to the catalytic subunits, deletion of CKB1 and/or CKB2 genes coding for regulatory subunits is not lethal for the yeast cells; it does not affect yeast growth under normal conditions but results in a phenotype of hypersensitivity to Na* and Li* cations [29].

The level of identity between plant, yeast and human CK2 β regulatory subunits is not as high as in the case of CK2 α subunits; for this reason complementation assays were needed in order to isolate CK2 β regulatory subunits (CK2 β 1 and CK2 β 2) in *Arabidopsis thaliana* [10]. Recently, a third CK2 β regulatory subunit (CK2 β 3) has been identified in *Arabidopsis* [11]. Data from the computational analysis of the Arabidopsis genome indicates the existence of a fourth CK2 β subunit. During several years, there was controversy about the existence of CK2 β subunits in maize. For instance, antibodies raised against chicken CK2 β failed to