

Erratum to: Characterization of transcription factor gene *SNAC2* conferring cold and salt tolerance in rice

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Due to an unfortunate mistake, an incorrect version of Fig. 2 has been used in the above mentioned publication.

The yeast-one-hybrid result presented was not for *SNAC2*. The correct representation of Fig. 2 is published on the following page and should be treated as definitive by the reader.

The online version of the original article can be found under doi:[10.1007/s11103-008-9309-5](https://doi.org/10.1007/s11103-008-9309-5).

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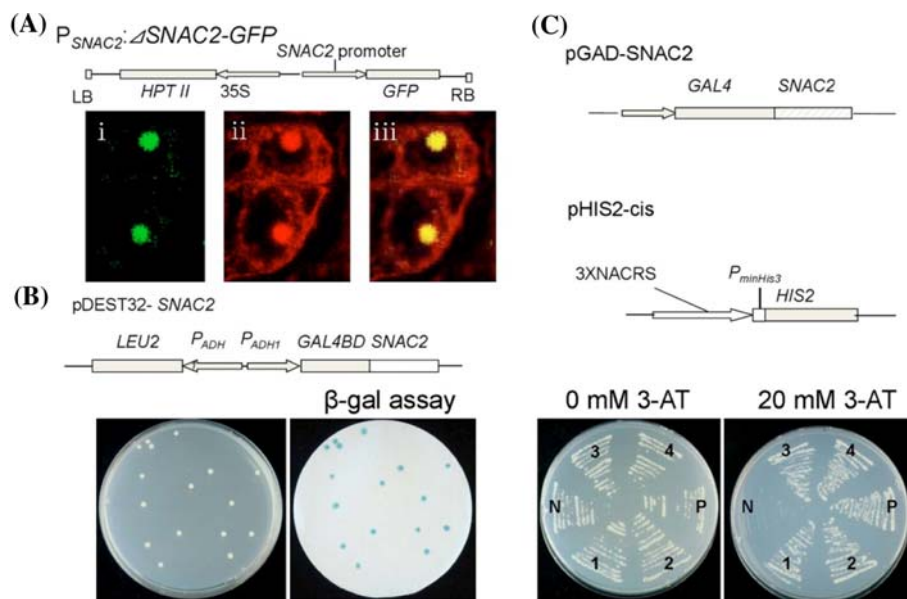


Fig. 2 SNAC2 features a transcription factor. **a** Construct of $P_{SNAC2}:\Delta SNAC2:GFP$ (The first 144 amino acids of SNAC2 was GFP under the control of *SNAC2* promoter) was transformed into rice and GFP signal was checked in calli cells with confocal microscopy. (i) Fluorescent image of GFP; (ii) fluorescent image stained with propidium iodide; (iii) merged image. **b** Transactivation assay of SNAC2 in yeast. Full SNAC2 protein was fused to the GAL4 binding domain (GAL4 BD) and transformed into yeast strain MV203, and

β -gal assay was performed to identify the transactivation activity (*LacZ* expression). **c** The pGAD-SNAC2 plasmid and the reporter construct pHIS-*cis* (Hu et al. 2006) were co-transformed into yeast strain Y187. The transformants were examined by growth performance on SD/Leu⁻/Trp⁻/His⁻ plates with or without 3-AT. *N*: negative control (p53HIS2 + pGAD-SNAC2); *P*: positive control (p53HIS2 + pGAD-Rec2-53); 1–4: four different colonies containing pGAD-SNAC2 and pHIS-*cis*