

**TABLE S11. Plasmid used in this study**

Plasmid	Genotype	Source
pINA443	Replicative form with <i>ARS68</i> , <i>URA3</i>	C. Gaillardin <sup>a</sup>
pINA445	Replicative form with <i>ARS68</i> , <i>LEU2</i>	C. Gaillardin <sup>a</sup>
pJCTGCAN18	Plasmid recovered from the genomic DNA library, complementing the fil <sup>-</sup> phenotype in CHY33188 strain. Replicative form with <i>ARS68</i> , <i>LEU2</i> .	This study
pJCTGCAN203	Plasmid constructed from pJCTGCAN18, <i>ZNC1</i> tagged with <i>GFP</i> in the <i>SpeI</i> site.	This study
pJCTGCAN227	The 2.2-kbp <i>Sall</i> - <i>XbaI</i> fragment from pJCTGCAN18 containing the <i>ZNC1</i> gene cloned into pBlueScript II SK(+).	This study
pJCTGCAN277	From pJCTGCAN227. Two cysteine residues within the Znc1p zinc finger region were replaced by two arginines (C <sub>22</sub> C <sub>25</sub> /R <sub>22</sub> R <sub>25</sub> ).	This study
pJCTGCAN282	From pJCTGCAN227. Two leucine residues within the Znc1p leucine zipper region were replaced by two arginines (L <sub>437</sub> L <sub>438</sub> /R <sub>437</sub> R <sub>438</sub> ).	This study
pJCTGCAN294	The 2.2-kbp <i>Sall</i> - <i>XbaI</i> fragment from pJCTGCAN203 expressing Znc1p-GFP was replaced with the 2.2-kbp <i>Sall</i> - <i>XbaI</i> fragment from pJCTGCAN277. (Znc1p C <sub>22</sub> C <sub>25</sub> /R <sub>22</sub> R <sub>25</sub> -GFP).	This study
pJCTGCAN297	The 2.2-kbp <i>Sall</i> - <i>XbaI</i> fragment from pJCTGCAN203 expressing Znc1p-GFP was replaced with the 2.2-kbp <i>Sall</i> - <i>XbaI</i> fragment from pJCTGCAN282. (Znc1p L <sub>437</sub> L <sub>438</sub> /R <sub>437</sub> R <sub>438</sub> -GFP).	This study

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