

CORRECTION

Correction: Velvet domain protein VosA represses the zinc cluster transcription factor SclB regulatory network for *Aspergillus nidulans* asexual development, oxidative stress response and secondary metabolism

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There is an error in panel D of [S8 Fig](#). Specifically, the upper panel should read ‘SclB-cYFP + nYFP’, not ‘SclA-cYFP + nYFP’. The authors have provided a corrected version here.

Supporting information

S8 Fig. GFP-fusion proteins of SclB are functional and phosphorylated and Bi-FC controls are negative. A) Strains expressing SclB either N- or C-terminally tagged with sGFP in $\Delta sclB$ background, $\Delta sclB$ and wildtype (WT) were point inoculated on solid MM and grown for 4 days in light. B) SclB-GFP and GFP-SclB fusion proteins expressed under native promoter are visualized in a western hybridization assay employing an α -GFP antibody (GFP) and Ponceau staining as loading control (Pnc). The black arrow indicates bands corresponding to full-length fusion proteins (*in silico* prediction 87.46 kDa). C) Protein crude extracts of GFP-SclB grown vegetatively were mixed with phosphatase inhibitor cocktail (-/PhoI), with Lambda phosphatase ($\lambda/-$), or Lambda phosphatase and phosphatase inhibitor cocktail ($\lambda/PhoI$). A control sample was left untreated (-/-). A subsequent western hybridization assay employing α -GFP antibody visualizes protein bands. D) Two strains, either expressing *sclB::cyfp* and the free second half of the split YFP (*nyfp*; upper part), or free *cyfp* and *rcoA::nyfp* (lower part), under control of a bi-directional nitrate promoter were constructed. Strains were inoculated in liquid MM and analyzed with fluorescence microscopy after 36 h at 30°C. (TIF)



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Reference

- Thieme KG, Gerke J, Sasse C, Valerius O, Thieme S, Karimi R, et al. (2018) Velvet domain protein VosA represses the zinc cluster transcription factor SclB regulatory network for *Aspergillus nidulans* asexual development, oxidative stress response and secondary metabolism. PLoS Genet 14(7): e1007511. <https://doi.org/10.1371/journal.pgen.1007511> PMID: 30044771