

APPENDIX

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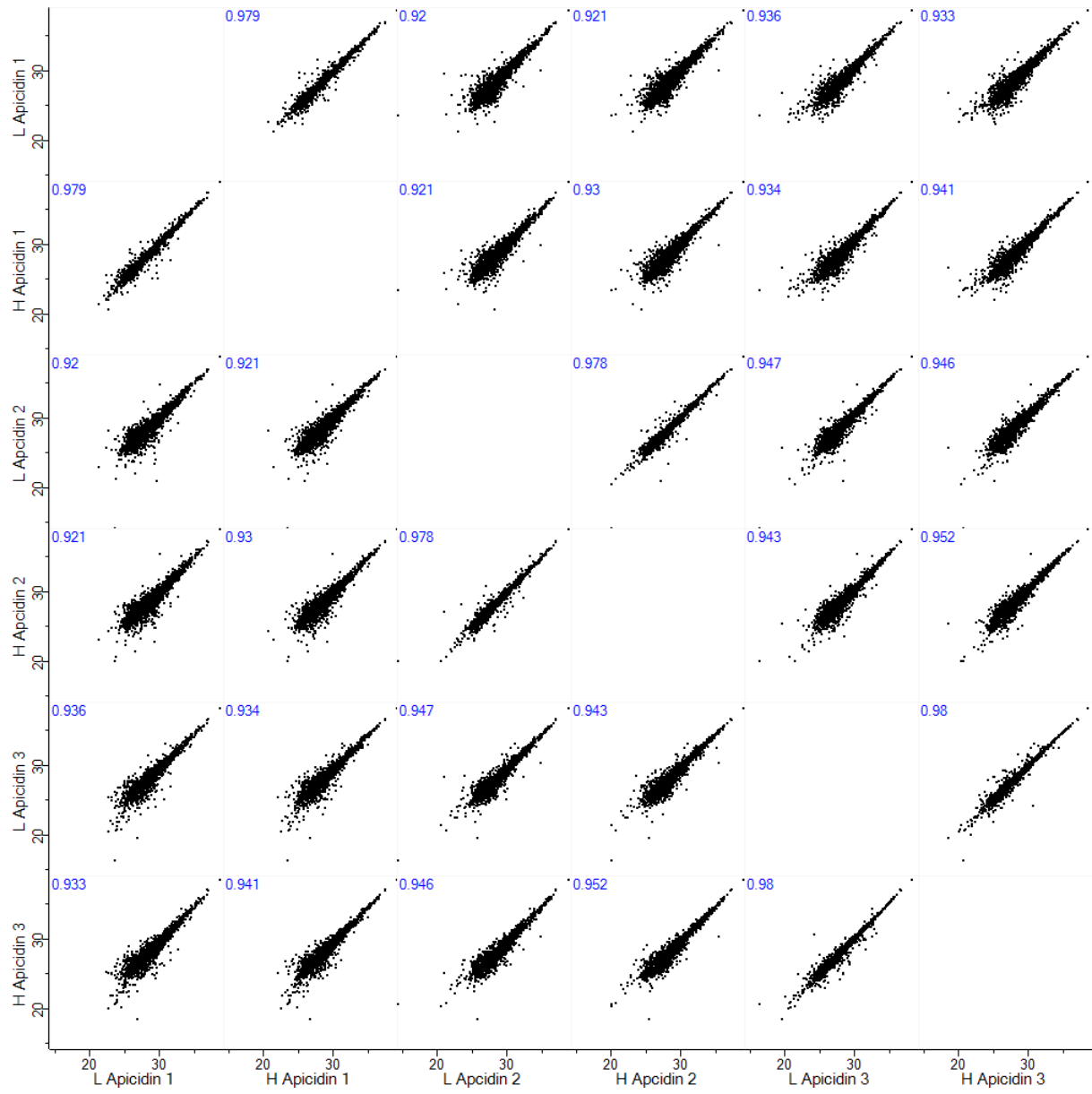
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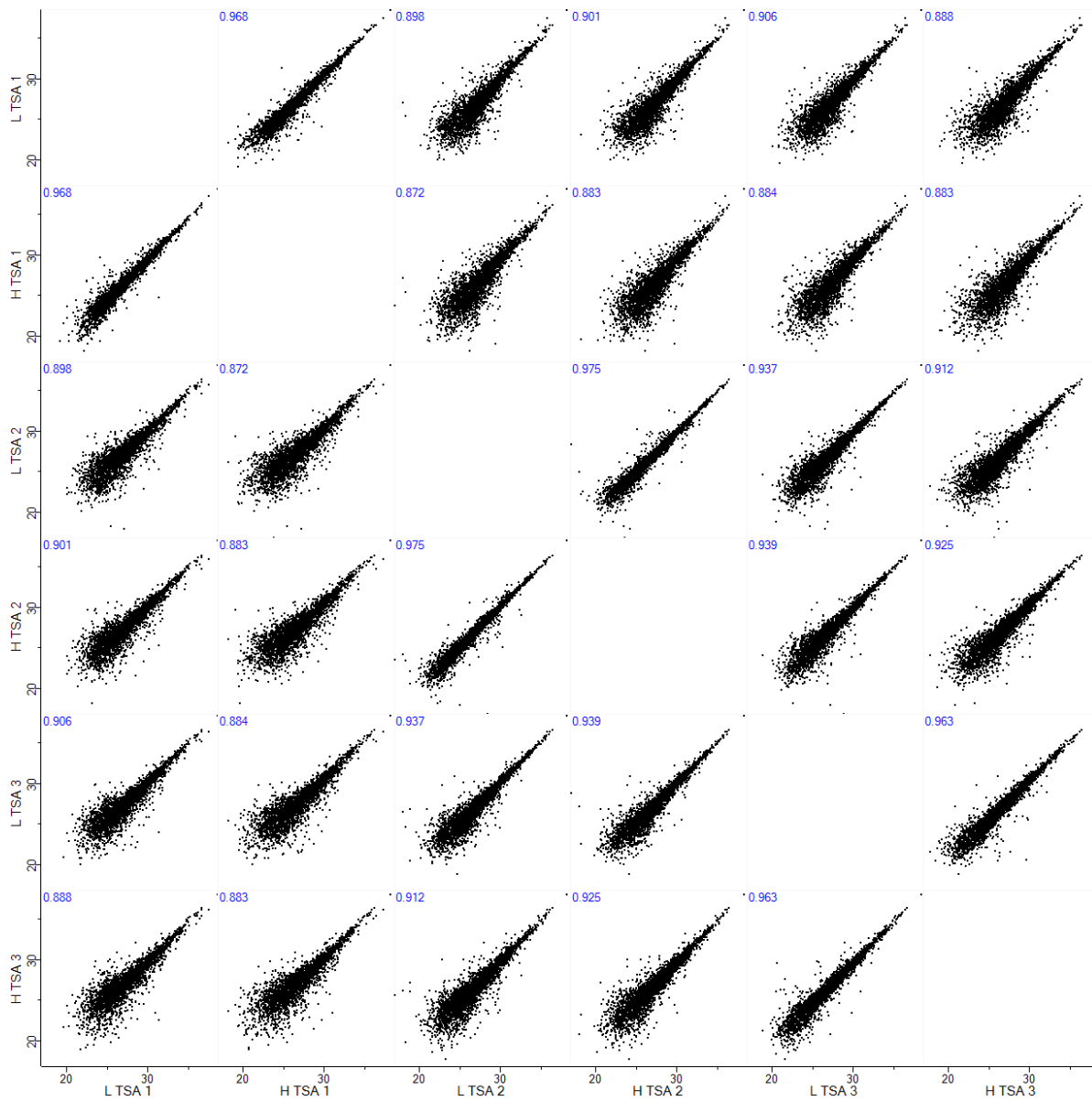
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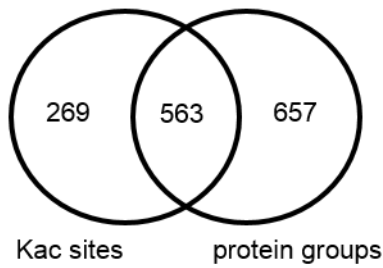
A



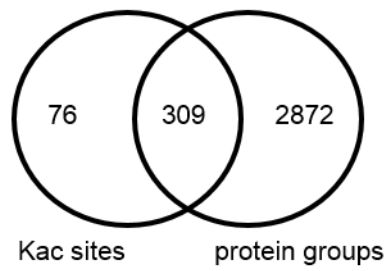
B

Appendix Figure S1: Multiscatter plots of light (L) and medium (H)-dimethyl-labeled non-normalized protein intensities from inhibitor experiments. (A) apicidin, (B) TSA (n = 3).

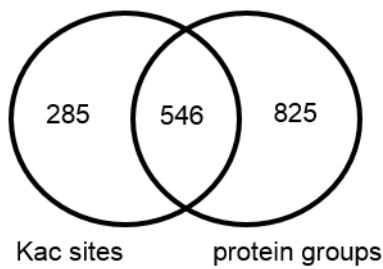
A: apicidin



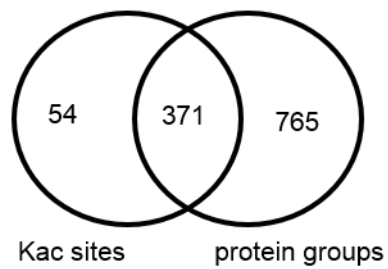
B: TSA



C: *hda14*



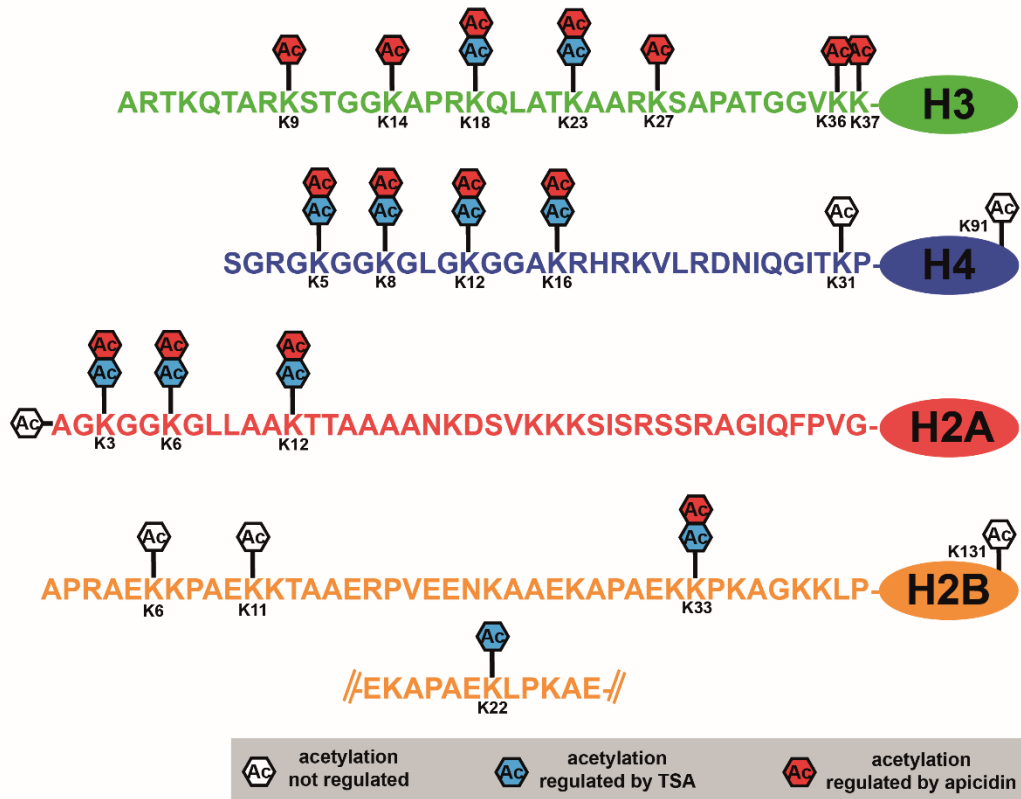
D: thylakoids



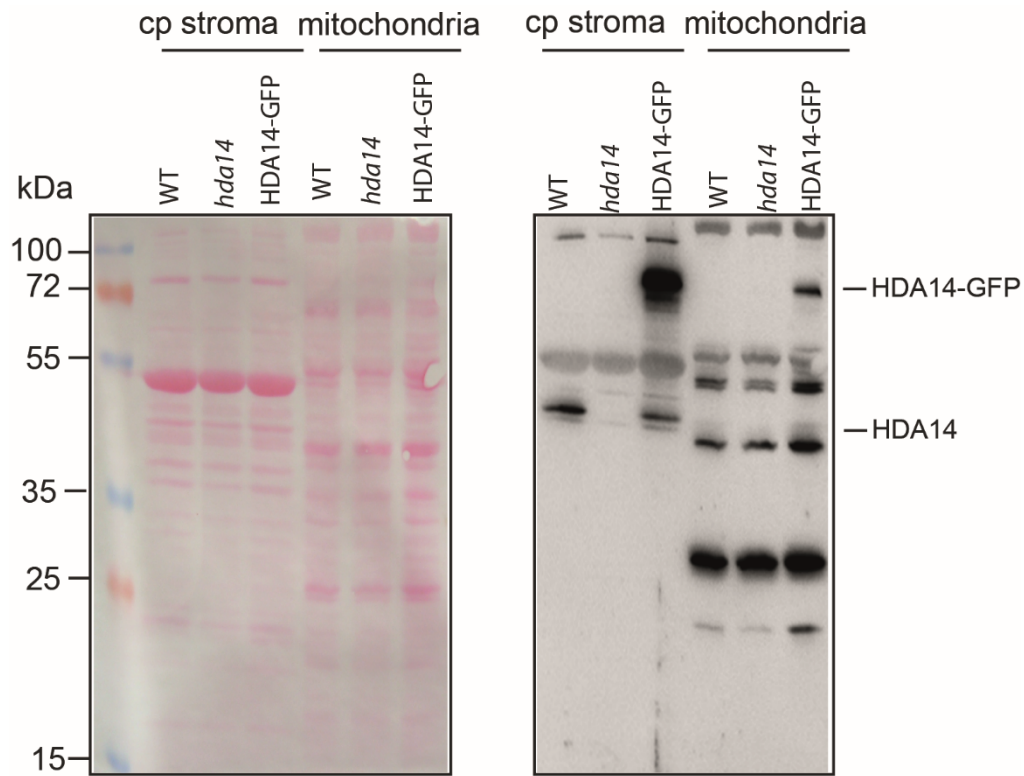
E: 2h low light



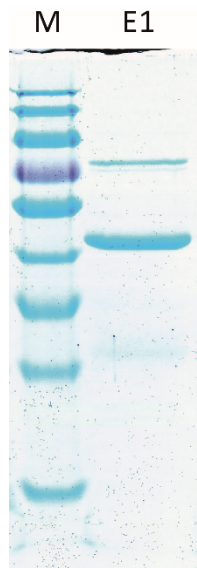
Appendix Figure S2: Venn Diagrams of overlapping quantified acetylation sites and protein groups. Experiments A-E refer to the supplementary datasets EV1-5.



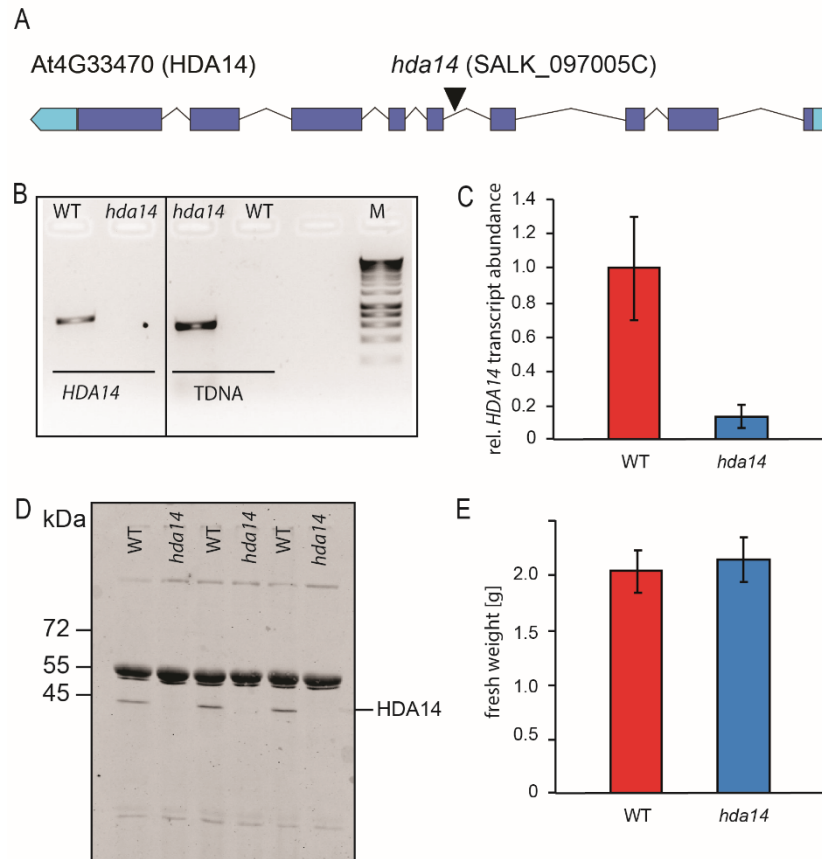
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Appendix Figure S4: Western-blot analysis of HDA14 on isolated chloroplasts and mitochondria. Ponceau S stain (left) Western blot analysis (right) of chloroplast (cp) stroma and mitochondria isolated from WT, *hda14* and 35S:HDA14:GFP seedlings. The endogenous HDA14 and HDA14-GFP fusion proteins are indicated. No HDA14 protein was detected in *hda14* seedlings.



Appendix Figure S5: Purified recombinant 6xHis- Δ 45-HDA14 protein. M: Marker, E1: elution fraction 1.



Appendix Figure S6: Genotyping and phenotyping of *hda14* line. (A) Scheme of the HDA14 gene with the TDNA insertion. (B) PCR with genomic DNA from WT and *hda14* with primers (HDA14_RP and HDA14_LP) which compose the TDNA insertion site (*HDA14*) and primers flanking the left border of the TDNA (SALK-Lb1.3) and gene specific HDA14_RP primers (TDNA). (C) qPCR with cDNA from WT and *hda14* using *HDA14* gene-specific primers. *Actin 2* primers were used as reference, (n = 3, mean \pm SD). For primers see materials and methods. (D) Western blot analysis with WT and *hda14* protein extracts (n = 3) using an antiserum raised against HDA14 protein (Tran et al., 2012). (E) Fresh weight of 5-week old WT and *hda14* plants (n = 21-27, mean \pm SD).