

Fig. S1: Mapman representation of the relative gene expression profile and related metabolic processes in *rpoTnp* dark-grown seedlings. Microarray data sets were processed using the RobiNA web tool and implemented into the MapMan visualization mask according to the recommendations on the website. Gene up-regulation is given by blue squares, gene down-regulation by red squares. A corresponding color code is indicated in the top right corner of the plot. Values refer to log₂ fold changes.

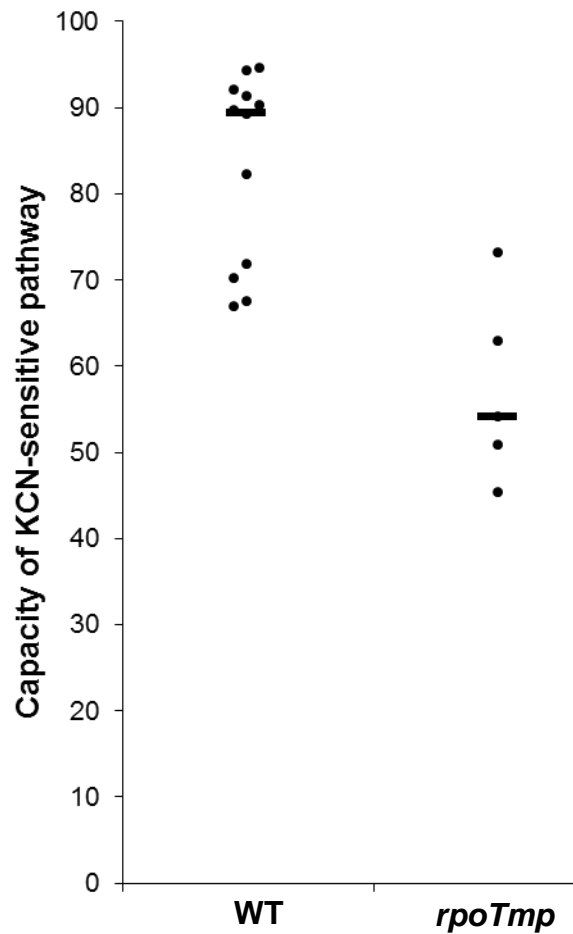


Fig. S2: Capacity of the KCN-sensitive pathway in etiolated seedlings. Capacity of the KCN-sensitive pathway was calculated upon addition of 1 mM SHAM into the measurement cell (final concentration). The figure presents the ratio in percentage of the KCN-sensitive O₂ consumption rate to the total consumption rate. Median values (thick horizontal lines) of N independent measurements (full circles) were scatter-plotted for WT (N=12) and *rpoTmp* (N=5).

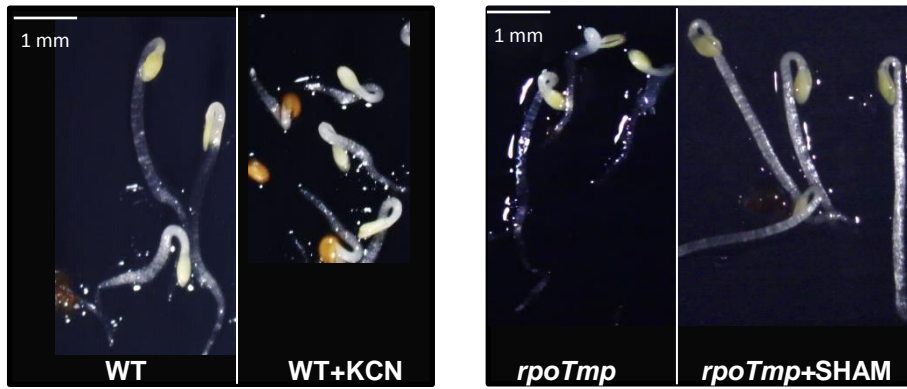


Fig. S3: Images of etiolated inhibitor-treated plants. Phenotype of the WT and *rpoTmp* grown in absence or in presence of 1 mM KCN (left panel) and 1mM SHAM (right panel), respectively. Scale bar corresponds to 1 mm.

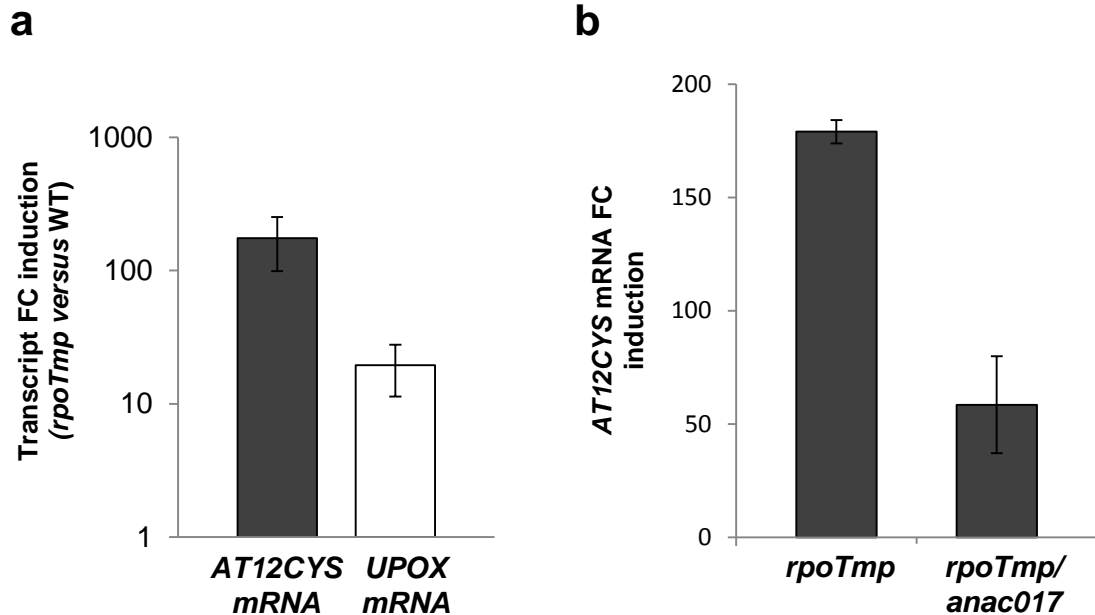


Fig. S4: Expression changes of oxidative stress marker genes in etiolated mutant plants compared to WT. a) Linear fold change (FC) induction of *AT12CYS* and *UPOX* transcripts in *rpoTmp* plants versus WT. Transcript expression levels were determined by RT-qPCR and normalized with the mean of *ACTIN 2-8* expression, used as reference gene. The mean values of three technical replicates are plotted. Error bars correspond to standard errors. **b)** Linear fold change (FC) induction of *AT12CYS* transcripts in *rpoTmp* and *rpoTmp/anac017* plants versus WT. The mean values of two biological RT-qPCR replicates are plotted.

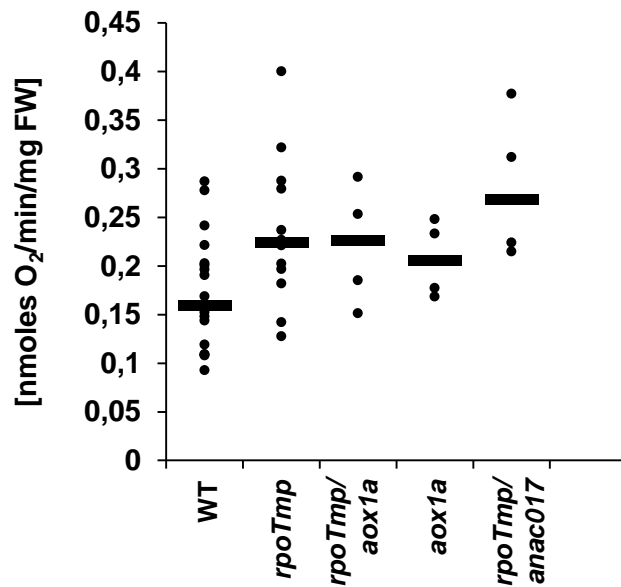


Fig. S5: Total O₂ consumption rate in etiolated mutant plants. Median values (thick horizontal lines) of N independent measurements (full circles) were scatter-plotted for WT (N=19), *rpoTmp* (N=12), *rpoTmp/aox1a* (N=5), *aox1a* (N=4), *rpoTmp/anac017* (N=4).

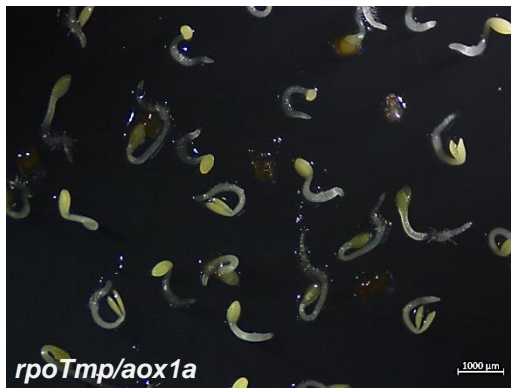
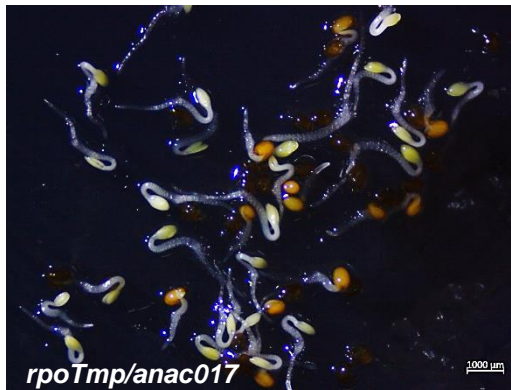


Fig. S6: Images of etiolated *rpoTmp*, *rpoTmp/anac017* and *rpoTmp/aox1a* mutant plants. Scale bar corresponds to 1 mm.

qPCR primers	
<i>AOX1a</i> FWD	GACGGTCCGTACGGTTTCG
<i>AOX1a</i> REV	CTTCTGATTCGCGTCCTCCTCCT
<i>AT12CYS2</i> FWD	GCCTCCTCAATCTGTGAACC
<i>AT12CYS2</i> REV	CCAAATGCAGTTCCAGTACC
<i>UPOX</i> FWD	CCGAGAACCCGCCAAAACC
<i>UPOX</i> REV	GCTTCTCTGCAACTGCCTC
<i>ACT 2.8</i> FWD	GGTAACATTGTGCTCAGTGGTGG
<i>ACT 2.8</i> REV	AACGACCTTAATCTTCATGCTGC

Table S1: Primers used in qPCR for expression analyses.