

Figure S1. RT-qPCR analysis of SS1, SS3, ISA3 and LDA expression in tobacco Wt, o/exTrxf and o/exNTRC homoplasmic plants (T₁ generation). Graphical representation of relative expression levels normalized to 16S rRNA expression that shows the fold change as the mean of the different biological repeats (n=3). Error bars represent SE.

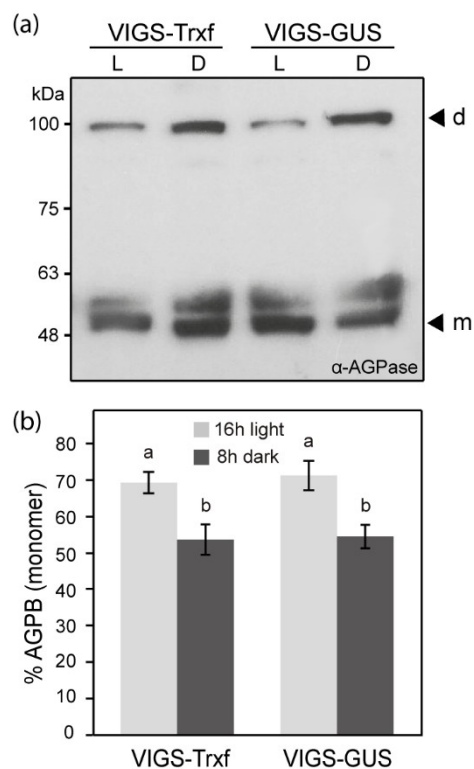


Figure S2. Redox activation of AGPase in VIGS-Trxf *N. benthamiana* leaves. Redox status of AGPase of mature leaves from Wt and VIGS-Trxf plants grown in phytotron sampled at the end of the light (L; 80 $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$) and dark (D) periods. (a) A representative non-reducing western blot is shown. Specific AGPB antibody was used. m: monomer; d: dimer. (b) Quantification of AGPB monomerisation by western blot analysis shown as the percentage of the 50 kDa monomer relative to the total amount of AGPB. Results are the mean \pm SE of four individual plants. Different letters above the bars represent significant differences ($P < 0.05$, ANOVA).

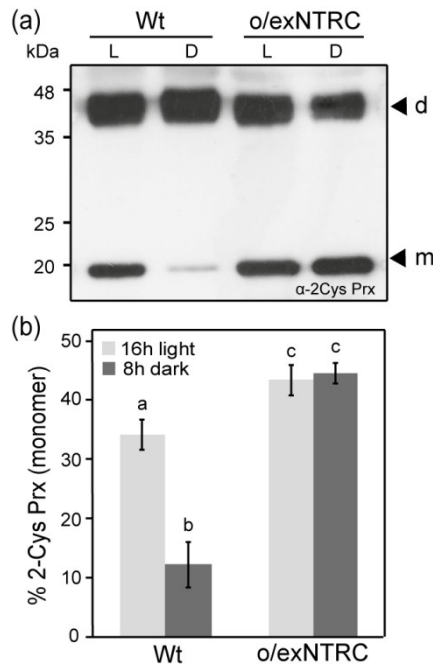


Figure S3. Redox status of 2-Cys Prx in the T₁ generation of o/exNTRC homoplasmic plants. Redox activation of 2-Cys Prx in leaves from 7-week-old o/exNTRC tobacco plants grown in phytotron and sampled at the end of the light (L) and dark (D) periods. (a) A representative non-reducing western blot of 2-Cys Prx in leaves of Wt and o/exNTRC plants is shown. m: monomer; d: dimer. (b) Quantification of 2-Cys Prx monomerization shown as the percentage of the 20 kDa monomer relative to the total amount of 2-Cys Prx. Results are the mean \pm SE for five plants. Different letters above each bar indicate significant differences ($P < 0.05$, ANOVA).

Table S1. Phenotypic characterization of the T₁ generation of o/exNTRC homoplasmic plants grown in phytotron under standard conditions at different developmental stages. Values are the mean \pm SE for 6 individual plants. Statistical significance compared to Wt plants is indicated by asterisks ($P < 0.05$, Student's t-test).

| | Wt | o/exNTRC |
|----------------------------|----------------|-----------------|
| <i>Flowering stage</i> | | |
| Height (cm) | 82.5 \pm 4.2 | 70.0 \pm 2.7* |
| Chlorophyll content (SPAD) | 32.8 \pm 1.5 | 30.5 \pm 0.9 |
| <i>Maturity</i> | | |
| Height(cm) | 87.0 \pm 4.5 | 86.7 \pm 4.5 |
| Chlorophyll content (SPAD) | 26.5 \pm 1.9 | 26.8 \pm 0.9 |

Table S2. List of primers used in these work.

| Primers used for NTRC amplification (5'-3')^{a,b} | | |
|------------------------------------------------------------------|----------------------------------------------------------|-----|
| <i>NTRC_for</i> | ccatgggt catcaccatcaccatcac GGATCCTTCTTCAGAGGCGAG | |
| <i>NTRC_rev</i> | gcggccgcTTATTTATGGCCTCAA | |
| Primers used for VIGS vector construction | Amplicon size (bp) | |
| <i>pTRV2-NTRC_for</i> | AGTgaattcCTGATACTCAGGAAGAATC | |
| <i>pTRV2-NTRC_rev</i> | CTAggatccACTGTACACATGGAGTAC | 585 |
| <i>pTRV2-trxf_for</i> | AGTgaattcGTTGCAGGAGATTATGGC | |
| <i>pTRV2-trxf_rev</i> | CTAggatccCTTTGGAGCAATCAC | 228 |
| Primers used for RT-qPCR analysis | Amplicon size (bp) | |
| <i>Actin_for</i> | CAGCAACTGGGATGATATGG | |
| <i>Actin_rev</i> | GGCGCTTCAGTTAAGAGGAC | 99 |
| <i>NTRC_for</i> | TGACTTAATGGATAGGATGAG | |
| <i>NTRC_rev</i> | GTTCACTACTTTGGATAGTAA | 116 |
| <i>trx_f_for</i> | CAAGTAAACGGCGTATCATT | |
| <i>trx_f_rev</i> | CAGACCACCGATTGCTTGCT | 81 |
| <i>16S rRNA_for</i> | CTTTTTAAGTCCGCCGTCAAA | |
| <i>16S rRNA_rev</i> | TCTTTCCGATCTCTACGCATTTT | 127 |
| <i>SS1_for</i> | GCTGAAGCAGCTCCGTATTC | |
| <i>SS1_rev</i> | GAGACGACCATTACACGATG | 101 |
| <i>SS3_for</i> | AAAACCCGATGAAGATGTCG | |
| <i>SS3_rev</i> | GCGCCAGTCATTA AAAAGCTC | 100 |
| <i>ISA3_for</i> | AGTGCTTCTCGGGAGTTCAA | |
| <i>ISA3_rev</i> | CGGGTTTTTGTTCATCAGCTT | 111 |
| <i>LDA_for</i> | GTGAAGGCAAAGTCATTGCT | |
| <i>LDA_rev</i> | TGTTTGCCACTTCACCAAAA | 109 |

^a Restriction sites are indicated in lowercase

^b 6xHis tag is indicated in bold.