

Supplementary data

Supplementary Tables

Sup. Table 1 Segregation analysis of F2 progeny of five independent WTxGFP6.4 F1 plants (L1-L5) for resistance to kanamycin and GFP fluorescence. Percentage of kanamycin resistant and GFP florescent plantlets out of n seedlings germinated and grown on agar-based Murashige and Skoog medium supplemented with 100 $\mu\text{g.mL}^{-1}$ kanamycin. Expected % of kanamycin resistant and GFP fluorescent plantlets for one and two unlinked T-DNAs carrying a functional nptII and mGFP4 transgenes are 75% and 93% respectively. n = number of germinated seeds.

Line	Kanamycin Resistant (%)	GFP Fluorescent (%)	n
WTxGFP6.4 L1 F2	92	92	223
WTxGFP6.4 L2 F2	92	92	90
WTxGFP6.4 L3 F2	92	92	106
WTxGFP6.4 L4 F2	93	93	60
WTxGFP6.4 L5 F2	90	90	70

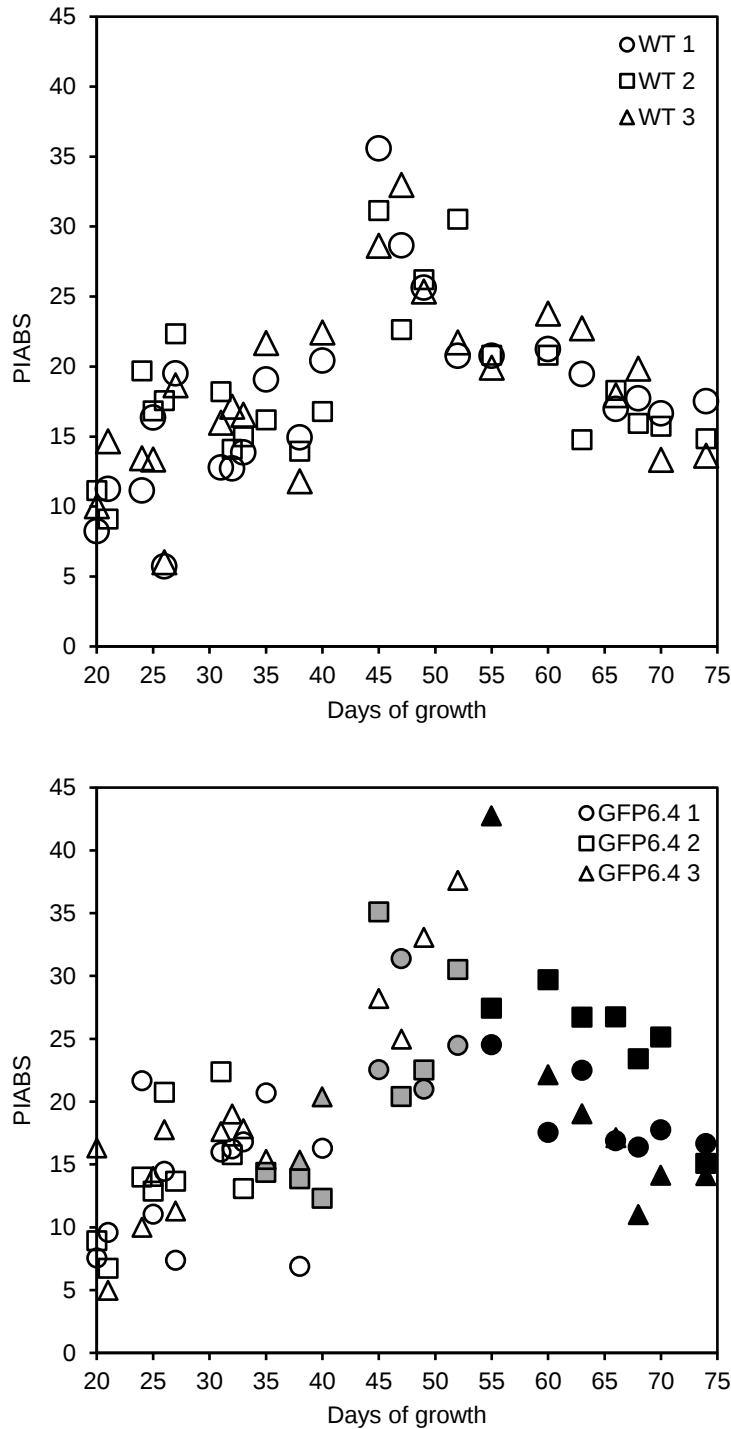
Sup. Table 2 Total number of reads sequenced in each sample. Four biological and three technical replicates were analyzed.

Sample ID	Total number of reads (fastq)	Platform	Experiment type
WT1	10578188	ILLUMINA	3'Quant seq
WT2	9052115	ILLUMINA	3'Quant seq
WT3	11095121	ILLUMINA	3'Quant seq
WT4	10320079	ILLUMINA	3'Quant seq
OS1	10352753	ILLUMINA	3'Quant seq
OS2	11306535	ILLUMINA	3'Quant seq
OS3	10098292	ILLUMINA	3'Quant seq
OS5	10466610	ILLUMINA	3'Quant seq
NS1	10328154	ILLUMINA	3'Quant seq
NS2	9625750	ILLUMINA	3'Quant seq
NS3	11977619	ILLUMINA	3'Quant seq
NS4	9627883	ILLUMINA	3'Quant seq
SS1	12241420	ILLUMINA	3'Quant seq
SS2	10833903	ILLUMINA	3'Quant seq
SS3	10438408	ILLUMINA	3'Quant seq
SS4	10522291	ILLUMINA	3'Quant seq

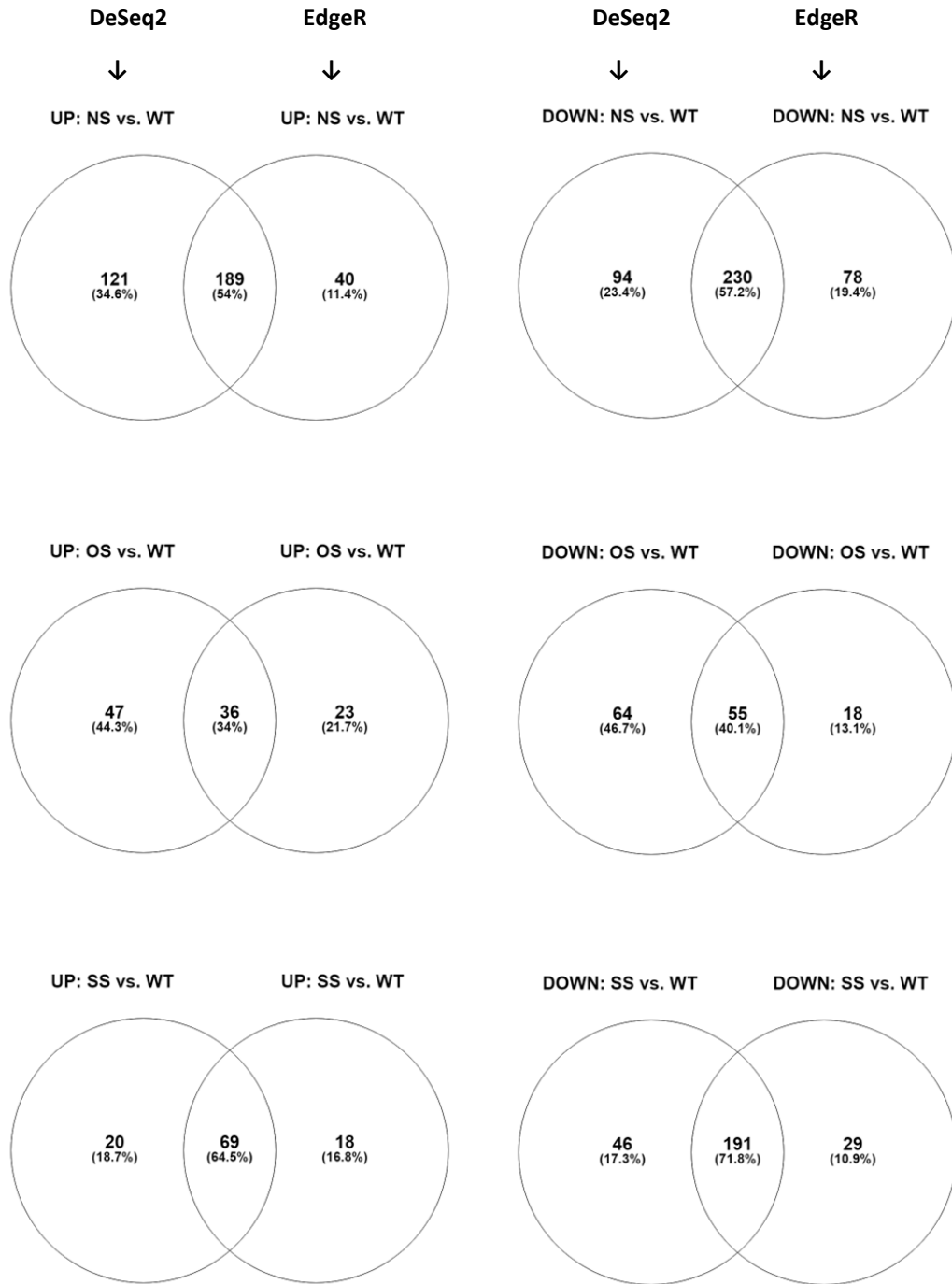
Sup. Table 3 DNA sequence of primers used for the amplification of target sequences for qPCR.

Target	Forward primer 5'-3'		Reverse primer 5'-3'		T _m	Amplicon size
CASP-like	CASP-F	AAGCAAACTCCACATCTCTCA	CASP-R	CGGATATTGGAGGCAGATTACTCA	F- 58.4 R- 61.0	113bp
Peroxidase 15	PEROX-F	TGGCTCATTCTTCTTCTTAGCCA	PEROX-R	ACTCAACTGAGCAATTTGATTTCATAG	F- 59.3 R- 58.1	120bp
BnaA05g03240D [Brassica napus]	BNA-F	CAGCTGGTCACCGACA	BNA-R	ACCACTGGCTTAACAGCTAAT	F- 54.3 R- 55.9	83bp
Tetratricopeptide repeat (TPR)-like superfamily	TPR-F	GAGAGAGAGGCGATGCTTTAT	TPR-R	TGATCCGAACAGCCCAATTCA	F- 57.9 R- 57.9	120bp
Cytochrome P450 superfamily	P450-F	AGATCTCCTCACTGAGTTCTGTA	P450-R	AGACACACAATAAAACTTCTTCTGG	F- 58.9 R- 58.1	107bp
C2 calcium/lipid-binding plant phosphoribosyltransferase	CAL-F	TGCCCTCTGAAGTTGCTAGTCT	CAL-R	GAAGAGATGTCAGTGTGTGTCGT	F- 57.9 R- 58.4	81bp
Protein kinase superfamily	KIN-F	CTCAGAAATGGGGGCTGTCA	KIN-R	TCTCCACCAAAATTTCTCTCCATC	F- 59.4 R- 58.4	98bp
UDP-Glycosyltransferase superfamily	UDP-F	TCCGTGACAGGTGGATTTC	UDP-R	GGTACCCCTTGTCCAATCGC	F- 59.4 R- 61.4	105bp
Stress responsive A/B Barrel Dom	BAR-F	TGAGGAATCTTGCAATAAAGGCT	BAR-R	TGGGCCCCATCTAAGAGGGTA	f- 57.1 r- 59.4	118bp
protein RESPONSE TO LOW SULFUR 1	RLS-F	TTGATCAGGCTCGGACCTAC	RLS-R	CGGTACGGGAAGCTGATTCCG	F- 59.4 R- 59.4	93bp
WRKY transcription factor 44	WRKY44-F	CCCCTGTGTGAAAGGATTGC	WRKY44-R	TCCAACCTTGTTTGGAGAGGT	F- 59.4 R- 57.9	118bp

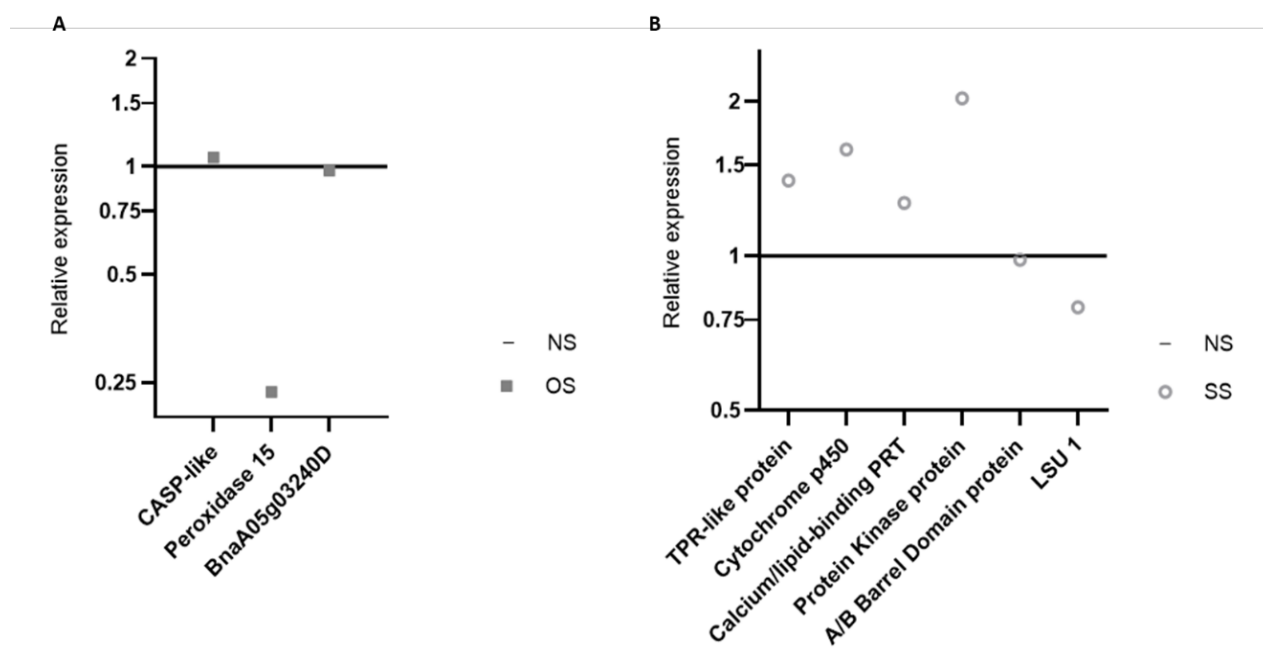
Supplementary figures



Sup. Fig. 1 Initiation and progression of GFP-PTGS do not affect photosynthetic activity in line GFP6.4. Performance index on absorption basis (PIABS) in function of time (number of days of growth) of three representative WT (a) and GFP6.4 plants (b) obtained from OJIP-test time series analysis. In (b), open, grey and closed markers indicate that leaf subjected to OJIP-test analysis was NS in NS plants, NS in plants presenting either OS or SS in distant leaf tissues, and SS in plants presenting either OS or SS in distant leaf tissues respectively.



Sup. Fig.2 Transcriptional reprogramming in lineGFP6.4 NS, OS and SS versus WT plants. Venn diagrams present up- and down-regulated gene sets.



Sup. Fig.3 Differentially expressed genes between: A) NS-OS GFP6.4 plants and B) NS-SS GFP6.4 plants. qPCR quantified data are presented as relative expression levels