

Transcriptional regulation of the glucose-6-phosphate/Phosphate translocator 2 is related to carbon exchange across the chloroplast envelope

Supplementary Tables and Figures

Sean E. Weise^{1,2}, Tiffany Liu³, Kevin L. Childs³, Alyssa L. Preiser¹, Hailey M. Katulski¹, Christopher Perrin-Porzondek¹, Thomas D. Sharkey^{1,2,4*}

¹MSU-DOE-Plant Research Laboratory, Michigan State University, East Lansing, MI USA

²Department of Biochemistry and Molecular Biology, Michigan State University, East Lansing, MI USA

³Department of Plant Biology, Michigan State University, East Lansing, MI USA

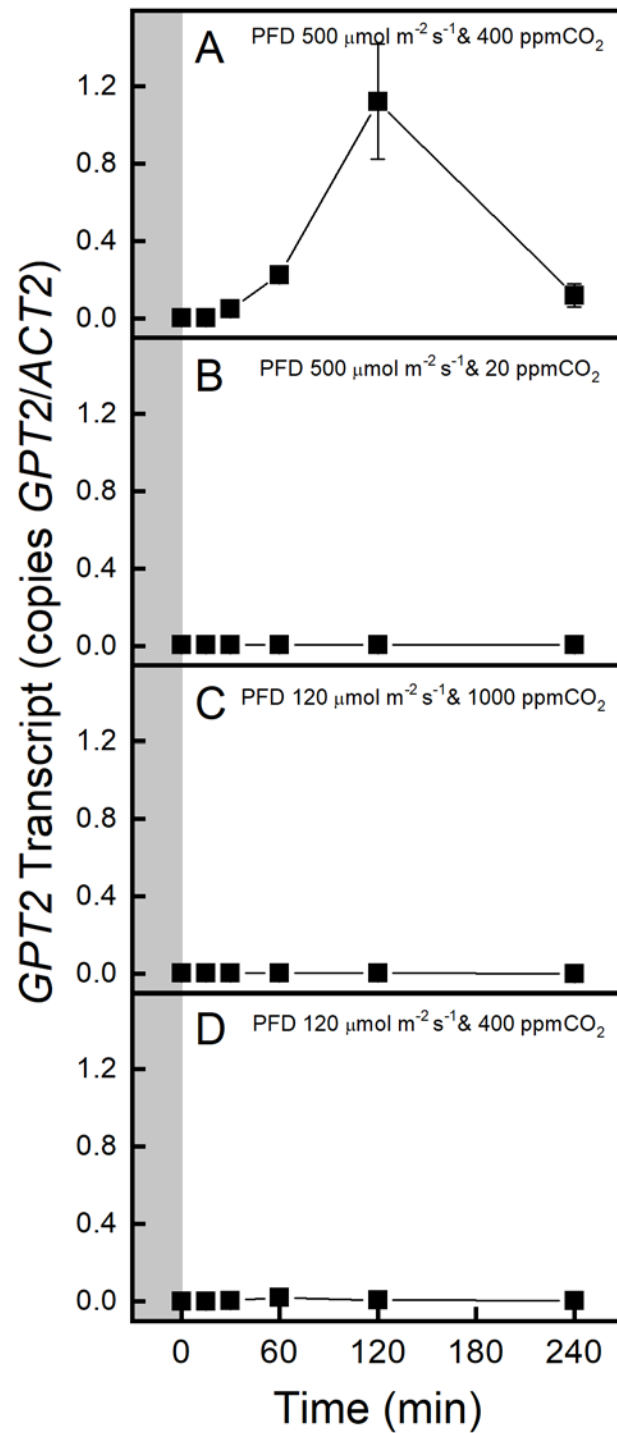
⁴Plant Resilience Institute, Michigan State University, East Lansing, MI USA

Supplementary Table S1. Primer sequences used in this study

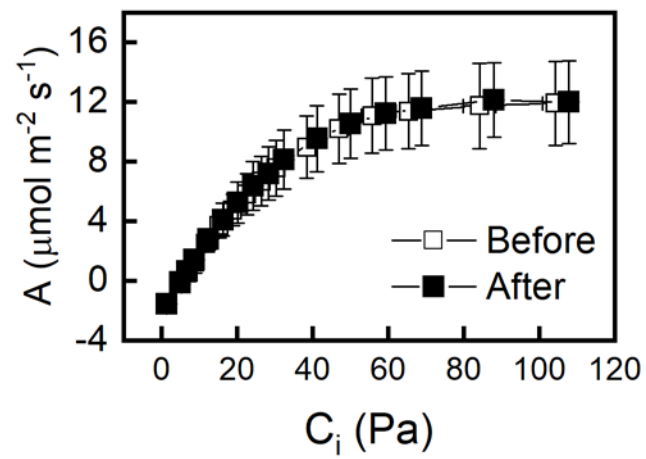
Gene	Gene Locus	Standard Generating Primers	qPCR primers
<i>ACT2</i>	At3g18780	GGTGATGAAGCACAATCCAA CAGTAAGGTCACGTCCAGCA	CAAAGGCCAACAGAGAGAAGA ATCACCAGAATCCAGCACAA
<i>IDI2</i>	At3g02780	CCGAATTTTCGTGCTTTCTC TGCAGCATTCTCACCACCTA	GTATGAGTTGCTTCTCCAGCAAAG GAGGATGGCTGCAACAAGTGT
<i>GPT2</i>	At1g61800	CGTGGAAGGTCCTCAAATGT TCACTGCTTCGCCTGTGAGT	AGACCAGATTTTCGCCGTTAACT ACTGCTTCGCCTGTGAGTAGAG
<i>XPT</i>	At5g17630	AGGATCACATTGGGTTCAG CTTTGCAGTGGCCTGAGAAT	CATTGCATCTGTTGGGACAC TGGACTGATCTCGTCGAGTG
<i>G6PDH1</i>	At5g35790	GAGAATGGCTGGACAAGGGT GCCGGTGTGAATAGATCCCA	ATTTCGGAACAGAAGGGCGT CAACCACATCTTCAAGCCGC
<i>G6PDH2</i>	At5g13110	TGGAAGAATCTCAAACCGCCT AGCCTCTCATATGCATCTGGT	ATTTCGGCACTGAAGGACGT GCGTTTCCATGGCAAAGAGG
<i>G6PDH3</i>	At1g24280	GAGTCTGATGGCGGTGAACA CTGCAAATGTCGGGGTCAGA	TGCATTGGACGAGAAGCTCA CACATTTACCGCGTCAACA
<i>RRTF1</i>	At4g34410	CCCAACCCGGTATCAAAAGGA CTTGGCCCACGGAATCCAAT	GCCTCGGTTGGATTTCAGACA TGATTTTCCCGCCACCTTCC

Supplementary Table S2. Specific differentially expressed *Arabidopsis* genes and loci used in heat maps in figure 4 and supplementary figure S5.

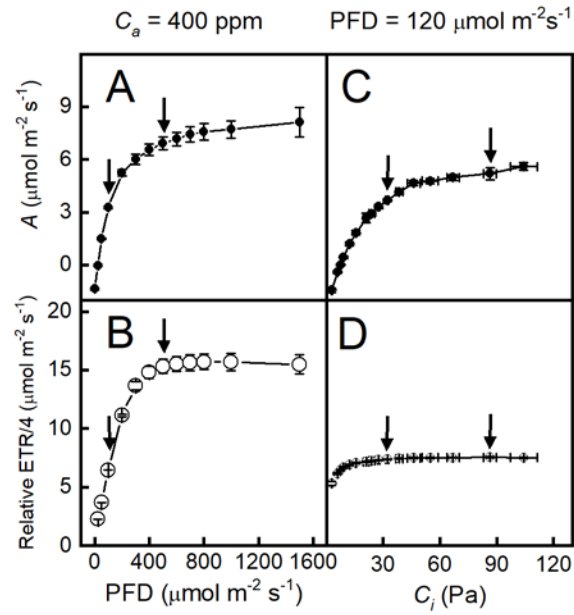
Gene	Locus	Gene	Locus
Calvin-Benson Cycle		TP to G6P	
<i>RBCS1A</i>	At1g67090	<i>TPT</i>	At5g46110
<i>RBCS1B</i>	At5g38430	<i>TPI</i>	At3g55440
<i>RBCS2B</i>	At5g38420	<i>FBA5</i>	At4g26530
<i>RBCS3B</i>	At5g38410	<i>FBA6</i>	At2g36460
<i>RCA</i>	At2g39730	<i>FBA7</i>	At4g26520
<i>GAPA</i>	At3g26650	<i>GPT2</i>	At1g61800
<i>GAPA2</i>	At1g12900	G6P to TP	
<i>FBA2</i>	At4g38970	<i>PFP81</i>	At4g04040
<i>CFBP2</i>	At5g64380	G6P to Sucrose	
<i>SBPASE</i>	At3g55800	<i>UGP2</i>	At5g17310
<i>PRK</i>	At1g32060	<i>SPS3</i>	At1g04920
<i>CP12-2</i>	At3g62410	<i>SPS4</i>	At4g10120
Starch Synthesis		<i>SPP2</i>	At2g35840
<i>APL2</i>	At1g27680	<i>G6PDH1</i>	At5g35790
<i>APL3</i>	At4g39210	<i>G6PDH2</i>	At5g13110
<i>APL4</i>	At2g21590	<i>G6PDH3</i>	At1g24280
<i>SS2</i>	At3g01180	<i>GAPN</i>	At2g24270
<i>SS3</i>	At1g11720		
<i>SS4</i>	At4g18240		
<i>GBSS1</i>	At1g32900		
<i>SBE1</i>	At3g20440		
<i>SBE3</i>	At2g36390		
<i>PHS1</i>	At3g29320		
<i>GWD</i>	At1g10760		
<i>PWD</i>	At5g26570		



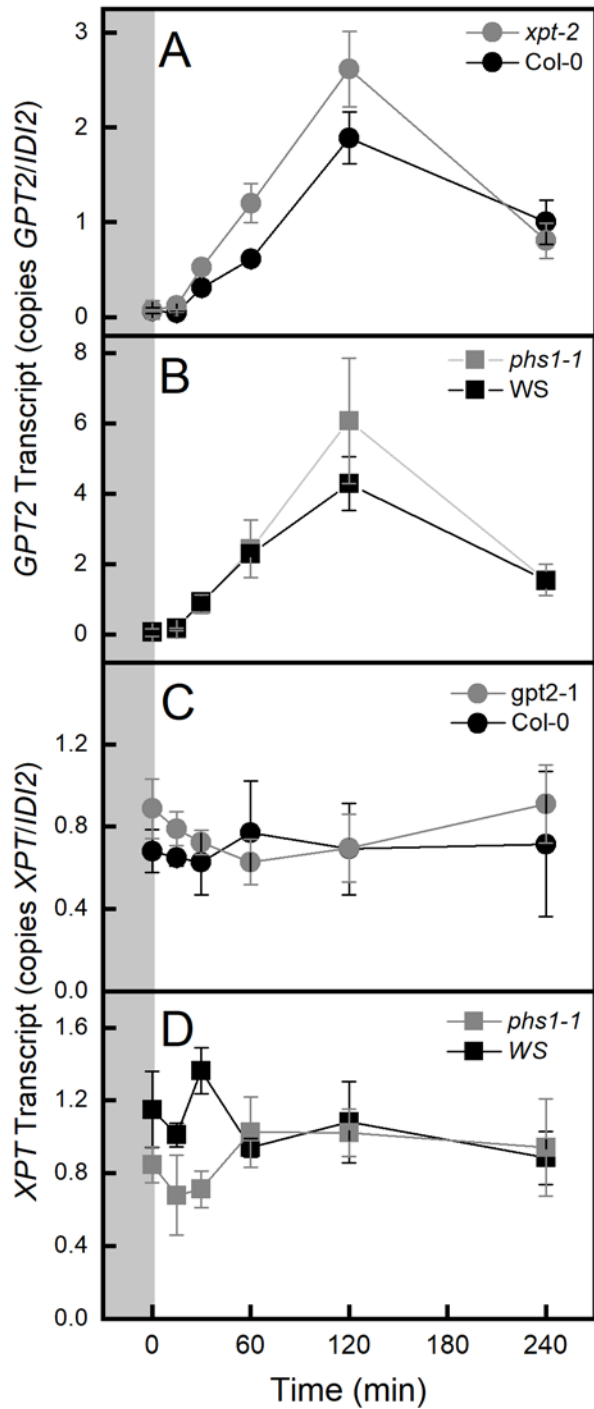
Supplementary Figure S1. Expression of *GPT2* in WS before and after transfer to an altered light or CO_2 environment. The grey block indicates sampling in growth conditions, PFD 120 $\mu\text{mol m}^{-2} \text{s}^{-1}$ and 400 ppm CO_2 . $n = 5$



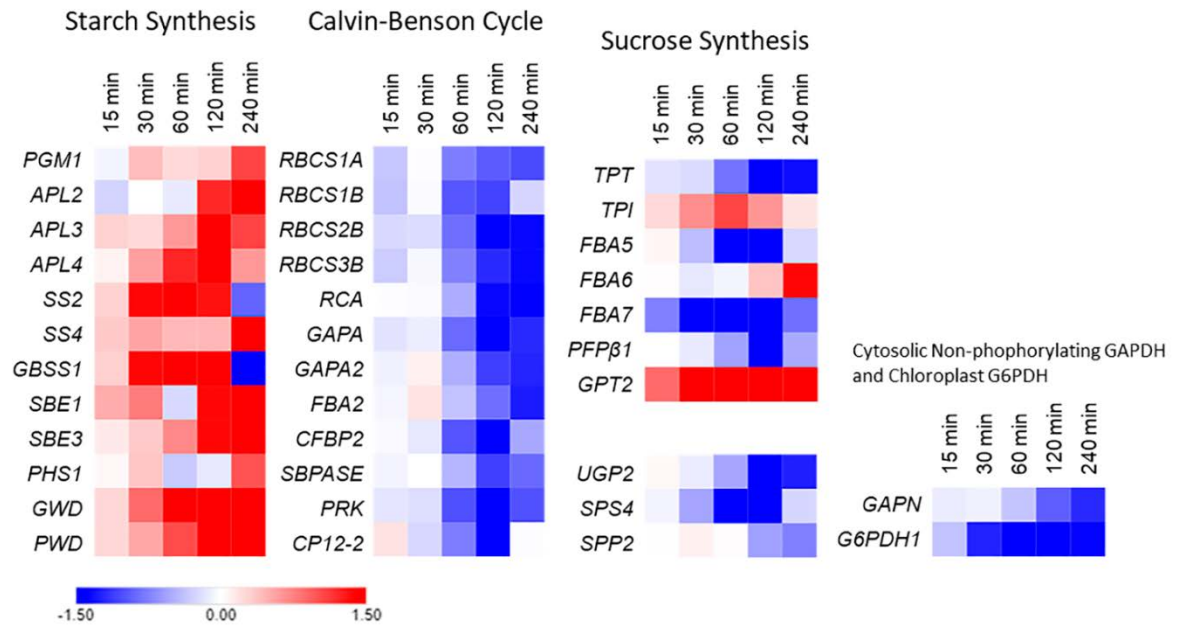
Supplementary Figure S2. A – C_i curves of WS taken before and after 4 hr incubation at a PFD of $500 \mu\text{mol m}^{-2} \text{s}^{-1}$ and C_a of 400 ppm. A – C_i curves were done at a PFD of $500 \mu\text{mol m}^{-2} \text{s}^{-1}$ n=5



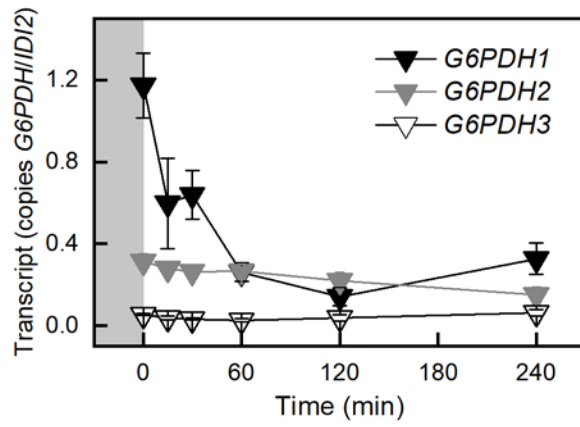
Supplementary Figure S3. Photosynthetic and relative electron transport (ETR) rates in Col-0 in varying light and CO_2 environments. Arrows indicate the PFD or average C_i values that correspond to the PFD or C_a values used in transcript experiments. ETR/4 is taken as the relative approximate rate of the Calvin-Benson cycle turnover. $n=5$



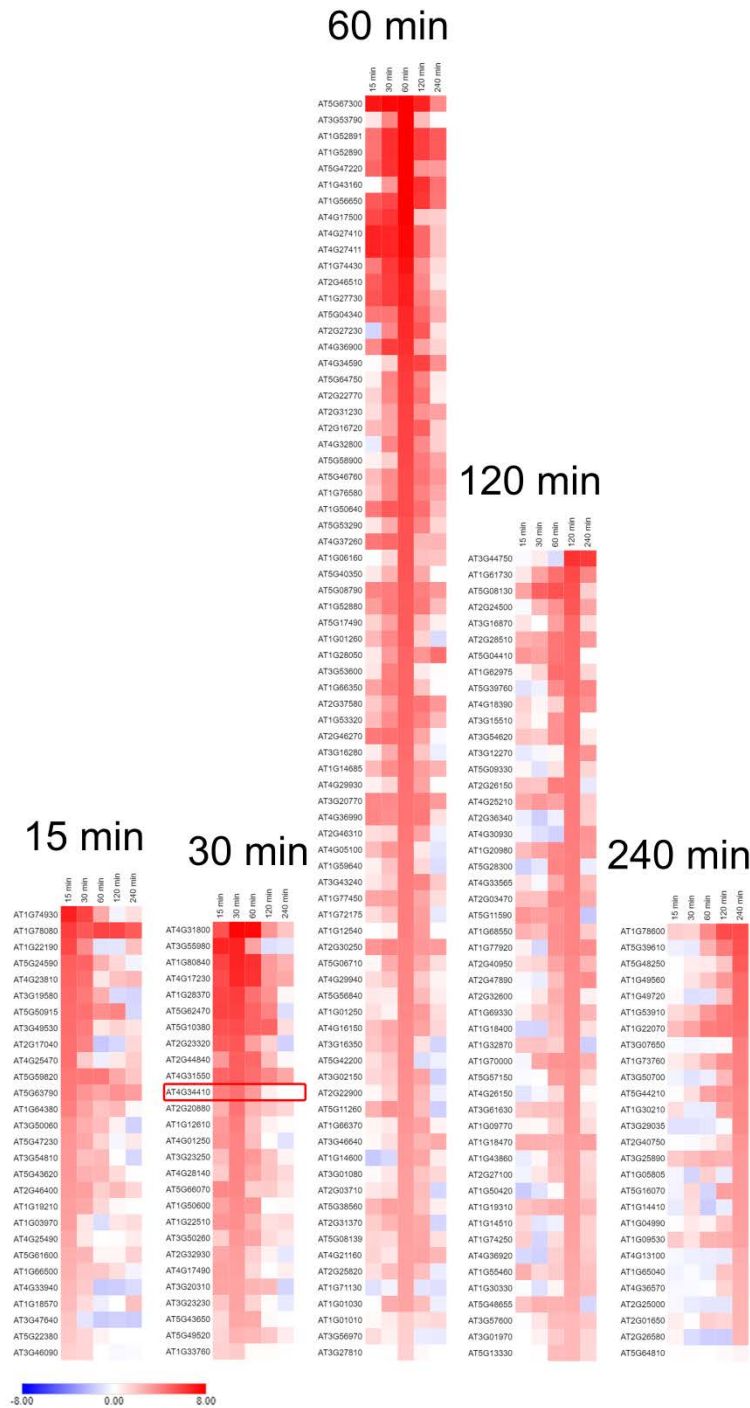
Supplementary Figure S4. Expression of *GPT2* and *XPT* in *xpt-2*, *gpt2-1*, and *phs1-1* mutants in response to a PFD of 500 $\mu\text{mol m}^{-2} \text{s}^{-1}$ C_a of 400 ppm. The grey block indicates sampling in growth conditions, PFD 120 $\mu\text{mol m}^{-2} \text{s}^{-1}$ and 400 ppm CO_2 . n = 5



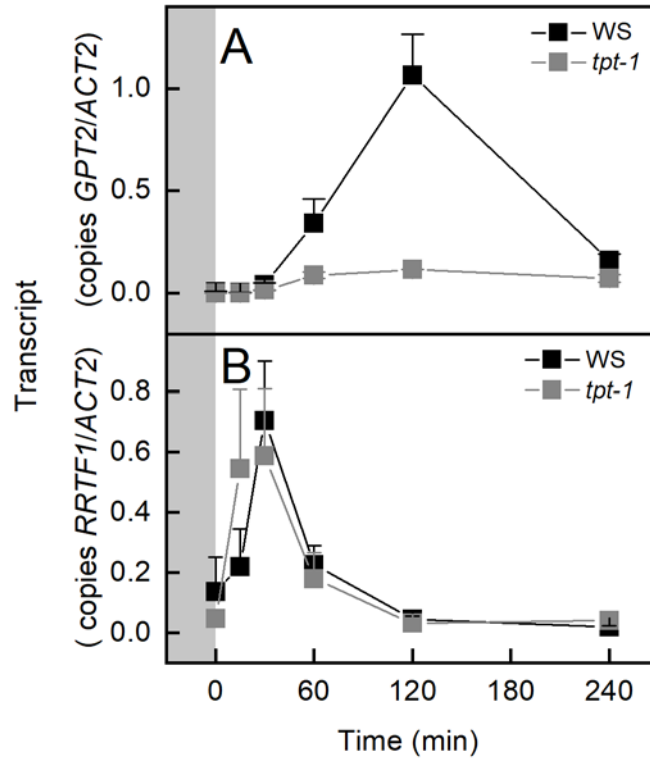
Supplementary Figure S5. Changes in transcription in WS in response to an increase in PFD from 120 to 500 $\mu\text{mol m}^{-2} \text{s}^{-1}$. Red boxes represent an increase in transcription and blue a decrease. Expression values shown had a \log_2 RPKM fold change greater than 1 and are on an absolute scale of -1.5 to 1.5 $n=3$



Supplementary Figure S6. Changes in transcription of the plastid *G6PDHs* in WS in response to a PFD of $500 \mu\text{mol m}^{-2} \text{s}^{-1}$ C_a 400 ppm. Grey block indicates sampling in growth conditions, PFD $120 \mu\text{mol m}^{-2} \text{s}^{-1}$ and 400 ppm CO_2 . $n=5$



Supplementary Figure S7. Transcription factors in WS whose expression was increased in response to an increase in PFD from 120 to 500 $\mu\text{mol m}^{-2} \text{s}^{-1}$. Transcription factors are grouped according to the time they had the highest transcript abundance after transfer to high light. The red box is around the *RRTF1* transcription factor. Expression values used for heat map generation are the $\log_2 \Delta\text{RPKM} + 1$ and are on an absolute scale of -8 to 8. $n=3$



Supplementary Figure S8. Expression of *GPT2* and *RRTF1* in response to an increase in PFD from 120 to 500 $\mu\text{mol m}^{-2} \text{s}^{-1}$ C_a 400 ppm in WS and the *tpt-1* mutant. Grey block indicates sampling in growth conditions, PFD 120 $\mu\text{mol m}^{-2} \text{s}^{-1}$ and 400 ppm CO_2 . $n=5$