Widening the landscape of transcriptional regulation of algal photoprotection

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Abstract

Availability of light and CO₂, substrates of microalgae photosynthesis, is frequently far from optimal. Microalgae activate photoprotection under strong light, to prevent oxidative damage, and the CO₂ Concentrating Mechanism (CCM) under low CO₂, to raise intracellular CO₂ levels. The two processes are interconnected; yet, the underlying transcriptional regulators remain largely unknown. By employing a large transcriptomics data compendium of *Chlamydomonas reinhardtii*'s responses to different light and carbon supply we reconstructed a consensus genome-scale gene regulatory network from complementary inference approaches and used it to elucidate the transcriptional regulation of photoprotection. We showed that the CCM regulator LCR1 also controls photoprotection, and that QER7, a Squamosa Binding Protein, suppresses photoprotection- and CCM-gene expression under the control of the blue light photoreceptor Phototropin. Along with demonstrating the existence of regulatory hubs that channel light- and CO₂-mediated signals into a common response, our study provides a unique resource to dissect gene expression regulation in this microalga.

Introduction

Photosynthetic microalgae convert light into chemical energy in the form of ATP and NADPH to fuel the CO₂ fixation in the Calvin–Benson cycle¹. They have evolved to cope with rapid fluctuations in light² and inorganic carbon³ availability in their native habitats. When absorbed light exceeds the CO₂ assimilation capacity, the formation of harmful reactive oxygen species can lead to severe cell damage; this is prevented by the activation of photoprotective mechanisms, collectively called non-photochemical quenching (NPQ). NPQ encompasses several processes that are distinguished in terms of their timescales², among which the rapidly reversible energy-quenching (qE) is, under most circumstances, the predominant NPQ component^{2,4}. The major molecular effector of qE in *Chlamydomonas* is the LIGHT HARVESTING COMPLEX STRESS RELATED protein LHCSR3, encoded by the *LHCSR3.1* and *LHCSR3.2* genes⁵ that slightly differ only in their promoters; LHCSR1 can also contribute significantly in qE under conditions where LHCSR3 is not expressed^{6,7}. PSBS, the key qE effector protein in higher plants⁸ is encoded as two highly similar paralogues *PSBS1* and *PSBS2* in *Chlamydomonas*⁹. They are only transiently expressed in *Chlamydomonas* under high light (HL)^{9,10} and accumulate under UV-B irradiation¹¹; their precise contribution in *Chlamydomonas* photoprotective responses is still unresolved¹².

Intracellular levels of CO₂ are modulated by the availability of its gaseous and hydrated forms³ in the culture media and the supply of acetate, that is partly metabolized into CO₂^{7,13}. Under low CO₂, *Chlamydomonas* activates the CO₂-concentrating mechanism (CCM) to avoid substrate-limitation of photosynthesis by raising the CO₂ concentration at the site of RuBisCO, where CO₂ is assimilated³. The CCM mainly comprises of carbonic anhydrases (CAHs) and of inorganic carbon transporters. Almost all CCM-related genes are under the control of the nucleus-localized zinc-finger type nuclear factor CIA5 (aka CCM1)¹⁴⁻¹⁶, including the Myb Transcription Factor LOW-CO₂ -STRESS RESPONSE 1 (LCR1) that controls expression of genes coding for the periplasmic CAH1, the plasma membrane-localized bicarbonate transporter LOW CO₂-INDUCED 1 (LCI1), and the low-CO₂ responsive LCI6, whose role remains to be elucidated¹⁷. CIA5 is also a major qE regulator activating transcription of genes encoding LHCSR3 and PSBS, while repressing accumulation of LHCSR1 protein⁷.

LHCSR3 expression relies on blue light perception by the photoreceptor phototropin (PHOT)¹⁸, on calcium signaling, mediated by the calcium sensor CAS¹⁹ and on active photosynthetic electron flow¹⁸⁻²⁰, likely via indirectly impacting CO₂ availability⁷. The critical importance of CO₂ in LHCSR3 expression is demonstrated by the fact that changes in CO₂ concentration can trigger LHCSR3 expression²¹⁻²³ even in the absence of light⁷. Accumulation of *LHCSR1* and *PSBS* mRNA is under control of the UV-B photoreceptor UVR8¹¹ and PHOT^{24,25}

and is photosynthesis-independent^{20,25}. While LHCSR1 is CO₂/CIA5 independent at the transcript level^{7,25}, PSBS is responsive to CO₂ abundance and is under partial control of CIA5⁷. A Cullin (CUL4) dependent E3-ligase^{24,26,27} has been demonstrated to post-translationally regulate the transcription factor (TF) complex of CONSTANS (CrCO)²⁶ and NF-Y isomers²⁷, which bind to DNA to regulate the transcription of *LHCSR1*, *LHCSR3*, and *PSBS*. The putative TF and diurnal timekeeper RHYTHM OF CHLOROPLAST 75 (ROC75) was shown to repress LHCSR3 under illumination with red light²⁸.

Here, we employed a large compendium of RNAseq data from *Chlamydomonas* to build a gene regulatory network (GRN) underlying light and carbon responses, and thus reveal the transcriptional regulation of qE at the interface of these responses. The successful usage of RNAseq data to infer GRNs has been demonstrated in many studies²⁹⁻³¹, although the data pose some challenges that require careful consideration. All of the developed approaches to infer a GRN quantify the interdependence between the transcript levels of TF-coding genes and their putative targets; the resulting prediction model serves as a proxy for the regulatory strength that the product of the TF-coding gene exerts on its target(s). It is usually the case that the number of observations (samples) is considerably smaller than the number of TFs used as predictors, leading to collinearity of the transcript levels and associated computational instabilities; further, as an artifact of the computational techniques, some of the inferred regulations may be spurious³²⁻³⁴. To address these issues, here we took advantage of combining the outcome of multiple regularization techniques and post-processing to increase the robustness of identified interactions³³⁻³⁵. In contrast to our approach, the existing predicted GRNs of *Chlamydomonas* either focused on nitrogen starvation³⁶ or used a broad RNAseq data compendium, not tailored to inferring regulatory interactions underlying responses to particular cues³⁷. Moreover, these GRNs were not obtained by combining the outcomes from multiple inference approaches, shown to increase accuracy of predictions³⁰, and their quality was not gauged against existing knowledge of gene regulatory interactions.

We used an RNAseq data compendium of 158 samples (**Supplementary Table 1**) from *Chlamydomonas* cultures exposed to different light and carbon supply as input to seven benchmarked GRN inference approaches that employ complementary inference strategies^{29,30}. We assessed the performance of each approach based on a set of curated TF-target gene interactions with experimental evidence from *Chlamydomonas*. Based on this assessment, we integrated the outcome of the five best performing approaches into a unique resource, a consensus network of *Chlamydomonas* light- and carbon-dependent transcriptional regulation. We used the consensus network to reveal regulators of qE genes and demonstrated the quality of predictions by validating two of the six tested candidates. We showed that LCR1

acts as a qE regulator by activating the expression of LHCSR3, and demonstrated that qE-REGULATOR 7 (QER7) is a repressor of qE and CCM-gene expression. Our work consolidates the extensive co-regulation of CCM and photoprotection⁷ based on the untargeted assessment of the obtained genome-scale GRN.

Results

Computationally inferred GRN recovered known regulatory interactions underlying qE and CCM in *Chlamydomonas*

We first aimed to employ published and here generated RNAseq data sets capturing the transcriptional responses of *Chlamydomonas* to light and acetate availability to infer the underlying GRN. To this end, we obtained data from two publicly available transcriptomics studies of synchronized chemostat wild-type (WT) cultures grown in a 12h/12h light dark scheme and sampled in 30 min to 2h intervals^{38,39}. We combined these with our RNAseq data generated from mixotrophically or autotrophically grown batch cultures of the WT and *phot* mutant acclimated to low light (LL) or exposed to HL (**Methods, Supplementary Table 1**). These data sets capture the expected expression patterns of the key genes involved on CCM and qE (**Fig. 1a**) in response to changes in acetate availability and light intensity. Specifically, we found strong up-regulation of these genes in the light^{7,25}, and a marked inhibition of *LHCSR3.1/2* and CCM genes by acetate as previously described^{7,40}.

We employed these data together with a list of 407 transcription factors from protein homology studies^{41,42} (**Methods, Supplementary Table 2**), as input to seven GRN inference approaches to robustly predict TF-target interactions, as shown in benchmark studies³⁰. Since there exists no study that experimentally probes TF binding to DNA on a genome-scale, we first curated a list of known, experimentally validated regulatory interactions underlying CCM and qE in *Chlamydomonas*^{22,26-28}, to assess the quality of GRNs inferred by the different approaches (**Fig. 1b**). As a negative control, we considered the lack of effect of the SINGLET OXYGEN RESISTANT 1 (SOR1) TF on *PSBS1* transcript levels in diurnal culture⁴³. The comparative analysis of the predicted and known interactions demonstrated that two of the applied approaches for GRN inference (i.e. ARACNE⁴⁴ and global silencing⁴⁵) are unable to recover any literature interactions when using a network density threshold of 10% of all possible TF-TF and TF-target interactions. This poor performance is due either to over-trimming or issues with the validity of the underlying assumptions, as seen in other case studies⁴⁶. We therefore considered only the remaining five approaches, namely: Graphical Gaussian Models (GGM), Context Likelihood of Relatedness (CLR), Elastic Net regression, Gene Network Inference with Ensemble of Trees (GENIE3), and Network Deconvolution to infer a consensus GRN by using the Borda count election method^{30,47} (**Methods**). We then ranked the inferred interactions within each approach and quantified the

variability of ranks for the known TF-target gene interactions. We found that the average standard deviation of the ranks of the TF-target gene interactions within an approach is larger than the average standard deviation for the rank of a TF-target gene interaction across the five approaches (**Extended Data Fig. 1, Fig. 1b**). This observation suggested that the properties of a given TF-target gene interaction have a stronger influence on its assigned rank than the inference approach used. More specifically, we noted that the regulation of *LHCSR* genes by the two NF-Y paralogues and the induction of *LCR1* by *CIA5* are not recovered by any of the used approaches; this is in line with reports showing that *CIA5* is constitutively expressed and regulated post-translationally¹⁶-not reflected in the transcriptomics data. Further, NF-Y factors that rely on complex formation with CrCO to regulate their targets²⁷, may also act via unresolved posttranslational mechanisms. Importantly, the regulatory interactions of the CCM effector genes *LCI1* and *CAH1* by *LCR1* are assigned very high ranks (top 1%) by the employed approaches (except for GGM) considered in the consensus GRN; moreover, the experimentally falsified interaction of *SOR1* and *PSBS1* transcripts⁴³ is correctly discarded by all approaches (**Fig. 1b**). In addition, we observed that CLR and GENIE3 demonstrated the best performance with respect to the set of known interactions. For instance, they identified the regulation of *LHCSR3.1* by CrCO^{26,27} and of *PSBS1* by NF-YB²⁷ (**Fig. 1b**). Generalization of this ranking beyond the known interactions underlying qE and CCM processes is challenging, due to the lack of genome-scale gold standard, and we therefore opted to combine the results of the five approaches, that showed comparable performance, in the consensus GRN (**Methods, Supplementary Table 3**) to increase robustness of the predictions. Our analyses of the overlap between the consensus and individual GRNs and the enrichment of TF-TF interactions demonstrated the robustness of the inferred interactions (**Supplementary Note 1, Extended Data Fig. 2**).

Consensus GRN pinpoints *LCR1* as a regulator of qE-related genes

Using the consensus GRN, we inferred direct regulators of *LHCSR* and *PSBS* genes) and ranked them according to the score resulting from the Borda method (**Methods**)^{30,47}. Mutants were available for four of the top ten of TFs with strongest cumulative regulatory effect on qE-related genes (**Fig. 2a, Supplementary Table 4**): Two knock-out mutants of previously uncharacterized genes were ordered from the CliP library⁴⁸, which we termed *qE-regulators 4* and *6* (*qer4*, *qer6*; see **Extended Data Fig. 3** for the genotyping of these mutants). Additionally, we obtained an over-expressor line of the N-acetyltransferase *LCI8*⁴⁹ and the knock-out strain of the known CCM regulator *LCR1*¹⁷. We tested for a regulatory effect by switching LL-acclimated mutant strains and their respective WT background to HL for 1h and quantified transcript levels of qE-related genes. For *qer4*, *qer6*, and *lci8-oe* we could not observe an effect on the transcript levels of investigated genes after HL exposure (**Extended Data Fig. 4**). Thus, *qer4* and *qer6* are considered false positive predictions of the GRN,

despite the fact that *qer4* accumulated 1.5 times more *LHCSR3.1* under LL than the WT. A review of the closest orthologs of LCI8 together with the experimental data indicate that it is likely involved in arginine synthesis⁴⁹ and wrongly included as histone acetylase in the list of TFs. Interestingly, LCR1, the highest ranking among the tested regulators showed significantly decreased expression of LHCSR3 at both the gene (3 times lower, **Fig. 2b**) and protein level (4 times lower, **Fig. 2c,d**) compared to the WT; as a result, *lcr1* developed very low NPQ and qE (**Fig. 2e**). Complementation of *lcr1* with the knocked-out gene (strain *lcr1-C*) restored LHCSR3 gene and protein expression as well as the qE phenotype (**Fig. 2b-e**). Interestingly, the *lcr1* mutant over-accumulated LHCSR1 and PSBS both at the transcript and at the protein level (**Fig. 2b-d**); Complementation of *lcr1* with the knocked-out gene (strain *lcr1-C*) restored LHCSR3 gene and protein expression as well as the qE phenotype (**Fig. 2b-e**). Because pre-acclimation conditions impact qE gene expression²⁵ we conducted independent experiments in which cells were acclimated to darkness before exposure to HL and we obtained very similar results. Our data demonstrated that *lcr1* showed significantly lower expression of LHCSR3 and higher expression of LHCSR1/PSBS at both the gene (**Extended Data Fig. 5a**) and protein level (**Extended Data Fig. 5b,c**), and had lower NPQ and qE (**Extended Data Fig. 5d**) than the WT, although the higher expression levels of *LHCSR1* gene were not rescued by the complementation with the missing *LCR1* gene (**Extended Data Fig. 5a**). Altogether, our data show that LCR1 is a regulator of qE by activating *LHCSR3.1* transcription and repressing LHCSR1 and PSBS accumulation.

Further, we revisited the role of LCR1 in regulating CCM genes¹⁷ by analyzing expression of selected CCM genes in WT, *lcr1* and *lcr1-C* cells shifted from LL or darkness to HL, conditions favoring CCM gene expression⁷. We first confirmed that under our experimental conditions *lcr1* could not fully induce *LCI1* (**Extended Data Fig. 6a, b**) in accordance to the report of the discovery of LCR1¹⁷. Our analyses further showed a statistically significant impairment of *lcr1* in inducing genes encoding the Ci transporters LOW-CO₂-INDUCIBLE PROTEIN A (LCIA), HIGH-LIGHT ACTIVATED 3 (HLA3), CHLOROPLAST CARRIER PROTEIN 1 (CCP1) and BESTROPHINE-LIKE PROTEIN 1 (BST1) as well as the carbonic anhydrase CAH4, when shifted from LL or dark to HL (**Extended Data Fig. 6a, b**), indicating that the role of LCR1 in low-CO₂ gene expression extends beyond the regulation of gene expression of *CAH1*, *LCI1* and *LCI6*¹⁷.

PHOT-specific GRN reveals a novel repressor of qE and CCM

The light-dependent induction of LHCSR3 is predominantly mediated by the blue light photoreceptor PHOT¹⁸. To analyze the PHOT-dependent transcriptional regulators, we first identified the interactions shared by the consensus network and a GRN inferred only by GENIE3, due to its good performance, using the RNAseq data from samples of *phot* and WT acclimated to LL and after 1h exposure to HL (**Methods**). This so-called PHOT-

specific GRN (**Methods, Supplementary Table 5**) comprises regulatory interactions underlying the transcriptomic changes observed in the *phot* mutant while burrowing the statistical power of the RNAseq compendium.

When investigating the top 10 regulators of qE genes in this PHOT GRN the interactions were weighted based on the importance score from GENIE3 (**Supplementary Table 6, Fig. 3a**). Among these interactions we recovered the previously discussed false positive LCI8 gene (see above, **Fig. 3a**) (**Extended Data Fig. 4**). In addition, two previously reported regulators of qE, ROC75²⁸ and CrCO^{26,27}, are also included in this list of PHOT-dependent regulators. These observations are in line with an existing hypothesis²⁴ suggesting that a CUL4-dependent E3-ligase targeting CrCO²⁶ acts downstream of PHOT. ROC75 has been previously reported to act independently of the PHOT signal based on qPCR studies of the mutant grown synchronously under different light spectra²⁸. In our RNAseq data, gathered under continuous white light, we observed a significant difference in expression levels of ROC75 between WT and *phot* (log2 fold-change = 1.03, adj. p-value = 1.80×10^{-7}).

The fact that several regulators showed larger regulatory strength than CrCO in the PHOT GRN indicates the existence of yet unreported regulators of qE effector genes in the PHOT signaling pathway. This is in line with existing results²⁶, showing that the knock-out of CrCO is insufficient to fully abolish light-dependent activation of LHCSR3. Following this reasoning we obtained *qer1* and *qer7*, the available regulator candidates mutants, from the CLiP library⁴⁸, (for genotyping see **Extended Data Fig. 3**). Our results show higher mRNA and protein levels of LHCSR1 in the *qer1* mutant (**Extended Data Fig. 7**); however, this could not be rescued by ectopic expression of the *QER1* gene in the *qer1* mutant background (**Extended Data Fig. 7**). We found significant upregulation of *LHCSR3.1* gene expression in the *qer7* mutant (1.7 times, **Extended Data Fig. 8a**) also reflected in higher NPQ (**Extended Data Fig. 8b**) and qE levels (**Extended Data Fig. 8c**) which we followed up in more detail. To this end, we ectopically expressed the WT *QER7* gene in the *qer7* mutant and generated the complemented strain *qer7-C* that expressed *QER7* to levels similar to those WT (**Extended Data Fig. 3c**). As a result, the *qer7-C* strain showed reduced *LHCSR3* gene expression, NPQ and qE levels as compared with the *qer7* mutant (**Extended Data Fig. 8a-c**). *LHCSR1* and *PSBS* seemed to be unaffected in the *qer7* in these LL to HL transition experiments (**Extended Data Fig. 8a**). As with LCR1, we also performed dark to HL experiments to further characterize the photoprotective responses of *qer7*; under these conditions, *qer7* accumulated significantly more *LHCSR1* (1.7 times) and *PSBS1* (2.2 times) while *LHCSR3* remained unaffected (**Extended Data Fig. 8d**). As in the LL to HL experiments (**Extended Data Fig. 8b, c**) *qer7* showed more NPQ and qE (**Extended Data Fig. 8e, f**). Complementation of *qer7* with the missing *QER7* gene restored all phenotypes

(*LHCSR1*, *PSBS*, NPQ, qE; **Extended Data Fig. 8d-f**). These data validate the prediction of QER7 as regulator of qE gene expression (**Fig. 3a**) and indicate that QER7 regulates different subsets of qE genes depending on the pre-acclimation conditions; *LHCSR3* when preacclimation is under LL, *LHCSR1* and *PSBS* when pre-acclimation occurs in darkness. Motivated by these findings and given the fact that most of the *Chlamydomonas* transcriptome, undergoes diurnal changes according to biological function³⁹ we decided to address the role of QER7 in regulating qE genes under light/dark cycles. We synchronized WT, *qer7* and *qer7-C* cells in 12h L/12h D cycle and exposed them to HL right after the end of the dark phase. Our results revealed that under these conditions QER7 functions as a repressor of all qE-related genes; the *qer7* mutant expresses significantly higher *LHCSR3*, *LHCSR1* and *PSBS* both at the gene (**Fig. 3b**) and protein (**Fig. 3c,d**) level, and exhibits higher NPQ and qE (**Fig. 3e**), with all phenotypes rescued in the *qer7-C* complemented line. Previous protein homology studies identified QER7 as Squamosa Binding Protein⁵⁰ or bZIP TF⁵¹, and here, we provide the first functional annotation of QER7 as a novel qE regulator.

QER7 co-regulates qE-related and CCM genes

The recent findings that the regulatory role of CIA5⁷ and LCR1 (**Fig. 2 and Extended Data Fig. 5**) extends beyond CCM to also control qE-related gene expression, prompted us to also inspect the expression levels of CCM genes in synchronized *qer7* cells (**Fig. 4a**). Indeed, for four of these transcripts (*CAH4*, *LCIA*, *CCP1*, *HLA3*) we observed a significant upregulation in *qer7* after HL exposure that was reversed by complementation with the *QER7* gene, indicating that QER7 suppresses expression of CCM genes; the suppression role of QER7 on CCM genes was only observable under HL, conditions that favor CCM gene expression⁷ and not under LL (**Fig. 4a**). Interestingly, and in line with QER7 acting as suppressor of CCM gene expression (**Fig. 4a**), we found LCR1 and QER7 among the top 10 regulators of the CCM genes probed by qPCR in the inferred GRNs (**Extended Data Fig. 9**).

After observing the inhibition of CCM gene transcription by QER7, we investigated the signaling pathway upstream of QER7. To this end, we quantified *QER7* gene expression in synchronized *phot* cultures. As expected, we observe a strong effect of the *PHOT* knock-out showing that PHOT suppresses *QER7* expression (**Fig. 4b**) implying that blue-light perception via PHOT is involved in the regulation of CCM related gene expression. Interestingly, *phot* expressed *LCR1* to levels similar to those in WT (**Fig. 4c**). Thus, while sharing part of their target genes, the two TFs mediate different signals. These data demonstrate that the PHOT-specific GRN successfully pinpoints TFs that depend on PHOT activity while retaining the statistical power to recover genuine regulatory interactions.

The two qE regulators that we validated in this study also regulate CCM genes. Therefore, we next investigated to what extent the observed coregulation pattern applies to the global, known transcriptional regulation of low CO₂ and light stress responsive genes. To this end, we took advantage of the size of the presented genome-scale GRNs and compiled a list of genes putatively involved in photoprotection (**Supplementary Table 7**) or the CCM (**Supplementary Table 8**); we then extracted the 10 TFs exhibiting the strongest regulatory strength on the genes in the compiled lists. We found six (empirical p-value<0.001, **Methods**) and four (empirical p-value < 0.01) of the top 10 regulators to be shared between these two responses in the consensus (**Fig. 5a,b**) and the PHOT-specific GRN, respectively (**Figure 5c,d**). The significant, large number of shared regulators is a strong indication that co-regulation of photoprotective and carbon assimilatory processes is a principal feature of *Chlamydomonas*' transcriptional regulatory program.

Discussion

The molecular actors and structure of the transcriptional regulatory mechanisms that shape *Chlamydomonas*' response to differential light and carbon availability are largely unknown, although they are paramount to survival of *Chlamydomonas* and offer valuable targets for biological engineering. Here we set out to elucidate the GRN underlying the response to light and carbon availability by combining the results from five complementary inference approaches and data from 158 RNAseq samples of cultures responding to these cues. The obtained GRN enabled us to identify novel regulators controlling qE. Experimentally testing available mutant strains for six of these candidates we were able to validate two TFs: QER7, suppressing qE- and CCM-gene expression, and LCR1, activating LHCSR3 and suppressing LHCSR1 and PSBS expression. The two TFs belong to independent signalling pathways: LCR1 has been known to controlled by CIA5 while we found that QER7 is controlled by Phototropin; this last finding adds a new layer of regulation of CCM by blue light sensing. We complemented the finding of the involvement of LCR1 in the regulation of qE related genes with an analysis of the genome-scale co-regulation of CCM and photoprotective genes. Indeed, we observed significant number (six out of the top 10) of regulators targeting both processes in the inferred consensus GRN. Together with previous studies, our results provide evidence for extensive signaling crosstalk between light and carbon sensing and name a set of TFs that likely integrate these disparate signals into a common transcriptional response. The GRN gives an unbiased representation of the genome-scale regulators acting in light and carbon sensing. Thus, it represents a powerful resource for future dissection of the transcriptional regulation of responses of *Chlamydomonas* to light and carbon availability.

Methods

Transcriptome analysis

We assembled a compendium of RNAseq data (**Supplementary Table 1**) that capture regulation of light-dependent processes by combining in-house produced RNAseq measurements with publicly available data from two studies of densely sampled diurnal cultures of *Chlamydomonas*^{38,39}. For the samples in the acetate time-resolved experiment, adapter sequences were specifically trimmed from raw reads using BBduk⁵² (ktrim=r k=30 mink=12 minlen=50). Raw reads of the diurnal transcriptome study from Strenkert et al.³⁹ were obtained from NCBI GEO database (GSE112394). Reads were aligned to the *Chlamydomonas* reference transcriptome⁵³ available from JGI Phytozome (Assembly version 5) using RNA STAR aligner. The BAM files obtained from these measurements were analyzed using HTSeq-count⁵⁴ (standed=reverse) to create raw read count files. The raw read counts from Zones et al.³⁸ were obtained as .tsv from NCBI GEO (GSE71469). The final data set consists of 158 samples from 62 experimental conditions or time points (**Supplementary Table 1**). Genes with less than 1 count per million in at least 9 measurements were discarded and the remainder were voom⁵³ transformed and normalized using library normalization factors based on the TMM⁵⁵ approach as implemented in the R Bioconductor package edgeR⁵⁶.

Transcription factor set from comparative genomics

To reduce the set of parameters in our network model, we compiled transcription factor (TF) annotations for the *Chlamydomonas* genome based on proteome homology studies. We obtained the proteomes and protein IDs of predicted *Chlamydomonas* TFs from Pérez-Rodríguez et al.⁴¹ Since these predictions were built based on the older *Chlamydomonas* assembly, we first used the conversion table provided by Phytozome to convert JGI4 to Crev5.6 IDs. For the TFs that could not be recovered by this approach we used the Phytozome BLAST tool to align these sequences against the Crev5.6 proteome (BLASTP, E threshold: -1, comparison matrix: BLOSUM62, word length: 11, number of reported alignments: 5). The reported hits were filtered for sequence identity > 97% and gaps ≤ 1. If sequences mapped multiple times to the same Crev5.6 gene ID, only the hit alignment closest to the N-terminus of the query sequence was kept. The hit was only accepted if the alignment started at least six residues from the N-terminus of the hit sequence. For Crev5.6 loci that had multiple JGI4 TF queries assigned to them the best hit was selected manually. This set was then extended by the TFs found in the study of Jin et al.⁴² and the regulators in the manually curated set of CCM and qE regulatory interactions (**Supplementary Tables 7 and 8**). Using this procedure, we compiled a list of 407 *Chlamydomonas* TFs (**Supplementary Table 2**) to be considered as regulators in the inferred networks.

312 **Gene regulatory network inference**

313 The CLR and ARACNE approach were based on all replicate measurements; for all other inference methods
314 the median from each condition was used as input. All input matrices were standardized gene-wise. If not
315 explicitly stated in the respective paragraph the implementations of all GRN inference approaches were
316 applied with their default settings.

317 **Graphical Gaussian Models**

318 The network inferred from a Graphical Gaussian model of gene regulation was obtained using the
319 implementation of the partial correlation estimate from Schäfer et al.³⁴ as implemented in the R GeneNet
320 package. All interactions between TFs and another gene/TF with non-zero partial correlations were included
321 as network edges.

322 **GENIE3**

323 The random forest-based network from GENIE3 was generated using the R Bioconductor implementation
324 provided by the authors⁵⁷. We used only expression levels of TFs as predictors.

325 **Elastic net regression**

326 A linear regression based network was obtained using the elastic net algorithm³⁵. A model was fit for each
327 gene using the expression levels of all TFs as predictors. The two hyperparameters λ_2 (quadratic penalty) and
328 s (fraction of L1 norm coefficients) were tuned for each gene model using 6-fold cross validation. The 2D
329 parameter space scanned was $\lambda_2=\{0,0.001,0.01,0.05,0.1,0.5,1,1.5,2,10,100\}$ and
330 $s=\{0.1,0.2,0.3,0.4,0.5,0.6,0.7,0.8,0.9\}$. The R2 value for each model was calculated as

$$R^2 = 1 - \frac{\sum(y - \hat{y})^2}{\text{var}(y)} \quad (1)$$

331 with y marking the vector of observed expression values and $\hat{y}$ the model predictions. Models with a negative
332 R2 value were discarded as regularization artifacts. The results of the remaining models were assembled into
333 a network in which interactions were ranked by regression coefficients β normalized by the maximum
334 absolute coefficient.

$$\beta_n = \frac{\beta}{\max|\beta|} \quad (2)$$

336 **CLR and ARACNE**

The implementation of mutual information (MI)-based network inference approaches from the R package *minet*⁵⁸ was used. Pairwise MI was estimated based on the Spearman correlation as proposed by Olsen et al.⁵⁹. Two networks were constructed based on these MI estimates. Using the CLR approach⁶⁰ non-significant interactions were removed based on the z-scores calculated from the marginal distributions of MI values for each gene pair. Alternatively, the ARACNE algorithm⁴⁴ was used to prune the network based on the data processing inequality. For both networks only interactions originating from a TF were taken into consideration and edges were ranked according to the assigned MI value.

Deconvolution and Silencing

For the two networks based on decomposition of the interaction matrix G the Pearson correlation matrix obtained from gene expression values was used as input.

The deconvolution approach introduced by Feizi et al.⁶¹ was implemented as previously described⁴⁶. The eigenvalue scaling factor β was initialized as $\beta=0.9$ and iteratively reduced in increments of 0.05 until the largest eigenvector of the direct interaction matrix generated by deconvolution was smaller than 1. Edges were ranked according to the deconvoluted interaction matrix.

The Silencing approach as described by Barzel et al.⁴⁵ was implemented in R. The proposed approximation of the direct interaction matrix S in which spurious interactions are silenced relies on the inverses of the observed correlation matrix G . In our implementation we used the Moore-Penrose pseudoinverse in case G was close to singular. In the resulting network edges were ranked according to the approximated silenced interaction matrix.

Consensus network construction

To improve network quality³⁰, we built a consensus network integrating the GRN models inferred by the different approaches introduced above. To this end, we used the Borda count election method⁴⁷ whereby the rank r of an interaction I in the consensus network built on the predictions from k approaches is given by arithmetic mean of the ranks in the individual networks

$$r_{consens}(I) = \frac{\sum_{i=1}^k r_i(I)}{k}. \quad (3)$$

Following the reasoning of Feizi et al.⁶¹) in this integration only the top 10% all possible edges in the GRN (625815) were considered from each individual ranking. For an edge that was not assigned a rank by some approaches, the missing ranks were set to 10% of all possible edges plus one.

Using this integration method, we assembled a consensus network based on all approaches to compare predictions from all GRN inference approaches (**Extended Data Fig. 1**). Due to this comparison and inability to recover known interactions (**Fig. 1b**) the rankings derived from ARACNE and Silencing were only considered in **Extended Data Fig. 1** and excluded from the final consensus network used for all other analyses. As with the individual networks returned by the different approaches the consensus network (**Supplemental Table 3**) was trimmed to the top 10% of all possible edges according to the integrated ranks. The weight of edges in the final network was set as $r_{consens}^{-1}$.

PHOT-specific network

To investigate the PHOT-specific regulatory interactions genes that are differentially expressed between phot mutant and wt under low and high light were inferred. To this end, transcript counts of genes with more than 1 count per million in at least four replicates from these conditions (**Supplementary Table 1**) were tested for differential expression using the R packages limma⁶², DeSeq2⁶³, and edgeR⁵⁶. Only genes deemed significant by all three tools after Benjamini-Hochberg correction for a false discovery rate of 0.05 were considered differentially expressed with respect to PHOT mutation

In the next step, we focused on the normalized and scaled expression levels from these differentially expressed genes and the previously mentioned conditions, to infer a PHOT-specific GRN using GENIE3. To improve robustness of this network, which was obtained from a comparably sparse data set, we only considered the edges in the intersection with the final consensus network. Again, for both networks only the top 10% of possible edges were taken into account. Therefore, the obtained PHOT-network represents a subnetwork of the final consensus in which edges are weighted by the “PHOT specific” GENIE3 importance measure (**Supplementary Table 5**).

Identification of major regulators

We compiled a manually curated list of possible target genes known to be involved in the processes of qE (*LHCSR1*, *LHCSR3.1/2*, *PSBS1/2*), photoprotection (**Supplementary Table 7**), and CCM (**Supplementary Table 8**). Based on the assumption that major regulators act on several genes important for a biological process, the regulatory strength of a candidate regulator (for the given process) was determined by the sum of edge weights w_{ij} between this regulator and the k genes in the respective target gene set

$$C(TF) = \sum_{j=1}^k w_{TFj}. \quad (4)$$

Empirical p-value calculation using Monte-Carlo simulation

The one-sided p-value for the overlap between the regulators of CCM and photoprotective genes was approximated by sampling the overlaps of random gene sets. To this end, we compiled two gene sets with the same cardinality as the curated CCM and photoprotective genes. The genes in these sets were randomly sampled without replacement from all targets in the respective networks. The 10 strongest regulators of these two gene sets were then obtained as previously described and the overlap was calculated as our sample statistic. This process was repeated 10,000 times and an empirical p-value was calculated from the number of iterations, r , where the overlap was higher or equal to the observed value, and the total number of iterations, n ⁶⁴:

$$p = \frac{r+1}{n+1}.$$

(5)

Strains and conditions

C. reinhardtii strains were grown under 15 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$ in Tris-acetate-phosphate (TAP) media⁶⁵ at 23 °C in Erlenmeyer flasks shaken at 125 rpm. For all experiments cells were transferred to Sueoka's high salt medium (HSM)⁶⁶ at 1 million cells mL^{-1} and exposed to light intensities as described in the text and figure legends. For the investigation of the impact of acetate on the genome-wide transcriptome, HSM was supplemented with 20 mM sodium acetate. *C. reinhardtii* strain CC-125 mt+ was used as WT. The *phot* (depleted from *PHOT1*; gene ID: Cre03.g199000), was previously generated⁶⁷ and recently characterized²⁵. For synchronized cultures, the cells were grown in HSM for at least 5 days under a 12h light/12h dark cycle (light intensity was set at 15 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$; temperature was 18 °C in the dark and 23 °C in the light). All CLiP mutant strains used in this study and their parental strain (CC-4533) were obtained from the CLiP library (REF); *qer1* (LMJ.RY0402.072278), *qer4* (LMJ.RY0402.202963), *qer6* (LMJ.RY0402.162350), *qer7* (LMJ.RY0402.118995). The *lcr1* (strain C44), *lcr1-C* (strain C44-B7) and its parental strain Q30P3 as described in¹⁷ were a kind gift from Hideya Fukuzawa. Before performing phenotyping experiments, we first confirmed that *lcr1* shows no expression of *LCR1* and that this is rescued in the *lcr1-C* strain (**Extended Data Fig. 10a**). The *lci8* overexpressing line was purchased from the Chlamydomonas Resource center; strain CSI_FC1G01, expressing pLM005-Cre02.g144800-Venus-3xFLAG in the CC-4533 background. Overexpression of LCI8-FLAG was verified by immunoblotting against FLAG (**Extended Data Fig. 10b**).

To complement *qer1*, a 1152 bp genomic DNA fragment from *Chlamydomonas* CC-4533 was amplified by PCR using KOD hot start DNA polymerase (Merck) and primers P11 and P12 (**Supplementary Table 9**). To complement *qer7*, a 5755 bp fragment DNA fragment from *Chlamydomonas* CC-4533 was amplified by PCR

with Platinum superfi DNA Polymerase (Thermo Fisher Scientific) and primers P13 and P14 (**Supplementary Table 9**). The PCR products were gel purified and cloned into pRAM118⁶⁸ by Gibson assembly⁶⁹ for expression under control of the *PSAD* promoter. Junctions and insertion were sequenced and constructs were linearized by EcoRV before transformation. Eleven ng/kb of linearized plasmid⁷⁰ mixed with 400 μ L of 1.0×10^7 cells mL⁻¹ were electroporated in a volume of 120 μ L in a 2-mm-gap electro cuvette using a NEPA21 square-pulse electroporator (NEPAGENE, Japan). The electroporation parameters were set as follows: Poring Pulse (300V; 8 ms length; 50 ms interval; one pulse; 40% decay rate; + Polarity), Transfer Pulse (20V; 50 ms length; 50 ms interval; five pulses; 40% decay rate; +/- Polarity). Transformants were selected onto solid agar plates containing 20 μ g/ml hygromycin and screened for fluorescence by using a Tecan fluorescence microplate reader (Tecan Group Ltd., Switzerland). Parameters used were as follows: YFP (excitation 515/12 nm and emission 550/12 nm) and chlorophyll (excitation 440/9 nm and 680/20 nm). Transformants showing high YFP/chlorophyll value were further analyzed by real time qPCR.

Unless otherwise stated, LL conditions corresponded to 15 μ mol photons m⁻² s⁻¹ while HL conditions corresponded to 300 μ mol photons m⁻² s⁻¹ of white light (Neptune L.E.D., France; see⁷ for light spectrum). All experiments were repeated at least three times to verify their reproducibility.

DNA Isolation and genotyping of CLiP mutants.

Total genomic DNA from CLiP mutants and corresponding wild-type strain CC-4533 was extracted according to the protocol suggested by CLiP website (<https://www.chlamylibrary.org/>). One μ L of the extracted DNA was used as a template for the PCR assays, using Phire Plant Direct PCR polymerase (Thermo Fisher Scientific). To confirm the CIB1 insertion site in the CLiP mutants, gene-specific primers were used that anneal upstream and downstream of the predicted insertion site of the cassette (primer pairs P3-P4, P7-P8, P9-P10 and P5-P6 for *qer6*, *qer1*, *qer7* and *qer4* respectively; **Supplementary Table 9**). While all these primers worked in DNA extracted from WT, they did not work in the DNA extracted from the mutants, with the exception of *qer4* (**Extended Data Fig. 3**), therefore primers specific for the 5' and 3' ends of the CIB1 Cassette were additionally used. All the primers used for genotyping were shown in **Supplementary Table 9**. We further confirmed the disruption of the genes of interest by quantifying their mRNA accumulation (**Extended Data Fig. 3**).

mRNA quantification

Total RNA was extracted using the RNeasy Mini Kit (Qiagen) and treated with the RNase-Free DNase Set (Qiagen). 1 μ g total RNA was reverse transcribed with oligo dT using Sensifast cDNA Synthesis kit (Meridian

Bioscience, USA). qPCR reactions were performed and quantitated in a Bio-Rad CFX96 system using SsoAdvanced Universal SYBR Green Supermix (BioRad). The primers (0.3 μ M) used for qPCR are listed in **Supplementary Table 10**. A gene encoding G protein subunit-like protein (GBLP)⁷¹ was used as the endogenous control, and relative expression values relative to *GBLP* were calculated from three biological replicates, each of which contained three technical replicates. All primers used for qPCR (**Supplementary Table 10**) were confirmed as having at least 90% amplification efficiency. In order to conform mRNA accumulation data to the distributional assumptions of ANOVA, i.e. the residuals should be normally distributed and variances should be equal among groups, Two-Way Analysis of Variance were computed with log-transformed data $Y=\log X$ where X is mRNA accumulation⁷².

Immunoblotting

Protein samples of whole cell extracts (0.5 μ g chlorophyll or 10 μ g protein) were loaded on 4-20% SDS-PAGE gels (Mini-PROTEAN TGX Precast Protein Gels, Bio-Rad) and blotted onto nitrocellulose membranes. Antisera against LHCSR1 (AS14 2819), LHCSR3 (AS14 2766), ATPB (AS05 085), CAH4/5 (AS11 1737) were from Agrisera (Vännäs, Sweden); antiserum against PSBS was from ShineGene Molecular Biotech (Shanghai, China) targeting the peptides described in Ref.⁹. ATPB was used as a loading control. An anti-rabbit horseradish peroxidase-conjugated antiserum was used for detection. The blots were developed with ECL detection reagent, and images of the blots were obtained using a CCD imager (ChemiDoc MP System, Bio-Rad). For the densitometric quantification, data were normalized with ATPB.

Fluorescence-based measurements

Fluorescence-based photosynthetic parameters were measured with a pulse modulated amplitude fluorimeter (MAXI-IMAGING-PAM, HeinzWaltz GmbH, Germany). Prior to the onset of the measurements, cells were acclimated to darkness for 15 min. Chlorophyll fluorescence was recorded during 10 min under 570 μ mol $m^{-2} s^{-1}$ of actinic blue light followed by finishing with 10 min of measurements of fluorescence relaxation in the dark. A saturating pulse (200 msec) of blue light (6000 μ mol photons $m^{-2} sec^{-1}$) was applied for determination of F_m (the maximal fluorescence yield in dark-adapted state) or F_m' (maximal fluorescence in any light-adapted state). NPQ was calculated as $(F_m - F_m')/F_m'$ based on⁷³; qE was estimated as the fraction of NPQ that is rapidly inducible in the light and reversible in the dark.

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Figures

Fig. 1: Characterization of the consensus GRN inferred by employing a compendium of RNA-seq data from diverse light and culture conditions. **a.** Expression levels of representative CCM and qE related genes are plotted over all samples used for network inference (z-scaled log values are depicted). The column annotation gives information on the culture conditions. The values of the rows of the heatmap correspond to the z-scores of representative CCM and qE genes plotted over all samples (columns) used in the network inference. **b.** The heatmap rows correspond to experimentally validated or falsified (neg) gene regulatory interactions involved in qE and CCM, curated from literature. The heatmap indicates ranking of these interactions by different approaches and the consensus network. Edges are considered highly ranked (depicted in blue) if they are above the 10% network density threshold. Edges ranked below this threshold are depicted in red. Edges that were not included in the given network are marked in grey. ARACNE and Silencing columns were only plotted for comparison and were not used in building the consensus GRN (see Methods).

Fig. 2: Consensus GRN for light and acetate responses pinpoints LCR1 as regulator of qE-related genes. **a.** Dot plot of the relative regulatory strength of the top 10 regulators of qE-related genes in the consensus GRN (see Methods). TFs are marked in green if qE transcript levels were affected in the respective knock-out strain and this effect was reversed by complementation with the missing gene. TFs for which no effect was observed are marked in red. TFs for which no mutant lines were available are plotted in grey. **b.** WT, *lcr1* and *lcr1-C* cells were acclimated for 16h in LL ($15 \mu\text{mol photons m}^{-2} \text{s}^{-1}$). After sampling for the LL conditions, light intensity was increased to $300 \mu\text{mol photons m}^{-2} \text{s}^{-1}$ (HL); samples were taken 1 h (RNA) or 4 h (protein and photosynthetic measurements) after exposure to HL. Shown are relative expression levels of qE-related genes at the indicated conditions normalized to WT LL ($n = 3$ biological samples, mean $\pm$ sd). **c.** Immunoblot analyses of LHCSR1, LHCSR3, PSBS and ATPB (loading control) of one of the three biological replicates, under the indicated conditions. **d.** Quantification of immunoblot data of all replicates in panel **c** after normalization to ATPB. Shown are the HL treated samples; WT protein levels were set as 1. **e.** NPQ and calculated qE (as an inset) 4h after exposure to HL ($n = 3$ biological samples, mean $\pm$ s.d). The p-values for the comparisons are based on ANOVA Dunnett's multiple comparisons test and as indicated in the graphs (*, $P < 0.005$, **, $P < 0.01$, ***, $P < 0.001$, ****, $P < 0.0001$). Statistical analyses for panel **b** and **d** were applied on log10- transformed values.

Fig. 3: A PHOT-specific GRN pinpoints QER7 as a suppressor of the expression of qE-related genes. **a.** Dot plot of the relative regulatory strength of the top 10 regulators of qE-related genes in the PHOT-specific GRN (see **Methods**). TFs are marked in green if qE transcript levels were affected in the respective knock-out strain and this effect was reversed by complementation with the knocked-out gene, in yellow, if the effect was not reversed by complementation, and in red, if no mutant effect was observed in the mutant. TFs for which no mutant lines were available are plotted in grey. **b.** WT, *qer7* and *qer7-C* cells were synchronized under 12h light ($15 \mu\text{mol m}^{-2} \text{s}^{-1}$)/12h dark cycles. After sampling for the dark conditions (end of the dark phase), cells were exposed to $300 \mu\text{mol photons m}^{-2} \text{s}^{-1}$ (HL); samples were taken 1 h (RNA) or 4 h (protein and photosynthetic measurements) after exposure to HL. Shown are relative expression levels of qE-related genes at the indicated conditions normalized to WT LL ($n = 3$

biological samples, mean $\pm$ sd). **c.** Immunoblot analyses of LHCSR1, LHCSR3 and ATPB (loading control) of one of the three biological replicate set of samples, under the indicated conditions. **d.** Quantification of immunoblot data of all replicates in panel **c** after normalization to ATPB. WT protein levels at HL were set as 1. **e.** NPQ and qE, measured 4h after exposure to HL ($n = 3$ biological samples, mean $\pm$ sd). The p-values for the comparisons are based on ANOVA Dunnett's multiple comparisons test and are indicated in the graphs (*, $P < 0.005$, **, $P < 0.01$, ***, $P < 0.001$, ****, $P < 0.0001$). Statistical analyses for panel **b** and **d** were applied on log10-transformed values.

Fig:4: QER7 suppresses transcription of CCM genes and depends on PHOT. WT, *qer7* and *qer7-C* cells were synchronized under 12h light ($15 \mu\text{mol m}^{-2} \text{s}^{-1}$)/12h dark cycles. After sampling at the end of the dark phase, cells were exposed to $300 \mu\text{mol photons m}^{-2} \text{s}^{-1}$ (HL) and samples were taken 1 h after HL exposure. **a.** Relative expression levels of CCM genes at the indicated conditions normalized to WT LL ($n = 3$ biological samples, mean $\pm$ sd). **b and c.** Relative expression levels of *QER7* (**b**) and *LCR1* (**c**) in synchronized WT, *phot* and *phot-C* (*phot* cells complemented with the *PHOT* gene) cells. The p-values for the comparisons are based on ANOVA Dunnett's multiple comparisons test on log10- transformed values and are indicated in the graphs (*, $P < 0.005$, **, $P < 0.01$, ***, $P < 0.001$, ****, $P < 0.0001$).

Fig. 5: Consensus and PHOT-specific GRNs indicate extensive co-regulation of CCM and photoprotective genes. Top: Venn diagram depicting the overlap of the top 10 predicted TFs of the curate CCM and photoprotective (PhPr) genes based on **a.** the consensus or **c.** PHOT-specific GRN. Bottom: Network representation of the top 10 TF sets of **b.** the consensus network or **d.** the *phot* GRN (center nodes, same color code as in panel **a** and the target genes used for prediction (left and right columns of nodes, photoprotection genes are shown in green, qE-related genes in blue, and CCM genes in red. Larger TF nodes indicate higher regulatory strength on target genes. The plotted edge width is proportional to the strength of the regulatory interaction.

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Fig. 3: A PHOT-specific GRN pinpoints QER7 as a suppressor of the expression of qE-related genes. **a.** Dot plot of the relative regulatory strength of the top 10 regulators of qE-related genes in the PHOT-specific GRN (see **Methods**). TFs are marked in green if qE transcript levels were affected in the respective knock-out strain and this effect was reversed by complementation with the knocked-out gene, in yellow, if the effect was not reversed by complementation, and in red, if no mutant effect was observed in the mutant. TFs for which no mutant lines were available are plotted in grey. **b.** WT, *qer7* and *qer7-C* cells were synchronized under 12h light ($15 \mu\text{mol m}^{-2} \text{s}^{-1}$)/12h dark cycles. After sampling for the dark conditions (end of the dark phase), cells were exposed to $300 \mu\text{mol photons m}^{-2} \text{s}^{-1}$ (HL); samples were taken 1 h (RNA) or 4 h (protein and photosynthetic measurements) after exposure to HL. Shown are relative expression levels of qE-related genes at the indicated conditions normalized to WT LL ($n = 3$ biological samples, mean $\pm$ sd). **c.** Immunoblot analyses of LHCSR1, LHCSR3 and ATPB (loading control) of one of the three biological replicate set of samples, under the indicated conditions. **d.** Quantification of immunoblot data of all replicates in panel **c** after normalization to ATPB. WT protein levels at HL were set as 1. **e.** NPQ and qE, measured 4h after exposure to HL ($n = 3$ biological samples, mean $\pm$ sd). The p-values for the comparisons are based on ANOVA Dunnett's multiple comparisons test and are indicated in the graphs (*, $P < 0.005$, **, $P < 0.01$, ***, $P < 0.001$, ****, $P < 0.0001$). Statistical analyses for panel **b** and **d** were applied on log10- transformed values.

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Supplementary Materials for

Widening the landscape of transcriptional regulation of algal photoprotection

Authors

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This PDF file includes:

Supplementary Note

Extended Data Figs. 1 to 10

Other Supplementary Materials for this manuscript include the following:

Tables S1 to S10 (to be downloaded as excel files)

Supplementary Note

Simulation studies³⁰ indicated that the top 10% of the edges in the consensus network are enriched in positive interactions if the underlying approaches perform better than guessing. Thus, a higher overlap with the consensus networks is expected to result in improved predictions. We found that the interactions from the network deconvolution approach exhibited the largest overlap with the consensus network, while GGM assigns the highest rank to consensus interactions (**Extended Data Fig. 2a**). Repeating this analysis by including the two worse performing approaches (i.e. ARACNE and Silencing) resulted in no qualitative changes in the overlaps (**Extended Data Fig. 2b**) and only 7.47% of difference to the original consensus network, demonstrating the robustness of the inferred interactions. Considering the 10% network density threshold, we also inspected the proportion of TF-TF interactions in the different networks and their consensus; we found that it ranges from 2.55% in the GENIE3 network to 2.77% in the network deconvolution approach (**Extended Data Fig. 2c**). Since the fully connected network containing all possible TF-TF and TF-target interactions has a relative TF-TF content of 2.52%, these findings suggest an enrichment of TF-TF interactions in GRNs inferred by all five approaches considered in the consensus.

Extended Data Fig. 1: Variation in ranks of known interactions across approaches and interactions from different inference approaches. Boxplots of the standard deviation of ranks calculated for the curated interactions plotted in **Fig. 1b** over all ranks assigned by one approach or over all ranks assigned to one interaction.

Extended Data Fig. 2: Overlap of consensus network with individual inference methods and proportion of TF-TF interactions in the obtained GRNs. **a.** The graph shows the overlap between the edges of the consensus network and the ranked interactions of the individual approaches normalized to the total number of edges in the consensus network used for regulator prediction. **b.** Same as in **a** but plotting the overlap with a consensus of all used approaches (including ARACNE and Silencing). **c.** Bar plots provide the proportion of TF-TF interactions in comparison to TF-target gene interactions contained in the GRNs inferred by the different approaches and the consensus GRN resulting from their integration. All depicted analyses considered only the interactions within in the 10% network density threshold (see **Methods**).

Extended Data Fig. 3: Genotyping of CLiP mutants affected in the predicted regulators of qE-related genes.

a. Insertion map of the CIB1 cassette in the different genes. Exons are shown in black, introns as interconnecting lines, 5'UTR and 3'UTR in light gray, and primers in arrows. The insertion site of the CIB1 cassette is indicated by the triangle; **b.** PCR-validation of the insertion site in the different CLiP mutants using genomic DNA. To confirm the CIB1 insertion site, gene-specific primers were used that anneal upstream and downstream of the predicted insertion site of the cassette (primer pairs P3-P4, P7-P8, P9-P10 and P5-P6 for *qer6*, *qer1*, *qer7* and *qer4* respectively; **Supplementary Table 9**). Pairs of primers used are indicated at the bottom of the agarose gels used to separate the PCR products. Note that the PCR product of P5-P6 is indicated by an arrow. **c.** Relative expression levels of predicted qE regulator genes in the different CLiP mutants after exposure to HL (300 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$) for 1h.

Extended Data Fig. 4: qE gene and protein expression in CLiP mutants bearing mutations in predicted qE regulator genes. WT, *lci8-oe*, *qer6* and *qer4* cells were acclimated for 16h in LL (15 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$). After sampling for the LL conditions, light intensity was increased to 300 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$ (HL); samples were taken 1 h after exposure to HL. Shown are relative expression levels of *LHCSR3.1*, *LHCSR1* and *PSBS1* at the indicated conditions normalized to WT LL ($n = 3$ biological samples, mean $\pm$ sd). The p-values for the comparisons between the mutants and the WT are based on ANOVA Dunnett's multiple comparisons test on log₁₀-transformed values and are indicated in the graphs (**, $P < 0.01$).

Extended Data Fig. 5: LCR1 is required for proper expression of qE-related genes during transitions from dark-to-light. WT, *lcr1* and *lcr1-C* cells were acclimated for 16h in darkness. After sampling for the dark conditions, light intensity was increased to 300 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$ (HL); samples were taken 1 h (RNA) or 4 h (protein and photosynthetic measurements) after exposure to HL. **a.** Relative expression levels of qE genes at the indicated conditions normalized to WT LL ($n = 3$ biological samples, mean $\pm$ sd). **b.** Immunoblot analyses of LHCSR1, LHCSR3, PSBS and ATPB (loading control) of one out of the three biological replicate samples, under the indicated conditions. **c.** Summary graph of immunoblots of all replicate samples of Extended Data Fig. 3b after normalization to ATPB. Shown are the HL treated samples; WT protein levels were set as 1. **d.** NPQ and calculated qE 4h after exposure to HL ($n = 3$ biological samples, mean $\pm$ s.d). The p-values for the comparisons are based on ANOVA Dunnett's multiple comparisons test and are indicated in the graphs (*, $P < 0.005$, **, $P < 0.01$, ***, $P < 0.001$, ****, $P < 0.0001$). Statistical analyses for panel **a** and **c** were applied on log₁₀- transformed values.

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Extended Data Fig. 6: LCR1 activates the transcription of CCM genes in LL and dark-acclimated cells. **a.** WT, *lcr1* and *lcr1-c* cells were acclimated for 16h in LL (15 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$). After sampling under LL conditions, light intensity was increased to 300 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$ (HL) and samples were taken 1 h after exposure to HL. Relative expression of CCM genes at the indicated conditions normalized to WT under LL. **b.** Relative expression of CCM genes acclimated in 16h of darkness ($n = 3$ biological samples, mean $\pm$ sd). The p-values for the comparisons are based on ANOVA Dunnett's multiple comparisons test on log₁₀-transformed values and are indicated in the graphs (*, $P < 0.005$, **, $P < 0.01$, ***, $P < 0.001$, ****, $P < 0.0001$).

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Extended Data Fig.7: Complementation of *qer1* with *QER1* (*qer1-C* strain) fails to rescue the *LHCSR1* and *PSBS* phenotypes. WT, *qer1* and *qer1-C* cells were acclimated for 16h in LL (15 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$). After sampling for the dark conditions, light intensity was increased to 300 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$ (HL); RNA samples were taken 1 h after exposure to HL. Shown are relative expression levels of qE genes at the indicated conditions normalized to WT LL ($n = 3$ biological samples, mean $\pm$ sd). The p-values for the comparisons are based on ANOVA Dunnett's multiple comparisons test on log10-transformed values and are indicated in the graphs (***, $P < 0.001$, ****, $P < 0.0001$).

Extended Data Fig.8: Relative expression of qE-related genes in asynchronous *qer7* cells. WT, *qer7* and *qer7-C* cells were acclimated for 16h in LL ($15 \mu\text{mol photons m}^{-2} \text{s}^{-1}$; **a-c**) or darkness (**d-f**). After sampling for the LL or dark conditions, light intensity was increased to $300 \mu\text{mol photons m}^{-2} \text{s}^{-1}$ (HL) and samples were taken 1 h (RNA) or 4 h (photosynthetic measurements) after exposure to HL. **a, d**. Relative expression of qE and CCM genes at the indicated conditions normalized to WT LL (**a**) or dark (**d**) respectively ($n = 3$ biological samples, mean $\pm$ s.d.). **b, e**. NPQ and **c, f**. calculated qE, 4h after exposure to HL ($n = 3$ biological samples, mean $\pm$ s.d.). The p-values for the comparisons are based on ANOVA Dunnett's multiple comparisons test and are indicated in the graphs (**, $P < 0.01$, ***, $P < 0.001$, ****, $P < 0.0001$). Statistical analyses for panel **a** and **d** were applied on $\log_{10}$ -transformed values.

Extended Data Fig. 9: Reconstructed GRNs recapitulate experimentally found regulators of qE and CCM.

Network representation of the top 10 regulators of CCM genes quantified via qPCR (**Fig. 4**), and qE genes as predicted from **a.** the consensus or **b.** PHOT-specific GRN. Regulators are represented by central nodes (same color code as in **Fig. 5**), target qE genes present in the respective network are shown in blue (*LHCSR1*, *LHCSR3.1*, *LHCSR3.2*, *PSBS1*, *PSBS2*) and target CCM genes are shown in red (*LCI1*, *LCI2*, *CAH4*, *CCP1*, *HLA3*, *BST1*). Larger TF nodes indicate higher regulatory strength on target genes. The plotted edge width is proportional to the strength of the regulatory interaction.

Extended Data Fig. 10: LCR1 mutation and LCI8 overexpression confirmation. a. Relative expression of *LCR1* in WT, *lcr1* and *lcr1-c* strains. Cells were acclimated for 16h in LL or darkness. After sampling for the LL or dark conditions, light intensity was increased to 300 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$ (HL), and samples for RNA purification were taken 1 h after exposure to HL ($n = 3$ biological samples, mean $\pm$ sd). *LCR1* gene expression was non-detectable (indicated as n/d in the graphs) in the *lcr1* mutant. **b.** Immunoblot analysis of LCI8-FLAG fused protein in WT and *lci8-oe* cells grown in TAP, LL. The p-values for the comparisons are based on unpaired t-tests on log10-transformed values and are indicated in the graphs (*, $P < 0.005$, ***, $P < 0.001$).

171 **Supplementary Table Captions**

172 **Supplementary Table 1:** RNAseq data sets used in this study; HSM: Sueoka's high salt medium (1); HSM –
173 new trace = modified Sueoka's high salt medium with trace elements as described in (2); Rep= biological
174 replicates

175 (1) Sueoka, N. Mitotic replication of deoxyribonucleic acid in *Chlamydomonas reinhardtii*. Proc. Natl.
176 Acad. Sci. U.S.A. 46, 83–91 (1960).

177 (2) Kropat, J. et al. A Revised Mineral Nutrient Supplement Increases Biomass and Growth Rate in
178 *Chlamydomonas Reinhardtii*. Plant J. 2011, 66 (5), 770.

179 **Supplementary Table 2:** List of Transcription factors used in this study; Family info was adapted from Pérez-
180 Rodríguez et al. (2010) and Jin et al. (2017)

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182 **Supplementary Table 3:** Edge list representation of the consensus network with mean ranks resulting from
183 Borda count election method (Borda (1781), Marbach et al. (2011)) as edge attributes.

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185 **Supplementary Table 4:** Top 10 predicted regulators of qE genes in the consensus GRN. Regulation strength
186 was calculated based on the edge rank in the consensus network.

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188 **Supplementary Table 5:** Edge list representation of the PHOT GRN with importance score from GENIE 3
189 (Huyn-Thu et al., 2010) as edge attributes.

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191 **Supplementary Table 6:** Top 10 predicted regulators of qE genes in the PHOT GRN. Regulation strength was
192 calculated based on the edge weight assigned by GENIE3.

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194 **Supplementary Table 7:** List of genes putatively involved in photoprotection used for regulator prediction

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196 **Supplementary Table 8:** List of genes putatively involved in CCM used for regulator prediction.

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198 **Supplementary Table 9:** PCR primers for CLiP mutant validation and complementation in this study

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200 **Supplementary Table 10:** RT-qPCR primers for all genes in this study

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