Cold priming uncouples light- and cold-regulation of gene expression in Arabidopsis thaliana

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Abstract

Background The majority of stress-sensitive genes responds to cold and highlight in the same direction, if plants face the stresses for the first time. As shown recently for a small selection of genes of the core environmental stress response cluster, pre-treatment of Arabidopsis thaliana with a 24 h long 4 °C cold stimulus modifies cold regulation of gene expression for up to a week at 20 °C, although the primary cold effects are reverted within the first 24 h. Such memory based regulation is called priming. Here, we show the effect of 24 h cold priming on cold regulation of gene expression on a transcriptome-wide scale and test if and how cold priming effects light regulation of gene expression.

Results 304 genes were differently regulated between cold-primed and non-primed plants after a second 24 h long 4 °C cold treatment. After triggering the plants with a heat-filtered highlight stimulus (800 μmol quanta m $^{-2}$ s $^{-1}$), 1011 genes showed priming dependent regulation. Only 32 of the priming-sensitive genes responded similarly to cold and light triggering. The majority of the priming-sensitive genes were regulated in a stressor-specific manner. 29 genes were even inversely regulated by the two triggering stimuli. Cold priming preferentially supported expression of genes involved in the defense against plant pathogens upon cold triggering. The regulation took place on the cost of the expression of genes involved in growth regulation and transport. On the contrary, cold priming resulted in stronger expression of genes regulating metabolism and development and weaker expression of defense genes in response to highlight triggering. qPCR in several independently cultivated and treated samples confirmed the trends observed by RNA-Sequencing.

Conclusion The 24 h long priming cold stimulus activates a several day lasting stress memory that controls cold and light regulation of gene expression and adjusts growth and defense regulation in a stressor-specific manner.

Background

Plants respond dynamically to a wide range of environmental signals and can adjust to many unfavourable conditions [1–3]. Performance optimization to persisting shifts in the conditions is called acclimation or acclimatization. It takes several days and involves cost-intensive changes in metabolism, gene expression and sometimes even in the anatomy and morphology [4, 5]. Specific signalling, such as by the cold-induced ICE (inducer of CBF expression)-CBF (C-repeat binding factor)-pathway [6] and e.g. ROS (reactive oxygen species)- and abscisic acid-signalling conjointly configure the plants towards activation and manifestation of higher stress tolerance [1, 7]. As soon as the conditions improve, most acclimation supporting reactions stop almost immediately and reverting regulation starts [8–10].

If the lag-phases between successive stress events, which are by themselves too short to establish protection, are short enough to maintain part of the acclimation responses, several short stimuli can lead to similar or higher stress tolerance than a continuous stress experience [11]. The phenomenon is called entrainment.

By contrast, priming is independent of the persistence of the stress or of accumulation of primary stress responses [12]. The stress memory (caused by the priming stimulus) uses information carriers that are set at low metabolic costs and modify the response to a later stress (triggering stress) [12−13]. Priming has been described for a wide range of biotic and abiotic stress stimuli [12−14]. However, in most cases the (precise) nature of the specific memory mechanism is still unknown. According to the first records, it can range from meta-stable metabolic imprints to trans-generation stable epigenetic marks [13, 15, 16].

In our earlier study on cold-priming, we showed that priming of Arabidopsis thaliana for 24 h at 4 °C differentially regulates genes of the core environmental stress response cluster, which are induced in response to various stressors, including cold [17, 18]. Cold priming weakened the induction of the zinc finger transcription factor ZAT10 (zinc-finger transcription factor 10; STZ; At1g27730) (and to a lesser extent BAP1 (BON1-associated protein 1; At3g61190)) upon a 5 day later cold stimulus and supported cold activation of CHS (chalcone synthase; TT4; At5g13930) and PAL1 (phenylalanine ammonium lyase 1; At2g37040) expression [17]. The same priming stimulus did not affect cold-induction of COR15A (At2g42540) [17], which is under control of the main cold acclimation regulating ICE-CBF-pathway [1].

In this small selection of genes, ZAT10 showed the strongest primability [17]. ZAT10 expression responds to a wide range of abiotic stresses, including high light intensities and cold [17−19]. The transcription factor mediates secondary gene expression regulation, such as induction of the non-plastid ascorbate peroxidase APX2 (At1g07890) and chloroplast iron superoxide dismutase FSD1 (At4g25100) and counteracts full activation of osmotic and salt tolerance [20].

ZAT10 is hardly expressed under non-stress conditions [17, 21]. In response to photooxidative stress, which occurs upon sudden cold or excess light [22−24], it is induced by reactive oxygen species (ROS), presumably by H_2O_2 [25]. In highlight, ZAT10 induction is supported by PAP (3'-phosphoadenosine 5'-phosphate) that accumulates upon photooxidative inhibition of the PAP-dephosphorylating chloroplast stroma localized phosphatase SAL1 (At5g63980) [26]. In the cold, CBF-dependent induction of the transcription factor CZF1 (At2g40140) activates ZAT10 expression [27, 28]. The various ZAT10 regulating pathways are differently controlled by chloroplast antioxidant protection. Whereas, for example, SAL1 regulation by ROS depends stronger on stromal ascorbate peroxidase (sAPX) function than on thylakoid ascorbate peroxidase (tAPX) activity, cold regulation of CBF genes is antagonized by tAPX [29] and cold priming of ZAT10 is solely mediated by transient post-cold accumulation of tAPX and can be antagonized by tAPX RNA silencing [17, 30].

Cold and excess light cause imbalances between light excitation of the thylakoid membrane and redox energy consuming chloroplast metabolism [22−24]. Consistent with the high similarity of the effects on photosynthesis, the two stress types regulate 87% of the responsive genes in the same direction in naïve plants [31]. Many cold-responsive genes, e.g. BAP1 and the ZAT (Zinc finger of Arabidopsis thaliana) transcription factors ZAT6 (At5g04340), ZAT10 and ZAT12 (At5g59820) [20, 32−34] belong to the group of "core environmental stress response genes" that are induced in response to various stresses and mediate stress response regulation and acclimation processes [18]. The high overlap between

transcriptome regulation in response to cold and light stress [31] suggests a strong trans-effect of cold-priming on light-regulation of gene expression. On the contrary, the complexity of regulation of primary stress responsive genes, like ZAT10, let assume cis- and trans-specific effects. In the present study, we compare the effect of 24-h cold priming on the response to a 5 day later applied 4 °C or temperature-controlled highlight (800 μmol photons $m^{-2} s^{-1}$) triggering stimulus, first, on frequently with ZAT10 co-regulated genes and, finally, in a transcriptome wide scale to investigate the specificity of cold-priming on future gene expression regulation.

Results

Cold priming results in decreased cold activation of specific ZAT genes

ZAT10 showed strongest primability in the previous study on selected cold-responsive genes [17]. To identify similarly regulated genes in Arabidopsis thaliana, publicly available data resources on transcript abundance regulation were scanned with GENEMANIA for ZAT10-like regulated genes [35]. The 15 highlighted genes (Fig. 1A) included BAP1, which is, like ZAT10, cold-priming sensitive and less inducible by cold 5 days after 24 h cold priming at 4 °C, as shown before [17]. Additionally, GENEMANIA highlighted the genes for the zinc-finger transcription factors ZAT6 (Zinc finger protein 6; At5g04340), ZAT11 (At2g37430), ZAT12 (At5g59820), ZAT5 (At2g28200), ZAT18 (At3g53600), the WRKY transcription factors WRKY33 (At2g38470) and WRKY40 (At1g80840), the AP2-type transcription factors ERF6 (Ethylene response factor 6; At4g17490), ERF13 (At2g44840) and ERF104 (At5g61600), the mitochondrial uncoupling protein PUMP4 (At4g24570) and the Ca^{2+}-binding protein encoding gene At4g272800. A similar analysis on the STRING v.11 platform [36] named also ACS6 (1-aminocyclopropane-1-carboxylate synthase 6; At4g11280), that is involved in ethylene biosynthesis, as a ZAT10 co-expressed gene (Fig. 1). All these genes respond, like ZAT10, to a wide range of abiotic stress stimuli and to oxidative stress [20, 37–43].

STRING v.11 further indicates protein-protein interactions (Fig. 1; orange lines). Via feed-back effects, they could impact on transcript abundance regulation. The ZAT10 transcription factor interacts with the MAP kinases MPK3 (At3g45640) and MPK6 (At2g43790), which are elements of a core plant stress signal transduction pathway responding to biotic and abiotic signals [44, 45]. MPK6 and MPK3 also phosphorylate ZAT6 [46], ERF6, ERF104 [41, 43], WRKY33 [47], WRKY40 [41] and ACS6 [48]. Additionally, ZAT10 interacts with the transcriptional co-repressors TOPLESS (TPL, At1g15750) and TOPLESS-RELATED-4 (At3g15880) [44, 49–51]. TPL binds also ZAT6 [50]. To test ZAT10-like regulated genes for the cold-primability of their cold regulation, we selected genes with different affinity to MPK6 / MPK3 and / or TPL, namely ZAT10, ZAT6, ACS6 and WRKY40 for a qPCR (quantitative polymerase chain reaction)-based priming analysis. We further included the gene for the bi-functional enolase LOS2, which is a negative upstream transcriptional regulator of ZAT10 [52]. The transcript levels of these genes were analyzed by qPCR immediately after triggering in previously naive plants (T) and plants that were cold-primed 5 days before cold triggering (PT). As controls, untreated plants (C) and plants (P) that perceived 5 days earlier the priming cold-treatment, but were not cold-triggered, were analyzed.

Like ZAT10, the transcript levels of ZAT12 and ZAT6 were significantly decreased in PT-plants as compared to T-plants, demonstrating priming-sensitivity (Fig. 1B). ACS6 and WRKY40 were not sensitive to the triggering stimulus, independent of whether the plants were cold-primed or not. Regulation of LOS2, which binds the ZAT10 promoter and controls ZAT10-mediated cold-induction of the cold and drought marker gene RD29 [52], was strongly cold-inducible (comparison of transcript levels in C- and T-plants), but not priming-regulated (comparison of transcript levels in T- and PT-plants). The analysis gave no indication that interaction with known ZAT10-interacting proteins controls priming, but demonstrated that cold-priming affects specific genes, even in a group of genes which are otherwise widely co-regulated with ZAT10 [17–19] (Fig. 1).

The effect of cold priming on the regulation of the ZAT genes upon highlight triggering

For comparison of the cold-priming effect on cold- and highlight triggering, we established a heat filtered highlight set-up (800 μmol quanta $m^{-2} s^{-1}$), which induces ZAT10 expression to a similar extent as the cold treatment used for cold priming and cold triggering does (documented in the response of T-plants in Fig. 2C). The set-up was evaluated by qPCR for its impact on regulation of well characterized light and heat regulated genes (Fig. 2A). After 2 h in highlight, the transcript levels of the light-inducible genes ELIP2 (early light induced protein 2, At4g14690; [53]), GPX7 (glutathione peroxidase 7, At4g31870; [54]) and PAL1 (phenylalanine ammonium lyase 1, At2g37040) were increased (Fig. 2B left). The heat filter was sufficient to counteract significant activation of the heat sensitive genes HSFA7a (At3g51910) and HSFA7b (At3g63350) [55, 56] (Fig. 2B right).

Besides induction of ZAT10, the light treatment increased the ZAT6 transcript levels almost as strong as the 24 hours cold treatment (T-plants in Fig. 2C). ZAT12 showed only a very weak (but also significant) response to the light treatment. In cold-primed plants, the mean transcript levels of ZAT6 were lower in PT-plants than in T-plants, indicating primability, although the effect was not significant. On the contrary, the transcript levels of ZAT10 and ZAT12 were more similarly regulated in primed and non-primed plants after light triggering. Consequently, cold-priming did not have any or has only very little effect on the light triggering response of these genes.

Photosynthetic performance after triggering

The differences between the cold- and the light triggering response of the ZAT genes (Fig. 1B and 2C), especially ZAT10, could result from effects of priming on the photosynthetic electron transport efficiency. To test this hypothesis, we compared the photosynthetic performance of photosystem-II in cold-primed plants after cold and light triggering by chlorophyll-a fluorescence analysis. Triggered (T) and primed + triggered (PT) plants were analysed site-by-site by 2-dimensional chlorophyll-a fluorescence imaging in middle-aged leaves, which show strongest priming sensitivity in 4-week-old plants [30] (Fig. 3).

After cold and light triggering, the maximal quantum yield of photosystem-II (F_V/F_M; 0 min in Fig. 3 top) was similar in dark-acclimated T- and PT-plants. Also, the quantum yield of photosystem II (Φ_{PS-II}) and photochemical and non-photochemical quenching (qP and NPQ) did not differ between primed and non-

primed plants (T- and PT) in both plant groups upon illumination with 185 µmol quanta m^{-2} s^{-1} actinic light (Fig. 3). The similarity of the response contradicts the hypothesis that the differences in gene expression regulation result from stress-induced damage or regulation of photosystem-II activity.

Effect of cold priming on cold- and highlight-regulated gene expression

For more insight into the effect of cold priming on the stress responses, we maximally widened the target gene spectrum and performed genome wide RNA-sequencing (RNASeq analysis) 2 h after cold (4 °C) and light triggering (800 µmol quanta m^{-2} s^{-1}) of 5 days earlier cold-primed and non-primed plants. RNA sequencing resulted in 23.76–24.14 million reads per sample (Suppl. Table 1). At minimum, 98.49% of the reads could be mapped to the TAIR10 genome (Suppl. Table 1). Sequences were recorded for 24085 different genes. The transcript levels of many well-known, highly cold and light-responsive transcription factors, e.g. CBF1 (At4g25490) and CBF3 (At4g25480) [57], ANAC078 (At5g04410) [58] and ZAT10 [17, 21] and ZAT6 [59], were 2 h after cold or light triggering already strongly decreased (Suppl. Table 2). At the same time, the transcript levels of secondarily cold regulated genes, such as the CBF3-regulated gene COR15A (At2g42540) and the ANAC078 target gene At1g56650, At3g01600 and At5g58610 [60] still were induced (Suppl. Table 2). Genes that are well characterized for their heat induction, such as HSFA2 (At2g26150), HSFA7a (At3g51910), and HSA32 (At4g21320), were only very weakly expressed in all samples (Suppl. Table 2). The transcript level of the senescence regulating NAC transcription factor ORE1 (ANAC092; At5g39610) [61] was not increased in any sample (Suppl. Table 2). The expression pattern confirmed high responsiveness of stressor-specific target genes and showed that the treatments did not induce heat signaling or activate senescence.

61.7% of the genes that were at least 2-fold up-regulated and 32.8% of the genes at least 2-fold down-regulated in response to light in unprimed plants, were also at least 2-fold regulated by the cold treatment. On the contrary, only 0.3 and 5.5% of the at least 2-fold regulated genes were inversely regulated by cold and light. Thus, our cold and light treatments widely regulated genes in the same direction in unprimed plants, similar as shown before by others [31].

Volcano plots (depicting the intensity of priming-dependent regulation based on the false discovery rate (FDR)) (Fig. 4 top) and blotting of the gene expression levels of primed plants (y-axes) against the gene expression levels of the respective unprimed plants (x-axes) (Fig. 4 bottom), showed that cold priming affected cold and light regulation of only specific genes. Cold triggering resulted in much less gene expression variability than light triggering in cold-primed plants (Fig. 4 bottom). In general, most significant priming-dependent regulation was observed for medium strongly expressed genes (Fig. 4 bottom).

Principal component analysis (PCA) (Fig. 5A) and clustering (Fig. 5B) of the relative transcript level in P- and PT-plants demonstrated that the priming effects on not, cold- and light-triggered plants differed in direction and intensity. Already from this first comparison, it can be concluded that the priming effects

observed after triggering did not result from prolonged gene dysregulation in response to the priming stimulus, but that priming affects the response to the triggering stimulus in a stressor-specific manner.

Long-term, not triggering-dependent gene expression effects of cold priming

For more stringent gene regulation analysis, the 13775 genes were selected that were detected in all samples and were recorded with FPKM (fragments per kilobase of exon per million reads mapped) values of 5 or higher in at least one data set. The effects of priming on the transcript levels were calculated by dividing the FPKM-values of primed and non-primed plants at the end of the lag-phase (P / C) and in cold-triggered (PT-C / T-C) and light-triggered plants (PT-L / T-L).

Transcriptome comparison between C and P plants at the end of the 5-day-long lag-phase demonstrated that the transcriptome was widely reverted prior to application of the triggering stimuli. Only for 12 genes more than 2-fold higher and only for 4 genes more than 2-fold lower transcript levels were recorded in primed plants as compared to control plants (Fig. 5C top, Suppl. Table 3). At1g53870 (encoding a LURP (Late/sustained Up-regulation in Response to Hyaloperonospora parasitica)-like protein, At1g73260 (putative trypsin inhibitor), At4g12490 and At4g12480 (two bifunctional inhibitor proteins, AZI3 and EARLI1), a cation exchanger (At3g51860) and a Haloacid dehalogenase-like hydrolase (HAD) superfamily protein (At5g36790) were strongest up-regulated. These genes were only weakly expressed under control conditions. Consequently, the absolute regulation of the transcript levels was low. On the contrary, the transcript levels of a transmembrane protein (At4g12495), the senescence and stress inducible gene SAG13 (At2g29350, encoding a short-chain alcohol dehydrogenase) and extensin-4 (At1g76930) were recorded with FPKM values higher than 10. Their transcript levels were more than 2-fold increased 5 days after cold priming reflecting a strong absolute effect (Supp. Table 3).

The four genes which were down-regulated in P compared to C encode lipid-transfer protein-4 (At5g59310), a glycine-rich protein (At1g04800), another LURP1-like protein (At1g53890) and an embryo development controlling gene (At4g29660) (Suppl. Table 3).

Analyzing the transcript abundance patterns at lower threshold (FPKM $\geq$ 5 in at least one of the treatments and $\log_2$ (primed / unprimed) $\geq$ I 0.5 I) (Fig. 5C bottom) showed only for two of the 365 potentially long-term regulated genes, namely a hypothetical gene (At5g23411) and At1g53870 (encoding a LURP1-related protein), co-upregulation in not, cold- and light-triggered plants. Only one hypothetical gene (At1g13470) was co-downregulation in all primed plant groups (Suppl. Table 4). The very low number of co-regulated genes shows that the priming memory is not stored in the regulation potential of individual genes, but affects gene regulation in a stressor-specific manner, such as by the control of the signal transduction pathways.

Common triggering-dependent effects of cold priming on cold and light triggering

Since cold and excess light regulate the majority of genes in the same direction [31], regulation of common signal transduction elements would result in high similarity between the effect of cold and light

triggering on priming sensitive genes. Already the analysis of a small selection of ZAT10-related genes showed differences (Fig. 1B and 2). On the transcriptome level, RNASeq analysis identified under the more stringent conditions used for analysis (FPKM $\geq$ 10 and $\log_2$(PT/T) $\geq$ I 1 I) only a gene for a not further characterized transmembrane protein (At4g22510) as potentially (at least 2-fold) priming co-regulated in cold- and light-triggered plants (Fig. 5C top).

Lowering the threshold to FPKM $\geq$ 5 and $\log_2$(PT/T) $\geq$ I 0.5 I showed 29 genes as being co-regulated in a priming-dependent manner after light and cold triggering (Suppl. Table 4). Eight of the 17 co-up-regulated transcripts map to the same chromosome region and several of the short genes overlap in sense and antisense orientation. Consequently, the FPKM values (as calculated for these genes) may overstate the actual transcript abundance and the regulation amplitudes of individual genes. The remaining co-up-regulated genes encode (besides hypothetical proteins and proteins of unknown function) with ERD6-like 1 (early response to dehydration-6 like-1; At1g08920), CC-NBS-LRR class immune receptor (At1g59218), the extensin OLE1 (At2g16630), a kinase inhibitor-like protein (At2g28870), plastome-encoded photoreceptor protein M (Atcg00220) and the plastid ribosomal subunit L32 (Atcg01020) a diverse spectrum of proteins.

In the group of the 12 genes, which are less expressed after light and cold triggering in primed plants (Suppl. Table 4), three encode disease associated genes, namely two β-glucanases (PR2 (BGL2; At3g57260) and BLG3 (At3g57240)) and one chitinase (At2g43570).

Specific effects of cold priming on cold and light triggering

Most priming-regulated genes were regulated by either cold or by light triggering (Fig. 5C). Under highly selective conditions (FPKM $\geq$ 10 and $\log_2$(PT/T) $\geq$ I 1 I), the transcript levels of only two genes, expansin-A8 (At2g40610) and glycin-rich protein 9 (At2g05440), were lower after cold triggering due to cold priming. In parallel, 13 genes were stronger expressed after cold triggering in cold-primed plants than in non-primed ones. Three of them, Kunitz trypsin inhibitor 1 (At1g73260), NIT2 (At3g44300) and SAG13 (At2g29350), were already induced prior to application of the triggering stimulus. Nine of the remaining 10 genes encode (hypothetical) lipid transfer proteins or are not characterized for their function (Suppl. Table 3). The remaining, trigger-specifically regulated gene was OLE1 (At2g16630) that encodes an extensin.

On the contrary, light triggering resulted in cold-primed plants in specific accumulation of transcript levels for 9 genes, of which three encode heat shock proteins. Various defense-related genes, such as PR2 (pathogen responsive gene 2, At3g57260), PR4 (At3g04720), a pathogen and circadiane controlled gene PCC1 (At3g22231) a chitinase (At3g12500) and five defensins, were less induced by highlight in primed plants than in naïve ones (Suppl. Table 3). Two genes, namely, At2g73260 and At4g12495, encoding a trypsin inhibitor and a transmembrane protein, showed inverse regulation in primed plants before and after light triggering. Inversion of the priming-effect by the triggering response demonstrates that priming actively affects gene regulation by the triggering light stress event.

The quantitative differences between the priming-impact on cold and light triggering were confirmed when the genes were filtered based on weaker criteria (FPKM $\geq$ 5 and $\log_2$(PT/T) $\geq$ I 0.5 I) (Fig. 5C): 130 genes were specifically induced and 121 down-regulated in cold-primed plants after cold triggering. Light triggering of cold-primed plants resulted in stronger induction of 613 and down-regulation of 334 genes in comparison to light-triggered non-primed plants.

Analysis of regulation patterns by qPCR

Regulation observed by RNASeq was evaluated by qPCR in 5 independently cultivated and treated biological replicates for 5 genes showing priming effects at the end of the lag-phase, for 5 genes which were regulated in a priming-dependent manner after cold triggering, and for 5 priming sensitive genes regulated by light (Fig. 6A). The priority was given to genes with high FPKM values. In the qPCR analysis, the transcript levels were normalized to the expression intensity of the constitutively expressed gene YLS8 (At5g08290) [62]. In all three gene sets, three genes were selected which are up-regulated in primed plants as compared to non-primed plants and two which were down-regulated. 13, out of the selected 15 genes, showed in the qPCRs significant regulation (Student t-Test, $p < 0.05$) consistent with the RNASeq data. The transcript levels of the other two genes, namely At5g59720 (encoding the heat-shock protein HSP18.2) and At1g73260 (encoding a Kunitz factor protein) were by average (although not significantly) more than 2-fold regulated in the same direction as in the RNASeq.

Of the five genes tested by qPCR for higher transcript levels 5 days after cold priming (Fig. 6A top), RNASeq analysis indicated only for SAG13 also higher transcript levels after cold triggering. qPCR in independently cultivated and treated biological replicates confirmed this effect (Fig. 6B). Additionally, it also showed down-regulation in primed plants after light triggering consistent with the RNASeq analysis (Suppl. Table 3; Fig. 6B). qPCR further confirmed the regulation observed by RNASeq for extensin-4 (At1g76930) and PR2 (At3g57260) before and after triggering (Fig. 6B). The ratios calculated from the FPKM values of primed and the respective unprimed plants (P/C; PT-C/T-C and PT-L/T-L) were for all treatments in the range of the values obtained by qPCR for the various biological replicates (Fig. 6B).

Functional categorization of the cold priming effect on the triggering response

Functional categorization of the priming-regulated genes based on analysing the enrichment of gene ontologies (GO) [63, 64] was performed with the wider data set ($\log_2$ (PT/T) >I 0.5 I; FPKM $\geq$ 5) on the AgriGO v2 platform (http://systemsbiology.cau.edu.cn/agriGOv2/). Data processing was evaluated using the Fischer test (F-test) and the Yekutieli method for α-level adjustment at a p-level of 0.05 [65]. The minimum threshold for statistical testing and multi-test adjustment was set to 5 genes per GO-term [66]. From the primary data, the subset of the most specific GOs within the hierarchical GO structure were extracted for the figures (Figs. 7 and 8). The full lists including information on the p-value and FDR (False Discovery Rate) and graphical images depicting all GO-terms in hierarchical order are provided in the supplements (Suppl. Table 5).

In the group of transcripts that were up-regulated in cold-primed plants after cold triggering, stress regulated genes were significantly enriched in comparison to non-primed cold stressed plants (Fig. 7). Especially genes responding to wounding, immune and programmed cell death regulation and / or genes under control of jasmonic acid signaling were over-represented. Additionally, priming preferentially affected the cold triggering response of genes involved in the starvation regulation and in flavonoid and anthocyanin biosynthesis (At4g22880, At4g09820, At2g02990, At3g29590, At5g17220, At5g42800, At4g14090, At5g54060). All eight genes of the latter group were also induced by excess light, but were less induced or even inversely regulated in primed plants after light compared to primed plants after cold triggering (Suppl. Table 5). CHS and PAL1, which regulate early steps of phenylpropanoid metabolism and were previously shown to be stronger activated in cold-primed plants upon cold triggering [17], were also stronger activated in cold-primed plants in response to cold triggering in the new dataset, although they did not pass the threshold criteria used here for the bioinformatics analysis. In parallel, cold triggering in cold-primed plants, resulted in weaker expression of genes involved in transport organization, growth and morphogenesis (Fig. 7). Various of the weaker expressed genes respond to auxin-activated signaling and response pathways.

After light triggering, genes involved in organelle organization, morphogenesis and nucleic acid metabolism were stronger induced in cold-primed plants than in non-primed ones. Genes responding to biotic stimuli, acids and oxygen-containing organic compounds (At5g44420, At3g15356, At3g22231, At2g14560 At1g73260, At4g10500, At3g16530) and genes involved in metabolic regulatory processes are less represented in primed plants (Fig. 8). In general, GO analysis showed that cold priming results in an inverse support of growth and biotic stress response upon cold and light triggering (Figs. 7 and 8, orange and dark green bars).

Sub-analysis of the priming-responsive genes inversely regulated by cold and highlight

In the group of 159 genes with higher transcript levels in PT-C plants than in T-C plants and the 379 genes down-regulated in PT-L plants as compared to T-L plants (FPKM values > 5 and $\log_2$ (PT/T) > |0.5 |) 17 genes were inversely regulated (Supp. Table 6). Additionally, 12 genes were inversely regulated between the group of 145 genes down-regulated PT-C plants and 633 genes up-regulated in PT-L (Supp. Table 6).

Six of these (in total) 29 inversely regulated genes were not annotated in TAIR10, which is the data background used for functional categorization with AgriGO v2. Only one biological function was identified as being overrepresented in the remaining group of 23 genes (Suppl. Table 7). Seven of the 23 genes, namely At2g29350, At4g37990, At1g73260, At2g43510, At3g22231, At3g04720 and At3g12500, respond to biotic stimuli. They all showed higher transcript levels after cold triggering and lower ones after light triggering if the plants were cold-primed before (Suppl. Table 6). Taking even slight regulation prior to triggering into account, all these genes show specific responses to light triggering (Suppl. Table 6). Three of them (At3g22231, At3g04720 and At3g12500) showed also specific up-regulation of the transcript levels after cold triggering and down-regulation in response to light. These three two-directionally regulated genes encode the plasma membrane protein Pathogen and Circadian Controlled 1

(PCC1; At3g22231), Pathogenesis Related 4 (PR4; At3g04720) and a basic chitinase (CHI-B; At3g12500). All three genes are associated with pathogen defense. Also CHS (At5g13930), but not the other core response gene PAL1 (At2g37040), showed stronger expression in cold-primed plants upon cold triggering and lower transcript levels after excess light triggering, although with lower amplitudes than PCC1, PR4 and CHI-B (Suppl. Table 2).

Expression network analysis on the GENEMANIA plattform indicated only very faint co-expression between PR4 and CHI-B and no co-regulation of the two genes with PCC1. The impression that these genes are hardly co-regulated in naïve plants was confirmed by comparison of transcript abundance regulation using the compare-mode of the eFP browser [67] on publicly available transcript abundance regulation data for developmental regulation in Arabiopsis thaliana and the response to biotic and abiotic stress. qPCR analysis confirmed the inverse regulation of pathogen related genes PCC1, PR4 and Kunitz 1 after cold and light triggering of cold-primed plants (Fig. 9). For CHI-B, the transcript levels were below the detection level of qPCR.

The other 16 genes, which responded inversely to cold and light triggering in a priming-dependent manner, have diverse functions. Five encode transmembrane proteins (At4g12495, At1g79170, At1g16916, At5g65580 and At1g53035), two protease inhibitors (At1g73260 and At2g43510), two protein phosphatases 2C (At5g02760 and At3g16800) (Suppl. Table 6).

Discussion

Stresses activate a sequence of events starting within seconds to minutes with the first measurable changes in transcriptional activity [68, 69]. After a period of massive regulation, the transcriptome gets adjusted to regulation of acclimation. Inactivation of primary stress regulation and secondary regulation dominates the post-stress phase [70]. In our experiment, 5 days after the plants perceived the priming cold stimulus, primary and secondary gene expression regulation was almost fully reset (Supp. Table 2; Fig. 5). At this stage, we exposed the plants either to cold or to excess light. The two stresses, if applied to naïve plants, regulate the majority of genes in the same direction [18, 31] (Suppl. Table 2). 5 days after cold priming, however, the same stimuli caused mainly specific effects and partly even inverse transcript abundance regulation (Figs. 1, 2, 5 and 6; Suppl. Table 2–4). Due to the low overlap between genes that were cold and light regulated in a priming-dependent manner (Figs. 1, 2 and 5 plus Suppl. Table 3–6), we conclude that cold priming uncouples cold and light regulation for specific genes.

The mechanisms, by which cold priming establishes the memory and how the priming-induced information is recorded in primed plants, are still under investigation [13, 15, 16]. Various studies suggest an epigenetic memory, such as histone and DNA (de-)acetylation or (de-)methylation, for storing information on thermal stress events [15, 71–73]. For example, COR15A (At2g42540) and COR47 (At1g20440) are stronger expressed, if the second cold stimulus quickly follows the first one [74]. The majority of the cold-induced histone marks, however, is metastable. Consequently, the effects on gene expression regulation get quickly lost. For example, the cold-priming effect on COR15A can be fully revert

within 24 h, if priming was performed with a short cold stimulus [72]. On the contrary, prolonged cold, such as 2 weeks at 4 °C, leads to higher transcript accumulation of COR15A upon a 5 day later applied 24 h 4 °C triggering stimulus [17]. Transformation of a metastable cold memory into a stable one, such as in the regulation of FLOWERING LOCUS C (FLC; At5g10140), requires several days or even several weeks of cold exposure [75]. Consistent with the previous qPCR-based analysis of COR15A regulation [17], 5 days after 24 h cold priming none of the reference genes for epigenetic cold memories, namely COR15A, COR47 or FLC (Suppl. Table 2) [74, 75], showed priming-dependent regulation in the present study.

Despite widely overlapping light and cold effects on the transcriptome of naïve plants [31] (Suppl. Table 2), co-regulation was only observed for 32 genes (of which several overlap and might reflect double annotations) in cold-primed plants (Fig. 5B). 29 genes were even inversely regulated. The latter group included well characterized genes of the core environmental stress response cluster [18], such as ZAT transcription factors, CHS and the pathogenesis associated genes PCC1, PR4 and CHI-B (Suppl. Table 2; Fig. 9). Cold-priming supported expression of stress (hormone) responsive genes upon cold triggering and resulted in lower expression of genes related to growth and metabolite transport (Fig. 7). Genes with functions in stress response regulation were down-regulated in cold-primed plants after light triggering and genes involved in growth, metabolism and development up-regulated (Fig. 8). Such inverse effects of cold priming on gene expression regulation demonstrate that priming affects cold- and light sensitivity and responsiveness in a stressor specific manner. With respect to biological function, our analysis highlighted two cold-priming effects:

(1) Cold priming supports cold-regulation of genes involved in anthocyanin and flavonoid metabolism

Eight genes involved in the biosynthesis of flavonoids and anthocyanins showed priming dependent regulation in response to cold triggering (Fig. 7; Suppl. Tables 2 and 5). Anthocyanin and flavonoid are broad spectrum protectants that not only filter ultra violet (UV) - and / or blue and red light, but have also antioxidant capacities [76, 77]. Their synthesis is activated by various stresses, including cold, UV light, drought, salt and highlight [78, 79]. CHS and PAL1, which were previously shown by qPCR to be stronger induced in cold-primed plants upon cold triggering [17], encode enzymes catalysing initial steps of phenylpropanoid metabolism and controlling the flux capacities into chalcone metabolism. Although regulation of CHS and PAL1 transcript levels did not pass the strict threshold criteria applied in this study, their transcript abundances were also higher in cold-primed plants upon cold triggering (and lower or unchanged upon highlight triggering) (Suppl. Table 2). Five of the eight cold- genes, namely At5g42800, At5g17220, At4g22880, At4g14090 and At4g09820, plus CHS and PAL1 can be activated by MYB75 (At1g56650) [80]. MYB75 expression is regulated by ZAT10 [20]. MYB75 transcript levels showed slight positive cold priming effects upon cold triggering, but not upon highlight triggering (Suppl. Table 2), consistent with selective priming-dependent regulation of ZAT10 upon cold, but not light triggering (Figs. 1 and 2). It could link priming regulation of CHS / PAL1 and ZAT10, which we previously hypothesized to be controlled by parallel induced, inversely acting pathways [17].

(2) Inverse cold-priming dependent regulation upon cold and light triggering

The most striking observation of our study was the inverse trade-off between the support of growth and defence upon cold- and light triggering after cold priming. Cold pretreatment is well known to decrease plant susceptibility to pathogens [81]. A recent transcriptome analysis, showing lower susceptibility of Arabidopsis against the pathogenic bacterium Pseudomonas syringae (Pst) strain DC 3000, explained the effect by cold-modulation of salicylic acid biosynthesis and signalling [82]. Salicylic acid levels, that increase in the cold [83], can activate local as well as systemic resistance [84]. Only 2 h after the 10 h long cold-priming treatment, cold-modification of defence signalling resulted in stronger expression of PAL1 and PR2, and weaker induction of PR4 upon infiltration with Pseudomonas syringae DC3000 [82]. In our study, 5 days after 24 h cold priming, PAL1 was stronger induced by cold and by highlight in cold-primed plants. However, the transcript levels of the more specific salicylic acid regulated gene PR2 were lower (Fig. 6 Suppl. Table 2) after both stress treatments if the plants were cold-primed. On the contrary, the gene for the chitin binding protein PR4 and PCC1 were not down-regulated, but strongly up-regulated by cold, and down-regulated by light in cold-primed plants (Fig. 9). In our opinion, such specific regulation of defence related genes upon cold- and highlight triggering can, like differential regulation of ZAT10 (Figs. 1 and 2), only be explained by specific modulation of the gene responsiveness to the specific trigger. In other words, we conclude that the cold priming memory uncouples core stress signalling, which co-regulates the genes upon cold and light stress in naïve plants, and deploys its regulatory potential on stressor specific gene regulation.

Conclusions

Controlling the balance between defence and growth is crucial for plants in a changing environment in order to optimize their fitness [85]. Our study demonstrated, that cold-priming differentially modifies regulation of specific genes and even uncouples regulation of genes of the core environmental response cluster. Transcriptome wide analysis of the consequences of cold priming demonstrated that cold triggering supports expression of various genes involved in defence and protection on the cost of the expression of transport and growth related genes. On the contrary, light triggering preferentially activates genes involved in metabolism and development, but down-regulates genes involved in the defence response. The overall pattern is manifested in the inverse regulation of 29 genes. From this, we conclude that cold-priming modifies stress signalling by differentiating cold and light induced regulation.

Methods

Plant growth and stress treatments

Arabidopsis thaliana (var. Col-0) plants were grown for 28 days individually in round pots (6 cm diameter) in soil at 20 ± 2 °C at a day - night regime of 10 h light / 14 h dark and an illumination rate of 100-110 µmol quanta$*m^{-2}*s^{-1}$. For priming, a 24 h cold stimulus was imposed to half of the 4-week-old plants by transferring them 2.5 h after the onset of light to a 4 ± 2 °C cold chamber with the same aeration, illumination and air humidity setting as the 20 °C chamber (Fig. 10). Afterwards the primed plants were placed back to 20 °C. The general settings were identical to those used in the previous study [17], except

that the temperature sensor in the cold chamber was exchanged to one shortening the phase length in the cooling rhythm, which better stabilizes the day and night temperatures. 1/3 of the primed and naïve plants was harvested 5 days after the end of the priming stimulus 2.5 h after onset of light. The control plants (C plants) were kept all time at the 20 °C.

Cold triggering was started after a lag-phase of 5 days at 20 °C with 1/3 of the primed and 1/3 of the control plants by transfer of the plants to 4 °C (cold triggering) (Fig. 10). For highlight triggering, 1/3 of the primed and 1/3 of the naïve plants were exposed for 2 h to a photon flux density of 800 µmol quanta m^{-2}·s^{-1} 30 minutes after the onset of light using halogen lamps (R7-s 500 W, Emil Lux GmbH Wermelskirchen, Germany). The heat emission of the halogen lamps was filtered through a water layer and additionally controlled by moderate ventilation. The leaf temperature was monitored on the upper leaf surface with an infrared thermometer. For each of at least 3 biological replicates, all rosette leaves of 5-7 individual plants were combined 2 h after the treatment and frozen in liquid N$_2$.

RNA-Isolation and RNA library construction

For RNA isolation, the plant material was harvested and ground to a fine powder in liquid N$_2$. RNA was extracted from 100 mg plant material using the Gene Matrix Universal RNA Purification Kit (EURx, Gdansk, Poland) including the DNase treatment recommended by the supplier. The RNA was precipitated from the solution overnight at -20 °C by adding 0.1 volume of 3 M sodium acetate (pH 5.2) and 2.5-volume absolute ethanol. After dissolving the RNA in 50 µl RNAse-frei H$_2$O, the RNA integrity was assessed by electrophoresis on a 2 % (w/v) agarose gel supplemented with 1 % formaldehyde.

For RNA library construction, the mRNA was enriched using oligo (dT) magnetic beads and depleted from rRNA by DNA/rRNA hybridization according to standard procedures of the Beijing Genomics Institute (Beijing, China). Afterwards, the mRNA was transcribed into cDNA and the second DNA strand was generated with random N$_6$ primer. The double stranded cDNA was then 5´-end repaired, 3´-poly-A-tailed and ligated with an oligo-dT-adapter. The ligation product was amplified with specific adapter primers by PCR. Single-end sequencing on the Illumina High-Seq4000 platform of the Bejing Genomics Institute lead to an average of 24 million (± 160.000) reads with a read length of 50 base pairs per treatment (Suppl. Tab. 1).

Bioinformatic analysis

The reads obtained by RNASeq, that did not contain adaptor sequences and less than 10 % unclear bases (= clean reads), were aligned to the Arabidopsis reference genome (TAIR10) using Bowtie (version 2.1.0; [86]) and HISAT (2.1.0; [87]). The number of aligned reads were normalized for each gene to the transcript length and the total number of reads per treatment by using the RSEM software package. For the 2000

highest expressed genes a principal component analysis was performed by using edgeR [88] and limma libraries [89] in R 3.5.1. Correlation analysis of differential expressed genes for the heat map was performed by using the R function pheatmap (https://CRAN.R-project.org/package=pheatmap). Gene co-regulation was analyzed with InteractiVenn [90]. Additional specific analyses were performed with GENEMANIA [35], STRING v.11 [36] and on the eFP browser platform [67]. The functional characterization by gene ontologies of significantly expressed and differentially regulated genes ($\log_2$-change > 1 and a FPKM value > 5) was performed on the AgriGO 2.0 platform [66].

Reverse transcription and quantitative real time PCR

For the real-time PCR analysis, the mRNA was transcribed into cDNA using the High Capacity Reverse Transcription Kit (Applied Biosystems, Carlsbad, CA, U.S.A.). Oligo $dT_{16}V$ primer were annealed to the polyA-tails of the mRNA during 10 min incubation at 25 °C. After 2 h at 37 °C, the reaction was stopped by 5 min heat inactivation of the enzyme at 85 °C. Real-time qPCR analysis was performed on the CFX96 real-time system (Biorad, Hercules, CA, U.S.A.) with 50 ng template cDNA and 0.2 µl 10xSYBR Green (Sigma-Aldrich, Germany) in 20 µl as described previously [17], except that the transcript levels were only standardized on *YLS8* (*yellow leaf specific protein 8*; At5g08290), since other typically in qPCR analysis used reference genes responded either to the cold or to the light treatment. All primers, if applicable, were designed to span exon-intron border by the QUANTPRIME software [91]. Primer sequences are list in the supplements (Suppl. Tab. 8).

Chlorophyll-a fluorescence analysis

After 20 min dark acclimatization, the maximal chlorophyll-a fluorescence (F_V/F_M) was determined with a saturating light flash (> 1000 µmol quanta m^{-2} s^{-1}) in primed (P) and primed + triggered (PT) plants in an Imaging PAM IMAG-K4B (Walz, Effeltrich, Germany). The effective quantum yield of photosystem II (Φ_{PS-II} = ($F_{M'}$- F)/$F_{M'}$)), photochemical quenching (qP = ($F_{M'}$ - F)/($F_{M'}$ - $F_{0'}$)) and non-photochemical quenching (NPQ = (F_M /$F_{M'}$) − 1) were analyzed with saturating light flashes spaced by 20 s time gaps before and during illumination with 185 µmol quanta m^{-2} s^{-1} actinic light.

Statistical analyses

The significance of difference was evaluated with Student t-test (p < 0.05) if two data sets were compared. Larger data sets were analysed with the Tukey posthoc test (*p* < 0.05) using the R 3.5.1 software package.

Abbreviations

C: control plants; C5: control plants at the end of the lag-phase; CBF: C-repeat binding factor; Col-0: *Arabidopsis thaliana* var. Columbia-0; FPKM: fragments per kilobase of exon per million reads mapped; F_V/F_M: quantum yield of photosystem II in dark acclimated plants; HSF: heat-shock factor; ICE: inducer of CBF expression; MAP: mitogen activated protein; MPK: MAP protein kinase; P: primed plants; P5: primed plants at the end of the lag-phase; PT: primed and triggered plants; PT-C: cold-primed and cold-triggered plants; PT-L: cold-primed and light-triggered plants; RNASeq: RNA-Sequencing; ROS: reactive oxygen species; sAPX: stromal ascorbate peroxidase; T: plants which only faced the triggering stimulus; T-C: cold-triggered plants; T-L: light-triggered plants; tAPX: thylakoid-bound ascorbate peroxidase; qP: photosynthetic quench; qPCR: real-time quantitative polymerase chain reaction; GO: gene ontology; NPQ: non photochemical quench; UV: ultra violet; Φ_{PS-II}: quantum yield of photosystem II in the light. Abbreviations for additional gene names are defined in the text by the Arabidopsis gene code (At-number).

Declarations

Ethics approval and consent to participate: Not applicable.

Consent for publication: Not applicable.

Availability of data and material: All the datasets generated and analysed during the current study were uploaded as with the manuscript as additional files. Primary data will be available on PrimeDB (https://primedb.mpimp-golm.mpg.de) by the time of publication.

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Author contributions: JB and AB designed and performed the experiment for the RNASeq analysis, AB performed all further experiments, drafted the figures and part of the manuscript. MB supervised the project and finalized the manuscript. All authors have read and approved the manuscript.

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Source of plant: The *Arabidopsis thaliana* var. Col-0 seed material was propagated from a seed stock originally obtained from INRA (stock 186 AV). The seed stock is identical to the one used in our previous studies [10, 17, 30].

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Figures
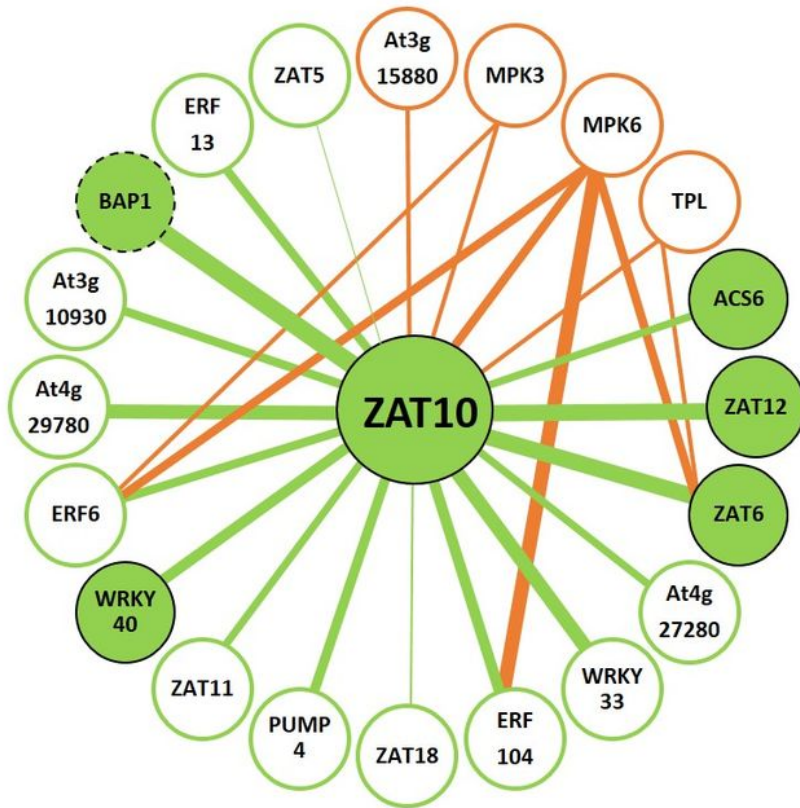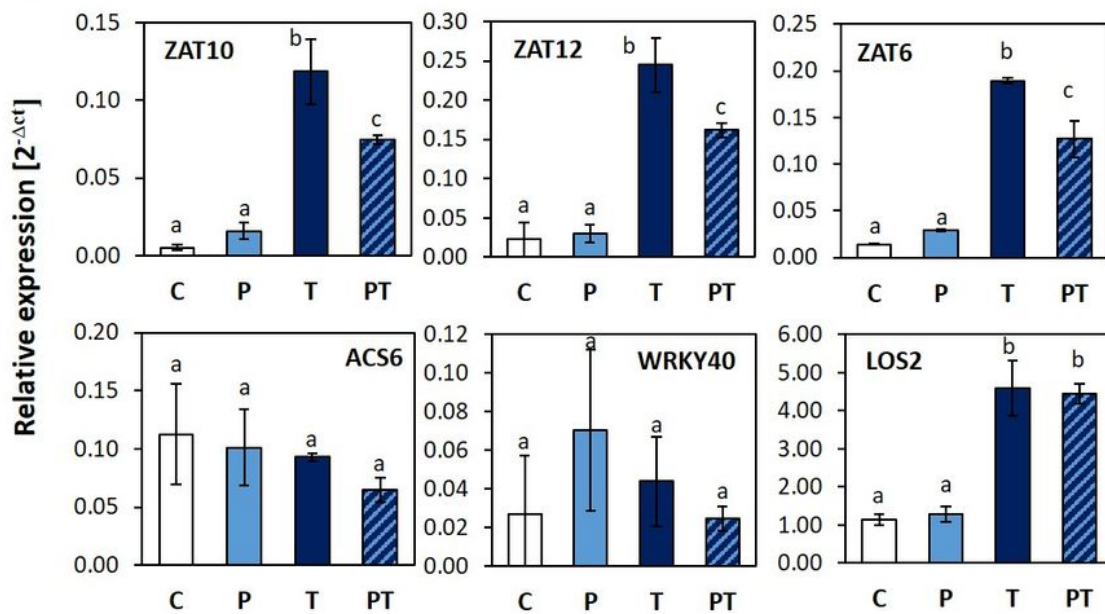
Figure 1

A: Genes frequently coregulated with ZAT10 (green) and proteins interacting with ZAT10 (orange) according to GENEMANIA and STRING. The thicker the connecting lines are drawn, the more studies reported co-regulation or interaction. Filled circles highlight the genes that were chosen for qPCR analysis. B: Effect of 24 h cold priming at 4 °C on cold-regulation of 4 genes co-regulated with ZAT10 in various studies and of the ZAT10 up-stream-regulator LOS2. Regulation of the relative transcript abundances

(standardized on YLS8; mean ± standard deviation) in control plants (C), only cold-primed (P), only cold-triggered (T) and cold-primed + cold-triggered plants (PT) immediately after triggering. Differ-ent letters label statistical significance of differences based on data obtained with 3 inde-pendently cultivated and treated biological replicates (Tukeys post hoc test; p < 0.05).
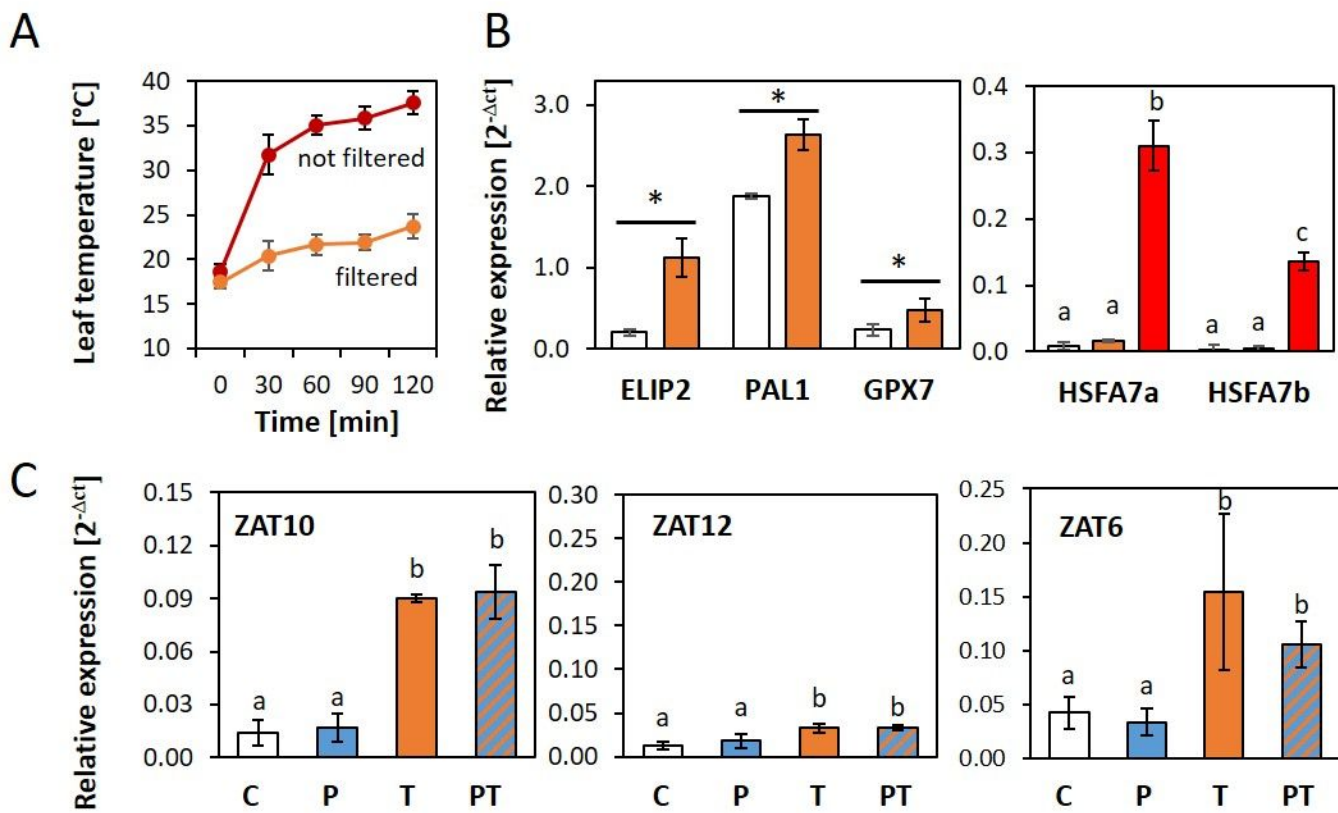
Figure 2

Effect of cold priming on light triggering. A: Leaf surface temperature in the heat-filtered (orange) and in the not heat-filtered illumination set-up (red). B: Relative tran-script abundance of light- and heat-responsive genes in control plants (white) and after 2 h heat-controlled illumination (orange) and not heat-controlled illumination (red). The transcript levels were standardized on the transcript levels of YLS8; Statistically significant dif-ferences in the relative transcript abundances are labelled with asterisks or different letters (n = 3 − 5; Tukeys post hoc test; p < 0.05). C: Effect of 24 h cold-priming at 4 °C on tran-script abundance regulation by a light stimulus. Regulation of the relative transcript abun-dances in control plants (C), only cold-primed (P), only light-triggered (T) and cold-primed + light-triggered plants (PT) immediately after light triggering. Different letters show statistical significance of differences in the relative transcript levels based on data obtained with 3-5 in-dependently cultivated and treated biological replicates (Tukeys post hoc test; p < 0.05).
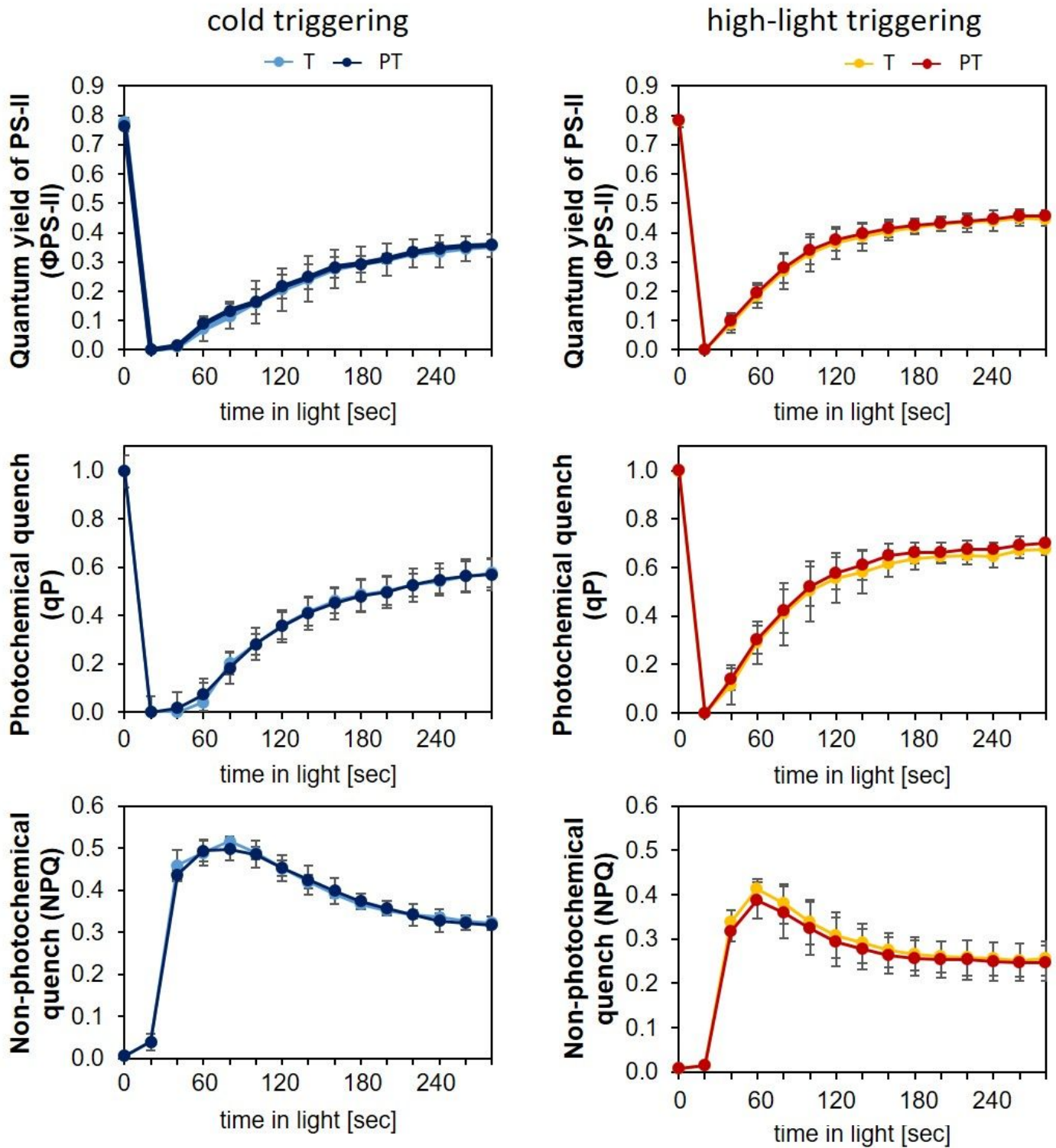
Figure 3

Effect of 24 h cold-priming at 4 °C on photosynthetic electron transport activity and regulation after cold (left) and light triggering (right). The means and standard deviations of the quantum yields of photosystem II (ΦPS-II), photochemical quenching (qP) and non-photochemical quenching (NPQ) as determined for each of the 4 biological replicates in paral-lel analysed only triggered (T) and primed + triggered plants (PT).
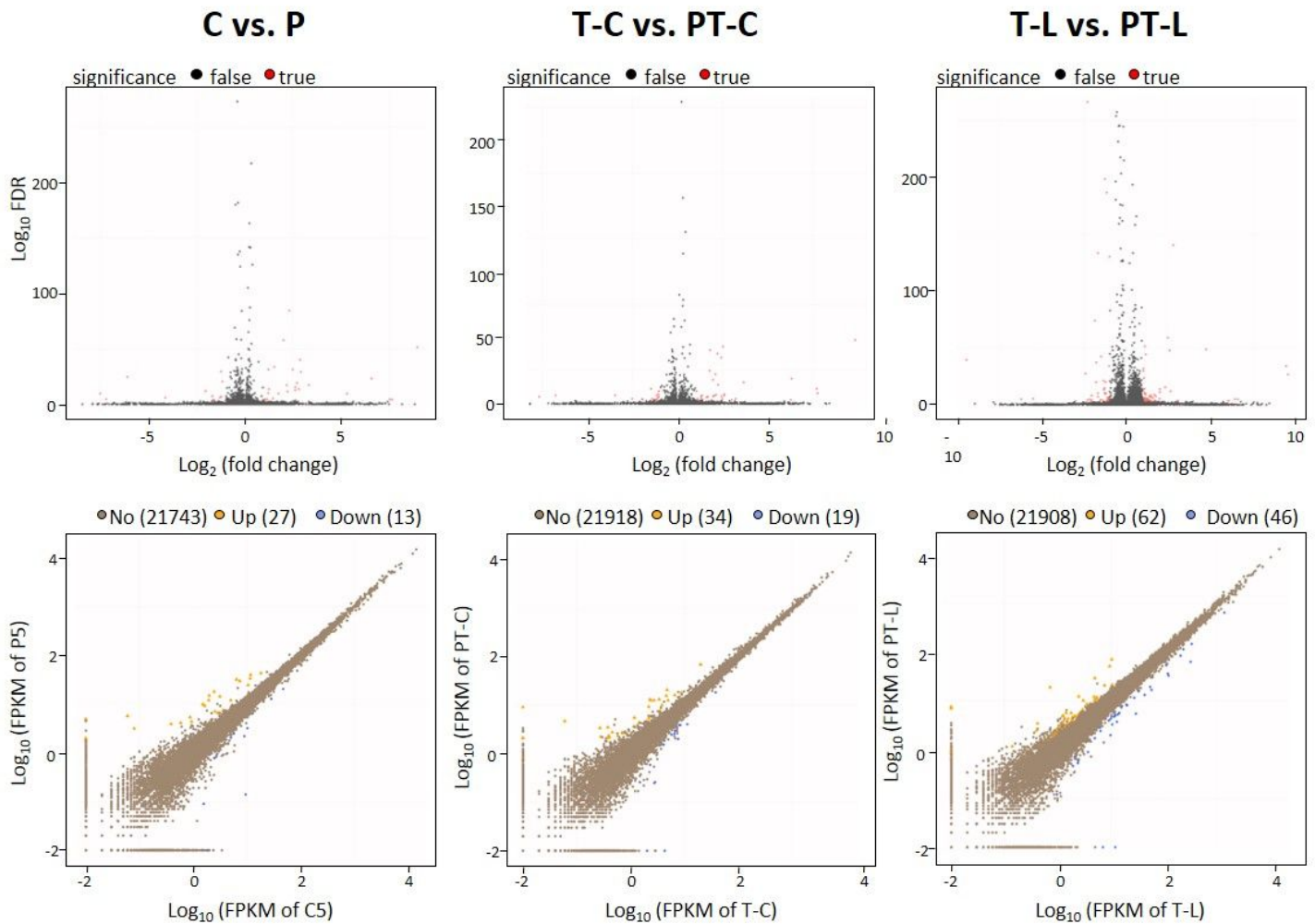
Figure 4

Statistical evaluation of priming-dependent regulation as obtained by RNASeq. Top: Volcano plots depicting genes with statistically significant regulation in red. Bottom: Com-parison of the regulation intensity in primed (y-axis) and non-primed plants (x-axes). Genes with an FPKM value > 0.001 and up-regulated at least with log2 (primed / unprimed) = I 1 I are labelled in yellow, down-regulated genes in blue. Data for non-triggered plants are shown on the left, for cold-triggered ones in the middle panels and for light-triggered plants to the right.
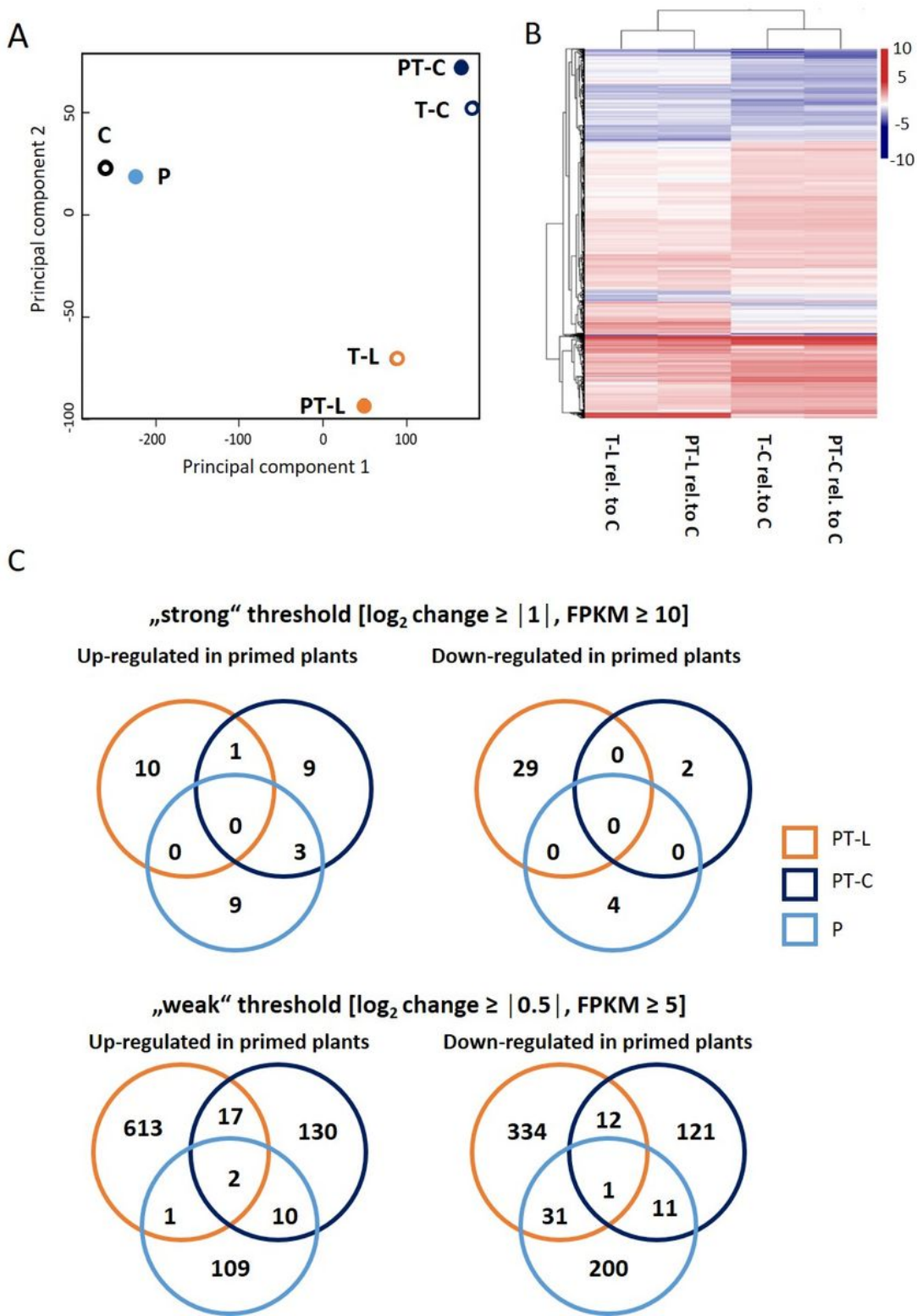
Figure 5

Transcript abundance regulation as observed by RNASeq. A: Principal component analysis separating the data sets of non-triggered (C, P), cold-triggered (T-C and PT-C) and light-triggered samples (T-L and PT-L) stronger according to the type of the triggering stimu-lus than to the priming effect (P or PT in comparison to C or T). B: Cluster analysis of tran-script abundance regulation in cold- or light-triggered, primed or unprimed samples relative to the transcript level in control plants. The heat map lists only

genes that were at least 2-fold stronger or less expressed in primed and / or triggered plants than in C-plants (FDR value < 0.001). 10-fold up-regulated transcripts are shown in dark red, 10-fold down-regulated tran-scripts are shown in blue. C: VENN-diagrams depicting the number of genes up- or down-regulated in a priming-dependent manner before (P) and after cold (PT-C) or light triggering (PT-L) at a strong threshold setting of log2 (primed/unprimed) ≥ I 1 I and FPKM ≥ 10 (top) or a weak threshold setting of log2 (primed/unprimed) ≥ I 0.5 I and FPKM ≥ 5 (bottom).
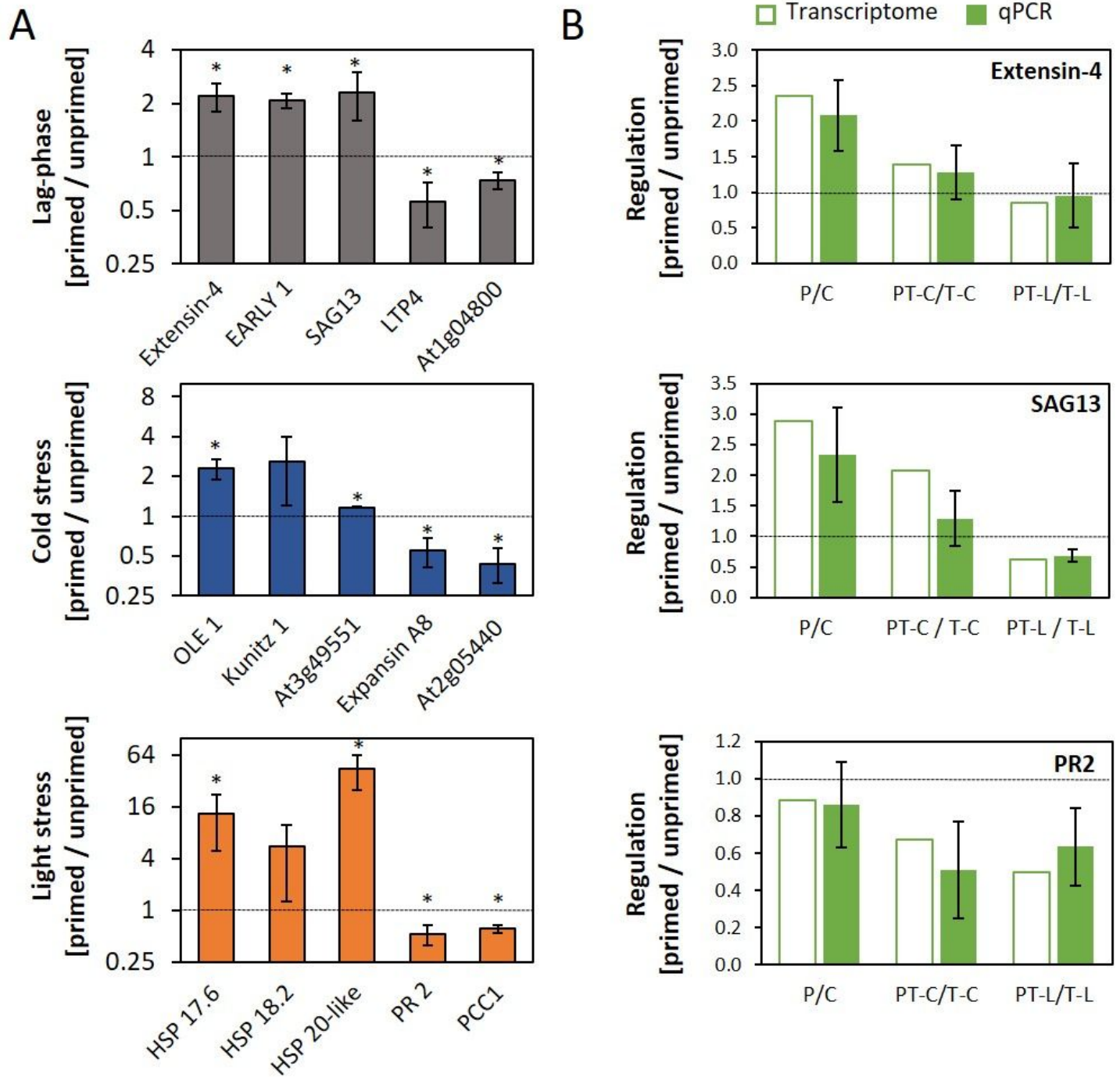
Figure 6

qPCR analysis of transcript abundance regulation. A: Consistency test on the regula-tion for genes showing strong regulation at the end of the lag-phase (top), after cold triggering (middle) or after light triggering. The transcript levels were quantified with gene specific primers and standardized on the transcript level of YLS8 in 3-5 independently culti-vated and treated biological replicates. Statistical significance of regulation (Student t-test; $p < 0.05$) is labelled with an asterisk. B: Testing for consistency of regulation of the RNASeq analysis throughout the experiment for three selected genes. For all samples, the transcript abundance ratio between primed and unprimed plants obtained by qPCR in four independently cultivated and treated biological replicates (green bars) was in the range of the ratio calculated based on the FPKM-values of the RNASeq analysis (white bars).
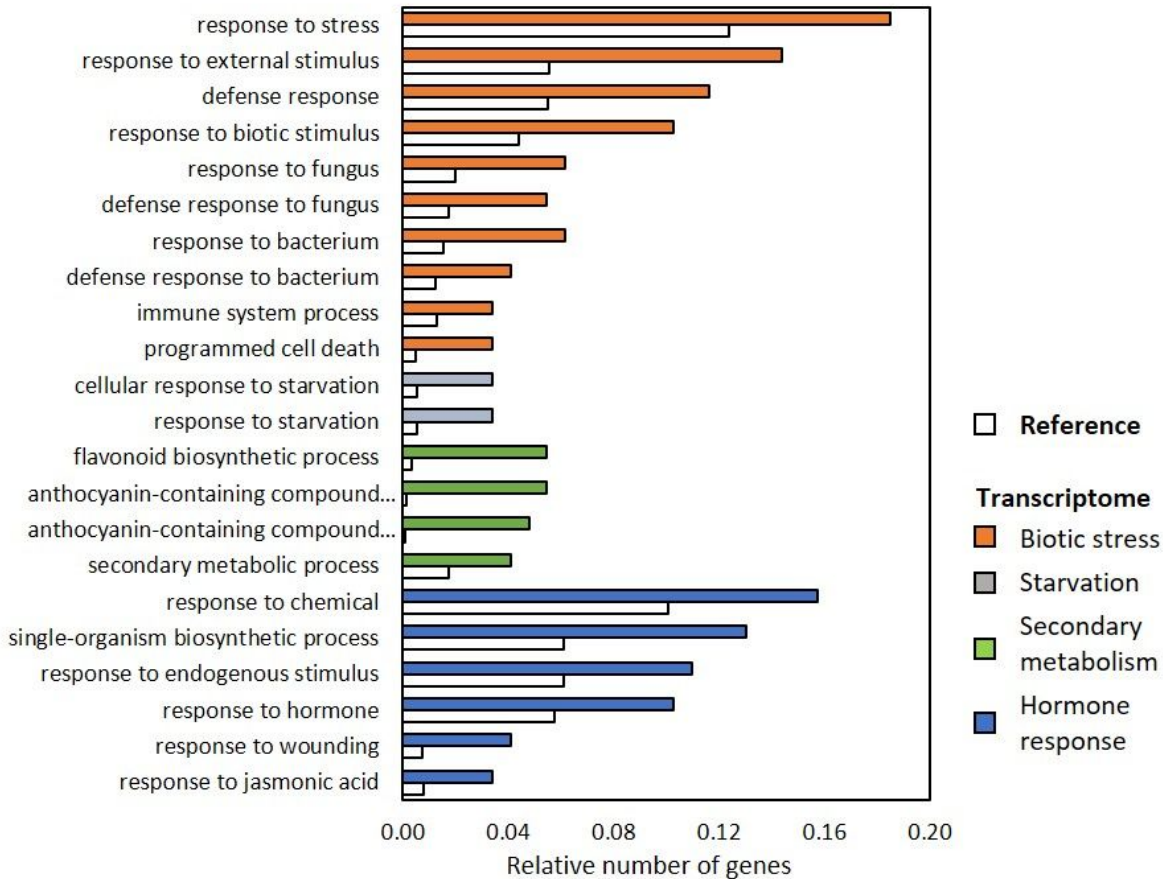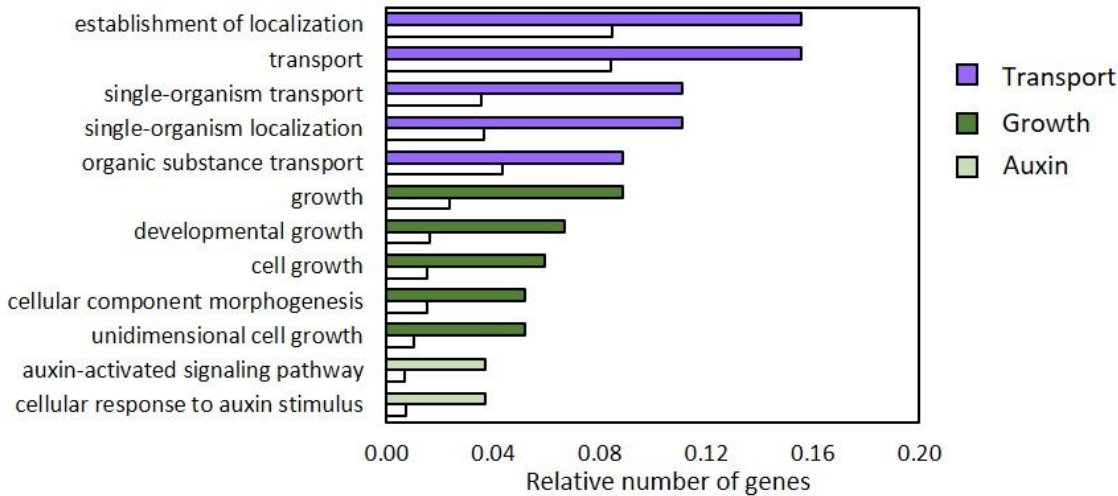
Figure 7

Functional characterization of genes regulated in priming-dependent manner after cold triggering. Enriched functional gene ontologies were identified with AgriGO using TAIR10 as background. The crude data including statistical information and the GO-term reference codes are summarized in Suppl. Tab. 5.
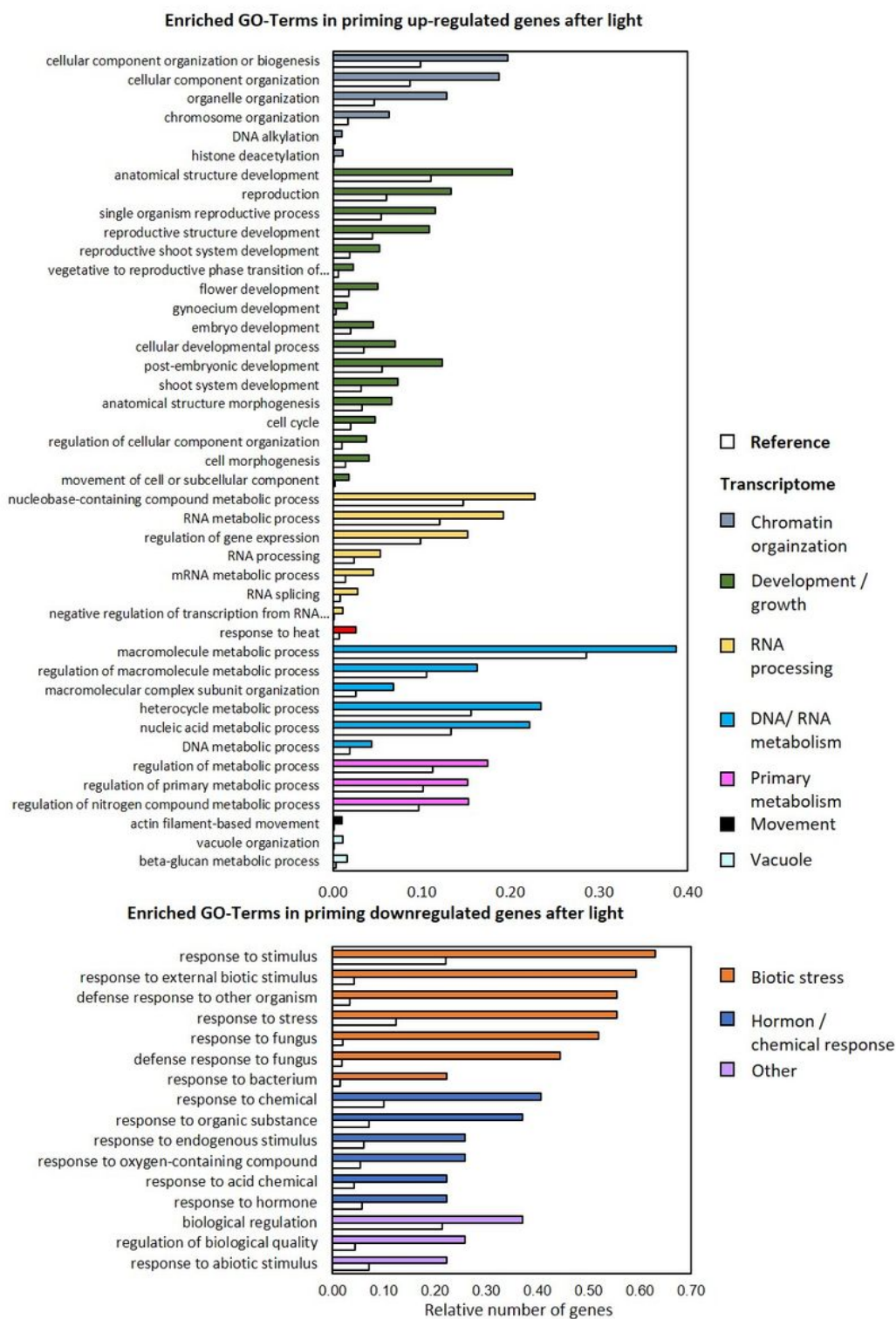
Figure 8

Functional characterization of genes regulated in priming-dependent manner after light triggering. Enriched functional gene ontologies were identified with AgriGO using TAIR10 as background. The crude data including statistical information and the GO-term reference codes are summarized in Suppl. Tab. 5.
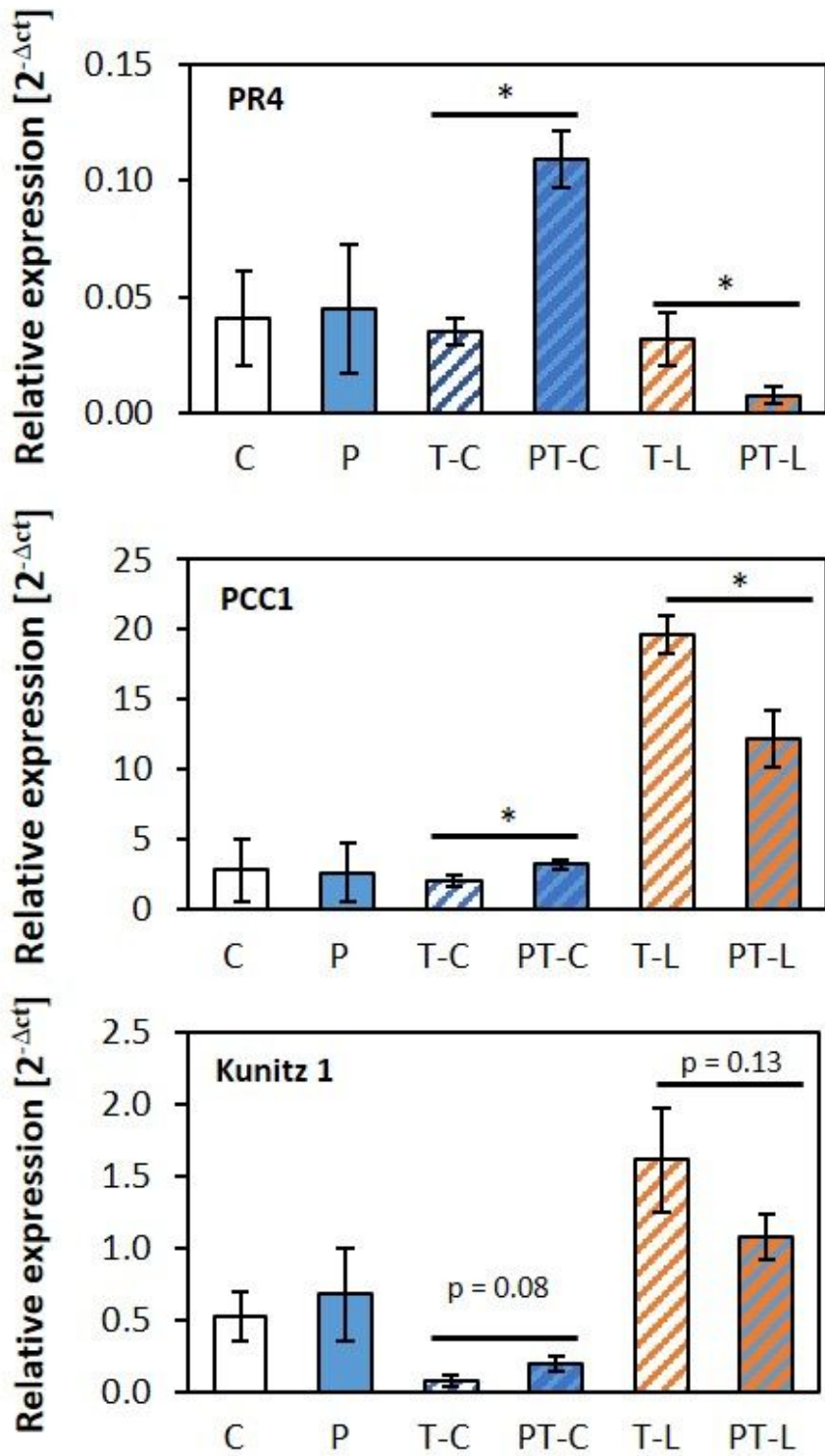
Figure 9

qPCR confirmation of inverse regulation of three pathogenesis-associated genes. The transcript levels were quantified with gene specific primers and standardized on the transcript level of YLS8 in 3 independently cultivated and treated biological replicates. Statistical signifi-cance of regulation (Student t-test; p<0.05) is labelled with an asterisk.
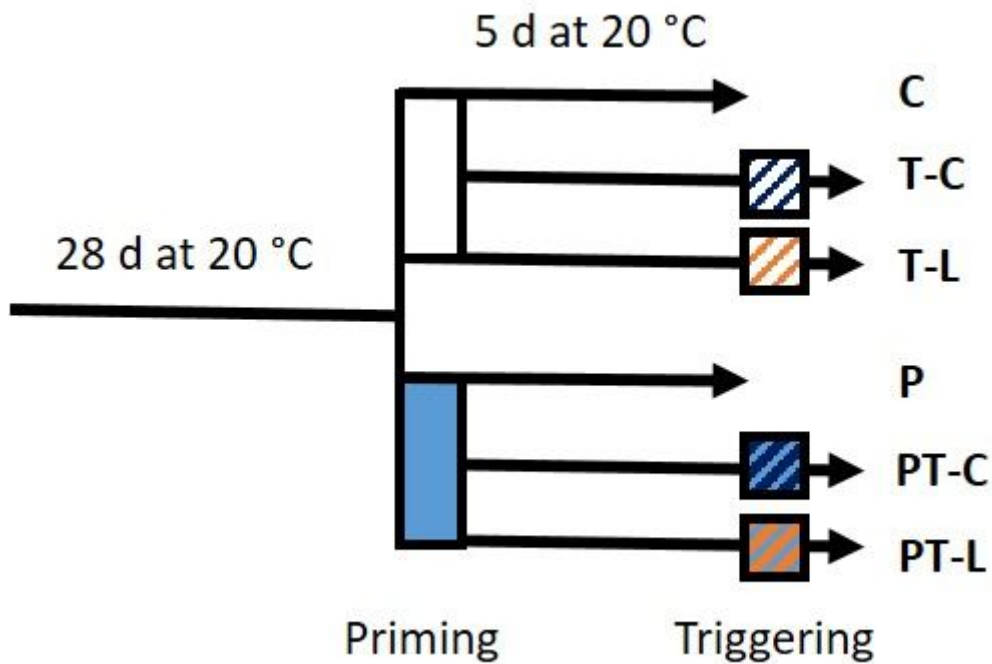
Figure 10

Experimental set-up. At an age of 28 days half of the plants were cold treated for 24 h at 4 °C (priming) and then retransferred to 20 °C and an illumination intensity of 100 - 110 µmol quanta m-2 s-1. 5 days later 1/3 of the plants of each set was cold-triggered for 24 h at 4 °C (T-C and PT-C), 1/3 light-triggered for 2 h at 800 µmol quanta m-2 s-1 (T-L and PT-L). The re-maining plants (control plants: C; only primed plants: P)) were harvested at the end of the lag-phase.