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Regulation of Arabidopsis gene expression by low fluence rate UV-B independently of UVR8 and stress signaling†

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UV-B exposure of plants regulates expression of numerous genes concerned with various responses. Sudden exposure of non-acclimated plants to high fluence rate, short wavelength UV-B induces expression *via* stress-related signaling pathways that are not specific to the UV-B stimulus, whereas low fluence rates of UV-B can regulate expression *via* the UV-B photoreceptor UV RESISTANCE LOCUS 8 (UVR8). However, there is little information about whether non-stressful, low fluence rate UV-B treatments can activate gene expression independently of UVR8. Here, transcriptomic analysis of wild-type and *uvr8* mutant Arabidopsis exposed to low fluence rate UV-B showed that numerous genes were regulated independently of UVR8. Moreover, nearly all of these genes were distinct to those induced by stress treatments. A small number of genes were expressed at all UV-B fluence rates employed and may be concerned with activation of eustress responses that facilitate acclimation to changing conditions. Expression of the gene encoding the transcription factor ARABIDOPSIS NAC DOMAIN CONTAINING PROTEIN 13 (ANAC13) was studied to characterise a low fluence rate, UVR8-independent response. ANAC13 is induced by as little as 0.1 $\mu\text{mol m}^{-2} \text{s}^{-1}$ UV-B and its regulation is independent of components of the canonical UVR8 signaling pathway COP1 and HY5/HYH. Furthermore, UV-B induced expression of ANAC13 is independent of the photoreceptors CRY1, CRY2, PHOT1 and PHOT2 and phytochromes A, B, D and E. ANAC13 expression is induced over a range of UV-B wavelengths at low doses, with maximum response at 310 nm. This study provides a basis for further investigation of UVR8 and stress independent, low fluence rate UV-B signaling pathway(s).

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Introduction

Ultraviolet-B radiation (UV-B; 280–315 nm) makes up less than 0.5% of the total solar spectrum but can have a major impact on all organisms. Plants, in particular, are dependent on sunlight as a source of energy and have evolved elaborate mechanisms to utilize solar energy in a positive manner and overcome the potentially negative effects of its most energetic wavelengths. Perception of specific wavelengths of light is carried out by photoreceptors, and the first UV-B photo-

receptor identified in plants was the UV RESISTANCE LOCUS 8 (UVR8) protein.^{1–3} Since its discovery, a growing body of research has revealed UVR8's function, and its relationship with other abiotic and biotic pathways.^{4–10}

Additionally, progress has been made in determining the crystal structure and mechanistic action of UVR8 in UV-B perception, including its intrinsic tryptophan-based chromophore.^{2,3,11–13} In essence, UVR8 senses UV-B photons *via* specific tryptophan residues within its structure, and this brings about monomerization of the homodimer. UVR8 monomers bind to CONSTITUTIVELY PHOTOMORPHOGENIC1 (COP1), ultimately triggering a network of transcriptional responses.^{7,10} The E3 ubiquitin ligase component COP1 is a master regulator of numerous proteins involved in photomorphogenesis and acts as a negative regulator in darkness. However, COP1 has a positive role in the UV-B signaling pathway and is required for expression of UVR8-regulated genes.^{14–17} ELONGATED HYPOCOTYL5 (HY5) and HY5 HOMOLOG (HYH) are both transcription factors that act downstream of UVR8 and COP1 to regulate expression of many UVR8 target genes.^{15,18} In addition,

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UVR8 has recently been found to interact directly with specific transcription factors to mediate responses to UV-B.^{19,20}

Overall, the UVR8 pathway gives the plant protection against potentially harmful UV-B wavelengths and initiates other processes, including morphogenic and physiological responses, entrainment of the circadian clock and protection against specific pathogens.^{5,9,10,15,18,21–23} Nevertheless, it has become apparent that other pathways mediate responses to UV-B that are independent of UVR8. Several studies, mainly involving microarrays, have shown that numerous genes are regulated by UV-B exposure, in some cases through the activation of stress pathways by relatively short wavelengths and high fluence rates of UV-B.^{14,18,24–27} However, UVR8-independent UV-B pathways remain poorly characterised, and the possible existence of additional UV-B photoreceptor(s) cannot be excluded.²⁸

The aim of this study was, firstly, to determine whether low, photomorphogenic exposures to UV-B initiate gene expression responses independently of UVR8 and to identify genes regulated by low fluence UV-B via UVR8-independent pathway(s), using transcriptomic analysis in *Arabidopsis thaliana*. Secondly, the UVR8-independent pathway regulating expression of a gene selected from the transcriptomic data was characterised. This gene encodes NAC DOMAIN CONTAINING PROTEIN 13 (ANAC13), a putative transcription factor.

Materials and methods

Plant material and light treatments

Seeds of wild-type *Arabidopsis thaliana* ecotypes Landsberg erecta (*Ler*), Columbia (Col-0), Wassilewskija (*Ws*) and the mutants *uvr8-1*,⁴ *uvr8-6*,¹⁵ *cry1cry2uvr8-2*,²⁹ *phyAphyBphyDphyE*,³⁰ *cry1cry2*,³¹ *phot1-5phot2-1*,³² *hy5-ks50hyh*³³ and *cop1-4*³⁴ were sown on compost, stratified at 4 °C for 48 h, and then grown in continuous white light of 20 μmol m⁻² s⁻¹ (warm white fluorescent tubes L36W/30, Osram) at 20 °C for 21 days. Plants were exposed to UV-B using either a broadband UVB-313 fluorescent tube (Q-Panel Co., USA) covered by cellulose acetate film (FLM400110/2925, West Design Products) or a narrowband tube (Philips TL20 W/01RS, λ_{max} 312 nm) at the fluence rates indicated in the figure legends. Control plants were kept in 20 μmol m⁻² s⁻¹ white light. The spectra of the UV-B sources, measured with a Macam Photometrics spectroradiometer model SR9910 are shown in Fig. S1.† Assays of expression at different UV wavelengths were undertaken using a pulsed Opolette 355 + UV II tuneable laser (Opotek Inc., USA) with a half bandwidth of 0.4 nm as described in ref. 35. Following UV exposure with the tuneable laser, plant material was left in darkness for 1 hour to allow transcripts to accumulate and then harvested and snap frozen in liquid nitrogen. All the data presented were obtained from 3 independent experiments with different sets of plants.

Transcript measurements

RNA extraction was performed using the RNeasy Plant Mini Kit (Qiagen) according to the manufacturer's instructions. cDNA

synthesis was performed as described in ref. 18. Quantitative PCR was performed using the MX4000 Stratagene real-time PCR system and a Brilliant III SYBR Green qPCR kit (Stratagene) following the manufacturer's instructions. A master mix was prepared of 1× SYBR Green Master Mix (Stratagene), 0.2 M of each primer, and appropriate volumes of cDNA and DEPC treated water. The PCR conditions were as follows: 3 min at 95 °C, 40 cycles of 10 s at 95 °C, 20 s at 60 °C, followed by a 60 to 95 °C dissociation protocol. Stratagene MX software was used to automatically calculate the cycle threshold (Ct) value for each reaction. Each reaction was performed in duplicate in three independent experiments. As a control for variation in RNA quantification, reverse transcription efficiency, and template preparation, the expression of *HY5*, *CHS* and *ANAC13* (At1g32870) transcripts was normalized to the mean of either *18S rRNA* or *ACTIN2* (as indicated in the figure legends). The relative levels of transcripts were calculated following the ΔΔCt method. The primers used for *HY5* were: 5'-CTGAAGAGGTTGTTGAGGAAC-3' and 5'-AGCATCTGGTTCTCGTCTGAAGA-3' (or 5'-GGCTGAAGAGGTTGTTGAGG-3' and 5'-CAGCATTAGAACCACCA CCA-3' for the data in Fig. 4); for *ANAC13*: 5'-AAGAAAGATCCGTCGGAAAA-3' and 5'-CCAATAGCCACGTTTCAGTAGC-3'; for *CHS*: 5'-CTACTTCCGCATCACCAACA-3' and 5'-TTAGGGACTTCGACCACCAC-3'; for *ACTIN2*: 5'-ACTAAAACGCAAAACGAAAGCGGTT-3' and 5'-CTAAGCTCTCAAGATCAAAGGCTTA-3'; and for *18S rRNA*: 5'-AAACGGCTACCACATCCAAG-3' and 5'-CCTC CAATGGATCCTCGTTA-3'.

For semi-quantitative PCR, to the appropriate volume of cDNA, a master mix was added consisting of 1× PCR Buffer (New England Biolabs), 0.2 mM dNTPs, 0.5 μM of each primer, 0.625 Units of Taq DNA Polymerase (New England Biolabs) and RNase free water to a final volume of 25 μl. The PCR cycle used was (step 1) incubation for 2 min 30 s at 95 °C, (step 2) a further 45 s at 95 °C, (step 3) incubation at 55–59 °C for 1 min, (step 4) elongation at 72 °C for 1 min and a final step of a further elongation at 72 °C for 5 min (step 5). Steps 2–4 were repeated 24–28 times depending on the primers used; the cycle number was selected to ensure that PCR product was quantitatively related to transcript level over a linear range of amplification. The primers used were as follows, *ACTIN2*: 5'-CTTACAATTTCCCGCTCTGC-3' and 5'-GTTGGGATGAACCAGAAGGA-3'; *ANAC13*: 5'-AGCTCGTTGTTTCGGCTAGT-3' and 5'-TCAGGAGACCAGAACC ATCC-3'; *CHS*: 5'-ATCTTTGAGATGGTGCTGC-3' and 5'-CGTCT AGTATGAAGAGAACG-3'; *At5g51440*: 5'-GCGGAAATGAAGAATG GTGT-3' and 5'-AAGTCAAAATCCCGAACACA-3'; *At2g41730*: 5'-GTCACCAAGGCATCGTAAGG-3' and 5'-ACTTGATAGCTGGCGACA CG-3'; *At3g22060*: 5'-ACAATGCGTTTCTCTTCCACA-3' and 5'-GCGAGTTGAATGTTGATGGAT-3'.

Transcriptomics

Three independent RNA samples were extracted as described above and transcriptomic analysis was undertaken by the Sir Henry Wellcome Functional Genomics Facility (SHWFGF, University of Glasgow). RNA quality was checked using an Agilent RNA BioAnalyzer 2100 (Austin, TX). The samples were then reverse transcribed and biotinylated cRNA hybridised to



Affymetrix Arabidopsis ATH1 GeneChips (High Wycombe, UK) as per the manufacturer's protocols. Subsequent washes and staining were performed using a Fluidics Station 400 (Affymetrix) according to manufacturer's instructions. The chips were scanned on a GeneArray Scanner 2500 (Affymetrix) and analysed by the SHWFGF using FUNALYSE version 2.0 (University of Glasgow, UK). Analysis involved normalization using Robust Multi Chip Average³⁶ and differentially expressed genes were determined using the Rank Products method.³⁷

Venn diagrams were constructed using GeneVenn software (<http://mcbc.usm.edu/genevenn/genevenn.htm>) and jvenn.³⁸ Microarray comparisons across studies were performed by incorporating all data into a FileMaker Pro 10® (Filemaker Inc., CA, USA) database. Data was then exported to Excel® (Microsoft, USA) and sorted according to frequency of occurrence for each gene. Gene Ontology (GO) term (biological processes) enrichment was performed using the R package clusterProfiler 3.10.1.³⁹ For the Venn analysis in Fig. 2, the gene lists were from Kilian *et al.* (2007)²⁶ Supplemental Table 2 (genes induced by multiple stresses) and Supplemental Tables 3 and 4 (genes induced by UV-B stress at different time points).

Results

Transcriptome analysis identifies genes regulated by low UV-B fluence rates independently of UVR8 and stress signaling

Several studies have been carried out previously with Arabidopsis to unravel UV-B regulated and UVR8 dependent pathways at the transcriptional level using microarrays and other techniques.^{6,15,18,24–26,40} Our initial research¹⁸ identified 639 genes induced in response to UV-B, of which 72 are dependent on UVR8. The implication is that many genes are induced by UVR8-independent pathway(s). However, the UV-B treatment used by Brown *et al.* (2005)¹⁸ (3 $\mu\text{mol m}^{-2} \text{s}^{-1}$ broadband UV-B for 4 hours) is sufficient to trigger several stress related genes in non-acclimated plants,²⁵ which could explain the preponderance of UVR8-independent genes in the microarray data. Hence, in this study we wanted to determine whether UVR8-independent pathways regulate gene expression at low fluence rates of UV-B. We therefore gave WT plants UV-B treatments (0.3 and 1 $\mu\text{mol m}^{-2} \text{s}^{-1}$ for 4 hours) that have been shown to initiate photomorphogenic gene expression rather than non-specific stress responses.²⁵ Microarray analyses were carried out similarly to Brown *et al.* (2005),¹⁸ comparing transcript profiles for each UV-B treatment to a non-UV-B-treated control. The gene lists are shown in Tables S1 (0.3 $\mu\text{mol m}^{-2} \text{s}^{-1}$) and S2 (1 $\mu\text{mol m}^{-2} \text{s}^{-1}$).† The data are presented with a false discovery rate (FDR), which represents the probability that a given gene in a list is a false positive. We use 5% FDR to facilitate comparison with previous published data, but are aware from transcript measurements that at this % FDR some genes in the lists may be only weakly differential. Comparisons at 2% or 1% FDR reduce the likelihood of false positives.

Fig. 1 shows Venn diagrams constructed from the microarray data at different false discovery rates (FDR) and depicts the

overlap between the microarrays. Intuitively one might expect that lower fluence rates of UV-B would up-regulate many fewer genes than higher fluence rates because the latter are likely to induce more stress related, non-specific genes. Conversely, this was not the case. At a FDR of 5%, 549 and 572 genes were up-regulated in the 0.3 and 1 $\mu\text{mol m}^{-2} \text{s}^{-1}$ microarrays respectively, which is comparable to the 639 genes induced at 3 $\mu\text{mol m}^{-2} \text{s}^{-1}$ in ref. 18. A similar outcome was obtained at FDRs of 2% and 1% (Fig. 1).

There was a degree of overlap between the genes induced by each fluence rate, with 169 genes (at 5% FDR) in common between the 3 treatments. However, there was a greater degree of overlap when comparing 0.3 and 1 $\mu\text{mol m}^{-2} \text{s}^{-1}$ (approximately 65% genes in common) and 1 and 3 $\mu\text{mol m}^{-2} \text{s}^{-1}$ (approximately 55% genes in common) than when comparing 0.3 and 3 $\mu\text{mol m}^{-2} \text{s}^{-1}$ (approximately 30% genes in common), which highlights the differences in signaling processes regulating gene expression at the lowest and highest fluence rates. The intermediate 1 $\mu\text{mol m}^{-2} \text{s}^{-1}$ UV-B treatment initiates expression of both 'low' and 'high' fluence rate genes and consequently less than 10% of the genes induced are specific to that treatment, whereas 33% and 49% of genes are specific to the 0.3 and 3 $\mu\text{mol m}^{-2} \text{s}^{-1}$ treatments, respectively.

The requirement for UVR8 for expression of specific genes was determined from previous analyses using a *uvr8* mutant.¹⁸ Most of the 72 UVR8-dependent genes induced by 3 $\mu\text{mol m}^{-2} \text{s}^{-1}$ UV-B (at 5% FDR¹⁸) were also detected in the present microarrays (64 and 61 genes for 0.3 and 1 $\mu\text{mol m}^{-2} \text{s}^{-1}$ respectively). Interestingly, UVR8-regulated genes show a greater representation at lower fluence rates than at 3 $\mu\text{mol m}^{-2} \text{s}^{-1}$ when the stringency is increased to 1% FDR. At 0.3 $\mu\text{mol m}^{-2} \text{s}^{-1}$ 28% of genes are UVR8-dependent at 1% FDR, compared to 11% at 5% FDR. Conversely, at 3 $\mu\text{mol m}^{-2} \text{s}^{-1}$ only 6% of genes are UVR8-dependent at 1% FDR, in contrast to 11% at 5% FDR. The above analysis indicates that a substantial number of genes induced by low fluence rates of UV-B are not regulated by UVR8. To extend the analysis, we compared the genes induced at low fluence rate UV-B with the set of UVR8 regulated genes identified by Favory *et al.* (2009),¹⁵ who listed a total of over 700 genes potentially regulated by UVR8 in the different UV-B exposures in their study. As shown in Fig. 2A, there is extensive overlap between the sets of UVR8 regulated genes identified in ref. 18 and 15. In addition, the Favory¹⁵ list increases the number of genes identified as UVR8 regulated at low fluence rate UV-B. Nevertheless, 44% of the 306 genes (at 2% FDR) induced by 0.3 $\mu\text{mol m}^{-2} \text{s}^{-1}$ UV-B are not dependent on UVR8 (Fig. 2A).

Since UVR8 independent UV-B signaling pathways are known to overlap with stress-related signaling pathways, at least at relatively high UV-B fluence rates,^{41,42} we examined whether any of the UVR8-independent genes induced by 0.3 $\mu\text{mol m}^{-2} \text{s}^{-1}$ UV-B are known targets of stress signaling pathways. We tested whether any of these genes were induced by UV-B stress in the microarray study of Kilian *et al.* (2007).²⁶ Of the 429 UV-B stress-induced genes,²⁶ only 11 were among the UVR8-independent genes induced by 0.3 $\mu\text{mol m}^{-2} \text{s}^{-1}$



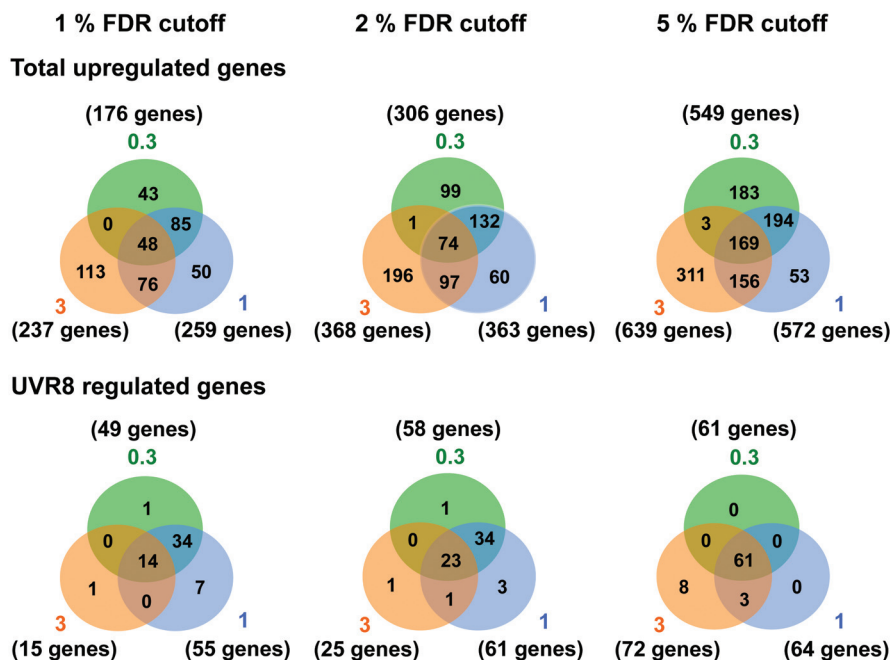


Fig. 1 Venn diagrams depicting the overlap in gene expression at different fluence rates of UV-B. Three-weeks-old *Arabidopsis* plants grown in a low fluence rate of white light ($20 \mu\text{mol m}^{-2} \text{s}^{-1}$) were treated with either 4 hours of 0.3 , 1 or $3 \mu\text{mol m}^{-2} \text{s}^{-1}$ broadband UV-B or were left in low white light as a control. This corresponds to a plant weighted UV^{51} of 0.08 , 0.27 , and 0.82 kJ m^{-2} , respectively, which compares with $4.8 \text{ kJ m}^{-2} \text{ day}^{-1}$ in Lund, Sweden, at midsummer under a cloudless sky.⁵¹ The numbers of genes that showed an increase in transcript level were calculated for each of three False Discovery Rates (FDR). Gene lists were then compared to those published by Brown *et al.* (2005)¹⁸ to determine overlap and dependence on UVR8. Numbers in orange circles denote transcripts identified in the Brown *et al.* (2005)¹⁸ $3 \mu\text{mol m}^{-2} \text{s}^{-1}$ UV-B microarray, those in blue and green are those found in this study to be induced by 1 and $0.3 \mu\text{mol m}^{-2} \text{s}^{-1}$ UV-B respectively.

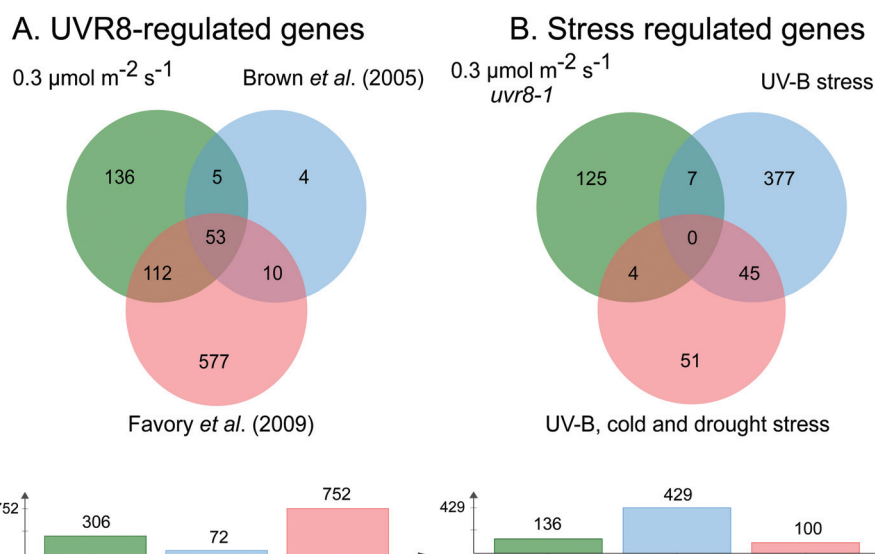


Fig. 2 Venn diagrams showing genes induced by low fluence rate UV-B independently of UVR8 and stress signalling pathways. (A) Genes induced by exposure of *Arabidopsis* to $0.3 \mu\text{mol m}^{-2} \text{s}^{-1}$ broadband UV-B for 4 hours (corresponding to a plant weighted UV^{51} of 0.08 kJ m^{-2}), cut at 2% FDR (as shown in Fig. 1) compared to genes shown to be regulated by UVR8 in microarray analyses of Brown *et al.* (2005)¹⁸ and Favory *et al.* (2009).¹⁵ (B) UVR8-independent genes identified in A compared to genes shown to be induced either by UV-B stress or in common by UV-B, cold and drought stresses in the microarray study of Kilian *et al.* (2007).²⁶ Venn diagrams are shown above with numbers of genes in each set below. The diagrams were constructed using jvenn.³⁸



UV-B (Fig. 2B). Four of these 11 genes were among a set of 100 genes induced in common by UV-B, cold and drought stresses²⁶ (Fig. 2B). Interestingly, 9 of the above 11 genes are among 52 UVR8-independent genes expressed at all 3 fluence rates employed in the present study, demonstrating that expression of this small set of putative 'stress-related' genes is not confined to stress conditions; it may be that these genes are expressed as a result of a change in environment rather than 'damage stress' *per se*.

According to the above analysis, 41% of the genes induced at 0.3 $\mu\text{mol m}^{-2} \text{s}^{-1}$ UV-B (125 genes in total at 2% FDR; listed in Table S3†) are not known targets of either UVR8 or stress signaling pathways, indicating that a distinct type of UV-B signaling operates at low UV-B fluence rates. To gain insights into the potential functions of genes induced *via* this UVR8 and stress independent pathway(s) we used GO enrichment analysis (Fig. S2†). This revealed that a substantial number of the genes are concerned with carbohydrate metabolism.

***ANAC13* expression is induced by low fluence rate UV-B independently of UVR8**

We examined a number of genes identified as UVR8-independent in the microarray analysis to test the validity of the results with respect to UV-B fluence rate: *ANAC13*, *At5g51440* (a putative HSP20-like chaperone), *At3g22060* (a gene expressed in response to abscisic acid) and *At2g41730* (a gene expressed in rosette leaves and responsive to high concentrations of Boron). As shown in Fig. 3, semi-quantitative RT-PCR confirmed the microarray result in that all four genes were induced by low fluence rate UV-B: *ANAC13*, *At2g41730*, and *At3g22060* were induced at 0.1 $\mu\text{mol m}^{-2} \text{s}^{-1}$, similar to *CHS*, whereas *At5g51440* was induced at 0.2 $\mu\text{mol m}^{-2} \text{s}^{-1}$. To further explore the potential low fluence rate UV-B, UVR8-independent pathway we decided to focus on one gene and selected *ANAC13*. Quantitative PCR analysis of the kinetics of *ANAC13* expression (Fig. 4) revealed that transcripts peak at around 6–12 hours after the start of UV-B exposure. This increase in expression is slower than that of the UVR8 dependent transcription factor *HY5* and *CHS*, which is downstream of the UVR8/*HY5* UV-B pathway. In addition, we wanted to test whether *ANAC13* is in fact UVR8-independent, as suggested by the microarray results and previously reported by Safrany *et al.* (2008).²⁷ Fig. 5 confirms that *ANAC13* is induced at low fluence rates (0.5 $\mu\text{mol m}^{-2} \text{s}^{-1}$) and, unlike *HY5*, this induction still occurs in the *uvr8* mutant. Fig. 5 further shows that *HY5* and *ANAC13* transcript levels are similar after 2 hours UV-B in WT, but at 6 hours *ANAC13* expression continues to rise with *HY5* starting to decline, consistent with the kinetics in Fig. 4. Overall, these results confirm that *ANAC13* expression is UVR8-independent and induced at low UV-B doses.

ANAC13* UV-B induction is independent of COP1 and *HY5*/*HYH

To further examine whether *ANAC13* is induced by the low fluence rate UV-B, UVR8-independent pathway, we tested *ANAC13* expression in *cop1* and *hy5hyh* mutants. Since both

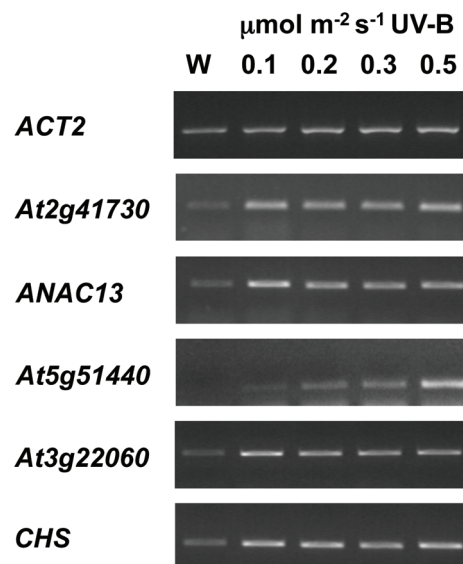


Fig. 3 Expression of UVR8-independent genes under very low fluence rates of UV-B. Three-weeks-old wild type plants grown in a low fluence rate of white light ($20 \mu\text{mol m}^{-2} \text{s}^{-1}$) were treated for either 4 hours with 0.1, 0.2, 0.3, or 0.5 $\mu\text{mol m}^{-2} \text{s}^{-1}$ broadband UV-B (corresponding to plant weighted UV⁵¹ of 0.03, 0.05, 0.08, and 0.14 kJ m^{-2}) or were left in low white light (W) as a control. Transcript levels were assayed using RT-PCR and compared with control *ACT2* transcripts.

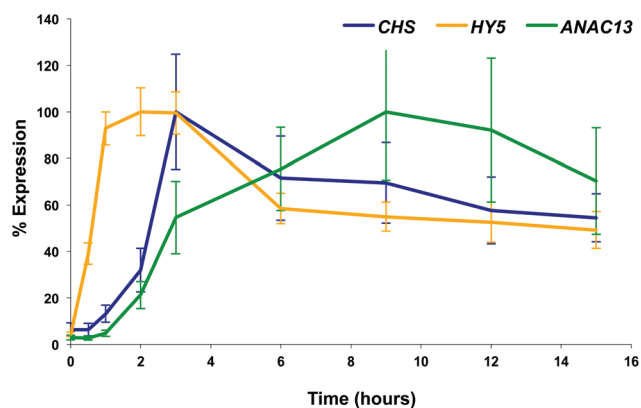


Fig. 4 Time course of expression of UV-B induced genes. Three-weeks-old wild type plants grown under $20 \mu\text{mol m}^{-2} \text{s}^{-1}$ white light were treated with $3 \mu\text{mol m}^{-2} \text{s}^{-1}$ broadband UV-B for the times shown before tissue was harvested and RNA extracted. Relative transcript levels (adjusted to *ACT2* transcript levels) were determined using qPCR and are shown as percentage of maximal expression for each gene. Bars represent S.E., $n = 6$.

COP1 and the transcription factors *HY5*/*HYH* are associated with UVR8 signaling, one would expect *ANAC13* expression to be unaffected in such mutants. In agreement with this notion *ANAC13* expression was induced by UV-B in *cop1-4* (Fig. 6A), as reported previously.²⁷ Furthermore, expression was similar in *hy5hyh* and WT plants (Fig. 6B), indicating that the UVR8 independent pathway regulating *ANAC13* expression is independent of *HY5*/*HYH* as well as *COP1*.



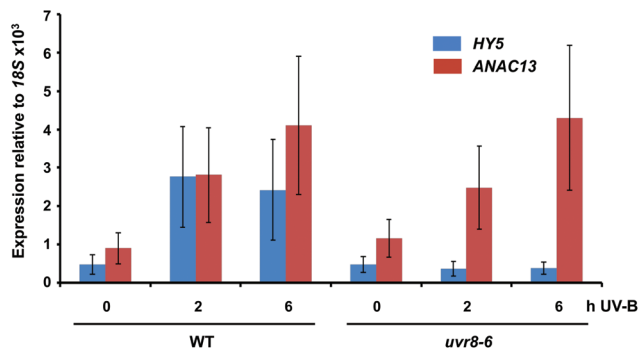


Fig. 5 *ANAC13* is UV-B induced and UVR8 independent. Quantitative RT-PCR assays of *HY5* and *ANAC13* transcripts, normalized to control 18S transcript levels, in wild-type Col-0 and *uvr8-6*. Plants were exposed (+) or not (–) to 0.5 $\mu\text{mol m}^{-2} \text{s}^{-1}$ of narrowband UV-B for 2 h or 6 h. The plant weighted UV⁵¹ was 0.005 kJ m^{-2} for the 2 h exposure and 0.015 kJ m^{-2} for the 6 h exposure, which compares with 4.8 $\text{kJ m}^{-2} \text{day}^{-1}$ in Lund, Sweden, at midsummer with a cloudless sky.⁵¹ Data are means of three experiments \pm S.E.

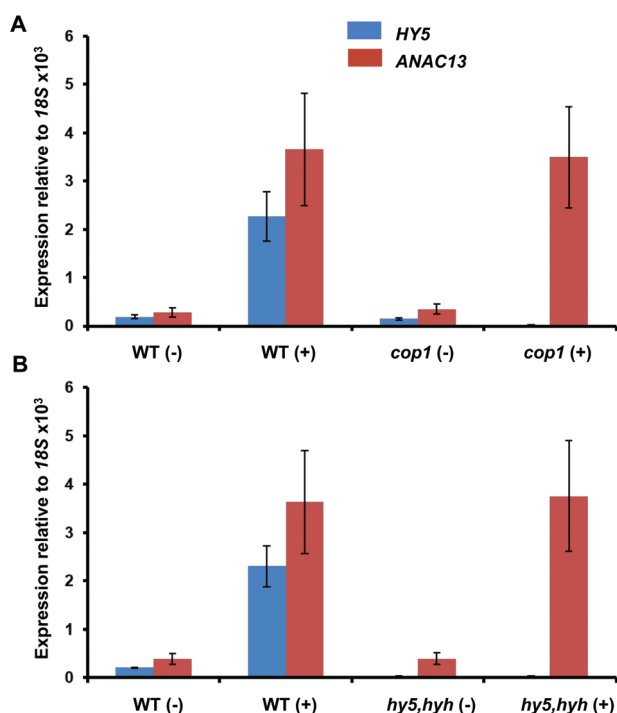


Fig. 6 UV-B induction of *ANAC13* is independent of *COP1*, *HY5* and *HYH*. Quantitative RT-PCR assays of *HY5* and *ANAC13* transcripts, normalized to control 18S transcript levels, in wild-type Ws, *cop1-4* (A) and *hy5/hyh* (B). 21-day old wild type, *cop1-4* or *hy5/hyh* mutant plants were grown in a low fluence rate of fluorescent white light (20 $\mu\text{mol m}^{-2} \text{s}^{-1}$) and then exposed (+) or not (–) to 0.5 $\mu\text{mol m}^{-2} \text{s}^{-1}$ of narrowband UV-B light for 2 h. The plant weighted UV⁵¹ was 0.005 kJ m^{-2} , which compares with 4.8 $\text{kJ m}^{-2} \text{day}^{-1}$ in Lund, Sweden, at midsummer with a cloudless sky.⁵¹ Data are means of three experiments \pm S.E.

ANAC13 UV-B induced expression is independent of several other known photoreceptors

To test the possibility that a photoreceptor other than UVR8 is able to detect low fluence UV-B and regulate *ANAC13*, or

perhaps act redundantly with UVR8, we tested several photoreceptor mutants for their ability to induce *ANAC13* in response to UV-B. To do this we used the *cry1cry2* mutant which lacks both cryptochromes, *phot1phot2* which lacks both phototropins, and *phyabde* which is deficient in all the phytochromes except PHYC. We also utilized the triple mutant, *cry1-cry2uvr8-2* to rule out the possibility of redundancy between the cryptochromes and UVR8. Fig. 7 shows that UV-B induction of *ANAC13* is similar to WT in all the mutants tested. Furthermore, in the triple mutant *cry1cry2uvr8-2*, *HY5* expression is not induced upon UV-B exposure, in contrast to *ANAC13* expression (Fig. 7D). Overall, these data indicate that *ANAC13* UV-B up-regulation is not activated *via* the known photoreceptors tested here and in addition *ANAC13* expression is not induced *via* an overlapping, redundant pathway operating between UVR8 and CRY1/CRY2.

ANAC13 shows maximum expression at longer wavelengths of UV-B in WT and *uvr8-6* plants

Finally, we wanted to determine which UV-B wavelengths produce the maximal expression of *ANAC13*. To do this we exposed both WT and *uvr8-6* plants to a range of UV-B wavelengths in increments of 10 nm from 280 to 320 nm for 1 hour, followed by 1 hour in darkness to allow transcripts to accumulate, then carried out qPCR to examine *ANAC13* and *HY5* transcript levels. Fig. 8 shows that both *ANAC13* and *HY5* expression are induced over the range of UV-B wavelengths in WT and only *ANAC13* expression is induced in the *uvr8-6* mutant. Fig. 8 also demonstrates that both *ANAC13* and *HY5* are maximally expressed at 310 nm and that the expression levels descend accordingly 300 > 290 > 280 > 320. The *p*-values (Table 1) show that *ANAC13* expression in WT and *uvr8-6* were not significantly different (>0.05) over the range of wavelengths tested. Overall, these data further support the claim that *ANAC13* is UVR8-independent and regulated by lower energy, longer wavelength UV-B.

Discussion

It is well established that UV-B exposure of plants regulates expression of a large number of genes.^{15,18,24,26} Such reprogramming of the transcriptome enables plants to re-adjust their metabolism, morphology and physiology to acclimate to changes in their radiation environment. While UVR8 regulates a substantial, functionally important subset of UV-B-regulated genes, it is evident that other processes can also mediate UV-B-induced gene expression. A number of studies have shown that sudden exposure of non-acclimated plants to high fluence rate, short wavelength UV-B induces expression of numerous genes that are not specific to the UV-B stimulus and are also induced by various stress treatments.^{24,26,41} The signal transduction processes initiated by UV-B stress treatments are reported to include DNA damage signaling, ROS signaling, MAPK kinase activity, and wound/defence signaling molecules.^{42–45} However, there is very little information about



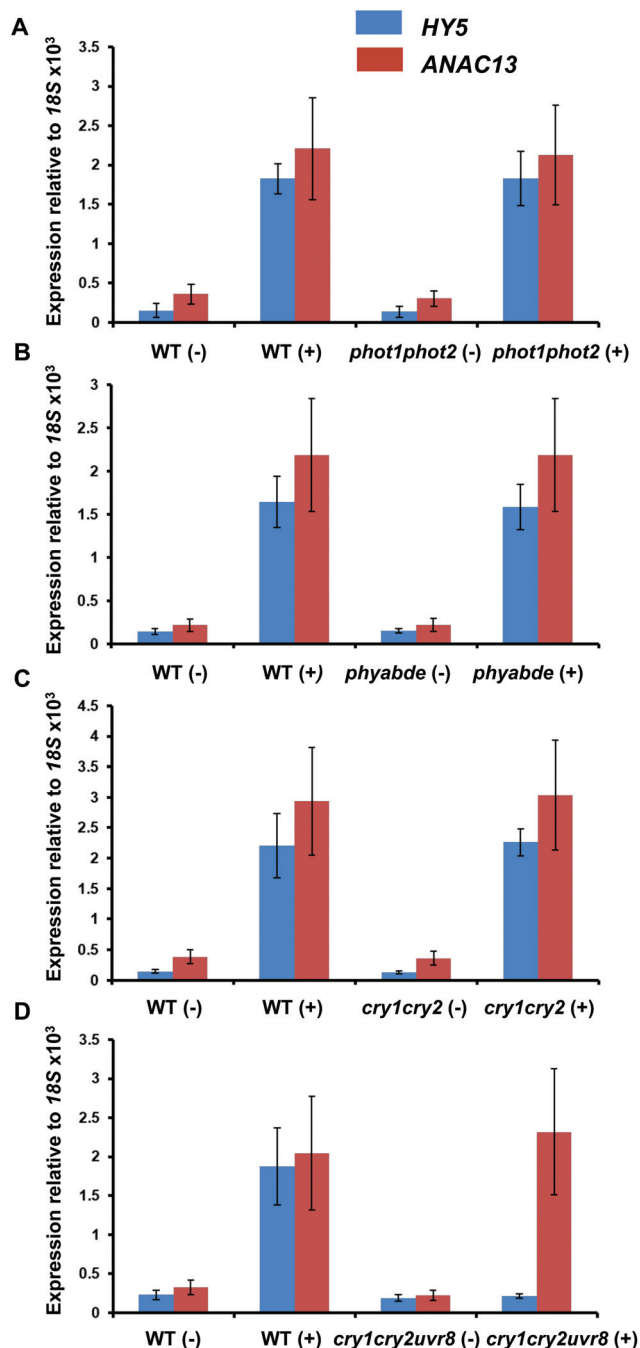


Fig. 7 UV-B induction of *ANAC13* is independent of multiple photoreceptors. Quantitative RT-PCR assays of *HY5* and *ANAC13* transcripts, normalized to control *18S* transcript levels, in wild-type *Ler* (or *Col-0* for *phot1phot2* comparison), *phot1phot2* (A), *phyABDE* (B), *cry1cry2* (C) and *cry1cry2uvr8-2* (D). Plants were exposed (+) or not (-) to $0.5 \mu\text{mol m}^{-2} \text{s}^{-1}$ of narrowband UV-B light for 2 h. The plant weighted UV⁵¹ was 0.005 kJ m^{-2} , which compares with $4.8 \text{ kJ m}^{-2} \text{ day}^{-1}$ in Lund, Sweden, at midsummer with a cloudless sky.⁵¹ Data are means of three experiments \pm S.E.

whether non-stressful, low fluence rate, longer wavelength UV-B treatments can regulate gene expression independently of UVR8, and that is the question addressed in this study.

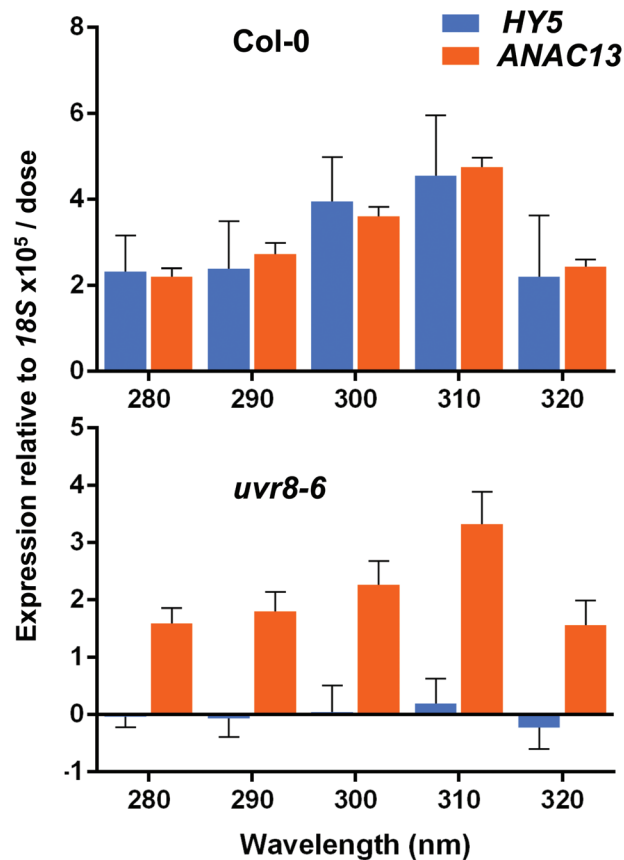


Fig. 8 *ANAC13* expression is maximal at 310 nm in WT and *uvr8-6* plants. Quantitative RT-PCR assays of *HY5* and *ANAC13* transcripts, normalized to control *18S* transcript levels, in wild-type *Col-0* and *uvr8-6*. Plants were exposed to a range of UV-B wavelengths for 1 h using a tuneable laser and after exposure left in the dark for 1 h before harvesting. Data are means of three experiments \pm S.E.

Table 1 *p*-Values obtained from ANOVA of response levels in each wavelength separately (*p* < 0.05 is significantly different). Columns 2 and 3: Comparison of *HY5* to *ANAC13* transcript levels in each genotype. Column 4: Comparison of *HY5* transcript levels in *Col-0* and *uvr8-6*. Column 5: Comparison of *ANAC13* transcript levels in *Col-0* and *uvr8-6*

	<i>Col-0</i>	<i>uvr8-6</i>	<i>HY5</i>	<i>ANAC13</i>
Comparison	<i>HY5/ANAC13</i>	<i>HY5/ANAC13</i>	<i>Col-0/uvr8-6</i>	<i>Col-0/uvr8-6</i>
280	0.991	0.015	0.002	0.491
290	0.926	0.046	0.011	0.402
300	0.888	0.012	0.0001	0.097
310	0.988	0.005	0.001	0.179
320	0.981	0.046	0.011	0.5

The UV-B exposures chosen for the microarray analysis were based on previous research that identified conditions for the induction of UVR8-dependent and UVR8-independent gene expression in *Arabidopsis*.²⁵ Plants were grown under identical conditions and given the same spectral quality and duration of UV-B exposure as previously,²⁵ however, the $0.3 \mu\text{mol m}^{-2} \text{s}^{-1}$ treatment used here is below the threshold of $1 \mu\text{mol m}^{-2} \text{s}^{-1}$



found to be required for UVR8-independent induction of stress related genes.²⁵ In the present study the UVR8-dependent genes were defined by both the Brown *et al.* (2005)¹⁸ and Favory *et al.* (2009)¹⁵ microarray analyses, which involved quite different plant growth and UV-B exposure conditions. It is therefore interesting that a substantial number of genes were induced by 0.3 $\mu\text{mol m}^{-2} \text{s}^{-1}$ UV-B independently of UVR8 (Fig. 1 and 2). Moreover, there is very little overlap between these UVR8-independent genes and genes reported to be induced by UV-B stress by Kilian *et al.* (2007).²⁶ This is not surprising because 0.3 $\mu\text{mol m}^{-2} \text{s}^{-1}$ UV-B is very unlikely to induce any of the known UV-B stress signaling pathways. Overall, the data indicate that a significant proportion of UV-B-responsive genes are regulated independently of both UVR8 and stress signaling.

The sets of genes induced by the 0.3 and 3.0 $\mu\text{mol m}^{-2} \text{s}^{-1}$ UV-B treatments had little overlap, but a small number of the UV-B regulated UVR8-independent genes were induced at all 3 UV-B fluence rates employed and several were also found to be induced under stress conditions.²⁶ At least some of these genes may be activated by 'change' rather than 'stress' *per se*. Hideg *et al.* (2013)⁴⁵ highlight the distinction between constructive stress, termed eustress, which promotes acclimation, and destructive stress in UV-B responses. It may be that transfer of plants that have not previously been exposed to UV-B to even a very low fluence rate of UV-B is sufficient to activate 'eustress' as opposed to 'damage stress' (distress), and that this UV-B induced eustress response is at least partly independent of UVR8.

It is not known how many different signaling pathways mediate the response to low fluence rate UV-B, and it is not clear why multiple mechanisms are employed. Redundancy is common in plant photoreception and signaling as it ensures that key stimuli, such as UV-B, will be perceived, and introduces flexibility in response. Characterisation of *ANAC13* gene expression provided information on one low fluence UV-B, UVR8-independent signaling pathway. *ANAC13* encodes a putative NAC domain containing transcription factor. The NAC domain (NAM, ATAF1/2 and CUC2) proteins are unique to plants and are thought to be involved in a wide range of processes including stress responses, development and growth.^{46–50} A previous study²⁷ reported that *ANAC13* could also be induced by short wavelength, relatively high fluence rate UV-B, but this is likely *via* a different signaling pathway. The kinetics of *ANAC13* induction by low fluence rate UV-B are different to those of the classic UVR8-regulated *HY5* gene and its downstream target *CHS*, supporting the notion that it is regulated *via* a distinct pathway. In addition, *ANAC13* is regulated by UV-B independently of COP1 and HY5/HYH, which are associated with UVR8 signaling. Moreover, at low fluence rates *ANAC13* is regulated by longer wavelength UV-B, maximally at 310 nm, further suggesting that it is not activated by stress signaling under these conditions. No detailed information is yet available on the nature of the low fluence pathway that regulates *ANAC13* expression. It does not appear to be regulated by the known photoreceptors, at

least based on our experiments with a range of photoreceptor mutants described above. However, the possibility of a novel photoreceptor cannot be excluded. There has been speculation about the existence of UV-B photoreceptors in addition to UVR8,²⁸ although no such molecule has yet been identified.

An important, but largely unanswered question is to what extent UVR8-independent pathways regulate gene expression in plants growing in natural environments where plants are not usually subject to UV-B stress. There is evidence that UVR8-independent pathways do operate,⁶ but they are not well defined. The present study highlights the potential for low, non-damaging fluence rates of UV-B, well within the wavelength range experienced by plants in nature, to regulate a substantial number of genes. Moreover, these pathways should not be considered as stress-related as they are evidently independent of classic stress signaling pathways. We therefore refer to these pathways as UVR8 and Stress-Independent (UASI) UV-B signaling pathways. Further research is now required to define the molecular mechanisms involved in these pathways and to assess the functional significance of gene expression responses that they mediate. The GO enrichment analysis suggests that carbohydrate metabolism may be an important function of genes regulated by UASI UV-B signaling, but no further insights are available at present.

Conflicts of interest

There are no conflicts of interest to declare.

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