

VIP1 response elements mediate mitogen-activated protein kinase 3-induced stress gene expression

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The plant pathogen *Agrobacterium tumefaciens* transforms plant cells by delivering its T-DNA into the plant cell nucleus where it integrates into the plant genome and causes tumor formation. A key role of VirE2-interacting protein 1 (VIP1) in the nuclear import of T-DNA during *Agrobacterium*-mediated plant transformation has been unravelled and VIP1 was shown to undergo nuclear localization upon phosphorylation by the mitogen-activated protein kinase MPK3. Here, we provide evidence that VIP1 encodes a functional bZIP transcription factor that stimulates stress-dependent gene expression by binding to VIP1 response elements (VREs), a DNA hexamer motif. VREs are overrepresented in promoters responding to activation of the MPK3 pathway such as *Trxh8* and *MYB44*. Accordingly, plants overexpressing VIP1 accumulate high levels of *Trxh8* and *MYB44* transcripts, whereas stress-induced expression of these genes is impaired in *mpk3* mutants. *Trxh8* and *MYB44* promoters are activated by VIP1 in a VRE-dependent manner. VIP1 strongly enhances expression from a synthetic promoter harboring multiple VRE copies and directly interacts with VREs in vitro and in vivo. Chromatin immunoprecipitation assays of the *MYB44* promoter confirm that VIP1 binding to VREs is enhanced under conditions of MPK3 pathway stimulation. These results provide molecular insight into the cellular mechanism of target gene regulation by the MPK3 pathway.

signal propagation | transcription factor | promoter motif | bZIP | *Arabidopsis*

The plant pathogen *Agrobacterium tumefaciens* transforms plant cells by delivering its transfer-DNA (T-DNA) into the plant cell nucleus, where it integrates into the plant genome. The subsequent growth of crown galls at the infection sites substantially interferes with plant development. There is ample experimental evidence demonstrating a key role of VIP1 (VirE2-interacting protein 1) in the nuclear import of T-DNA during *Agrobacterium*-mediated plant transformation (1, 2). VIP1 exhibits a stress-dependent subcellular localization; and phosphorylation of VIP1 at serine 79 by the stress-activated MAPK MPK3 (3) triggers its translocation from cytoplasm to the nucleus (2). Analysis of the *vip1-1* mutant line, which produces only a truncated VIP1 protein, has shown that the N-terminal portion of VIP1 is sufficient to bind to and target VirE2 to the nucleus to facilitate transient genetic transformation by *Agrobacterium* (4). However, this is insufficient for stable genetic transformation and *Agrobacterium*-induced tumorigenesis (4). Apart from *Agrobacterium* transformation, no in planta function has yet been ascribed to VIP1 in *Arabidopsis*. Considering that the only purpose of a plant protein would not merely lie in assisting pathogen invasion, we aimed to investigate the in planta function of VIP1 and the functional relevance of its stress-dependent nuclear translocation.

In this work, we focus on the molecular events immediately downstream of stress-activated VIP1. We report on the role of VIP1 as a transcriptional regulator of genes that are targets of the stress-activated MPK3 cascade. VIP1 mediates transcriptional induction of target genes by binding to a DNA motif, termed VRE for VIP1 response element. Multiple copies of VRE motifs are found in the promoters of various stress-responsive genes, and evidence is provided that VIP1 directly

targets VRE motifs in vitro and in vivo. This work gives insight into the molecular mechanism how stress-induced MPK3 activation is linked to target gene induction.

Results and Discussion

PR1 is an Indirect Target of VIP1. In transiently transformed *Arabidopsis* protoplasts, VIP1 shows a predominantly nuclear localization, and we found that nuclear VIP1 can activate the transcription of the pathogenesis-regulated protein 1 gene (PR1) (2), a major stress marker gene. A bZIP domain located in the C-terminal region classifies VIP1 as a putative transcription factor (Fig. S1). First, we addressed the question whether PR1 was directly regulated by VIP1. VIP1 is one of the 75 members belonging to the bZIP family of *Arabidopsis thaliana*. A phylogenetic study places VIP1 into the so far uncharacterized subgroup I (5). The 12 members of this family carry a characteristic lysine (K) instead of the otherwise highly conserved arginine residue (K212 in VIP1) within the bZIP domain. The nature of this particular residue is a determinant for the binding preference of bZIP family members, with arginine-type members preferentially binding to palindromic, and lysine-type members preferentially binding to nonpalindromic DNA elements (6, 7). If VIP1 encodes a functional bZIP transcription factor, mutation at K212 might affect its transactivating activity. To test this idea, K212 of VIP1 was replaced by the related residue arginine (R), or the nonsimilar threonine (T) residue, to potentially alter the DNA-binding preference or DNA-binding capacity of the protein, respectively. In protoplast transient expression experiments, only wild-type VIP1, but not its mutant derivatives K212R or K212T, efficiently activated the PR1 promoter::GUS (β -glucuronidase) reporter construct (Fig. 1A).

PR1 gene expression is regulated through binding of proteins belonging to the TGA subfamily of bZIP transcription factors to specific promoter motifs, including the as-1 element (consisting of subelements LS5 and LS7) and LS4 (8). Mutations at LS4, LS7, and/or LS5 did not affect VIP1 activation of PR1 promoter variants in protoplast cotransfection studies (data not shown). Moreover, VIP1 was unable to induce an as-1 element-containing synthetic promoter or the cauliflower mosaic virus CaMV 35S promoter, which harbors as-1 elements (data not shown). Electrophoretic mobility shift assays (EMSA) also showed no detectable binding of recombinant VIP1 to fragments of the PR1 promoter or to the as-1 element alone (Fig. 1B).

Together, these findings suggest that VIP1 is a functional bZIP

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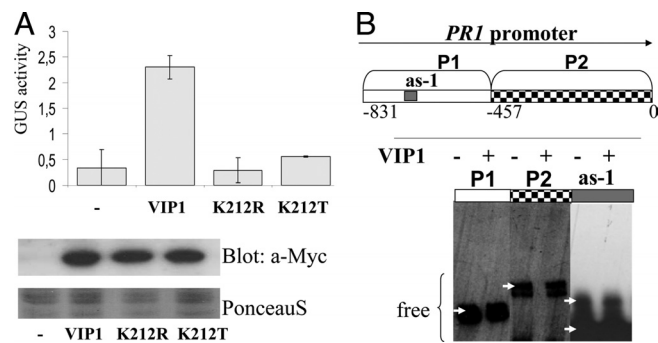


Fig. 1. Indirect induction of PR1 by VIP1. (A) bZIP residue Lys-212 is essential for VIP1-induced PR1 expression. Protoplasts were transfected with the PR1::GUS reporter construct alone or in combination with constructs for overexpression of VIP1-Myc or VIP1 K212-Myc mutant variants. Given are mean values and standard deviations of GUS activity (nmol 4-MU min⁻¹ (mg protein)⁻¹; $n = 6$). Transgene expression was visualized by immunoblotting with anti-Myc antibody. Equal loading was visualized by staining of the membrane with PonceauS. (B) VIP1 does not bind to the PR1 promoter (EMSA). The indicated PR1 promoter fragments or the as-1 element were biotin-labeled and incubated with or without recombinant VIP1 protein. Nonbound DNA fragments are indicated by arrows. No band shift was observed.

transcription factor, but that, unlike TGA proteins, VIP1 most probably acts on PR1 expression in an indirect manner.

PR1 is known as a late stress-responsive gene. Considering the rapid activation of MPK3 upon stress that is followed by VIP1 phosphorylation and nuclear translocation within minutes (2), good candidates for VIP1 direct target genes might therefore rather be found among early stress-responsive genes.

Identification of VRE as a DNA Motif Binding to VIP1. In contrast to the subgroup of TGAs and several other family members in *Arabidopsis* (5, 9) little is known about the subfamily I of bZIPs. Unlike most other bZIP proteins, members of subfamily I are predicted to form homo- rather than heterodimers (10). Co-immunoprecipitation experiments (Fig. S2) and bimolecular fluorescence complementation (BiFC) analysis (4), confirm homodimerization activity of VIP1. Consequently, additional proteins may not necessarily be required for VIP1 to bind to DNA. The target DNA motifs of bZIP transcription factors related to VIP1 (subfamily I) have been identified in other species, such as tomato VSF-1 (GCTCCGTTG) (11) and tobacco RSG (TCCAGCTTGA, TCCAAGTTGGA) (6). However, the dissimilarity in VSF-1 and RSG target DNA motifs suggests that, despite high homology in the bZIP domains, the DNA binding preferences of VSF-1 and RSG are not conserved. Moreover, no target DNA motifs are yet known for any *Arabidopsis* bZIP subfamily I member.

In an attempt to identify DNA elements targeted by VIP1, we carried out random DNA binding selection assays (RDSAs). To this end, chitin-immobilized recombinant VIP1 protein was incubated with random 17- or 18-bp-long DNA fragments flanked by defined primer annealing sites (*SI Text*). VIP1-bound DNA fragments were PCR-amplified and used as input for another RDSA cycle with a second aliquot of VIP1 protein. By raising the concentration of nonspecific competitor, double-stranded DNA poly(dIdC), the stringency was increased in each subsequent purification cycle. A total of five cycles were performed, and the amplified DNA fragments were subsequently cloned and sequenced.

Of the seven independent candidate elements isolated, five (E1–E5; Fig. 2A) contained the common motif ACNGCT. Moreover, E2 carries two copies of ACNGCT, and two candidate fragments were isolated in duplicates, indicating the selectivity of the RDSA procedure.

To test whether VIP1 can directly interact with the consensus motif, EMSA was performed using recombinant VIP1 protein and

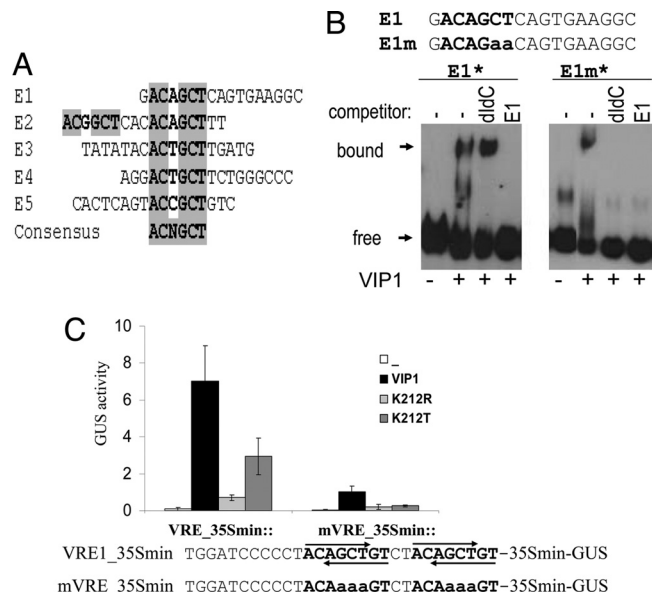


Fig. 2. Identification and binding of VIP1 to VREs. (A) Alignment of potential VIP1-target DNA elements isolated by RDSA. The consensus motif shared by five candidate fragments is highlighted. (B) VIP1 binds to VRE sequences (EMSA). Binding of VIP1 to candidate motif E1 isolated by RDSA (Left) or element E1m, carrying a mutation at the ACNGCT motif (Right). Biotin-labeled (*) E1 or E1m were incubated with or without recombinant VIP1 in the absence or presence of an excess of either nonspecific (dIdC) or specific inhibitor (nonlabeled E1). (C) VIP1-induced expression from a synthetic multiple-VRE-containing promoter. Protoplasts were transfected with synthetic promoter-GUS reporter gene constructs containing four copies of VRE (or mutated VRE-mVRE), alone or in combination with 35S::VIP1-Myc or 35S::VIP1 K212-Myc mutant variants and analyzed as described in Fig. 1A. Lower: Sequence of the promoter region. VRE sequences are shown in bold, and their orientation is indicated by arrows.

biotin-labeled DNA fragments. A clear band shift indicated the DNA-binding capability of VIP1 to the E1 fragment (Fig. 2B). The binding specificity of VIP1 for the ACNGCT core sequence was tested in EMSA with the E1m DNA fragment, which is identical to E1 except for a mutation in the ACNGCT motif (ACNGaa). EMSAs showed that although VIP1 can also bind to biotin-labeled E1m, this binding is abolished in the presence of the nonspecific competitor poly(dIdC) as well as by a 20-fold excess of nonlabeled E1 (Fig. 2B). In contrast, binding of VIP1 to labeled E1 is only substantially reduced in the presence of the specific competitor (nonlabeled E1) but not by the nonspecific competitor poly(dIdC). These results strongly indicate that VIP1 directly targets the ACNGCT DNA motif, which we hitherto call VRE. The VRE sequence does not match any characterized regulatory DNA motif, nor does it contain the TGAC core sequence commonly targeted by various bZIP proteins (12). Consistent with the predicted preference for nonpalindromic DNA elements (7), VRE is devoid of a palindromic core.

VIP1 Induces Transcription from a Synthetic Promoter Carrying VRE Motifs. To test whether VIP1 can directly regulate VRE-containing promoters, we generated two synthetic promoter-reporter gene constructs. Four tandem copies of VRE, two in either orientation, were fused to a minimal promoter fragment derived from the CaMV35S promoter to give VRE::GUS. A mutated derivative containing two mismatches in each of the four VRE copies was generated to give mVRE::GUS (Fig. 2C). Protoplasts transfected with the reporter gene constructs alone showed no detectable GUS activity (Fig. 2C). However, constitutive coexpression of VIP1 strongly induced GUS expression from the VRE promoter, but only poorly from the mVRE promoter (Fig. 2C). VIP1 mutant variants

Table 1. VRE1 overrepresentation in promoters of stress-responsive genes

Experimental condition	Total number of differentially expressed genes	Proportion of promoters with two or more VRE	P value	Reference
Total genome	33,282	1,946 (5.8%)	1.0	
Stress datasets (under conditions when MPK3 is activated)				
Transient response to flg22	42	(6) 14.3%	0.0198*	17
Early systemic wound response	14	(4) 28.6%	0.0003***	18
Early response to cold, mannitol and NaCl	66	(11) 16.7%	0.0002***	19
Common response to cold, osmotic stress, wounding and biotic stress	197	(18) 9.1%	0.049**	20
Common response to 9 or more abiotic or biotic stress treatments	182	(16) 8.8%	0.0905*	21
Altered expression in <i>mkp1</i>	21	(3) 16.7%	0.0002***	22
Control datasets (transcriptome studies of conditions not affecting MPK3 activity)				
Pollen-enriched	456	27 (5.9%)	0.9463	23
Response to methyl jasmonate	975	50 (5.1%)	0.3388	24
Response to ACC (ethylene)	314	18 (5.7%)	0.9311	24

The proportion of genes containing at least two VRE copies in their 500 bp promoter regions was compared between all *Arabidopsis* genes and datasets from published microarrays. The statistical significance, as calculated by Chi² test, is indicated as * (significant), ** (highly significant), and *** (extremely significant).

in which the characteristic lysine residue (K212) that confers DNA-binding specificity had been replaced by an arginine (K212R) or a threonine (K212T) did not activate expression from either promoter. These findings substantiate the results from RDSA and EMSA experiments and suggest that VIP1-binding to VRE boxes is directly responsible for inducing gene expression. In line with the notion that PR1 expression is regulated by VIP1 only indirectly, no VRE sequences are found in the PR1 promoter.

VRE Defines a DNA-Binding Motif in Promoters of Stress Genes. We then wondered whether the VRE motif could be a bona fide regulatory motif. MPK3 is activated by multiple biotic and abiotic stresses (reviewed in references 3 and 13). Based on the stress-activated protein kinase MPK3-triggered rapid nuclear translocation of VIP1 in planta (2), we hypothesized that VIP1 might bind to promoters of genes involved in the early stress response. Apart from a small number of uncharacterized candidates isolated from a protein chip-based screen for targets of MPK3 in vitro (14, 15), little is known about the steps of early stress signaling immediately downstream of MPK3. Thus, VIP1 is so far the only transcription factor with in vivo evidence for a direct regulation by MPK3. If VIP1 was a major mediator of MPK3-mediated gene expression and if VREs were the major motifs targeted by VIP1, one might expect to find an accumulation of VRE motifs in the promoters of genes responsive to the signals activating the MPK3 pathway. To test this theory, we performed a statistical analysis of *A. thaliana* promoters, exploiting publicly available microarray data resources.

Although in eukaryotic genes regulatory elements can be distributed anywhere within approximately 1,500 bp upstream of the transcription start sites, they are commonly found within the proximal 500 bps (16). We therefore compared the abundance of VRE sequences (in both forward and reverse orientations) in the 500-bp promoter regions of *A. thaliana* genes with reported stress-regulated transcript abundance in published microarrays to the overall abundance in all *Arabidopsis* genes. Because bZIP proteins form dimers, with both monomers binding to DNA, we expected VIP1 target promoters to carry at least two copies of VREs. Indeed, there were no significant differences in the abundance of stress- or nonstress-regulated promoters carrying a single VRE copy (data not shown), whereas the proportion of promoters carrying two or more VRE copies was significantly higher in stress-regulated genes retrieved from several independent transcriptome studies (Table 1). There was no apparent bias for the orientation of the motif (ACNGCT or AGCNGT).

A gene expression study of the early and late wound response in local and systemic leaves identified 14 genes that are induced systemically early after wounding (18). Four of these genes (28.6%)

carry multiple VRE copies in their promoters, a significantly higher proportion than would be expected from the overall abundance of such promoters in the *Arabidopsis* genome (5.8%). Since this is only a small dataset, one should be careful to generalize from these findings. However, we also found proportionally more multiple VRE-containing promoters among the dataset of genes with reported early transient induction by the bacterial elicitor flagellin-derived peptide flg22 (17). Of the 42 genes whose expression is induced after 30 min, but declined after 60 min of treatment, six (14.3%) harbor multiple VRE sequences in their promoters. It is interesting to note the time correlation between MPK3 activity and the expression of VRE-containing genes. MPK3 activity increases rapidly upon flg22 treatment and sharply declines after 30 min (25). Transcription factors that are directly activated by MPK3 are therefore likely to contribute to transient flg22-triggered changes in gene expression.

Three independent genome-wide studies have established clusters of genes that commonly respond to multiple stress treatments (19–21). Among these clusters, genes with multiple VREs in their 500-bp promoter region are clearly overrepresented (Table 1). VREs are also more abundant in promoters of genes differentially expressed in *mkp1* plants, which are impaired in the MPK3-inactivation via dual-specific phosphatase MKP1 (22). To assess the significance of the overrepresentation of multiple VRE-containing promoters in the statistical analysis of these stress-related transcriptome datasets, we also included datasets obtained under conditions when MPK3 is not activated or found to play a prominent role (23, 24) (Table 1). Unlike their overrepresentation in datasets of stress-responsive genes (8.8–28.6%), multiple VRE-containing promoters in these control datasets (5.1–5.7%) show an abundance that is close to that expected for the entire *Arabidopsis* genome (5.8%).

Investigation of VRE Regulation in Two Potential VIP1 Target Genes.

Based on the above observations, we hypothesized that VREs may serve as transcriptional regulatory motifs in at least a subset of stress-responsive genes. For the selection of putative target genes of the MPK3-VIP1 pathway, we applied two criteria: such genes should be (i) responsive to conditions that correlate with MPK3 activation and (ii) carry at least two VRE copies in their promoter regions. Clearly, this approach excludes promoters which are bound by possible heterodimers of VIP1 and other bZIP-like proteins recognizing distinct DNA motifs.

For a more detailed analysis of the contribution and mechanism how VRE motifs function in the context of MPK3-mediated signaling by VIP1, we selected two genes, thioredoxin *Trxh8* (At1g69880) and the transcription factor *MYB44* (At5g67300). *Trxh8* and/or *MYB44* gene expression correlates with activation of

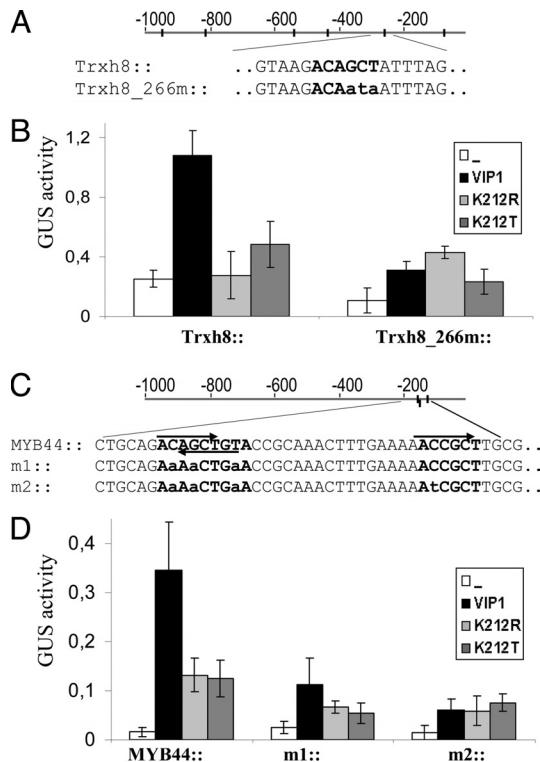


Fig. 3. VIP1 target genes. (A and C) Schematic presentation of the *Trxh8* and *MYB44* promoter regions. Promoter regions matching the VRE1 consensus (ACNGCT or its reverse complement, AGCNGT) are highlighted, respective mutations of the promoter constructs are shown in lowercase. Note that in the *MYB44* promoter two oppositely-oriented copies of VRE1 are overlapping. (B and D) Protoplast cotransfection. Protoplasts were transfected with *Trxh8* (B) or *MYB44* (D) native and mutated promoter::GUS constructs in the absence or presence of VIP1 or Lys-212 mutant variants K212R, K212T, and analyses as described in Fig. 1A.

MPK3 in the response to a diversity of stresses, e.g., treatment with the plant pathogen *Pseudomonas syringae*, various pathogen-associated molecular patterns, wounding, and oxidative stress (18, 25–27).

The 1,000-bp promoter region of *Trxh8* harbors six VRE copies (Fig. 3A). Protoplasts transfected with a 1,100-bp *Trxh8* promoter-driven GUS construct displayed higher reporter gene activity when cotransfected with a construct for constitutive overexpression of VIP1 (35S::VIP1) (Fig. 3B). We then tested whether VIP1 activated the *Trxh8* promoter in a VRE-dependent manner. As statistical analyses indicated the presence of regulatory DNA elements to be positioned most often at regions 200–300 bp upstream of the translation start site (28), we mutated the VRE motif that is located 266 bp upstream of the *Trxh8* transcription start site to give *Trxh8_266m*::GUS. Compared with protoplasts transfected with a wild-type *Trxh8*::GUS construct, *Trxh8_266m*::GUS activity was significantly compromised upon cotransfection with 35S::VIP1 (Fig. 3B). Consistent with our concept that the lysine residue K212 is essential for VIP1 DNA binding, overexpression of VIP1 K212 mutant variants (K212R or K212T) did not activate the *Trxh8* promoter (Fig. 3B).

The second potential VIP1 target gene, *MYB44*, is one of the 197 core stress responsive genes reported by Ma and Bohnert (20). The *MYB44* promoter region harbors three VRE copies, two of which are overlapping in opposite orientation (Fig. 3C). Protoplast cotransfection studies revealed that a 137-bp *MYB44* promoter fragment containing the three VRE copies was sufficient to drive VIP1-induced GUS reporter gene expression. VRE mutation (*MYB44m1*, *MYB44m2*) rendered the promoter nonresponsive to activation by VIP1 (Fig. 3D). Similar to *Trxh8* promoter activation,

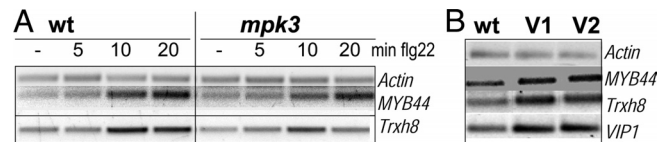


Fig. 4. MPK3-dependent stress responsiveness of VIP1 target genes *Trxh8* and *MYB44*. (A) Flg22-induced expression of VIP1 target genes in wild-type and *mpk3* mutant plants. Fourteen-day-old seedlings grown in Petri dishes were adjusted for 24 h in liquid medium before application of water or 2.5 μ M flg22. Samples were frozen after 0, 5, 10, and 20 min and assessed by semiquantitative RT-PCR. *Actin* and *MYB44* transcripts were amplified in a multiplex PCR. (B) VIP1 target gene expression in VIP1 overexpressing plants. Semiquantitative RT-PCR analysis of expression levels of the indicated genes in 14-day-old seedlings of wild-type and two stably transformed VIP1-HA/e lines (V1, V2).

induction of *MYB44* expression by VIP1 depends on VIP1 residue Lys-212. Taken together, these results are consistent with the assumption that *Trxh8* and *MYB44* are direct VIP1 targets and that direct binding through the bZIP domain of VIP1 to its cognate VRE sequences is necessary for promoter activation.

Defining the VIP1 Signaling Pathway. According to our hypothesis, *MYB44* and *Trxh8* should be stress-regulated target genes of VIP1, whose nuclear localization depends on prior phosphorylation by stress-activated MPK3. Consequently, mutant plants lacking MPK3 or VIP1 would be expected to be impaired in the stress-induced expression of these genes. We analyzed *MYB44* and *Trxh8* gene expression at 5, 10, and 20 min of treatment with flg22. In wild-type plants, *MYB44* and *Trxh8* transcripts rapidly accumulated upon flg22 treatment (Fig. 4A). Similar stress gene induction was observed in the previously characterized *vip1* mutant line, *vip1-1* (SALK_001014) (Fig. S3). [The only other available putative *vip1* T-DNA insertion line (SALK_148460) turned out to express wild-type levels of the complete VIP1 transcript.] *vip1-1* forms a C-terminally truncated protein in which the MPK3 phosphorylation site, as well as the entire bZIP domain, are retained and is apparently not affected in its transcriptional activation capacity. Thus, *vip1-1* mutants were found unsuitable for testing our hypothesis. We therefore focused on the analysis of *mpk3* null mutants.

Compared with wild-type plants, we consistently observed lower flg22-induced *MYB44* and *Trxh8* transcript accumulation in *mpk3* mutant plants (Fig. 4A). Stress gene induction was not completely blocked in *mpk3*, most probably because MPK3 can be partially compensated for by the highly homologous MAPK MPK6. Previous in vitro kinase assays had revealed that VIP1 can, although very poorly, also be phosphorylated by MPK6 (data not shown). In addition, MPK3-VIP1-mediated stress gene induction might be partially masked by the contribution of additional regulatory elements targeted by other stress-related transcription factors.

Complementary to studying VIP1 target gene expression in *vip1* or *mpk3* mutant plants, we sought further confidence of VIP1 being an activator of *MYB44* and *Trxh8* expression by a gain-of-function approach. To this end, transgenic plants constitutively overexpressing hemagglutinating-tagged VIP1 (VIP1-HA/e) were generated. In all six lines tested (shown for two lines in Fig. 4B), overexpression of the VIP1-HA transgene correlated with pronounced accumulation of *MYB44* and *Trxh8* transcripts.

Taken together, the observed impairment of stress-triggered induction of two candidate VIP1 target genes, *MYB44* and *Trxh8*, in *mpk3* null mutants as well as the elevated transcript levels of these genes in VIP1-HA overexpressing plants suggest a direct regulation via the following pathway [stress→MPK3→VIP1→VRE promoter activation→stress gene induction]. That *MYB44* and *Trxh8* transcript levels are enhanced in nontreated VIP1-HA overexpressing plants is likely to be because of a basal nuclear level of VIP1. This, in turn, may be explained by the strong activity of the constitutive promoter driving VIP1 expression or a basal level of

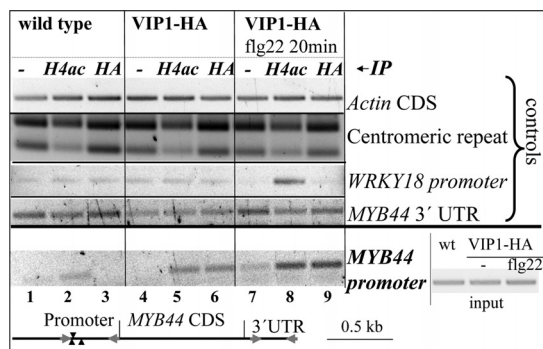


Fig. 5. ChIP of the *MYB44* promoter region comprising the VRE motifs. Chromatin of wild-type (wt) or VIP1-HA overexpressing plants (-/+ application of a 20-min treatment with flg22) was immunoprecipitated without antibody (mock, -) or with antibodies directed against histone H4ac or HA. PCR with primers binding to the actin coding region, a 180-bp repetitive centromeric region, or a WRKY18 promoter region served as controls for equal DNA content, for successful ChIP, and effective flg22 response, respectively. *Lower:* relative position of the primers used for *MYB44* promoter amplification. *Right:* "Input," PCR amplification products of "input" DNA samples [chromatin isolated from wt, VIP1o/e (-/+ flg22) before immunoprecipitation] with *MYB44* promoter-specific primers.

MPK3 activity. Upon stress treatment, additional VIP1 molecules are recruited from the cytoplasm to the nucleus. This assumption is consistent with our previously reported microscopy analysis, where a fraction of the transgenic VIP1 protein was found in the nucleus already in nontreated VIP1-YFP-expressing plants (2).

Chromatin Immunoprecipitation Reveals Stress-Dependent *In Vivo* Binding of VIP1 to the *MYB44* Promoter. To investigate whether VIP1 directly interacts with target promoters in planta, VIP1-HA overexpressing plants were subjected to chromatin immunoprecipitation (ChIP) analysis and compared with wild-type plants (Fig. 5).

ChIP performed on wild-type plants without (mock control) or with anti-HA antibody yielded no detectable enrichment of a *MYB44* promoter region (spanning the VRE1-containing region) (Fig. 5, lanes 1 and 3, respectively). In contrast, we observed *MYB44* promoter amplification products when ChIP was performed with anti-histone H4ac antibody. This antibody recognizes acetylated histone H4 and enriches for expressed genes [as evidenced by the shift of amplification products of a repetitive centromeric region (29) in the anti H4ac samples] (Fig. 5, lane 2).

In contrast, ChIP with either anti-HA or anti-histone H4ac antibody yielded detectable *MYB44* promoter amplification products in VIP1-HAo/e plants (Fig. 5, lanes 5 and 6, respectively), consistent with the elevated *MYB44* transcript levels in these plants (Fig. 4B).

Because MPK3-dependent phosphorylation of VIP1 results in enhanced activation of transcription by VIP1, we also tested whether the VIP1 interaction with the *MYB44* promoter will increase under MPK3-activating conditions. VIP1-HAo/e plants were stimulated with flg22 for 20 min before chromatin isolation. ChIP analysis with anti-HA antibody resulted in stronger accumulation of *MYB44* promoter-specific amplification products in flg22-treated compared with nontreated samples of VIP1-HAo/e plants (Fig. 5, lanes 9 and 6, respectively), indicating more efficient binding of VIP1 to the *MYB44* promoter. Interestingly, when ChIP was performed with anti-histone H4ac from flg22-treated VIP1-HAo/e plants and compared with nontreated conditions, a much stronger amplification was equally obtained (Fig. 5, lanes 8 and 5, respectively). These results indicate that activation of the *MYB44* promoter by flg22 is correlated with enhanced association of acetylated histone H4 with the *MYB44* promoter. The enhanced association of histone H4ac or VIP1-HA with the *MYB44* promoter upon flg22 treatment was not because of unequal reaction conditions as

evidenced by ChIP with the same antibodies, but PCR performed with primers for the actin coding region or a 180-bp centromeric region (Fig. 5). PCR amplification of a WRKY18 promoter region served as an additional control. The activity of the early stress-responsive WRKY18 promoter is regulated via W-boxes recognized by WRKY proteins (30). Enrichment of a WRKY18 promoter region was observed in samples of flg22-treated VIP1-HAo/e plants immunoprecipitated with anti-histone H4ac-antibody, but not with anti-HA antibody. This indicates that VIP1o/e plants responded to flg22 treatment with activation of the WRKY18 promoter, but that this activation was not mediated through binding of VIP1.

In summary, our ChIP experiments provide evidence that VIP1 can directly bind to the *MYB44* promoter *in vivo*. Furthermore, binding is enhanced under conditions of flg22-induced MPK3 activation. These data are consistent with the observed enhanced *MYB44* transcript levels in VIP1 overexpressing plants and the inducibility of *MYB44* expression by flg22.

MAPK-targeted transcription factors are good candidates for regulating gene expression of components involved in the primary/immediate stress response (e.g., redox-regulatory enzymes such as thioredoxins) and of additional transcription factors, which in turn activate respective subsets of target genes encoding components involved in stress adaptation.

By acting directly downstream of MPK3, VIP1 is a potent transducer of the stress signaling MAPK cascade. Through the transcriptional induction of genes encoding transcription factors themselves, such as *MYB44*, VIP1 may also be involved in mediating later than the immediate early responses characterized by the transient activation pattern of the MAP kinase. The presence of multiple VREs in the promoters of several stress-responsive genes suggests that VREs might also serve to coordinately activate a larger set of genes by VIP1. It is likely that the signaling pathways for responses to individual stresses bifurcate downstream of VIP1, a possible scenario could be through heterodimerization with additional transcriptional activators and/or repressors that are regulated by separate signaling pathways.

Since even in nontreated plants, a minor pool of VIP1 is found in the nucleus (2), VIP1 may be contributing to the background levels of activation of promoters harboring VREs.

VIP1 joins the group of *Arabidopsis* bZIP proteins with known DNA target motifs. VRE is a DNA element. VIP1 is the so far only characterized member of the bZIP subfamily I. For few *Arabidopsis* bZIP proteins, target motifs and putative target genes have been identified (8, 9). Unlike VIP1, these proteins display a constitutive nuclear localization and are not subject to any known posttranslational control.

A recent study reports on a B3-DNA binding protein, NtWIF, whose transactivation activity is regulated through phosphorylation by WIPK, the putative MPK3 ortholog in tobacco (31). A number of stress-related putative target genes, including *WIPK*, have been identified by microarray analysis. MPK3 is also subject to transcriptional control (13). However, a self-amplifying circuit of MPK3 signaling via VIP1—similar to NtWIF-induced *WIPK* gene expression (31)—appears unlikely, since no VRE sequences are found in the MPK3 promoter.

A previous study (32) has shown that heterologous overexpression of VIP1 in tobacco results in developmental abnormalities, including growth retardation and impaired differentiation. These effects may be related to aberrant activation/repression of VIP1-targeted promoters, since VIP1 in tobacco locates exclusively to the nucleus. As no transcriptional analysis has been performed on these plants, the identity of VIP1-targeted tobacco promoters remains elusive. If VIP1 overexpression activated stress gene expression also in tobacco, this might account for the developmental abnormalities displayed by these plants, since constitutive activation of stress responses is often accompanied by developmental defects (30). Alternatively, the phenotype found in VIP1 overexpressing tobacco plants might be the consequence of an imbalance of endogenous

bZIP proteins (subfamily I-type) that compete with VIP1 for their true dimerizing partners

Moreover, Avivi et al. (32) observed that VIP1 gene expression is enhanced in de-differentiating *Arabidopsis* cells (undergoing protoplasting); and we found ectopically expressed VIP1-YFP predominantly in the nucleus (2). VIP1 may thus be involved in the maintenance of enhanced basal expression of stress-responsive genes in protoplasts and thereby confer a certain preparedness against challenging conditions, as de-differentiating cells are (because of the lack of a cell wall) particularly susceptible to stress.

In summary, VIP1 is a protein incorporating two distinct functions. It mediates the nuclear import of *Agrobacterium* T-DNA, thereby assisting plant transformation. The second function involves transcriptional regulation of stress-responsive genes. Both VIP1 functions rely on stress-triggered nuclear translocation of VIP1 and are mediated through phosphorylation by MPK3. In our study, we have also identified VRE as a DNA motif that is bound by VIP1 in vivo and whose abundance in stress-responsive promoters correlates with MPK3 activation. These results provide evidence that one response to stress-triggered MPK3 activation is mediated by VRE-dependent stress gene expression through phosphorylation-dependent control of VIP1. The potential impact of VIP1-mediated stress gene activation in stress tolerance will be addressed in future studies.

The question arises how *Agrobacterium* can take advantage of the stress-triggered nuclear translocation of VIP1 if nuclear VIP1 activates the defense response. In fact, numerous stress-response genes are induced in the early response to *Agrobacterium* infection (33). Subsequent expression of these genes is suppressed later during infection with strains that can transfer Vir proteins and T-DNA (33). Once it has fulfilled its function as T-DNA transporter, VIP1 undergoes proteasomal degradation triggered by *Agrobacterium* VirF (34).

Thus, the initial stress gene activation upon *Agrobacterium* contact might—at least partially—be attributed to the action of nuclear VIP1, whereas because of its proteasomal degradation later in the infection process, VIP1 cannot activate defense genes any more.

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Methods

Plant Material and Treatment. *A. thaliana* Col-O and CaMV35S::VIP1-HA over-expressing plants were grown on plates containing ½ Murashige and Skoog (MS) medium (Duchefa), 1% sucrose and 0.8% agar at 24 °C and a 16 h photoperiod. For flg22 treatment, 14-day-old wild-type seedlings were adapted overnight in liquid ½ MS medium containing 1% sucrose. Flg22 was added to a final concentration of 2.5 μM. Seedlings were frozen at 0, 5, 10, and 20 min after treatment.

Protoplast Transfection and GUS Quantification. *Arabidopsis* protoplasts preparation, transfection, protein extraction, GUS activity quantification, and immunoblotting were performed as described in ref. 2.

Labeling of RDSA Candidates for EMSA. Inserts of sequenced clones were PCR-amplified with primers RDSA_1fo/re or RDSA_2fo/re (Table S1), respectively, and labeled using the biotin 3' end labeling kit (Pierce). Efficiency of labeling was checked by dot blot analysis according to the manufacturer's instructions. The following labeled oligonucleotides were annealed E1fo/E1re; E1mfo/E1mre; as-1fo/as-1re.

Electrophoretic Mobility Shift Assay. Recombinant VIP1 was incubated with biotin-labeled DNA fragments in RDSA binding buffer for 40 min at 4 °C. The reactions were loaded on a 6% TBE gel that had been prerun for 1 h at 4 °C. Blotting and detection were performed using the Light Shift Chemiluminescence kit (Pierce). For competition analysis, 500 ng poly(dIdC) or a 20-fold excess of nonlabeled RDSA candidate fragment E1 were added to the binding reactions.

Chromatin Immunoprecipitation. Four-week-old wild-type and VIP1-HAo/e plants were subjected to ChIP analysis with rabbit polyclonal antibody directed against acetylated Histone H4 (Upstate) or monoclonal mouse anti-hemagglutinin antibody following a protocol described in <http://www.epigenome-noe.net/researchtools/protocol.php?protid=13>.

Additional Experimental Procedures. A detailed description of plasmids and cloning, RT-PCR analysis, expression and purification of VIP1 recombinant protein, the Random DNA Selection assay, and VRE motif abundance calculation can be found in the *SI Text*.

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