Destabilization of interaction between cytokinin signaling intermediates AHP1 and ARR4 modulates *Arabidopsis* development

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Summary

- Eukaryotic two-component signaling involves His-Asp-His-Asp multi-step phosphorelay (MSP). In *Arabidopsis thaliana*, cytokinin-mediated MSP signaling intermediates include histidine kinases (HKs), histidine phosphotransfer proteins (Hpts) and response regulators (RRs). The structure-function relationship of interaction between Hpt (e.g., AHP1) and RR (e.g., ARR4) is poorly understood.
- Using a homology model and yeast two-hybrid analysis, we identified key amino acids of ARR4 at AHP1- Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ interaction interface. Mutating them in *Arabidopsis* (*arr3,4,5,6,8,9* hextuple mutant background) and performing root length assays provided functional relevance, and co-immunoprecipitation (Co-IP) assay provided biochemical evidence for the interaction.
- The homology model mimics crystal structures of Hpt-RR complexes. Mutating selected interface residues of ARR4 either abolished or destabilized the interaction. D45A and Y96A mutations weakened interaction with AHP1, and exhibited weaker rescue of root elongation in the hextuple mutants. Co-IP analysis using cytokinin-treated transgenic *Arabidopsis* seedlings provided biochemical evidence for weakened AHP1-ARR4 interaction. The relevance of the selected residues for the interaction was further validated in two independent pairs of Hpt-RR proteins from *Arabidopsis* and rice (*Oryza sativa*).
- Our data provide evidence for a link between Hpt-RR interaction affinity and regulation of downstream functions of RRs. This establishes a structure-function relationship for the final step of a eukaryotic MSP signal cascade.

Keywords: AHP1, Arabidopsis histidine phosphotransfer protein 1; ARR4, Arabidopsis response regulator 4, Two-Component Signaling; Co-immunoprecipitation; Cytokinin Signaling Intermediates; Structural basis of AHP1-ARR4 interaction; Multi-Step Phosphorelay (MSP); Root Elongation Assay; Plant Hormone Signaling

Introduction

Two-component signaling (TCS) systems mediate a wide spectrum of signaling events in prokaryotic and eukaryotic organisms by sensing and responding to various signals. The

canonical TCS consists of a membrane-bound sensor histidine kinase (HK) that senses the signals and gets autophosphorylated on the conserved His residue in the kinase domain (Stock *et al.*, 2000). The signal is transmitted as a phosphoryl group to the conserved Asp residue in the receiver domain (RD) of a response regulator (RR). Compared to the prokaryotic TCS systems, the eukaryotic TCS system is more complicated because of the presence of a multi-step phosphorelay (MSP) (Appleby *et al.*, 1996). This is necessitated by the presence of RRs in the nucleus while the receptors occur on outer membranes. Therefore, the MSP signaling system follows a sophisticated His-Asp-His-Asp phosphorelay among the multiple signaling intermediates.

The signal transduction pathway of cytokinins, a major class of plant hormones, is an example of the MSP signaling system in plants (To & Kieber, 2008; Hwang et al., 2012). In Arabidopsis thaliana, the hybrid sensor kinase family consists of ARABIDOPSIS HISTIDINE KINASE 2 (AHK2), AHK3 and AHK4/CRE1/WOL1 that function as cytokinin receptors (Inoue et al., 2001; Suzuki et al., 2001), AHK1 which is a putative osmosensor (Tran et al., 2007) and CKI1 and AHK5 that are cytokinin-independent HKs (Desikan et al., 2008; Deng et al., 2010). Autophosphorylation of AHKs at the conserved His in the kinase domain initiates MSP, which is then relayed intramolecularly to the conserved Asp in the RD (Hwang et al., 2012). The RD transfers the phosphate group to Histidine phosphotransfer proteins (Hpts), namely, the Arabidopsis Histidine phosphotransfer Proteins (AHPs), which in turn, transmit the phosphoryl group to conserved Asp in the RD of Arabidopsis Response Regulators (ARRs) located mainly in the nucleus. The phosphorylation of ARRs results in their activation, which mediate cytokinin-regulated responses. Two families, namely, type-A and type-B ARRs are involved in this MSP. Arabidopsis has 10 type-A ARRs (ARR3-9 and ARR15-17) (Muller & Sheen, 2007), which are primary transcriptional targets of cytokinin signaling, being rapidly upregulated upon cytokinin treatment (Hwang & Sheen, 2001). There are 11 type-B ARRs (ARR1, ARR2, ARR10-14 and ARR18-21) that consist of an RD at the N-terminus and a DNA-binding domain at the C-terminus (Hosoda et al., 2002). They are transcriptional activators of cytokinin-regulated genes, including type-A ARRs, thereby functioning as positive regulators of cytokinin signaling (Hwang & Sheen, 2001). Furthermore, in depth molecular characterization of different cytokinin signaling intermediates helped to identify cognate Hpt, type-A RR and type-B RR proteins in rice (Tsai et al., 2012). Characterization of selected rice RRs has shown that they function in a manner similar to their Arabidopsis counterparts (Hirose et al., 2007).

The phosphorylated (activated) type-A ARRs negatively regulate cytokinin signaling and phosphorylation at the conserved Asp is a prerequisite for their function (Lee *et al.*, 2008). This also highlights that interaction of AHPs with type-A ARRs and phosphorelay from the former to the latter is a critical step for cytokinin signaling cascade. Several studies have validated the interaction and phosphotransfer between AHPs and type-A ARRs using yeast two-hybrid assay and monitoring the transfer of radioactively labeled PO_4^{3-} group (Imamura *et al.*, 1998; Mira-Rodado *et al.*, 2007).

Structural snapshots of the mechanistic basis of interaction and phosphotransfer between RDs and Hpts were obtained from protein complex crystal structures of TCS intermediaries. Examples are available from organisms belonging to various kingdoms, such as, CheA₃P1•P-CheY₆ from *Rhodobacter sphaeroides* (Bell et al., 2010), SLN1_{RD}-YPD1 and SLN1_{RD}-YPD1•Mg²⁺•BeF³⁻ from Saccharomyces cerevisiae (Xu et al., 2003; Zhao et al., 2008), as well as from the recently published AHK5_{RD}-AHP1•Mg²⁺ complex from *Arabidopsis* (Bauer et al., 2013). Importantly, AHK5_{RD}-AHP1 complex structure is the first crystal structure of a plant HK_{RD}-Hpt complex. However, it is noteworthy that the two complex structures from eukaryotes mentioned above only represent the 'Asp-His' interaction, which corresponds to the middle portion of His-Asp-His-Asp phosphorelay. The structural details of the final His-Asp step i.e., interaction and phosphotransfer between Hpts and RR_{RD} have not been studied so far. This could be partly because of the problems associated with the procurement of high yields of recombinant ARR proteins with significant purity (Verma et al., 2013). An alternative approach to address this knowledge gap could be generation of homology models using available structural information followed by structure-function analysis. A critical comparison of the structures of different receiver domains, e.g., CheY; RR (Escherichia coli) (Lee et al., 2001), SLN1_{RD}; HK_{RD} (S. cerevisiae) (Xu et al., 2003) and CKI1_{RD}, AHK5_{RD} (Arabidopsis, both are HK_{RD}) (Muller-Dieckmann et al., 1999; Pekarova et al., 2011) revealed that RDs from HKs and RRs possess similar $(\alpha/\beta)_5$ fold across kingdoms. This signifies that the available crystal structures can serve as templates for building computational models of HK_{RD} and RR_{RD}.

In this study, we generated an *in silico* model of AHP1 complexed with 16-175 amino acid region of ARR4 (henceforth, this region will be referred to as Δ ARR4⁽¹⁶⁻¹⁷⁵⁾) to interpret the final step (Hpt–RR) of cytokinin signal transduction. Mutations in key amino acid residues of

ARR4 identified from AHP1– Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ interaction interface resulted in either abolition or weaker interactions with AHP1 in a yeast two-hybrid assay. Interactions of cognate protein pairs from *Arabidopsis* and rice were also tested. *In planta* analyses of two mutants of ARR4, which showed weakened interaction with AHP1, also showed weakened cytokinin signaling. The mutants showed weaker rescue of root elongation, a cytokinin-mediated developmental event, as compared to wild-type ARR4. Co-IP analysis provided a biochemical explanation of the observed differences in root elongation. Our results help to explain the structure-function relationship of AHP1-ARR4 interaction, which is a critical step in cytokinin signaling.

Materials and Methods

Plant materials and growth conditions

Arabidopsis thaliana ecotype Columbia-0 (Col-0) plants were used as wild-type control for *in planta* experiments. Plants were grown at 23 °C under long-day conditions (16 h light/8 h dark). All transgenic plant lines were generated in *arr3,4,5,6,8,9* hextuple mutant background in which six type-A ARRs (*ARR3, ARR4, ARR5, ARR6, ARR8* and *ARR9*) were knocked-out (To *et al.*, 2004). Seeds of the hextuple mutant (CS25279) were obtained from the Arabidopsis Biological Resource Centre (http://www.abrc.osu.edu).

For seedling assays, the seeds were surface-sterilized and sown on Murashige and Skoog (MS) semi-solid medium (Caisson LABS) containing 1X MS, 0.05% MES, 1% sucrose and 0.6% GelriteTM (<u>https://www.plantmedia.com/</u>), unless stated otherwise. They were subsequently stratified at 4 °C for 3 d in the dark followed by incubation at 23 °C under constant white light (~50 μ E/m²/s) (To *et al.*, 2004).

Plasmid construction

Full-length cDNAs encoding AHP1, ARR4, Δ ARR4⁽¹⁶⁻¹⁷⁵⁾, AHP2, ARR5, OsHP1 and OsRR6 were amplified by PCR and cloned into pJET vector (Thermo Scientific). The various mutant versions of ARR4, ARR5 and OsRR6 were generated by site-directed mutagenesis approach. All clones were verified by sequencing. For yeast two-hybrid assay, AHP1, AHP2 and OsHP1 were cloned into HA tag containing pGADT₇ vector (Clontech), whereas, ARR4, Δ ARR4⁽¹⁶⁻¹⁷⁵⁾, ARR5, OsRR6 and all the mutants of ARR4, ARR5 and OsRR6 were cloned into myc tag containing pGBKT7 vector (Clontech). For Bimolecular Fluorescence

Complementation (BiFC), modified pSAT1 vectors were used in which the expression cassette of pSAT1 including the 35S promoter and the N/C-EYFP was fused to pGreen binary vector HY105. AHP1 was cloned at the C-terminal of cEYFP, while, ARR4, ARR4^{D45A} and ARR4^{Y96A} were cloned at C-terminal of nEYFP. For GFP localization, ARR4, ARR4^{D45A} and ARR4^{Y96A} were fused at the N-terminus of GFP driven by 35S promoter. For generation of transgenic plants, wild-type ARR4 and the two mutants were cloned into the pGreen 35S vector possessing an HA tag at the 3' end.

Generation of transgenic plants

The hextuple mutant (*arr3*, *4*, *5*, *6*, *8*, *9*) seeds obtained from ABRC were germinated in soil and PCR-based screening was done to confirm homozygosity and presence of T-DNA inserts in the six genes. The hextuple mutant did not show phenotypic changes compared to the wild-type. Seeds from homozygous mutant plants were collected and used for subsequent experiments. Transgenic plants were generated by introducing relevant constructs into the hextuple mutant plants via *Agrobacterium tumefaciens*-mediated floral dip method (Clough & Bent, 1998). Selection was done using BASTA (2 ml/l) spray followed by genotyping-PCR of the survivors for confirmation. The transgenic lines were taken to T3 generation for homozygosity before they were used for analyses.

Homology modeling and identification of AHP1 - ARR4 contact points

Full-length protein sequence of AHP1, Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ and OsRR6 were submitted to SWISS MODEL (http://www.swissmodel.expasy.org) and the coordinates were generated. The models of the individual proteins were superimposed on the SLN1_{RD}-YPD1 complex crystal structure (PDB id: 2R25) (Zhao *et al.*, 2008) using COOT software (Emsley & Cowtan, 2004) to generate the complex model coordinates. Subsequently, these coordinates were energy minimized. These model coordinates were used to calculate the interaction interface of the two proteins by the CCP4 program.

Yeast two-hybrid assay

The yeast two-hybrid experiment was performed as per the manufacturer's protocol for Matchmaker GAL4-based two-hybrid system (Clontech). Equal amounts of AD constructs were mixed with the corresponding BD constructs in separate reactions and the mixtures were introduced into AH109 yeast strain. The transformed yeast cells were selected on Leu⁻/Trp⁻/His⁻ and also on Leu⁻/Trp⁻/His⁻/Ade⁻ in order to screen for stronger interactions.

Aliquots plated on Leu⁻/Trp⁻ medium were used as transformation control. Simultaneously, the cells were diluted 1:10 and 1:100 fold using 0.9% NaCl and plated on Leu⁻/Trp⁻ and Leu⁻/Trp⁻/His⁻ media. The plates were incubated for 3 to 4 d at 30 °C and then photographed.

For AHP1 and ARR4 interaction, the presence of both the proteins in the respective yeast cells were detected by Western blot analysis of the transformed cells using anti-HA and antimyc antibodies (Santa Cruz) for AD and BD vector clones, respectively. Total proteins were extracted from overnight yeast cultures (3 ml, with cell density normalized to the culture with lowest OD_{600}) as described (Riezman *et al.*, 1983). To ensure equal loading, each extract was subjected to 50% trichloroacetic acid (TCA) precipitation and the resultant protein pellets were resuspended in 100 µl of 2x SDS loading dye prior to SDS PAGE.

Bimolecular fluorescence complementation (BiFC)

The fusion constructs of cEYFP-AHP1, nEYFP-ARR4, nEYFP-ARR4^{D45A} and nEYFP-ARR4^{Y96A} were introduced into *Agrobacterium tumefaciens* strain GV3101. The colonies were grown overnight and the next day 1 ml culture pellet for each construct was resuspended in infiltration buffer (10 mM MES pH 5.6, 10 mM MgCl₂ and 100 μ M acetosyringone) to get a final OD₆₀₀ of 0.6. Equal volumes of infiltration solution of the pair of constructs to be tested for interaction were mixed and incubated for 3 h at room temperature with gentle shaking. After incubation, leaves from three-week-old *Nicotiana benthamiana* plants were infiltrated with the *Agrobacterium* mixture on their abaxial surfaces using a syringe (Walter *et al.*, 2004). The leaves were examined for YFP signal three days post-infiltration using Carl Zeiss 510 Meta laser scanning confocal microscope (http://www.zeiss.de/axiovert200) with excitation at 514 nm. All images were recorded with the same settings. The signal intensity was measured using ImageJ software (National Institute of Health, USA).

Protoplast isolation and transfection for GFP localization

The protoplasts were extracted from leaves of 3- to 4-week-old *Arabidopsis* (Col-0) plants (Yoo *et al.*, 2007). For each reaction, approximately 2×10^5 protoplasts were transfected with $15 - 20 \mu g$ of plasmid DNA corresponding to 35S::ARR4-GFP, $35S::ARR4^{D45A}-GFP$, $35S::ARR4^{Y96A}-GFP$ and then incubated for 12 to 16 h at 25 °C in the dark. The GFP signals were monitored by laser scanning microscopy as above, but with excitation at 488 nm. Expression of 35S::GFP was used as a control.

RNA extraction, cDNA synthesis and quantitative real time-PCR (qRT-PCR)

Total RNA was extracted from10-day-old seedlings of wild-type (Col), hextuple mutant and selected transgenic lines of ARR4 and the two mutants using TRIzol[®] reagent (Life Technologies) as per the manufacturer's protocol. About 1 μ g of extracted RNA for each sample was used for reverse transcription as per the manufacturer's instructions using Maxima First strand cDNA synthesis kit (Thermo Scientific). The 10 μ l qRT-PCR reaction mixture included 1 μ l cDNA (diluted five folds), 0.2 μ l of each primer, 5 μ l 2x KAPA SYBR[®] Master Mix (KAPA Biosystems) and sterile water. PCR was performed using StepOneTM Real-Time PCR systems (v2.1; Applied Biosystems) and the PCR conditions were: denaturation at 95 °C for 20 s; 40 cycles of denaturation at 95 °C for 3 s and annealing and extension at 60 °C for 30 s. Amplification of *TUB2* gene was used as an internal control for normalization. StepOne TM software (v2,1; Applied Biosystems) was used for data analysis from two independent biological replicates.

Root elongation assay

Arabidopsis seedlings were grown in culture petri plates containing MS medium supplemented with the indicated concentrations of BA or 0.1% DMSO (solvent). The plates were incubated vertically for 10 d in continuous light (To *et al.*, 2004). The positions of roots were marked on the plates on 4th and 9th day and the plates were photographed on the 10th day. The root growth of root between days 4 and 9 was measured using ImageJ software. Data presented are means \pm SE from at least 30 seedlings per transgenic line per treatment with at least two independent transgenic lines for each construct.

ARR7 response to cytokinin treatment

For treatment with cytokinin, the seedlings were grown on horizontal MS plates with 0.5% Gelrite. 10-day-old seedlings were transferred to 1X liquid MS supplemented with 50 nM BA and samples were collected at 0, 30 and 60 min of treatment (To *et al.*, 2004). RNA extraction, cDNA synthesis and qRT PCR were performed as mentioned above. Expression of *ARR7* was analyzed from two independent biological replicates.

Analysis of protein levels in transgenic lines

Total protein was extracted from 10-day-old seedlings of the different transgenic lines used for the study using 100 mM Tris HCl (pH 8.0), 150 mM NaCl, 5 mM EDTA, 5 mM EGTA, 10 mM DTT, 0.5% Triton X-100 and 1X Complete protease inhibitors (Roche Applied

Sciences) (To *et al.*, 2007). Protein extracts were separated by SDS-PAGE and transferred to PVDF membrane (BIORAD). HA-tagged proteins were detected using anti-HA antibody (Santa Cruz) and visualized by chemiluminescent detection (Thermo SCIENTIFIC) by autoradiography. Subsequently, the membranes were stained with Ponceau and Rubisco protein band was used as loading control.

For examining the protein stability, 10-day-old seedlings were treated with 200 μ M cycloheximide for different time points (0, 30, 60, 90, 120 min) and western blot was done as mentioned above. Signals were quantified using Image J (National Institute of Health).

Co-IP of ARR4-HA, ARR4^{D45A}-HA and ARR4^{Y96A}-HA using recombinant GST-AHP1

14-day-old seedlings of one representative transgenic line each of wild-type ARR4 and two mutant versions of ARR4 were treated with 10 nM BA for 45 min. Subsequently, total protein was extracted in 400 mM sucrose, 50 mM Tris HCl (pH 7.5), 10% glycerol and 2.5 mM EDTA. Recombinant GST-AHP1 was expressed and purified as described (Verma *et al.*, 2013), but the GST tag was not cleaved. The protein concentration for recombinant GST-AHP1 and seedling extracts were estimated using Direct Detect[®] Spectrometer (MERCK MILLIPORE). 2 µg of GST-AHP1 protein was immobilized onto 30 µl of Glutathione Sepharose 4B (GE Healthcare) beads. Subsequently, about 20 µg of seedling protein extracts from representative transgenic lines were added to the immobilized GST-AHP1 in independent reactions and incubated for 2 h at 4 °C in the Co-IP buffer (50 mM Tris HCl, 100 mM NaCl, 10% glycerol, 5 mM EDTA, 0.1% Triton X-100). Subsequently, the beads were washed 3 times with Co-IP buffer for 10 min and subjected to SDS-PAGE. Proteins were transferred to PVDF membrane (BIORAD) and probed with anti-HA (Roche) antibody and reprobed with anti-GST (Sigma) antibody.

Results

Generation of a homology model for AHP1-ΔARR4⁽¹⁶⁻¹⁷⁵⁾ complex

In order to understand the structural basis of AHP1-ARR4 interaction, we generated a homology model for AHP1- Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ using SLN1_{RD}-YPD1•Mg2⁺•BeF³⁻complex crystal structure (PDB id: 2R25) as the template (Zhao *et al.*, 2008). The sequence similarity of AHP1 with YPD1 is ~44% (Fig. 1a). The sequence similarity of 16-175 amino acid region of

ARR4 (putative receiver domain) with the receiver domain (R1; 1086-1221) of SLN1 is ~42% (Fig. 1b). The histidine residue required for phosphotransfer (His79 for AHP1; highlighted in yellow) and all the functionally critical amino acid residues of ARR4 including the two aspartates (Asp41; highlighted in blue and Asp95; highlighted in yellow) and a lysine (Lys147; highlighted in purple) required for phosphorylation are highly conserved. Further, amino acid 32 to 172 of ARR4 aligned with SLN1_{RD}. This was in agreement with Imamura *et al.* (1998) who depicted that Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ was capable of receiving phosphoryl group from bacterial Hpt domain under *in vitro* conditions. Together, this prompted us to use Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ for subsequent analyses.

For generating the complex model AHP1- Δ ARR4⁽¹⁶⁻¹⁷⁵⁾, we employed the crystal structure of AHP1 (PDB id: 4EUK) (Bauer *et al.*, 2013). It consists of six α -helices in which four helices bundle to form a central core (shown in red in Fig. 1c). His79, the conserved histidine required for phosphorylation extends from one of the helices of the bundle (inset in Fig. 1c). However, for $\Delta ARR4^{(16-175)}$, a predicted 3-dimensional structured was used. The predicted structure depicted a central core of five parallel β -strands enveloped by five α -helices in groups of two and three, giving an $(\alpha/\beta)_5$ topology (shown in green in Fig. 1c). The conserved residues such as Asp41, Asp95 and Lys147 form a pocket (inset in Fig. 1c) that resembles the phosphate-binding pocket of other response regulators (Bourret, 2010). Moreover, the $(\alpha/\beta)_5$ fold exhibited by $\Delta ARR4^{(16-175)}$ model is similar to the structures of other receiver domains, such as, CKI1_{RD} from Arabidopsis (PDB id: 3MMN; ~49% sequence similarity) (Pekarova et al., 2011), CheY₃ from Vibrio cholerae (PDB id: 3TO5; ~46% sequence similarity), AHK5_{RD} from Arabidopsis (PDB id: 4EUK; ~39% sequence similarity) (Bauer et al., 2013) and SLN1_{RD} from S. cerevisiae (PDB id: 2R25; 27% sequence similarity) (Zhao et al., 2008). The AHP1- Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ complex model clearly depicted that residues from the two proteins that are required for phosphotransfer are in close proximity at the interface (inset in Fig. 1c).

AHP1-ΔARR4⁽¹⁶⁻¹⁷⁵⁾ complex model mimics natural Hpt-RR complexes

The computational model of AHP1– Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ complex showed significant alignment with the complex crystal structures of CheA₃P1-CheY₆ (PDB id: 3KYI; rmsd of 3.2 Å for 191 C α atoms) (Bell *et al.*, 2010), SLN1_{RD}-YPD1 (PDB id: 2R25; rmsd of 1.49 Å for 223 C α atoms) (Zhao *et al.*, 2008) and AHK5_{RD}–AHP1 (PDB id: 4EUK; rmsd of 1.72 Å for 252 C α atoms) (Bauer *et al.*, 2013) (Fig. 2a). Interestingly, the independent proteins better superimpose than the complex. It indicates possible differences in the relative disposition of the two molecules in the complex compared to the complexes of its homologs. Fig. 2(b) shows the structure based alignment which reveals the conserved position of the key amino acids among these structural homologs. Taken together, it suggests that the *in silico* AHP1- Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ complex model mimics the natural complexes.

We next investigated if the Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ region is functional by examining its ability to interact with AHP1 in the yeast two-hybrid system. Both full-length ARR4 and Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ were able to interact with AHP1, because yeast cells harboring AD-AHP1 and BD-ARR4 or BD- Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ grew on Leu⁻/Trp⁻/His⁻/Ade⁻ medium (Fig. 2c). On the contrary, no growth was observed for yeast cells containing empty AD vector with BD-ARR4 or BD- Δ ARR4⁽¹⁶⁻¹⁷⁵⁾. Normal growth was observed on Leu⁻/Trp⁻ confirming efficient yeast transformation.

Identification of amino acid residues of ARR4 which affect its interaction with AHP1

To identify the key amino acids involved in AHP1–ARR4 interaction, intermolecular interaction analysis was conducted (<3.8 Å) with the AHP1– Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ complex (Table S1). Based on this analysis, 12 residues of ARR4 were identified to interact with AHP1 in which 5 amino acids namely, Asp45, Arg51, Tyr96, Cys97 and Pro148 has more interactions, and were selected for subsequent studies [Fig 3(a) shows mapping of these residues on AHP1– Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ complex model]. It is noteworthy that the selected interacting residues (except Cys97) are conserved among all the 10 members of type-A ARR family (Fig. S1). Cys97 was conserved in 6 of the 10 members.

To test if the ARR4 amino acid residues selected using *in silico* approaches are indeed involved in interaction with AHP1, each of the five amino acids was individually mutated to Ala in full-length ARR4. The individual mutants were tested for interaction with AHP1 in the yeast two-hybrid system (Fig. 3b). Full-length wild-type ARR4 showed significant interaction with AHP1 as the yeast cells harboring the two plasmids grew well on Leu⁻/Trp⁻/His⁻/Ade⁻. On the contrary, ARR4^{D45A}, ARR4^{R51A} and ARR4^{Y96A} mutants showed reduced interaction with AHP1, since yeast cells grew only on Leu⁻/Trp⁻/His⁻ and not on Leu⁻/Trp⁻/His⁻/Ade⁻. Besides, among the three mutants, the strength of interaction was in the following sequence: ARR4^{R51A} > ARR4^{Y96A} > ARR4^{D45A}, as revealed by serial dilution (Fig. 3b). However, Cys to Ala mutation at 97th position had no major impact on ARR4 interaction with AHP1 because ARR4^{C97A} grew on Leu⁻/Trp⁻/His⁻/Ade⁻. Interestingly, an abolition of interaction between the two proteins was observed with ARR4^{P148A} mutant. The presence of

AHP1, ARR4 and different ARR4 mutant proteins in yeast cells were detected by western blot analysis using anti HA (for AD:AHP1) and anti c-myc (for BD:ARR4) antibodies (Fig. 3c).

We further explored the impact of weakened AHP1-ARR4 interaction on the downstream functions of ARR4 using ARR4^{D45A} and ARR4^{Y96A} mutants for further analyses. These two mutants showed reduced interaction affinity towards AHP1 in the yeast two-hybrid assays. ARR4^{P148A} was not used, because it lead to abolition of interaction. To investigate if the mutations have affected the subcellular localization of the protein, Arabidopsis mesophyll 35S::ARR4^{D45A}-GFP with 35S::ARR4-GFP, protoplasts transfected and were 35S::ARR4^{Y96A}-GFP and examined for GFP signals. For all three proteins, namely, wild-type ARR4, ARR4^{D45A} and ARR4^{Y96A}, we observed a strong signal emerging predominantly from the nucleus, with weak or negligible GFP signal in the cytoplasm (Fig. S2). This clearly revealed that point mutations have not affected the protein localization, since GFP signal of the two mutants overlapped completely with that of wild-type ARR4. Moreover, the localization pattern of proteins coincides with the published information (Sweere et al., 2001), thereby further validating our observations. The expression of 35S::GFP was used as the transformation control.

To further verify the weakening of interaction due to mutation of ARR4 at Asp45 and Tyr96 to Ala, ARR4^{D45A} and ARR4^{Y96A} mutants were tested for interaction with AHP1 in *Nicotiana benthamiana* leaves using BiFC. The YFP signals were observed for all three, namely, wild-type and two variants of ARR4, but the signal intensity was maximum for ARR4 followed by ARR4^{Y96A} and then ARR4^{D45A} (Fig. 3d,e). This further confirmed that the two proteins were able to interact despite the mutations, however, interaction affinity was in the following declining order ARR4 > ARR4^{Y96A} > ARR4^{D45A}, thereby validating the yeast two-hybrid results. No signals were observed when the two vectors containing only the YFP fragments were infiltrated ensuring that the signals obtained for AHP1-ARR4 interaction were not due to non-specific contacts (Fig. 3d).

The selected Asp and Tyr residues of ARR4 were conserved in ARR5 (Fig. S1) as well as in OsRR6, a rice type-A response regulator (Fig. S3). Further, the superposition of the computational model of OsRR6 onto ARR4 model clearly showed that the selected residues of OsRR6 exhibit similar three dimensional orientation as observed for the corresponding

ARR4 residues (Fig. 3f). Hence, to further examine if the mutation of Asp36 and Tyr88 in ARR5 and Asp61 and Tyr104 in OsRR6 can affect the interaction with their corresponding Hpt proteins, we tested the interaction of ARR5, OsRR6 and their mutants with AHP2 and OsHP1, respectively. Wild-type ARR5 and OsRR6 showed significant interaction with AHP2 and OsHP1, respectively, because yeast cells harboring the plasmid pairs showed proper growth on Leu⁻/Trp⁻/His⁻ (Fig. 3g). ARR5^{D36A} or ARR5^{Y88A} did not show any interaction with AHP2. On the other hand, OsRR6^{D61A} showed weakened interaction with OsHP1 (exhibited by serial dilution), whereas OsRR6^{Y104A} failed to interact with OsHP1. These data further validated the involvement of the selected amino acids in mediating the interaction with Hpts.

D45A and Y96A point mutations did not alter ARR4 protein stability in Arabidopsis

As a definitive validation of the functional relevance of weakened AHP1–ARR4 interaction in the native environment, we generated transgenic *Arabidopsis* lines overexpressing 35S::ARR4-HA, 35S::ARR4^{D45A}-HA and 35S::ARR4^{Y96A}-HA in the type-A ARR hextuple mutant (*arr3,4,5,6,8,9*) background (To *et al.*, 2004). For each construct, we generated at least five independent transgenic lines. The hextuple mutant background was chosen to overcome the functional redundancy shown by type-A ARRs (Fig. S4). Two homozygous transgenic lines were selected each for 35S::ARR4-HA: #2-2-ARR4-HA, #6-2-ARR4-HA, , 35S::ARR4^{Y96A}-HA: #1-3-ARR4^{Y96A}-HA, #3-1-ARR4^{Y96A}-HA, and 35S::ARR4^{D45A}-HA: #3-1-ARR4^{D45A}-HA, #5-1-ARR4^{D45A}-HA. Comparable levels of relative *ARR4* transcripts and corresponding proteins were detected in all the selected transgenic lines (Fig. 4a,b).

To test if the mutations have altered the relative protein stability of the HA-tagged fusion proteins *in planta*, we examined the protein turnover rates in one representative transgenic line each for #2-2-ARR4-HA, #3-1-ARR4^{D45A}-HA and #3-1-ARR4^{Y96A}-HA. Detection of relative protein levels at 30, 60, 90 and 120 min after cycloheximide treatment showed that the protein turnover rates for the three proteins were comparable, thereby confirming that mutations have not affected the stability of the protein (Fig. 4c-f).

D45A and Y96A mutations weakened ARR4-mediated cytokinin response

To investigate the effect of mutations on ARR4-mediated cytokinin functions, we employed the root elongation bioassay, which is one of the best-characterized responses of cytokinin (To *et al.*, 2004). Examination of wild-type (Col-0) and hextuple mutant seedling roots at 0, 5, 10, 50 and 100 nM N^6 -benzyladenine (BA) concentrations showed that the root length of

hextuple mutants was significantly inhibited at 5 and 10 nM BA, compared to Col-0 (Fig. S5). Nevertheless, at 50 and 100 nM concentrations, even wild-type root growth was retarded. The data are consistent with earlier reports (To et al., 2004). Consequently, homozygous transgenic lines corresponding to 35S::ARR4-HA (#2-2-ARR4-HA, #6-2-(#3-1-ARR4^{D45A}-HA, #5-1-ARR4^{D45A}-HA) 35S::ARR4^{D45A}-HA ARR4-HA,), and 35S::ARR4^{Y96A}-HA (#1-3-ARR4^{Y96A}-HA, #3-1-ARR4^{Y96A}-HA,) were tested for primary root elongation at 5 nM and 10 nM BA concentrations (Fig. 5a,b). An additional transgenic line for each of the three constructs, viz. #8-1-ARR4-HA, #4-1-ARR4^{D45A}-HA and #5-1-ARR4^{Y96A}-HA was also included for the assay (Fig. S6). We observed that in the absence of exogenous cytokinin i.e., 0 nM BA, primary root growth was similar for all seedlings including wild-type (Col-0) and hextuple mutants. On the contrary, at 5 nM and 10 nM BA, 35S::ARR4-HA exhibited longer primary roots than the two weaker interacting mutants and the hextuple knock-out seedlings.

To further examine the impact of mutations on ARR4 functions, we treated the transgenic seedlings of #6-2-ARR4-HA, #3-1-ARR4^{D45A}-HA, #5-1-ARR4^{D45A}-HA, #1-3-ARR4^{Y96A}-HA and #3-1-ARR4^{Y96A}-HA with 50 nM BA and analyzed the expression levels of *ARR7*, an early cytokinin response gene. To *et al.*, (2004) have shown that upon cytokinin treatment transcript levels of *ARR7* were much higher in the hextuple mutant than the wild-type reaching to maximal levels at 30 min. This is due to the lack of negative feedback regulation offered by other type-A ARRs. *ARR7* levels were significantly lower in #6-2-ARR4-HA as compared to the hextuple mutant, #3-1-ARR4^{D45A}-HA, #5-1-ARR4^{D45A}-HA, #1-3-ARR4^{Y96A}-HA and #3-1-ARR4^{Y96A}-HA at 30 min of BA treatment (Fig. 5c). This clearly showed that wild-type ARR4 was able to significantly suppress *ARR7* expression, whereas the two mutants did not significantly suppress *ARR7* transcript levels. Together, these results highlighted that wild-type ARR4 was able to rescue hextuple mutant more efficiently, as compared to the minimal rescue exhibited by the two weak interacting mutants of ARR4.

D45A and Y96A mutations weakened ARR4 binding to AHP1

To investigate if the observed phenotypic differences between the wild-type and mutant versions of ARR4 can be attributed to the weak interaction between AHP1 and ARR4 mutants, we performed a Co-IP experiment using recombinant GST-AHP1 and HA-tagged ARR4 from a representative transgenic line for each (#2-2-ARR4-HA, #3-1-ARR4^{D45A}-HA and #3-1-ARR4^{Y96A}-HA). Since the phenotypic differences were observed in the presence of

cytokinin, 14-day-old seedlings were treated with 10 nM BA for 45 min. GST-AHP1 showed reduced binding to mutated forms of ARR4 compared to the wild-type ARR4 (Fig. 6). The efficiency of pull-down was in the following declining order: ARR4-HA, ARR4^{Y96A}-HA and ARR4^{D45A}-HA, which is consistent with our yeast two-hybrid and BiFC results (Fig. 3b,e). The Co-IP data clearly indicate that a tight AHP1-ARR4 binding is critical for efficient progression of cytokinin signaling.

Discussion

Characterization of cytokinin signaling components and analyses of associated transcriptional networks have enhanced our understanding of the molecular functioning of the phosphorelay. Nevertheless, structural knowledge of the key signaling steps is essential for a better understanding of the signaling pathways. Although, crystal structures of cytokinin signaling intermediaries, such as CKI_{RD} (Pekarova *et al.*, 2011) and AHK5_{RD}-AHP1 complex (Bauer *et al.*, 2013) contributed significantly by providing structural snapshots of the HK_{RD} and mode of phosphotransfer from AHKs to AHPs, interaction between AHPs and ARRs have not been studied structurally. The fact that transfer of phosphoryl group from AHPs to type-A ARRs is an essential step for negative regulation of cytokinin signaling further necessitates the structural examination of the interaction. Our study provides a computational model of AHP1– Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ complex and highlights how it can be used to understand the structure-function relationship of the interactions between an Hpt and RR_{RD} in higher eukaryotes.

The AHP1– Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ complex model provides significant insights into the mechanistic aspects of interaction between AHP1 with ARR4. It demonstrated that similar to other Hpt-RD complex structures, the conserved His79 from AHP1 and the 'phosphate-binding pocket' from ARR4 are present in close proximity and hence can efficiently execute the phosphotransfer process (Fig. 1). Model-based identification of amino acid residues of AHP1 from the interaction interface of AHP1– Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ complex revealed the presence of amino acids that are involved in hydrophobic interactions and hydrogen bond formation (Table S1). Furthermore, the destabilization of AHP1-ARR4 interaction caused by mutations in ARR4 at Asp45, Arg51, Tyr96, Pro148 and Lys150 positions have helped in highlighting the amino acid residues of ARR4 critical for its interaction. Furthermore, OsHP1-OsRR6

interaction was similarly destabilized when the conserved Asp and Tyr residues were mutated (Fig. 3), thereby extending the validity of the study to a monocotyledonous species as well.

The poor rescue of hextuple knock-out plants by ARR4^{D45A} and ARR4^{Y96A} in the root elongation assay and *ARR7* expression assay (Fig. 5) clearly indicated that the mutations, besides affecting the interaction ability of ARR4 with AHP1, also perturbed ARR4 functions. The possibility that observed malfunctioning of ARR4 mutants could be due to structural aberrations caused by the mutations was invalidated by the ability of ARR4^{D45A} and ARR4^{Y96A} to interact with AHP1 (Fig. 3), suggesting that mutations had not affected the overall fold of ARR4. Also, predominant nuclear localization of ARR4^{D45A}-GFP and ARR4^{Y96A}-GFP, similar to ARR4-GFP (Fig. S2) showed that mutations had not disturbed the subcellular localization of ARR4. Moreover, comparable protein turnover rates of the wild-type and mutant forms of ARR4 clearly highlighted that mutations have not affected the stability of ARR4 protein (Fig 4).

Our data of weakened interaction between AHP1 and ARR4 leading to altered root elongation response supports the view that a close and tight contact between the two signaling intermediates is critical for efficient phosphotransfer and mediating downstream functions. Similarly, earlier studies of SLN1_{RD}-YPD1 complex structures of yeast in the absence and presence of Mg²⁺ and BeF³⁻ showed that YPD1 undergoes a rigid-body shift for alignment of conserved His within ideal distance of, and in linear O-P-N geometry with, conserved Asp of SLN1_{RD} for efficient phosphotransfer (Zhao *et al.*, 2008). Furthermore, substitution of a key Met in bacterial CheY₆ at the interface of CheA₃P1–CheY₆ showed reduction in interaction and rate of phosphotransfer between them (Bell *et al.*, 2010). The Co-IP analysis in the present study showed that even after cytokinin treatment, there was only weak interaction between the mutant versions of ARR4 and AHP1 (Fig. 6), suggesting that this might be occurring *in vivo* as well and hence the resultant phenotype in ARR4^{D45A} and ARR4^{Y96A} mutants (Fig. 5). Based on this, we can speculate that weaker interaction of ARR4 with AHP1 could result in slow rate of phosphotransfer resulting in the altered root elongation phenotype as shown in the model (Fig. 7).

In conclusion, our data provide evidence for a link between AHP1–ARR4 interaction and ARR4-mediated cytokinin signal progression.. These findings highlight the intricacies of the mechanistic basis of phosphoryl group transfer from Hpt to RR and regulatory functions of

RRs. Hence, our study establishes a structure-function relationship for the final step of a eukaryotic MSP signal cascade.

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Supporting Information

Fig. S1. Multiple sequence alignment of ARR4 with other members of type-A ARR family.

Fig. S2. Subcellular localization of GFP fusion proteins of ARR4, ARR4^{D45A} and ARR4^{Y96A} in *Arabidopsis* mesophyll protoplasts.

Fig. S3. Sequence alignment of $\triangle ARR4^{(16-175)}$ with rice type-A RR, OsRR6.

Fig. S4. Validation of type-A ARR hextuple knock-out plants.

Fig. S5. Hextuple mutants are more sensitive to cytokinin than wild-type (Col-0).

Fig. S6. Root elongation assay data for additional transgenic lines (Supplement to Fig. 5).

Table S1. Amino acid residues involved in AHP1- Δ ARR4⁽¹⁶⁻¹⁵⁾ interaction at 3.8 Å intermolecular distance.

Figure Legends

Figure 1. Generation of a homology model of AHP1- Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ complex. (a) Sequence alignment of AHP1 with its homolog from *S. cerevisiae*, YPD1. The conserved His required for phosphorylated is highlighted in yellow. (b) Sequence alignment of 16-175 amino acid region of ARR4 with SLN1_{RD}, receiver domain of *S. cerevisiae* HK protein SLN1. The conserved phosphate-binding pocket residues i.e., the two aspartates (in blue and yellow) and lysine (in purple) are highlighted. (c) Overall view of the computational model of AHP1- Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ complex, in which AHP1 is depicted in red and Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ is depicted in

green. The inset shows 3-dimensional orientation of conserved residues required for phosphorylation i.e., His79 from AHP1 and Asp41, Asp95 and Lys147 from ARR4. Color coding is the same as in sequence alignment (a) and (b).

Figure 2. Validation of the AHP1- Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ homology model by its comparison with the existing natural complex structures. (a) The C α superposition of the AHP1- Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ *in silico* complex model with its homologues CheA₃P1-CheY₆ (orange), SLN1RD-YPD1 (blue) and AHK5RD–AHP1 (magenta). The structural alignment were carried out in PyMol (DeLano, 2002). (b) The structure-based sequence comparison of ARR4 and its homologs. Most of the substitutions are carried with ARR4 and thus we compared its sequences with its structural homologs. The structural alignment was performed using the program Coot (Emsley & Cowtan, 2004). Highly conserved and conserved residues are highlighted. The secondary structures of ARR4 are provided on the top. This figure was prepared using the program ESPript (Gouet *et al.*, 1999). (c) Both full-length ARR4 and Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ showed interaction with AHP1 in the yeast two-hybrid system. Growth of yeast cells expressing AD:AHP1 and BD:ARR4 or BD: Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ was examined on selection medium lacking Leu, Trp, His and Ade. Selection medium lacking Leu and Trp was used for checking transformation efficiency. The empty pGADT7 vector was used as a negative control. AD: Gal4 activation domain; BD: Gal 4 DNA-binding domain.

Figure 3. Identification of ARR4 residues present at AHP1- Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ complex interface. (a) ARR4 residues Asp45, Arg51, Tyr96, Cys97, Pro148 mapped onto AHP1- Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ model. Cys97 is not shown for clarity (AHP1: red, Δ ARR4⁽¹⁶⁻¹⁷⁵⁾: green). (b) Yeast two-hybrid analysis of AD:AHP1 interaction with BD:ARR4 mutants. pGBKT7 vector was used as the control. 10⁰, 10⁻¹ and 10⁻² represent no dilution, 10x, 100x dilutions, respectively. (AD: Gal4 activation domain; BD: Gal4 DNA-binding domain). (c) Western blot detection of AD:AHP1 and BD:ARR4 in yeast cells using anti-HA and anti-c-myc antibodies. Line numbers from one to six correspond to yeast cells harboring AHP1 and ARR4 or different mutants of ARR4, *viz*. ARR4^{D45A}, ARR4^{R51A}, ARR4^{Y96A}, ARR4^{C97A} and ARR4^{P148A}, respectively (similar to the numbering in Fig 3b). (d) BiFC interactions of ARR4 and mutants with AHP1. Confocal images of *N. benthamiana* leaf cells coexpressing YFP-C:AHP1 with either YFP-N:ARR4 or YFP-N:ARR4^{D45A} or YFP-N:ARR4^{Y96A}. Column headings: Yellow Channel - reconstituted YFP fluorescence, Transmitted - bright-field images, Overlay – superimposition of both channels. (e) Relative quantification of the YFP fluorescence from AHP1 interaction with ARR4, ARR4^{D45A} and ARR4^{Y96A} using ImageJ (NIH, USA). Data are mean \pm SE of signal intensities (28-30 spots per sample), with ARR4 set as 100%. (f) (f) Overlay of the computational model of OsRR6 (cyan) onto Δ ARR4⁽¹⁶⁻¹⁷⁵⁾ (green). The Asp45, Tyr96 and Pro148 of ARR4 selected for analysis were also conserved in OsRR6 (Asp61, Tyr104 and Pro156, highlighted in red). (g) Yeast two-hybrid interaction analysis of AD:AHP2 and AD:OsHP1 with wild-type and mutants of BD:ARR5 and BD:OsRR6, respectively. 10⁰, 10⁻¹ and 10⁻² represent no dilution, 10x and 100x dilutions, respectively. (AD: Gal4 activation domain; BD: Gal4 DNA-binding domain).

Figure 4. Detection of relative transcript and protein levels and analysis of protein turnover rates in transgenic lines. (a) The transcript levels of *ARR4*, *ARR4*^{D45A} and *ARR4*^{Y96A} were detected in the different transgenic lines used in the study. Data presented are mean \pm SD from two independent biological replicates. (b) The protein levels of ARR4-HA, ARR4^{D45A}-HA and ARR4^{Y96A}-HA were detected in the respective transgenic lines used for the study. Rubisco protein was used as the loading control. (c) The protein levels of ARR4-HA, ARR4^{D45A}-HA and ARR4^{Y96A}-HA were detected in 10-day-old seedlings treated with cycloheximide for the indicated time points using anti-HA antibody. One representative line of each transgenic was used for the study. Rubisco protein was used as the loading control. (d-f) Relative protein levels were normalized to the loading control and to their respective levels at time 0 min. Results from two independent experiments were averaged and shown with error bars representing SD. An exponential best-fit curve was fitted through the data points. Correlation coefficient (R²) values are indicated as a measure of curve fit. The half-life was estimated from the curve assuming first-order kinetics.

Figure 5. ARR4^{D45A} and ARR4^{Y96A} showed weaker rescue of hextuple knock-out than ARR4 in *Arabidopsis* root elongation assay and *ARR7* expression in response to exogenous cytokinin. (a) A representative snapshot of the root elongation of *Arabidopsis* seedlings corresponding to two independent transgenic lines each of ARR4-HA, ARR4^{D45A}-HA and ARR4^{Y96A}-HA on 0 nM BA and 10 nM BA. (b) The quantification of primary root elongation of two independent transgenic lines for each of the three constructs at 0 nM, 5 nM and 10 nM BA concentrations. The data represent the mean \pm SE of primary root growth between 4th day and 9th day from at least 30 individual seedlings for each transgenic line at each BA concentration. Two-tailed Student's *t*-test was performed among means of root lengths under each BA concentration. Transgenic lines with different letters were significantly different from others (P < 0.001). Scale bar = 1 cm. (c) 10-day-old seedlings of wild-type, hextuple mutant and different transgenic lines were treated with 50 nM BA for indicated time points and *ARR7* expression levels were analyzed. Data presented are mean \pm SD from two independent biological replicates and normalized to *TUB2* expression.

Figure 6. Co-IP of ARR4-HA, ARR4^{D45A}-HA and ARR4^{Y96A}-HA with GST-AHP1 after cytokinin treatment. Total protein was extracted from 14-day-old cytokinin-treated seedlings for 45 min of one representative transgenic line for each of the three constructs. The protein was allowed to bind to recombinant GST-AHP1 immobilized on glutathione resin for 2 h. Resin was washed 3 times and samples were subjected to SDS-PAGE and subsequently probed with anti-HA and anti-GST antibodies. GST protein was used as control to show the binding specificity of the HA-tagged proteins towards AHP1. Input shows the amount of the different proteins at the start of the experiment, whereas, pull-down shows the amount of proteins detected at the end of the experiment after the washing steps. BA: N^6 -benzyladenine.

Figure 7. A schematic representation of the root elongation response mediated by AHP1-ARR4 interaction strength. The mechanistic basis of the modification of cytokinin signal strength via AHP1-ARR4 interaction is depicted in the model. 'H' represents the conserved His79 of AHP1 and 'D' represents the conserved Asp95 of ARR4, the two amino acids required for phosphorylation. Grey rectangles on ARR4 indicate the amino acids residues at the interaction interface of ARR4 (Asp45, Arg51 and Tyr96) involved in interaction with AHP1. Two of these rectangles were replaced by yellow boxes in D45A/Y96A to indicate the mutations of Asp45 and Tyr96 to Ala. The red circle harboring 'P' represents the phosphoryl group. Representative photographs of the seedling phenotypes observed are taken from Figure 4. Scale bar = 1 cm.

(a)			
	Name	Organism	Protein Sequence
	AHP1 YPD1	A. thaliana S. cerevisiae	MDLVQKQKSLQDYTKSLFLEGILDSQFTQLQQLQDESNPDJVSQVVTLGFQDSDRIINDL 60 NSTIPSEINWTINEIISMDDDSDSSGLIIQGIDQAQTTFAQM 46
	AHP1 YPD1	A. thaliana S. cerevisiae	SLSIDQQVVDFKKV <mark>OPHVHQHXGSS</mark> SSICAQRVKNAQVVFRSGCEQQNVEACH 113 QRQ <mark>HD</mark> GEK-NLTELDNLG <mark>HFLKGSSAALCLQRIAWVO</mark> ERIQNLGRKMEHJFPNKTELVNT 105
	AHP1 YPD1	A. thaliana S. cerevisiae	RCLQQVKQEYYLVKNRLGTLFKLEQQIVASGGMIPAVEHGG
	AHP1 YPD1	A. thaliana S. cerevisiae	NL 167 (ld: 15.58 %; Sm: 44.15 %)
(h)			

(b)

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Name	Organism	Protein Sequence
ARR4	A. thaliana	MSVGGIGGIESAPLDLDEVHVIAVODSLVDRIVTERIURITSCKVTAVDSGWRALZFIGL 60
SLN1-R1	S. cerevisiae	TSVKILVVDDNHVNOEVIRRMINLEGIENIELACDGODAFDK 42
ARR4	A. thaliana	DNERASAEFDRIKVDIHIT <mark>D</mark> YC <mark>YE</mark> GMTCYELLRRUKESSNFREVEV <mark>U</mark> INSSENVLTRIDR 120
SLN1-R1	S. cerevisiae	VRELTSKGENYNMHFMDVC <u>H</u> KVDCLLSTRMERRDIGY-TSEIVALTAFADDSNHRE 98
ARR4	A. thaliana	CH2ECAODELLAEVKLADVKRIRSHITKDVKLSNGNKRKI 160
SLN1-R1	S. cerevisiae	CH2SCMNGFISKEIKRPKIKTIITEFCAAYQGKKNNK 135 (ld: 21.25 %; Sm: 42.5 %)



Figure 1. Generation of a homology model of AHP1-ΔARR4(16-175) complex. (a) Sequence alignment of AHP1 with its homolog from S. cerevisiae, YPD1. The conserved His required for phosphorylated is highlighted in yellow. (b) Sequence alignment of 16-175 amino acid region of ARR4 with SLN1RD, receiver domain of S. cerevisiae HK protein SLN1. The conserved phosphate-binding pocket residues i.e., the two aspartates (in blue and yellow) and lysine (in purple) are highlighted. (c) Overall view of the computational model of AHP1-ΔARR4(16-175) complex, in which AHP1 is depicted in red and ΔARR4(16-175) is depicted in green. The inset shows 3-dimensional orientation of conserved residues required for phosphorylation i.e., His79 from AHP1 and Asp41, Asp95 and Lys147 from ARR4. Color coding is the same as in sequence alignment (a) and (b).

154x160mm (300 x 300 DPI)



Figure 2. Validation of the AHP1-ΔARR4(16-175) homology model by its comparison with the existing natural complex structures. (a) The Ca superposition of the AHP1-ΔARR4(16-175) in silico complex model with its homologues CheA3P1-CheY6 (orange), SLN1RD-YPD1 (blue) and AHK5RD-AHP1 (magenta). The structural alignment were carried out in PyMol (DeLano, 2002). (b) The structure-based sequence comparison of ARR4 and its homologs. Most of the substitutions are carried with ARR4 and thus we compared its sequences with its structural homologs. The structural alignment was performed using the program Coot (Emsley & Cowtan, 2004). Highly conserved and conserved residues are highlighted. The secondary structures of ARR4 are provided on the top. This figure was prepared using the program ESPript (Gouet et al., 1999). (c) Both full-length ARR4 and ΔARR4(16-175) showed interaction with AHP1 in the yeast two-hybrid system. Growth of yeast cells expressing AD:AHP1 and BD:ARR4 or BD:ΔARR4(16-175) was examined on selection medium lacking Leu, Trp, His and Ade. Selection medium lacking Leu and Trp was used for checking transformation efficiency. The empty pGADT7 vector was used as a negative control. AD: Gal4 activation domain; BD: Gal 4 DNA-binding domain.

134x115mm (300 x 300 DPI)



Figure 3. Identification of ARR4 residues present at AHP1-ΔARR4(16-175) complex interface. (a) ARR4 residues Asp45, Arg51, Tyr96, Cys97, Pro148 mapped onto AHP1-ΔARR4(16-175) model. Cys97 is not shown for clarity (AHP1: red, ΔARR4(16-175): green). (b) Yeast two-hybrid analysis of AD:AHP1 interaction with BD:ARR4 mutants. pGBKT7 vector was used as the control. 100, 10-1 and 10-2 represent no dilution, 10x, 100x dilutions, respectively. (AD: Gal4 activation domain; BD: Gal4 DNA-binding domain). (c) Western

blot detection of AD:AHP1 and BD:ARR4 in yeast cells using anti-HA and anti-c-myc antibodies. Line numbers from one to six correspond to yeast cells harboring AHP1 and ARR4 or different mutants of ARR4, viz. ARR4D45A, ARR4R51A, ARR4Y96A, ARR4C97A and ARR4P148A, respectively (similar to the numbering in Fig 3b). (d) BiFC interactions of ARR4 and mutants with AHP1. Confocal images of N. benthamiana leaf cells coexpressing YFP-C:AHP1 with either YFP-N:ARR4 or YFP-N:ARR4D45A or YFP-N:ARR4Y96A. Column headings: Yellow Channel - reconstituted YFP fluorescence, Transmitted - bright-field images, Overlay – superimposition of both channels. (e) Relative quantification of the YFP fluorescence from AHP1 interaction with ARR4, ARR4D45A and ARR4Y96A using ImageJ (NIH, USA). Data are mean ± SE of signal intensities



Figure 4. Detection of relative transcript and protein levels and analysis of protein turnover rates in transgenic lines. (a) The transcript levels of ARR4, ARR4D45A and ARR4Y96A were detected in the different transgenic lines used in the study. Data presented are mean \pm SD from two independent biological replicates. (b) The protein levels of ARR4-HA, ARR4D45A-HA and ARR4Y96A-HA were detected in the respective transgenic lines used for the study. Rubisco protein was used as the loading control. (c) The protein levels of ARR4-HA, ARR4D45A-HA and ARR4Y96A-HA were detected in 10-day-old seedlings treated with cycloheximide for the indicated time points using anti-HA antibody. One representative line of each transgenic was used for the study. Rubisco protein was used as the loading control. (d-f) Relative protein levels were normalized to the loading control and to their respective levels at time 0 min. Results from two independent experiments were averaged and shown with error bars representing SD. An exponential best-fit curve was fitted through the data points. Correlation coefficient (R2) values are indicated as a measure of curve fit. The half-life was estimated from the curve assuming first-order kinetics. 159x177mm (300 x 300 DPI)

(b)



Figure 5. ARR4D45A and ARR4Y96A showed weaker rescue of hextuple knock-out than ARR4 in Arabidopsis root elongation assay and ARR7 expression in response to exogenous cytokinin. (a) A representative snapshot of the root elongation of Arabidopsis seedlings corresponding to two independent transgenic lines each of ARR4-HA, ARR4D45A-HA and ARR4Y96A-HA on 0 nM BA and 10 nM BA. (b) The quantification of primary root elongation of two independent transgenic lines for each of the three constructs at 0 nM, 5 nM and 10 nM BA concentrations. The data represent the mean ± SE of primary root growth between 4th day and 9th day from at least 30 individual seedlings for each transgenic line at each BA concentration. Two-tailed Student's t-test was performed among means of root lengths under each BA concentration. Transgenic lines with different letters were significantly different from others (P < 0.001). Scale bar = 1 cm. (c) 10-day-old seedlings of wild-type, hextuple mutant and different transgenic lines were treated with 50 nM BA for indicated time points and ARR7 expression levels were analyzed. Data presented are mean ± SD from two independent biological replicates and normalized to TUB2 expression. 170x192mm (300 x 300 DPI)



Figure 6. Co-IP of ARR4-HA, ARR4D45A-HA and ARR4Y96A-HA with GST-AHP1 after cytokinin treatment. Total protein was extracted from 14-day-old cytokinin-treated seedlings for 45 min of one representative transgenic line for each of the three constructs. The protein was allowed to bind to recombinant GST-AHP1 immobilized on glutathione resin for 2 h. Resin was washed 3 times and samples were subjected to SDS-PAGE and subsequently probed with anti-HA and anti-GST antibodies. GST protein was used as control to show the binding specificity of the HA-tagged proteins towards AHP1. Input shows the amount of the different proteins at the start of the experiment, whereas, pull-down shows the amount of proteins detected at the end of the experiment after the washing steps. BA: N6-benzyladenine. 80x65mm (300 x 300 DPI)



Figure 7. A schematic representation of the root elongation response mediated by AHP1-ARR4 interaction strength. The mechanistic basis of the modification of cytokinin signal strength via AHP1-ARR4 interaction is depicted in the model. 'H' represents the conserved His79 of AHP1 and 'D' represents the conserved Asp95 of ARR4, the two amino acids required for phosphorylation. Grey rectangles on ARR4 indicate the amino acids residues at the interaction interface of ARR4 (Asp45, Arg51 and Tyr96) involved in interaction with AHP1. Two of these rectangles were replaced by yellow boxes in D45A/Y96A to indicate the mutations of Asp45 and Tyr96 to Ala. The red circle harboring 'P' represents the phosphoryl group. Representative photographs of the seedling phenotypes observed are taken from Figure 4. Scale bar = 1 cm. 80x50mm (300 x 300 DPI)