

Review

# Multiple Links between HD-Zip Proteins and Hormone Networks

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**Abstract:** HD-Zip proteins are unique to plants, and contain a homeodomain closely linked to a leucine zipper motif, which are involved in dimerization and DNA binding. Based on homology in the HD-Zip domain, gene structure and the presence of additional motifs, HD-Zips are divided into four families, HD-Zip I-IV. Phylogenetic and bioinformatics analysis of *HD-Zip* genes using transcriptomic and genomic datasets from a wide range of plant species indicate that the HD-Zip protein class was already present in green algae. Later, *HD-Zips* experienced multiple duplication events that promoted neo- and sub-functionalizations. HD-Zip proteins are known to control key developmental and environmental responses, and a growing body of evidence indicates a strict link between members of the HD-Zip II and III families and the auxin machineries. Interactions of HD-Zip proteins with other hormones such as brassinolide and cytokinin have also been described. However, it is striking that among the genes regulated by REV, a HD-Zip III protein playing a key role in apical development, are genes that mediate ABA signaling. Furthermore, HAT1 and HAT3, two HD-Zip II proteins involved in key developmental processes, repress ABA biosynthesis and signaling, indicating an essential role of these factors in adjusting development to changing environment.

**Keywords:** Arabidopsis; developmental pathways; environmental responses; HD-Zip transcription factors; hormones

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## 1. The HD-Zip class of proteins

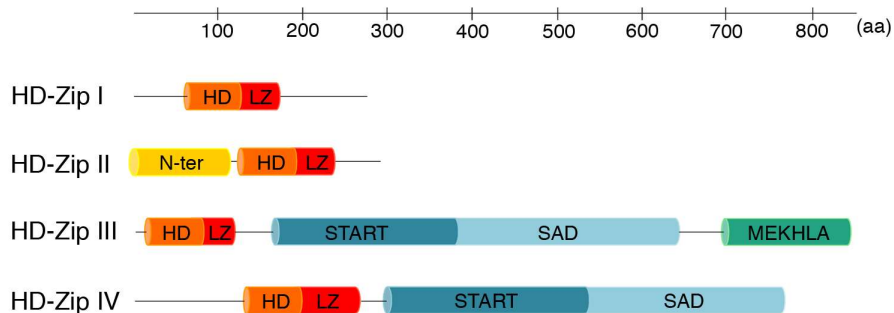
The homeodomain-leucine zipper (HD-Zip) class of proteins appears to be present exclusively in the plant kingdom, and is characterized by the presence of a homeodomain closely linked to a leucine zipper motif [1]. The Arabidopsis genome codes for 48 HD-Zip proteins that, on the basis of sequence homology in the HD-Zip domain, the presence of additional conserved motifs, and specific intron and exon positions, have been grouped into four families: HD-Zip I (17 members), HD-Zip II (10 members), HD-Zip III (5 members) and HD-Zip IV (16 members) [2-7].

*HD-Zip genes* are evolutionary highly conserved and there is evidence that they were already present in green algae [8-11]. Later in evolution, the HD-Zip class experienced multiple duplication events that promoted neo- and sub-functionalizations for terrestrial life [11].

Experimental work has demonstrated that the HD-Zip domain, but not the HD by itself, interacts with DNA [12], and it has been shown that a correct spatial relationship between the HD and the leucine zipper motif is crucial for DNA binding [12]. Binding-site selection analysis and subsequent chromatin immunoprecipitation sequencing experiments have determined that the HD-Zip proteins recognize pseudo-palindromic DNA elements [3, 12-14]. HD-Zip I proteins interact with the CAAT(A/T)ATTG motif [12, 15, 16] whereas HD-Zip II proteins preferentially bind the CAAT(C/G)ATTG motif [12, 13]. Binding-site selection analysis identified GTAAT(G/C)ATTAC as the sequence preferentially recognized by HD-Zip III proteins [3]; however, more recent genome-wide binding site experiments suggest that the AT(G/C)AT central core is sufficient for DNA binding [14]. For HD-Zip IV proteins, the CATT(A/T)AATG motif was shown to be required for DNA-binding [13], and found in the promoters of true target genes [17-19]. Interestingly, the identified cis-elements are very similar, particularly the HD-Zip II and III binding sites which share the same core sequence [AAT(G/C)ATT] [3, 12], thus suggesting that members of the different families of HD-Zip proteins may regulate common target genes [20, 21].

Beside the homeodomain-leucine zipper motif, HD-Zip I proteins have no other established functional domain; conversely, most of the HD-Zip II transcription factors contain an LxLxL type of ERF-associated amphiphilic repression (EAR) motif [7, 22] (Figure 1), and there is evidence that they function as negative regulators of gene expression [7,20, 23-25]. Furthermore, it was recently shown that HOMEBOX ARABIDOPSIS THALIANA (HAT) 1 and HAT22 interact with the TOPLESS (TPL) co-repressor protein [26]. HD-Zip III and HD-Zip IV proteins, in addition to the HD-Zip domain, possess a steroidogenic acute regulatory protein-related lipid-transfer (START) domain motif with putative lipid-binding capability and a Small body size–mothers against decapentaplegic homolog 4 (Smad4) activation domain (SAD) [27, 28] (Figure 1). Finally, HD-Zips III, and not HD-Zips IV, share a MEKHLA domain (Figure 1). A region within this domain shows sequence similarity to the PAS (Per-Arnt-Sim)-domain which has been shown to act as intracellular sensor of light, oxygen or redox-potentials [28]. It has been shown that this domain is involved in the dimerization of HD-Zip III proteins with DORNROSCHEN (DRN) and DORNROSCHEN-like (DNRL) transcription factors [29].

HD-Zip proteins are known to control key developmental and environmental responses [21, 30-32]. Here we report recent advances on the understanding of the interactions between HD-Zip transcription factors and hormone signaling networks.



**Figure 1.** Schematic representation of the protein domains possess by each HD-Zip family. *ATHB1*, *ATHB2*, *ATHB8*, and *GLABRA2* were chosen as representative members of the HD-Zip I, II, III and IV families, respectively. N-term, N-terminus consensus; HD, Homeodomain; Zip, Leucine zipper; START, steroidogenic acute regulatory protein-related lipid transfer domain; SAD, Small body size–mothers against decapentaplegic homolog 4 (*Smad4*) activation domain; MEKHLA, named after the identification of the highly conserved amino acids Met, Glu, Lys, His, Leu, Ala.

## 2. HD-Zips I

Sequence comparison and phylogenetic analysis indicated that members of the HD-Zip I protein family can be classified into six different clades, I to VI, and that the presence and the position of conserved sequences may be related to specific function(s) [4, 33]. Genome-wide expression studies revealed that several *Arabidopsis HD-Zip I* genes show transcriptional changes in response to treatments with abscisic acid (ABA) [4], and there is evidence that members of clades I and II of the HD-Zip I family have roles related to drought stress and ABA-signaling in different plant species [34-40]. For example, *ARABIDOPSIS THALIANA HOMEBOX (ATHB) 7* and *ATHB12* (belonging to the clade I) [33] are both strongly induced by water-deficit and ABA. Chromatin immunoprecipitation and gene expression analyses have demonstrated that *ATHB7* and *ATHB12* positively regulate the expression of five genes encoding clade A protein phosphatases type 2C (PP2C), acting as central negative regulators of ABA signaling [41-43]. Furthermore, it has also been shown that *ATHB7* and *ATHB12* act to repress the transcription of two members of the *PYRABACTIN RESISTANCE1 (PYR1)/PYR1-LIKE (PYL)* gene family, encoding the ABA receptors [44-46]. Together the data indicate that *ATHB7* and *ATHB12* function as negative regulators of the ABA response in *Arabidopsis* [47]. Evidences exists that *HOMEBOX (HB) 6* (a.k.a *ATHB6*, clade II) may also act as a negative regulator of ABA signaling [48].

The *HELIANTHUS ANNUUS HOMEBOX4 (HAHB4)* gene, encoding a protein homologous to *ATHB7* and *ATHB12*, is also regulated by ABA and drought, as well as by methyl-jasmonic acid (MeJa) or ethylene (ET) or biotic stresses [49, 50]. The ectopic expression of *HAHB4* in *Arabidopsis* negatively affects the synthesis of ET and resulted in plants more resistant to drought [49]. In addition, functional analysis of transgenic *Arabidopsis* and maize plants constitutively expressing *HAHB4* suggested that this HD-Zip I protein acts as an integrator of MeJa and ET pathways [50].

It is worth to mention that in some plants there is evidence that ABA synthesis and signaling it is relevant to fully activate defence responses against insect herbivores and, in general, hormonal interactions are important to regulate plant responses to abiotic stresses and growth-defense tradeoffs [51]. Of interest is the finding that the *ATHB13* (clade V) gene, positively regulated by

low temperature, drought, and salinity, can confer cold, drought and broad-spectrum disease resistance when overexpressed [38, 52, 53]. The results point to a role of some HD-Zip I proteins as integrators of internal and external signals in the regulation of abiotic and biotic stresses.

Together with the effects on stresses described above, the overexpression of HD-Zip I proteins very often resulted also in alterations of the shape and growth of the plant, including cotyledon, leaf and supporting organs [53-55], suggesting a role of some HD-Zip I proteins in specific growth and/or developmental pathways. It is worth to mention that at least *ATHB12* and some ABA signaling components are regulated by *KANADI1* (*KAN1*) [56], a factor controlling organ polarity, including the patterning of leaf primordia in *Arabidopsis* [57]. In addition, the *Medicago truncatula* *HB1*, highly related to *ATHB7* and *ATHB12*, represents an example of a cross-talk between ABA and auxin in the regulation of organ development. The *MHB1* gene is strongly regulated by salt, osmotic and ABA stresses. There is evidence that *HB1* controls the emergence of lateral roots likely by repressing the auxin-regulated *LOB-Binding Domain 1* (*LBD1*) gene [58], a member of a family of plant-specific transcription factors involved in lateral organ development [59].

In the post-embryonic development, *HB5* (clade II) behaves as a growth-promoting transcription factor of the hypocotyl. In particular, it promotes the gibberellin acid (GA)-mediated expansion of the epidermal and cortex cells by a positive direct modulation of *EXPANSIN3* (*EXP3*) gene expression [60]. *ATHB1* (Clade III) is a direct target of PHYTOCHROME INTERACTING FACTOR 1 (*PIF1*) and plays a role in hypocotyl growth under short-day regime likely through a positive regulation of genes involved in cell elongation [61]. Interestingly, *ATHB1* expression is positively regulated by ethylene in *Arabidopsis* [G.M. unpublished data; [62], known to regulate the elongation of the hypocotyl in low light and shade [63], whereas the tomato ortholog *HB1* directly regulates the expression of *1-AMINOCYCLOPROPANE-1-CARBOXYLATE OXIDASE* (*ACO*) gene encoding a key enzyme in ethylene biosynthesis [64].

Very recent work implicated three related HD-Zip I proteins belonging to clade VI [33] in the control of shoot branching [65]. Indeed, it was found that TEOSINTE BRANCHED1, CYCLOIDEA, PCF (TCP) transcription factor *BRANCHED1* (*BRC1*) binds to and positively regulates the transcription of the *HB21*, *HB40* and *HB53* genes, all belonging to clade VI. These HD-Zip I proteins are necessary and sufficient to enhance the expression of *9-CIS-EPOXICAROTENOID DIOXIGENASE 3* (*NCED3*), a key ABA biosynthesis gene, and for ABA accumulation inside axillary buds in conditions of Low Red/Far-Red (R/FR) ratio light or short photoperiod. This, in turn, causes suppression of bud development. Relevantly, the *BRC1/HB21/40/53* regulatory module appears to be conserved in monocot and dicot species [65].

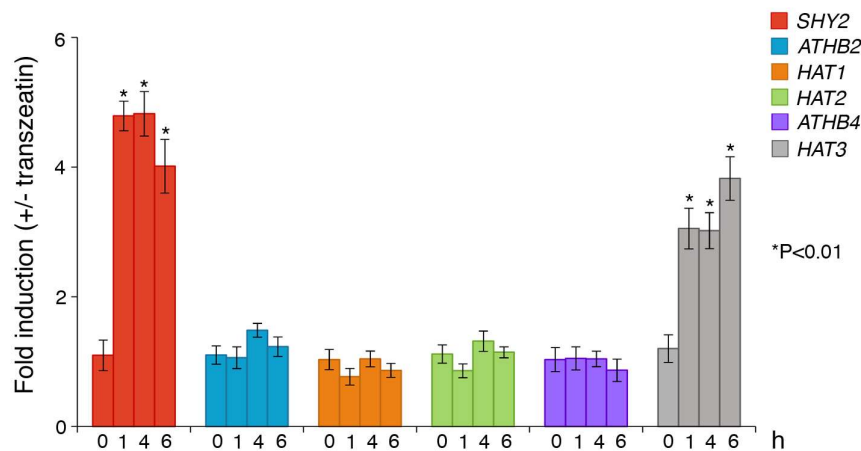
Besides the multiple links found between HD-Zip I proteins and ABA, evidence of a direct interaction between HD-Zips I and auxin also exists [66]. Auxin has a central role during embryogenesis and post-embryonic development. The transcriptional auxin response is controlled by AUXIN RESPONSE FACTOR (ARF) transcription factors and AUXIN/INDOLE-3-ACETIC ACID (AUX/IAA) proteins. AUX/IAAs function as repressors by forming dimers with ARFs, and their degradation releases the inhibition on ARF transcription factors [67, 68]. *BODENLOS* (*BDL*)/*IAA12* and its interacting ARF partner *MONOPTEROS* (*MP*)/*ARF5* play a key role during the early stages of embryo development. Both a dominant mutant of (*BDL*)/*IAA12* and a loss-of-function mutant of *MP/ARF5* lack a seedling root and display cotyledon defects [69-72]. Interestingly, it was found that the *HB5* protein (clade II) directly negatively regulates *BDL/IAA12* expression.

Overexpression of HB5 during embryogenesis transcriptionally suppresses the expression of *BDL/IAA12* and rescues the rootless phenotype of the *bdl/iaa12* dominant mutant. Together the data lead to the hypothesis that HB5 may contribute to spatially restrict *BDL/IAA12* expression during embryogenesis [66]. Evidence that HB6, a close homolog of HB5, may act redundantly with HB5 in the negative regulation of *BDL/IAA12* have also been provided [66].

Finally, the cross-talk between ethylene and auxin in the control of root elongation mediated by *ATHB52* has been recently uncovered. It is very well established that root elongation is inhibited by ethylene in *Arabidopsis* and other species through the action of auxin [73-76]. The *ATHB52* gene is positively regulated by ETHYLENE-INSENSITIVE3 (EIN3), a key transcription factor of the ethylene signal transduction pathway. A molecular and genetic analysis has shown that *ATHB52* binds the promoters of *PIN FORMED2 (PIN2)*, coding for a polar auxin carrier, and of *WAVY ROOT GROWTH1 (WAG1)* and *WAG2*, encoding PIN polarity regulators. The positive modulation by ethylene of the *PIN2/WAG1/WAG2* module exerted through *ATHB52* could affect the local polar auxin transport in the root tip resulting in the inhibition of primary root elongation [77].

### 3. HD-Zips II

The HD-Zip II protein family contains 10 members which can be divided into four clades ( $\alpha$ - $\delta$ ) [7]. Remarkably, all the *HD-Zip II*  $\gamma$  and  $\delta$  (*HAT3*, *ATHB4*) genes are rapidly induced by changes in the R/FR ratio light that promote shade avoidance in the Angiosperms [7, 78] and several evidence exist that HD-Zip II  $\gamma$  and  $\delta$  proteins act as positive regulators of this response [7, 23, 25, 79-83]. By contrast, *HD-Zip II*  $\gamma$  and  $\delta$  genes display distinct responses to plant hormones. For example, *HAT2* is induced by auxin [25] whereas *HAT3*, and not the other  $\gamma$  and  $\delta$  genes, is up-regulated in the tip of roots treated with cytokinins (CKs) (Figure 2).



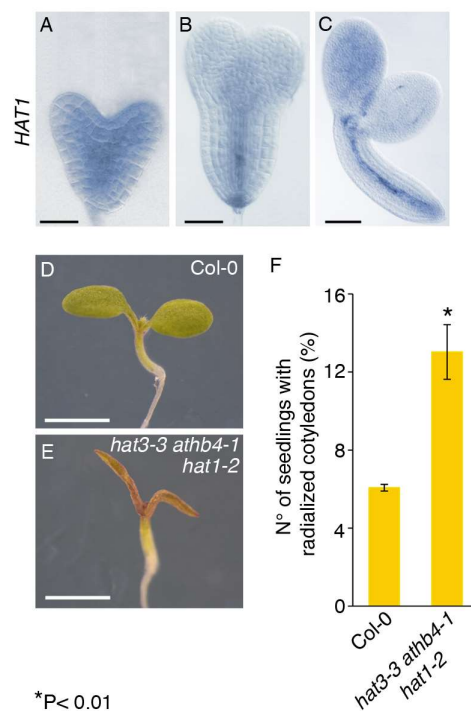
**Figure 2.** *HAT3*, and not *ATHB2*, *HAT1*, *HAT2* and *ATHB4*, is positively regulated by cytokinins in the root tip. RT-qPCR analyses of *SHORT HYPOCOTYL 2 (SHY2)/IAA3 (SHY2)*, positive control [84], *ATHB2*, *HAT1*, *HAT2*, *ATHB4* and *HAT3* in Col-0. Seedlings were grown for 5 days in a light/ dark cycle (16/8 h), and then treated with transzeatin (5  $\mu$ m) for the indicated times. The graph shows the relative expression levels of *SHY2*, *ATHB2*, *HAT1*, *HAT2*, *ATHB4* and *HAT3*. Each value is the mean of three biological replicates normalized to *EF1 $\alpha$*  expression ( $\pm$ SD). Statistical significance was assessed as previously described [83]. \* $P < 0.01$ .

Interestingly, several evidence demonstrate that, besides their function in shade avoidance, HD-Zip II  $\gamma$  and  $\delta$  transcription factors play a crucial role in embryo apical development and essential developmental processes in sunlight, including shoot apical meristem (SAM) activity, organ polarity and gynoecium development [20, 30, 85-88] (Figure 3).

A number of links have been established between HD-Zip II  $\gamma$  and  $\delta$  proteins and auxin. Plants with elevated levels of ATHB2 displays a constitutive shade avoidance response, and it was shown that ATHB2-induced elongation of the hypocotyl depends on the auxin transport system, as it is abolished by auxin transport inhibitors [23]. Furthermore, the lateral root phenotype observed in ATHB2 overexpressing seedlings is rescued by IAA [23]. Finally, it was found that both auxin synthesis and transport are affected in *hat3 athb4* and *hat3 athb4 athb2* mutant embryos [20, 30]. The enhanced cotyledon phenotype of *hat3 athb4 hat1* with respect to *hat3 athb4* strongly suggests that HAT1 works redundantly with HAT3 and ATHB4 in the control of auxin synthesis and transport during embryogenesis (Figure 3).

Interestingly, HAT1 has also been linked to brassinosteroid (BR) signaling pathway. BRs signal through plasma membrane-localized receptor and other components to modulate the BES1/BZR1 (BRI1-EMS SUPPRESSOR 1/BRASSINAZOLE RESISTANT1) family of transcription factors that regulate thousands of target genes [89-94]. Relevantly, it was recently found through chromatin immunoprecipitation (ChIP) experiments that *HAT1* is a direct target of BES1 [95]. HAT1 functions redundantly with its close homolog HAT3, as the double loss-of-function mutant *hat1 hat3* displayed a reduced BR response stronger than that of the *hat1* and *hat3* single mutants. Expression levels of several BR-repressed genes are increased in *hat1 hat3* double mutant and reduced in HAT1 overexpressing lines, thus strongly suggesting that HAT1 functions to repress the expression of a subset of BR target genes. Consistently, it was found that HAT1 binds to DNA elements in BR-repressed gene promoters and functions as a BES1 corepressor [95]. Furthermore, it was shown that HAT1 can be phosphorylated and stabilized by GSK3 (GLYCOGEN SYNTHASE KINASE 3)-like kinase BIN2 (BRASSINOSTEROID-INSENSITIVE 2), a negative regulator of the BR pathway [95-98].

Furthermore, HAT1 was identified among the transcription factors interacting with the GIBBERELLIN INSENSITIVE (GAI) DELLA protein, a master negative regulator in gibberellin (GA) signaling [99, 100]. Further work is needed to establish the specific GA-response(s) in which HAT1 is involved.



**Figure 3.** HAT1 is redundant to HAT3 and ATHB4 in regulating cotyledon development. (A–C) *In situ* hybridization of Col-0 embryos with *HAT1* probe. Heart (A), torpedo (B) and mature (C) embryos. (D, E) Col-0 (D) and *hat3-3 athb4-1 hat1-2* (E) seedlings (*hat3-3 athb4-1* is described in [20]; *hat1-2*, line SALK\_059835). (F) The graph represents the percentage of seedlings displaying two radialized cotyledons in *hat3-3 athb4-1* and *hat3-3 athb4-1 hat1-2* mutants. Error bars represent s.d.; \* $P < 0.01$ , Student's t-test. Scale bars: A, B, 30  $\mu\text{m}$ ; C, 100  $\mu\text{m}$ ; D, E, 1 mm.

Recent work has demonstrated that HAT1, apart from its role in auxin-regulated cotyledon development (Figure 3), in BR-mediated growth responses [95], in GA signaling, and in viral defense response in a manner dependent on salicylic acid (SA) [101], it also negatively regulates, redundantly with HAT3, ABA-mediated drought responses through suppression of ABA biosynthesis and signaling [102]. The expression of both *HAT1* and *HAT3* is indeed repressed by ABA. Evidence have been provided that HAT1 can bind to specific DNA sequences on the promoters of *NCED3* and *ABA DEFICIENT (ABA) 3*, two key ABA biosynthesis genes, and negatively regulate their expression, thus resulting in a reduction of ABA synthesis. In addition, it was observed that HAT1 overexpressing plants display reduced sensitivity to ABA and less tolerance to drought stress, whereas the double loss-of-function *hat1 hat3* mutant show opposite phenotypes. Finally, it was found that Sucrose non-fermenting 1-related protein kinase (SnRK) 2.3, a positive component of ABA signaling, physically interacts with and phosphorylates HAT1, decreasing its protein stability and binding activity [102].

Relevantly, at least other two HD-Zip II proteins, ATHB17 and HAT22/ABA INSENSITIVE GROWTH 1 (ABIG1), are linked to ABA [103, 104]. *ATHB17* expression is induced by ABA, and evidence have been provided that *athb17* loss-of-function mutants are ABA-insensitive and drought-sensitive whereas lines overexpressing ATHB17 display opposite phenotypes.

Interestingly, the effect of *ATHB17* on seedling growth in the presence of ABA is stage-specific. Indeed, it is observed exclusively during the post-germination seedling establishment stage [103]. Recent work identified *HAT22/ABIG1* as a transcription factor required for ABA-mediated growth inhibition, but not for seed dormancy and stomatal closure. It has been proposed that drought acts through ABA to increase *HAT22/ABIG1* transcription which, in turn, inhibits new shoot growth and promotes leaf senescence [104].

#### 4. HD-Zips III

The HD-Zip III family contains five members: *ATHB8*, *CORONA* (*CNA*), *PHABULOSA* (*PHB*), *PHAVOLUTA* (*PHV*), and *REVOLUTA* (*REV*). It is well established that HD-Zip III proteins act as master regulators of embryonic apical fate [105], are required to maintain an active SAM and to establish lateral organ polarity [106-108], and are necessary for xylem formation and specification [109-114]. Recent work has also implicated HD-Zip III proteins in the regulation of the shade avoidance response [14, 21, 31].

The pattern of HD-Zip III expression largely coincides with that of auxin distribution [8, 9, 115-119]. Furthermore, *HD-Zip III* genes are regulated at the post-transcriptional level by the microRNAs *miR165/166*, which negatively affect their expression through mRNA cleavage [107, 120]. Relevantly, *REV* directly positively regulates the *HD-Zip II* genes *HAT3*, *ATHB4*, *ATHB2*, and *HAT2*, and evidence exists that *PHB* and *PHV* are involved in the control of *HAT3* expression [14, 20]. It has been recently shown that *REV* physically interacts with *HAT3* and *ATHB4* to directly repress *MIR165/166* expression in the adaxial side of the leaf [88].

The interconnection between HD-Zip III transcription factors and auxin began to be clarified especially through the molecular-genetic analysis of the vascular system [121]. *REV* directly positively regulates the auxin biosynthetic genes *TRYPTOPHAN AMINOTRANSFERASE OF ARABIDOPSIS 1* (*TAA1*) and *YUCCA5* (*YUC5*) [14, 122]. Furthermore, it has been demonstrated that genes implicated in auxin transport, including the influx carriers *LIKE AUXIN RESISTANT 2* (*LAX2*) and *LAX3*, and response are also direct targets of *REV* [119, 122, 123]. Interestingly, *REV* also up-regulates the expression of *NAKED PINS IN YUC MUTANTS 1* (*NPY1*) and *WAG1*, encoding an AGC protein kinase highly homologous to PINOID (*PID*) [124]. *NPY* genes encode proteins with a Broad-Complex, Tramtrack, and Bric-a-brac (*BTB*)-Poxvirus and Zinc Finger (*POZ*) domain that together with AGC kinases determine the subcellular polar targeting of the PIN efflux carriers, thus establishing the direction of auxin transport [125-128].

A recent study reinforces the interconnection between HD-Zip III transcription factors and auxin signaling [129]. HD-Zip III proteins are known to determine xylem patterning in the Arabidopsis root [113], and it has been shown that *PHB* directly interacts with the promoter of both *MP/ARF5*, a transcription factor gene playing a major role in vascular development, and *IAA20*, encoding an IAA protein that is stable in the presence of auxin and able to interact with *MP*. The double mutant of *IAA20* and its closest homolog *IAA30* forms ectopic protoxylem, whereas elevated levels of *IAA30* result in discontinuous protoxylem, analogous to a weak *mp* mutant. It has therefore proposed a mechanism in which *PHB* stabilizes the auxin response within the xylem axis by activating both *MP* and its repressors *IAA20* and *IAA30* to ensure correct vascular patterning and differentiation of xylem cells [129].

Cross-talk between auxin and cytokinin (CK) is crucial during several developmental processes,



including vascular development. A number of studies have indicated that ARABIDOPSIS HISTIDINE PHOSPHOTRANSFER PROTEIN 6 (AHP6), an inhibitory pseudophosphotransfer protein, is positively regulated by auxin and counteracts CK signaling, allowing protoxylem formation in the root. Conversely, CK signaling negatively regulates the spatial domain of AHP6 expression [130, 131]. Interestingly, it has been shown that *PHB* acts redundantly with other HD-Zip III transcription factors to repress *AHP6* expression in a dose-dependent manner [113].

The interaction between HD-Zip III proteins and cytokinin network is further strengthened by the finding that *PHB* directly activates *ISOPENTENYLTRANSFERASE 7 (IPT7)*, a gene coding for a rate-limiting component of the cytokinin biosynthesis pathway. This in turn promotes cell differentiation and regulates root length. It was also shown that CK represses both *PHB* and *miR165* [132]. The authors referred to these interactions as an incoherent regulatory loop in which CK represses both its activator and a repressor of its activator, and proposed that this circuit might provide robustness against CK fluctuations [132].

A recent study on the mechanism underlying shoot regeneration in Arabidopsis provided an additional link between HD-Zip III transcription factors and CK signaling [133], somehow predicted also on the basis of protein-protein interaction studies between HD-Zip III and DNR/DNRL transcription factors [29, 134, 135] and the analysis of the regeneration capacity of a *cna* mutant encoded by the *hoc* locus [136].

Four B-type ARABIDOPSIS RESPONSE REGULATOR (ARR) transcription factors, ARR1, ARR2, ARR10, and ARR12, have essential roles in shoot regeneration. Indeed, the shoot regenerative capacity is impaired in the *arr1 arr10 arr12* mutant with respect to wild type [137-139]. The A-type ARRs play opposite roles in shoot regeneration, and it has been shown that overexpression of ARR7 or ARR15 results in a marked reduction of the regeneration capacity [140]. Remarkably, it was found that ARR1, ARR2, ARR110 and ARR12 interact with *PHB*, *PHV* and *REV* HD-Zip III transcription factors, and that these complexes in turn activates *WUSCHEL (WUS)* expression [133].

Beside the evidence of molecular interactions between HD-Zip III proteins and key components of the auxin and cytokinin networks, strikingly, it has been recently reported that *REV* positively directly regulates the expression of the gene encoding the ABA receptor protein *PYL6* [123, 124]. Remarkably, *PYL6* is oppositely directly regulated by *KAN1*, a key determinant of abaxial cell fate in the leaf [57, 124, 141]. Furthermore, microarray data revealed that the expression of *REV*, *PHB* and *PHV* significantly decreases upon ABA application, as a consequence of ectopic induction of *miR165* expression [21, 123]. It has been therefore proposed that the connection between ABA perception and signaling and HD-Zip III transcription factors may be required to adapt leaf development to alterations in water availability [21, 123].

Finally, it is worth to mention that recent work has revealed that the expression of *miR165/166* is regulated by a complex hormonal cross-talk during root development in Arabidopsis [142].

#### HD-Zips IV

Many of the *HD-Zip IV* genes were shown to be specifically or preferentially expressed in the epidermis of developing embryos and/or other plant organs [143]. It is worth to mention that the epidermis plays a critical role also in plant defence against pathogens and in protection from environmental stresses [143, 144].

*GLABRA2 (GL2)*, the first identified member of the HD-Zip IV protein family, promotes

trichome differentiation [145], and suppresses root hair formation in the root epidermis [19, 146]. In particular, *GL2* controls cell fate determination of N-cells (non-hair cells; atrichoblast) [147], through a negative regulation of the phospholipase D zeta 1 (*PLDz1*) [18], and of several basic helix-loop-helix (bHLH) transcription factors including *ROOT HAIR DEFECTIVE 6* (*RDH6*) involved in root hair initiation [146]. There is strong evidence that BRs are required to maintain position-dependent cell fate specification in root epidermis. Indeed, BRs are required for normal expression levels and patterns of *WEREWOLF* (*WER*) and *GL2*, master regulators of epidermal patterning. Loss of BR signaling results in loss of H cells, likely as a consequence of reduced expression of *CAPRICE* (*CPC*), a direct downstream target of *WER* [148].

Recent work has also established a link between HD-Zip IV proteins and GA signaling [149]. It was indeed shown that *DELLA* proteins interact directly with *MERISTEM LAYER 1* (*ATML1*) and its paralogue, two HD-Zip IV proteins required for epidermis specification and binding to the L1 box present in the promoters of epidermis-specific genes [17, 150]. Silencing of both *ATML1* and *PDF2* inhibits epidermis-specific gene expression and delays germination [149]. Evidence were provided that, upon seed imbibition, increased GA levels reduce *DELLA* protein levels, thus releasing *ATML1/PDF2* to activate epidermis-specific expression and promote seed germination [149].

Apart from their role in development, interactions between HD-Zip IV transcription factors and environmental responses have also been reported [151]. For example, *HOMEODOMAIN GLABRA 11* (*HDG11*) was identified via activation tagging as a gene involved in drought tolerance. The mutant has higher levels of ABA than the wild type, and displays enhanced root growth with more lateral roots and reduced stomatal density. Overexpression of *HDG11* also conferred drought tolerance associated with augmented lateral roots and reduced leaf stomatal density in both *Arabidopsis* and tobacco [152]. Higher level of ABA and improved drought tolerance have been observed also in cotton, poplar and rice transgenic plants overexpressing *HDG11* [151, 153]. It has been suggested that *HDG11* positively regulates the expression of cell wall loosening protein genes, including *EXP5*, resulting in a well-developed root system [154].

Intriguingly, *HDG11* acts as a maternal regulator of zygote asymmetry through a direct activation of the *WUSCHEL RELATED HOMEODOMAIN 8* (*WOX8*) gene whose product leads to asymmetric division of the zygote [155].

## 6. Conclusions

A fundamental question in plant biology is how plants integrate environmental signals with intrinsic developmental programs and how coordinate the growth of different organs depending on resource availability. Over recent years remarkable progress has been made, and the molecular mechanisms controlling these processes are being elucidated. The functional analysis of the HD-Zip proteins revealed that they are part of complex networks involved in the integration of external signals through the regulation of hormonal pathways involved in the control of fundamental developmental processes. Although many factors belonging to each HD-Zip family are implicated in specific processes, it is interesting to note that the simple overexpression of *ATHB2* (and other members of the HD-Zip II family) and *HDG11* is sufficient to generate plants looking for optimal light and water for growth, respectively. This could be explained by the existence of organ- and/or tissue-specific hubs that stimulated by external and/or internal (hormonal) signals

converge to coherently adjust the development and growth to a specific environmental signal. These hubs may have been generated during evolution through multiple duplication events that have promoted neo and sub-functionalization of factors operating on specific pathways. The identification of these hubs it has the potential to lead to a more unified vision of the development and growth of the plant according to environmental stresses that could be applied for the improvement of the cultivated plants.

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#### Abbreviations

ABA	Abscisic acid
ABA	ABA DEFICIENT
ABIG1	ABA INSENSITIVE GROWTH 1
ACCO	1-AMINOCYCLOPROPANE-1-CARBOXYLATE OXIDASE
AHP6	HISTIDINE PHOSPHOTRANSFER PROTEIN 6
AUX/IAA	AUXIN/INDOLE-3-ACETIC ACID
ARF	AUXIN RESPONSE FACTOR
ARR	ARABIDOPSIS RESPONSE REGULATOR
ATHB	ARABIDOPSIS THALIANA HOMEBOX
ATML1	MERISTEM LAYER 1
BDL	BODENLOS
BES1/BZR1	BRI1-EMS SUPPRESSOR 1/BRASSINAZOLE RESISTANT1
PHB	PHABULOSA
PHV	PHAVOLUTA
b-HLH	basic Helix-Loop-Helix
BIN2	BRASSINOSTEROID-INSENSITIVE 2
BR	Brassinosteroid
BRC1	BRANCHED1
ChIP	Chromatin Immunoprecipitation

CK	Cytokinin
CNA	CORONA
CPC	CAPRICE
DRN	DORNROSCHEN
DNRL	DORNROSCHEN-like
EAR	ERF-Associated Amphiphilic Repression
EIN3	ETHYLENE-INSENSITIVE 3
EXP3	EXPANSIN 3
ET	Ethylene
HAHB4	HELIANTHUS ANNUUS HOMEobox 4
HB	HOMEobox
HAT1	HOMEobox ARABIDOPSIS THALIANA 1
HDG11	HOMEODOMAIN GLABRA 11
HD-Zip	Homeodomain-leucine zipper
IPT7	ISOPENTENYLTRANSFERASE 7
GA	Gibberellin Acid
GAI	GIBBERELLIN INSENSITIVE
GL2	GLABRA2
GSK3	GLYCOGEN SYNTHASE KINASE 3
KAN1	KANADI 1
LAX2	LIKE AUXIN RESISTANT 2
LBD1	LOB-Binding Domain 1
MeJa	Methyl-Jasmonic acid
MP	MONOPTEROS
NCED3	9-CIS-EPOXICAROTENOID DIOXIGENASE 3
NPY1	NAKED PINS IN YUC MUTANTS 1
PDF2	PLANT DEFENSIN 2
PID	PINOID
PIF1	PHYTOCHROME INTERACTING FACTOR 1
PIN2	PIN FORMED2
PLDz1	Phospholipase D zeta 1
POZ	Broad-Complex, Tramtrack, and Bric-a-brac (BTB)-Poxvirus and Zinc Finger
PP2C	Protein Phosphatases type 2C
PYL	PYRABACTIN RESISTANCE1 (PYR1)/PYR1-LIKE
RDH6	ROOT HAIR DEFECTIVE 6
R/FR	Red/Far-Red
REV	REVOLUTA
SAD	Small body size-mothers against decapentaplegic homolog 4 (Smad4) activation domain
SAM	Shoot Apical Meristem
SHY2	SHORT HYPOCOTYL 2
SnRK	Sucrose non-fermenting 1-related protein kinase
START	Steroidogenic acute regulatory protein-related lipid-transfer

TAA1	TRYPTOPHAN AMINOTRANSFERASE OF ARABIDOPSIS 1
TCP	TEOSINTE BRANCHED1, CYCLOIDEA, PCF
TPL	TOPLESS
WAG1	WAVY ROOT GROWTH 1
WER	WEREWOLF
WOX8	WUSCHEL RELATED HOMEODOMAIN 8
WUS	WUSCHEL
YUC5	YUCCA5

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