

# Shoot meristem maintenance is controlled by a GRAS-gene mediated signal from differentiating cells

Jeroen Stuurman, Fabienne Jäggi, and Cris Kuhlemeier<sup>1</sup>

Institute of Plant Sciences, University of Bern, Altenbergrain 21, CH-3013 Bern, Switzerland

**Plant shoot development depends on the perpetuation of a group of undifferentiated cells in the shoot apical meristem (SAM). In the *Petunia* mutant *hairy meristem* (*ham*), shoot meristems differentiate postembryonically as continuations of the subtending stem. *HAM* encodes a putative transcription factor of the GRAS family, which acts non-cell-autonomously from L3-derived tissue of lateral organ primordia and stem provasculture. *HAM* acts in parallel with *TERMINATOR* (*PhWUSCHEL*) and is required for continued cellular response to *TERMINATOR* and *SHOOTMERISTEMLESS* (*PhSTM*). This reveals a novel mechanism by which signals from differentiating tissues extrinsically control stem cell fate in the shoot apex.**

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Plants differ from animals in that they continuously form new organs (stems, leaves, or flowers) during post-embryonic shoot development. This depends on the perpetuation of a shoot apical meristem (SAM) at the very summit of the growth axis. Within the SAM, a few stem cells are specified whose daughters give rise to the entire aerial plant body (for review, see Weigel and Jürgens 2002). A primary question in plant development is how the SAM is maintained and how stem cells are specified.

A major signaling pathway that controls stem cell identity is mediated by the homeobox gene *WUSCHEL* (*WUS*). *WUS* is critical for SAM maintenance as judged by mutation (Laux et al. 1996) or by genetically programmed elimination during flower development (Lenhard et al. 2001; Lohmann et al. 2001). *WUS* expression is under negative feedback control of one of its target genes, *CLAVATA3* (*CLV3*), which keeps stem cell maintenance and differentiation in dynamic equilibrium (Brand et al. 2000; Schoof et al. 2000). Other major factors that control SAM maintenance are the homeobox genes of the *KNOTTED* (*KNOX*) family. One of these, *SHOOTMERISTEMLESS* (*STM*), is thought to distinguish the SAM from lateral organs (Byrne et al. 2000). Ectopic *KNOX* expression can induce adventitious

SAMs at sites that normally are fully differentiated (e.g., see Sinha et al. 1993).

Factors like *WUS*, *STM*, and *CLV* are all expressed within undifferentiated cells of the SAM and are meristem intrinsic. There is evidence, however, that SAM development depends on extrinsic activities present in differentiating tissues. For example, SAM function can be affected by failures in lateral organ development or in the establishment of their ad-abaxial polarities (Waites et al. 1998; Lynn et al. 1999). Clearly, the SAM is intricately linked to its differentiating environment by non-cell-autonomous control systems, the extent and molecular mechanisms of which are important issues.

Here, we describe the GRAS gene *HAIRY MERISTEM* (*HAM*) of *Petunia*. *HAM* mediates a signal from lateral organ primordia and stem provasculture that is essential and specific for maintaining the SAM. This defines a novel pathway that links stem cell perpetuation to differentiation.

## Results and discussion

### *HAIRY MERISTEM* is required for meristem maintenance

The recessive *hairy meristem* (*ham-B4281*) mutation was found in a screen for meristem defects in a population of ~60,000 *dTph1* insertions in *Petunia* (Koes et al. 1995). All *ham* mutants (100%, n = 60, stable allele *hamFT-7*; Fig. 3B, below) ceased organ formation during vegetative growth after initiating a variable number of leaves ( $9.9 \pm 3.9$ , n = 60), whereas wild-type plants produced  $17.9 \pm 1.2$  (n = 20) leaves before transition to flowering (Fig. 1A,C). Terminating vegetative SAMs developed a differentiated epidermis with trichomes (Fig. 1E-G), a feature normally found only on leaf primordia and subapical stem. *ham* axillary meristems differentiated likewise (data not shown) but without any organ formation. If axillary shoots did grow out, they arose exclusively from early vegetative nodes and terminated after several leaves. Occasionally, *ham* mutants developed inflorescence nodes from adventitious shoots, as judged by a nearly opposite pair of bracts, and their apices terminated with trichomes on a differentiated epidermis (Fig. 1H,I). Flowers were rare and produced fewer, although normal, floral organs per whorl [3–5 sepals (s), 3–5 petals (p), 1–3 stamens (st), 0 carpels (c), n = 10] than wild type (invariably 5s, 5p, 5st, 2c; Fig. 1B,D). Intact *ham* flowers have never been observed. Floral meristems terminated as flat structures (Fig. 1J) at the cost of inner whorls.

Before termination, *ham* plants could not be distinguished from wild type by shoot morphology, leaf histology, or root growth (data not shown). Thus, *HAM* is specific for postembryonic maintenance of all shoot and floral meristems, reflecting a shared and essential property. The differentiation of epidermis with trichomes on the SAM is a unique feature of *ham*, pointing toward a developmental mechanism that has hitherto not been uncovered.

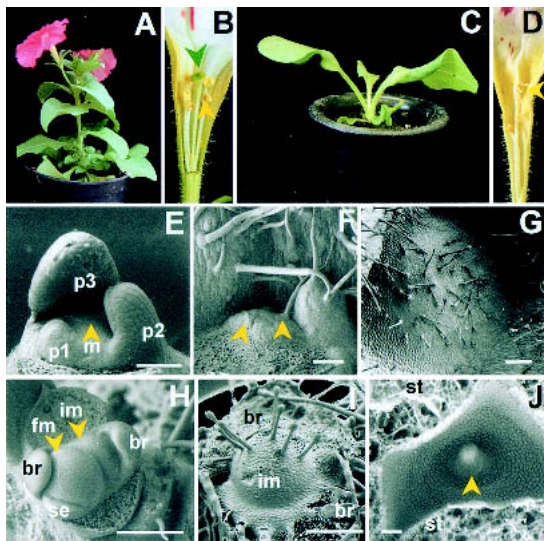
### *ham* meristems differentiate as stem

The wild-type vegetative SAM contains two outer cell layers (L1 and L2; Fig. 2A) with cell divisions predomi-

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<sup>1</sup>Corresponding author.

E-MAIL [cris.kuhlemeier@ips.unibe.ch](mailto:cris.kuhlemeier@ips.unibe.ch); FAX 41-31-332-2059.

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**Figure 1.** Phenotype of *ham* mutants. (A) Wild-type W138 *Petunia*. (B) Wild-type W138 flower with the internal whorls of organs. Five stamens (yellow arrow) surround two fused carpels (green arrow). (C) *ham-B4281* plant, terminated during vegetative rosette growth. (D) *ham* mutant flower lacking two stamens and both carpels. (E) Wild-type vegetative apex. p3, p2, and p1 = leaf primordia in order of decreasing age; m = meristem. (F) Vegetative *ham* apex shortly after termination. Primordium initiation has ceased. The meristem displays ectopic trichomes (arrows). (G) As in F, 2 wk after termination, the central dome has increased in size and is covered with trichomes. (H) Wild-type inflorescence apex. fm = floral meristem, im = inflorescence meristem, br = bract, se = sepal. (I) *ham* inflorescence apex with ectopic trichomes. br = last initiated bracts. (J) *ham* floral meristem, showing termination after initiation of three stamens (st). In place of carpels, a flat apex is visible with a small outgrowth (arrow). Bars, 100  $\mu$ m.

nantly in the anticlinal plane, leading to a gain in surface area. The subepidermal L2 layer shows more frequent periclinal division in the SAM's periphery at sites of leaf initiation (Fig. 2A), which are marked by disappearance of *PhSTM* expression (Fig. 2B). *PhSTM* was isolated in this study as a class-1 *KNOX* gene, most similar to tobacco *NTH15* (Nishimura et al. 1999; data not shown). Its expression pattern is typical for *STM*-like genes (e.g., see Long et al. 1996).

In *ham* seedlings with 2–4 leaves, no obvious histological deviations from wild type could be observed in the SAM (data not shown), consistent with the general absence of a *ham* phenotype at this stage. At termination, *ham* was best recognized by the absence of new leaf primordia (Fig. 2C). In L2, periclinal divisions in the central zone of the SAM were observed more frequently (Fig. 2C), and *PhSTM* expression was weak, although its pattern was normal (Fig. 2D). Two weeks after overt termination, *ham* meristems showed differentiated cell types of epidermis (trichomes), several layers of highly vacuolated subepidermal cells, and vascular differentiation (Fig. 2F). This histological structure is strikingly similar to the radial pattern of stem tissue underneath a wild-type SAM: epidermis, cortex, vasculature, and pith (Fig. 2E). At this stage, *PhSTM* expression could no longer be detected (data not shown).

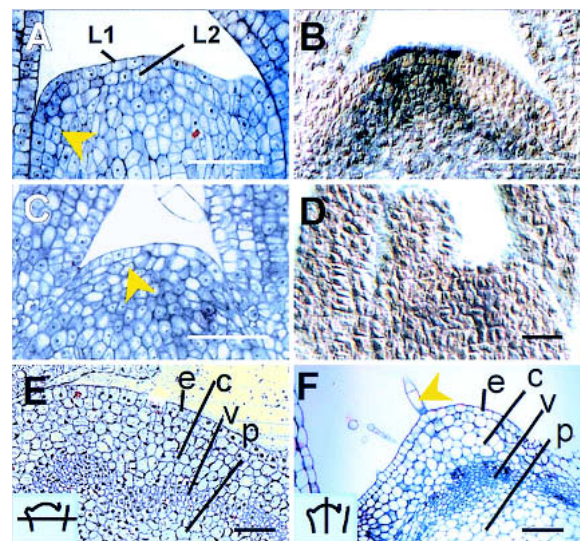
On the basis of these data and the idea that leaf initiation requires local auxin maxima whereas stem develops by default (Reinhardt et al. 2000), we conclude that *ham* meristems differentiate as continuations of the stem.

More importantly, *PhSTM* expression continues for some time after meristem termination, indicating that cells at the *ham* apex lose the ability to respond to *PhSTM* but not the potential to express it in a normal pattern.

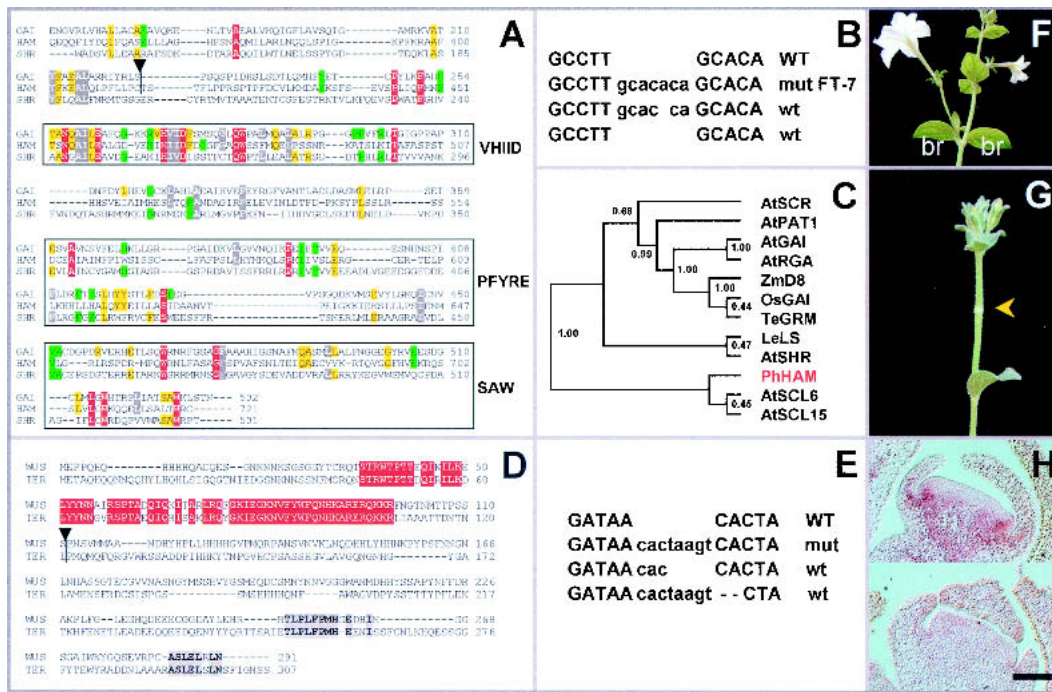
#### *HAM* encodes a GRAS protein

*ham-B4281* was genetically unstable. We differentially displayed *dTph1* transposon insertions using selective inverse PCR (siPCR; see Materials and Methods). A single siPCR product fully cosegregated with the mutation (data not shown). The *ham* phenotype reverted to wild type whenever excision of *dTph1* restored the translational reading frame (Fig. 3B). This shows that the siPCR product is part of *HAM*.

*HAM* cDNA contained a 721 amino acids open reading frame with strong homology with the GRAS family (Fig. 3A; Pysh et al. 1999). The homology extended over all the conserved residues of this family, which are located primarily in the C-terminal half of the protein. GRAS proteins involved in gibberellin signal transduction share a functional amino acid motif in the N-terminal half (DELLA; Dill et al. 2001). *HAM* lacks this motif. Comparative sequence analysis of a variety of GRAS proteins, ranging from gibberellin signaling proteins to root patterning factors, indicated that *HAM* falls into a distinct group with the putative *Arabidopsis* proteins AtSCL6 and AtSCL15 (SCARECROW-like; Fig. 3C). Bootstrap analysis strongly supported this group (frequency 1.00, 1000 samplings), making AtSCL6 and AtSCL15 good candidates for *HAM* orthologs. The *dTph1* element was inserted 413 amino acids down-



**Figure 2.** Histology of *ham* apices. (A) Wild-type vegetative meristem. The arrow indicates a periclinal division in the L2 layer of an initiating leaf primordium. (B) In situ localization of *PhSTM* transcript in a wild-type vegetative apex. The signal (blue) is excluded from the leaf primordia. (C) *ham* vegetative apex showing a cessation of organ initiation and periclinal division in the central zone (arrow). (D) In situ localization of *PhSTM* transcript in a *ham* apex shortly after termination. (E) Transverse section of developing stem, just below a wild-type meristem (section schematized, inset). e = epidermis, c = cortex, v = vasculature, p = pith. (F) Older *ham* apex in longitudinal section showing a layered structure of differentiated tissue. e, c, v, p as in E. Arrow = trichome. Bars, 50  $\mu$ m.



**Figure 3.** Cloning and structure of *HAM* and *TER*. (A) Protein sequence alignment of the C-terminal portion of *HAM* with *SHR* and *GAI*. Absolutely conserved positions are red, and conserved residues are grey (>90%), yellow (>80%), or green (>70%), based on alignments of 12 GRAS sequences as in the cladogram of C. Atypical residues at conserved positions are not colored. VHIID, PFYRE, and SAW are domains as defined in Pysh et al. (1999). Triangle = *dTph1* insertion in *ham-B4281*. (B) *ham* excision alleles. WT = wild-type sequence flanking the insertion (ClustalG software) showing the similarities of 12 GRAS sequences. Numbers indicate bootstrap frequencies of each branchpoint in the cladogram. GenBank accession nos. AtGAI (CAA75492), AtSHR (NP195480), AtPAT1 (AAF73237), AtRGA (CAA75493), LeLS (AAD05242), AtSCL6 (NP191926), AtSCR (AAB06318), AtSCL15 (NP191622), ZmD8 (AAL10319), TeGRM (CAB51555), OsGAI (BAA90749), and PhHAM (AF481952). (D) Full protein alignment of *TER* (GenBank accession no. AF481951) and *WUS* (GenBank accession no. CAA09986). Conservation is given on the basis of these two orthologs only. Red residues indicate positions in the homeobox, and grey residues denote blocks of conspicuous colinearity. (E) As in B but for *ter-B1382*. (F) Wild-type *Petunia* inflorescence producing two bracts (br) and a flower per node. (G) *HAM* cosuppressor showing a node without bracts and flower (arrow). (H) *In situ* hybridization of *HAM* RNA on wild-type (upper) and cosuppressed (lower) floral meristems. Wild type shows a signal in the initiating petal primordia. The cosuppressor lacks this signal. Bar, 50  $\mu$ m.

stream from the start codon in the coding region of *HAM* (Fig. 3A), probably resulting in a null phenotype.

Conclusive evidence for the identity of *HAM* was obtained by cosuppression, using the full cDNA expressed in the sense orientation from the 35S promoter in transgenic plants. One cosuppression line was selected on the basis of absence of endogenous *HAM* expression (Fig. 3H) and analyzed in detail. Although vegetative development was mostly normal, axillary shoots typically terminated in hairy meristems (data not shown). Interestingly, 52% of inflorescence nodes (n = 138, 14 plants) skipped one or more organs, resuming organ initiation in the node that followed (Fig. 3F,G). This is probably a weak *ham* phenotype, with SAM cells differentiating into stem before acquiring organ identity. The recovery of these SAMs indicates that some self-maintaining properties of the plant's stem cell population can compensate for a partial loss of *HAM* function.

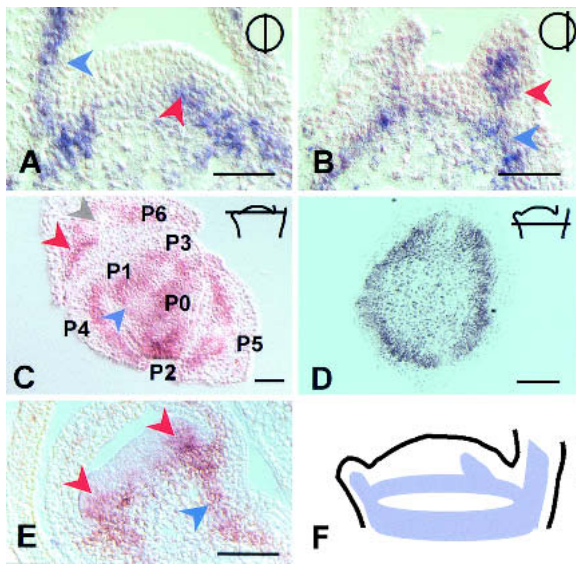
*HAM acts non-cell-autonomously from lateral organ primordia and stem provasculature*

RT-PCR detected *HAM* cDNA in all shoot tips and in roots but not in expanding leaves (data not shown). *HAM* expression was analyzed in detail by RNA *in situ* hybridization. In vegetative apices, *HAM* was expressed in

deeper layers of the meristem at the presumptive site of organ initiation (Fig. 4A), as well as in the developing stem vasculature (Fig. 4B). In transverse sections, a signal was obtained in the developing primordia at least until P6 (Fig. 4C). *HAM* was expressed strongly in the L3-derived ground tissues in the inner part of the primordia and weakly in the main vascular bundle of older primordia (Fig. 4C). Beneath the vegetative meristem, *HAM* was observed in a ring that corresponds to the provasculature of the stem (Fig. 4D). In the inflorescence and floral meristems, similar *HAM* expression was detected consistently in all organ primordia and all floral whorls (Fig. 4E), as well as in provascular tissue (Fig. 4E). Figure 4F summarizes this pattern for a vegetative meristem.

These data indicate that *HAM* expression from the L3 layer of the SAM may be sufficient for function in the overlying layers L1 and L2. To test this, we analyzed the *HAM:ham* segregation ratios for four independent revertant branches on unstable *ham-B4281* homozygotes. These branches developed wild-type flowers indeterminately. Within-flower self-fertilizations from three branches gave rise to nearly normal Mendelian 3:1 (*HAM:ham*) patterns. However, one revertant branch did not segregate a single wild-type progeny among 80 siblings derived from the eighth and tenth flower on the branch. As gametes are derived mostly from L2 (e.g., see





**Figure 4.** Expression pattern of *HAM*. (A) In situ localization of *HAM* transcript in a near median (*top right inset*) longitudinal section through a vegetative apex. Signal is in the developing primordia (blue arrow) and at the presumptive position of a newly initiating primordium (red arrow). (B) As in A, with a section located more peripherally (*top right inset*). The signal is seen in a developing primordium (red arrow), as well as in a ring-shaped pattern that corresponds to the developing stem vasculature (blue arrow). (C) As in A, but in a transverse section. The position of the section is indicated in the *top right inset*. The signal is observed in the inner ground tissues of the primordia (red arrow) and is weaker in the main vascular bundle of older primordia (grey arrow). The blue arrow indicates *HAM* expression in a ring-shaped pattern that merges with primordia P1 and P0 and corresponds to provascular tissue of the differentiating stem. P6, P5, and so forth indicate the consecutive order of primordium initiation with decreasing age. (D) As in C, but at a position just below the meristem (indicated in *top right inset*). *HAM* expression is seen as a ring that corresponds to the provascularization of the stem. (E) *HAM* localization during development of the floral meristem, as exemplified for initiating petal primordia. Expression is observed in inner cell layers at the site of petal initiation (red arrows) and in subtending provascular tissue of the developing pedicel (blue arrow). (F) Schematic representation of the *HAM* expression pattern as exemplified for a wild-type vegetative meristem. Bar, 50  $\mu$ m.

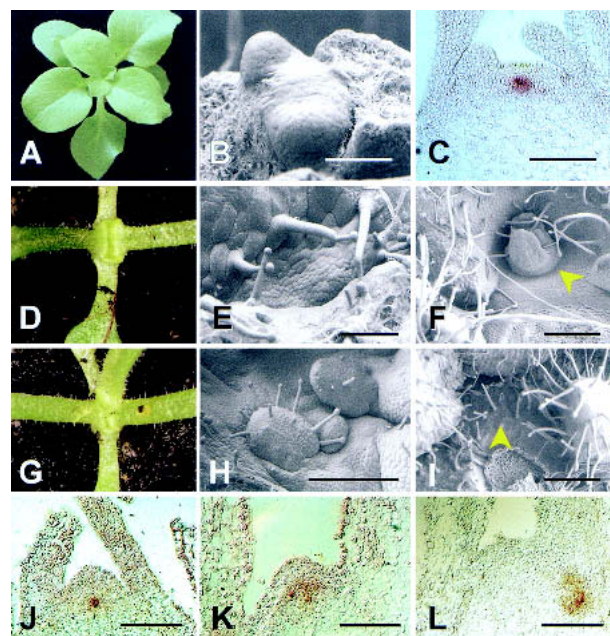
Stewart and Burk 1970), the wild-type phenotype of this branch was conferred by cells that were not related clonally to L2. Given the expression pattern of *HAM*, this branch was most likely a periclinal chimera with a revertant L3. Taken together, this essentially shows a non-cell-autonomous mode of action of *HAM*. Because revertant branches did not influence the mutant phenotype in other parts of the same plant, the signal must be short range.

#### *HAM* acts in parallel with TERMINATOR

*ham* has some important similarities with *wuschel* mutants of *Arabidopsis* (the meristem is not maintained in both). To investigate the relationships between *HAM* and *WUS*, we isolated the *Petunia WUS* ortholog from a mutant *terminator* (*ter-B1382*). Like *wus*, *ter* ceased shoot development after the two first true leaves (Fig. 5A,D), continuously reinitiating ectopic leaves and defective meristems from flat apices (stop-and-go growth; Fig. 5E,F). This led to increasingly bushy plants that

rarely flowered. Occasional flowers had fewer floral organs per concentric whorl (data not shown), strongly resembling *wus* mutants (Laux et al. 1996). A single siPCR product cosegregated with *ter* (data not shown). The predicted *TER* protein was highly similar to *WUS* (Fig. 3D), mainly in the homeobox and with conspicuous blocks of homology in the C terminus. *ter* could be reverted to wild type whenever excision of *dTph1* restored the open reading frame (Fig. 3E). *TER* gene expression patterns were identical to *WUS* (Fig. 5C). *dTph1* was inserted 12 amino acids downstream from the homeobox, probably resulting in a null mutation. We refer to *TER* as *PhWUS* and to its mutation as *ter*.

During early vegetative growth, *ter ham* double mutants showed an initial stop-and-go growth characteristic for *ter* but subsequently started to display *ham* phenotypes (Fig. 5F,I). We compared the structure of 15 meristems of both *ter* single mutants and *ter ham* double mutants in mature plants. In *ter* single mutants, we found ectopic leaves and meristems on a flat apex in 13 of 15 cases (Fig. 5F). In contrast, in *ter ham* double mutants, the typical *ham* phenotype of a trichome-covered



**Figure 5.** Relations between *ham* and *ter*. (A) Wild-type *Petunia* during vegetative rosette growth. (B) Wild-type shoot apical meristem (SAM) with the first two true leaf primordia. Cotyledons have been removed. (C) In situ localization of *TER* (*PhWUS*) transcripts in a wild-type vegetative apex. (D) *ter-B1382* seedling. Growth has ceased after production of the first two true leaves. (E) *ter* seedling apex after initiation of the two first leaves. A flat, disorganized structure replaces the SAM. (F) *ter* apex with an ectopic meristem (arrow, stop-and-go growth). (G) *ter ham* double mutant seedling. An additional leaf, compared with D, occurred with a low frequency in *ter* single mutants as well. (H) *ter ham* double mutant seedling apex. Initiation of ectopic leaves is observed (stop-and-go). (I) *ter ham* double mutant apex on an older plant. The SAM displays a trichome covered surface characteristic for *ham* single mutants. (J) In situ localization of *PhWUS* transcripts in a *ham* mutant apex shortly after termination. The signal is essentially normal. (K) In situ localization of *PhWUS* transcripts in a later *ham* mutant apex. Expression occurs in a disorganized pattern. (L) As in K. *PhWUS* expression in the main apex has disappeared. In the axillary position, expression is disorganized and deeply internal. Bars: B, 25  $\mu$ m; C,E,I, 50  $\mu$ m; F,I, 200  $\mu$ m; H,K,L, 100  $\mu$ m.

surface was found in 10 of 15 cases (Fig. 5I). In the other five cases, the surface was flat and lacked trichomes but showed no sign of resumption (data not shown). Thus, in early seedling stage *ter ham* double mutants closely resemble *ter*, but at later stages they behave like *ham*. In situ hybridization on *ham* single mutants showed that, at termination, *PhWUS* expression was essentially normal (Fig. 5J). In later stages, we observed no expression (Fig. 5L) or a patchy pattern (Fig. 5K). *PhWUS* expression in axillary *ham* meristems did the same (Fig. 5L).

We conclude that specification and maintenance of stem cells by *HAM* and *WUS* are largely parallel processes, although maintenance of spatially correct *PhWUS* expression indirectly depends on *HAM*. Like *PhSTM*, *PhWUS* expression continues for some time after meristem termination. Thus, cells in a differentiating *ham* shoot apex lose the ability to respond to *PhWUS* activity but not the potential to express it.

#### Role of HAM in shoot meristem and stem cell maintenance

By acting from L3 of organ primordia and provasculation of the stem, *HAM* affects communication among differentiating and nondifferentiating cells of the SAM. *HAM* is likely to signal cell fate decisions in the shoot apex, promoting the undifferentiated state as a distinct cellular identity. Judging from the nonpleiotropic nature of the *ham* mutation, this may be the single most important function of *HAM*. It might work in a similar way as its homolog *SHORTROOT* (*SHR*) of *Arabidopsis*, which controls cell fate by intercellular movement (Nakajima et al. 2001).

The antidifferentiation function of *HAM* is distinct from that of *STM* and *WUS*. The loss of responsiveness of apical cells to *PhSTM* and *PhWUS* in *ham* apices establishes a hitherto unknown cellular precondition on which these homeobox factors must act. We propose that, by means of *HAM*, cells that exit the meristematic domain and initiate differentiation actively maintain a field of uncommitted apical cells, protecting against a default path of development into stem. In the presence of a factor like *WUS*, these apical cells may acquire stem cell identity (Schoof et al. 2000). In the absence of *STM*, conversely, they may develop into lateral organs (e.g., see Byrne et al. 2000).

In root and shoot systems of plants, the differentiation and division patterns of stem cell daughters are directed by positional information from surrounding cells (Stewart and Burk 1970; van den Berg et al. 1995). In the root, initial cells for radial cell files differentiate like their maturing neighbors unless inhibited by the quiescent center (van den Berg et al. 1997). Our data indicate that, also in the shoot, stem cell daughters differentiate according to a prepattern present in more mature (stem-) tissue, and that maintenance of stem cell activity depends on extrinsic antidifferentiation factor(s). Whereas the molecular identities of antidifferentiation activities in the root are unknown, *HAM* is a likely candidate for those of the shoot.

*HAM* adds a new level of complexity to the developmental interactions in the meristem by showing a specific, essential role of differentiating cells in maintaining the undifferentiated state. Its biological meaning would be the coupling of stem cell fate to differentiation, effec-

tively assuring indeterminate growth potential by a positive feedback mechanism on tissue level.

#### Materials and methods

##### Transposon mutagenesis and mutant construction

For transposon mutagenesis, ~2000 M1 plants (*Petunia hybrida* strain W138) from 600 different families were self-fertilized, and M2 families were inspected visually for developmental abnormalities. Mutants were verified in segregation studies and manually cross-pollinated for double mutant construction. *ham ter* double mutants were identified on the basis of their *dTph1* insertion genotype using PCR. All plants were grown in a greenhouse under standard conditions.

##### Structural analysis and histology

Scanning electron microscopy was performed on fresh plant material with Hitachi S-3500N equipment. Histological sections were made by fixing tissue in 4% formaldehyde in 50-mM sodium phosphate buffer (pH = 7.2), dehydrating through an ethanol series, embedding in Epon (Fluka), sectioning to 3  $\mu$ m, and staining with 0.1% toluidine blue. All sections shown were selected for near-median plane, unless indicated otherwise. For sections, only stable *ham* mutants were used that carried a translational frameshift allele (*hamFT-7*; see Fig. 3B).

##### Gene isolation and analysis

Full-length *PhSTM* (GenBank accession no. AY112704) and *HAM* (GenBank accession no. AF481952) cDNAs were isolated by filter screening of a ZAPII cDNA library of young inflorescence shoot tips (gift of R. Koes). *PhSTM*: heterologous probe from tomato LeT6 (Janssen et al. 1998, gift of N. Sinha). *HAM*: siPCR fragment as probe. The full coding region of *PhWUS* (GenBank accession no. AF481951) was amplified and sequenced with primers *ter*5'F (5'-GCACGAGGCCATATTCTCTTCACT-3') and *ter*3'R (5'-CCATGGTCCAAAGATACGTTAGTAC-3') from the above cDNA library.

For selective iPCR (siPCR), genomic DNA was digested with *Mbo*I, self-ligated with T4 ligase, and religated with *Ssp*I. Such templates were selectively preamplified using *dTph1*-specific primers *out4* (5'-GAACGGTTGTCCTCTTGAACC-3') and *out6* (5'-GTGGCAGCCCA GATTCGATCnn-3'). *Out6* flanks an *Mbo*I site in *dTph1* and carries any of 16 possible combinations of 3'-terminal bases (NN) extending into plant DNA. Preamplifications were reamplified with *out6* + NN primers and a nested *dTph1-out1* primer (5'-GGGAATTCGCTCCGCCCTG-3'). Products were separated on 5% native polyacrylamide gels and stained with SYBR Gold (Molecular Probes, Inc.). Candidates were cut from the gel, eluted in water, reamplified, and sequenced. A detailed protocol is available from the authors.

To analyze footprint alleles for *ham* and *ter*, genomic DNA was isolated from revertant branches on homozygous insertion mutants or from their progeny. DNAs were PCR amplified with *ham3* (5'-CAAGGAG GCTCTGCAGTTGC-3') and *ham4* (5'-CTTCAGAAAAGGACTTATA AGCATCC-3') or *terF* (5'-GAAAAGGCTTATTGCTGCTGCTACC-3') and *terR* (5'-CCTAAAGCTCTTCTCCATAGCTAAAG-3'). Products were separated on 5% polyacrylamide gels, and bands with small size increases relative to wild type were eluted, reamplified, and sequenced.

##### Expression analyses

RT-PCR was performed on first strand cDNA using gene-specific primers for the 3' untranslated region of *HAM* cDNA (*HAM*3'F, 5'-AGCTTTAT CTAATGAAGCCAGAAGAGG-3'; *HAM*3'R, 5'-ACAGGGAAATGAG ACAAGATACATCAC-3'). In situ hybridization was performed using DIG-labeled antisense riboprobes according to the protocol at <http://www.wisc.edu/genetics/CATG/barton/protocols.html>. *PhWUS* and *PhSTM* probes were from full-length cDNAs. *HAM* probes were either full-length cDNA or a 976-bp probe from the nonconserved 5' part of the cDNA (corresponding to amino acids 1–325). Both *HAM* probes gave identical results.

##### Transgenics

The full *HAM* coding region was amplified with *Pfu* DNA polymerase and primers *hamATG* (5'-ATCTAGAGTTTAAAGATGATTGCAATAC C-3') and *hamTGA* (5'-AAGATCTCTCCTCAGCACCTCCAAGTC-3'), digested with *Bgl*II and *Xba*I, and inserted in the sense orientation into a pBI121 (Clontech)-derived vector to replace the GUS cassette. Transfor-

mation was into *P. hybrida* W115 using *Agrobacterium tumefaciens* LBA4404.

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## References

- Brand, U., Fletcher, J.C., Hobe, M., Meyerowitz, E.M., and Simon, R. 2000. Dependence of stem cell fate in *Arabidopsis* on a feedback loop regulated by CLV3 activity. *Science* **289**: 617–619.
- Byrne, M., Barley, R., Curtis, M., Arroyo, J.M., Dunham, M., Hudson, A., and Martienssen, R.A. 2000. *Asymmetric leaves1* mediates leaf patterning and stem cell function in *Arabidopsis*. *Nature* **408**: 967–971.
- Dill, A., Jung, H.-S., and Sun, T.-P. 2001. The DELLA motif is essential for gibberellin-induced degradation of RGA. *Proc. Natl. Acad. Sci.* **98**: 14162–14167.
- Janssen, B.J., Williams, A., Chen J.J., Mathern, J., Hake, S., and Sinha N. 1998. Isolation and characterisation of two knotted-like homeobox genes from tomato. *Plant Mol. Biol.* **36**: 417–425.
- Koes, R.E., van Houwelingen A., Mur, L., Spelt, C., Quattrocchio, F., Wing, J., Oppedijk, B., Ahmed, S., Maes, T., Gerats, T., et al. 1995. Targeted gene inactivation in *Petunia* by PCR based selection of transposon insertion mutants. *Proc. Natl. Acad. Sci.* **92**: 8149–8153.
- Laux, T., Mayer, K.F.X., Berger, J., and Jürgens, G. 1996. The *WUSCHEL* gene is required for shoot and floral meristem integrity in *Arabidopsis*. *Development* **122**: 87–96.
- Lenhard, M., Bohnert, A., Jürgens, G., and Laux, T. 2001. Termination of stem cell maintenance in *Arabidopsis* floral meristems by interactions between *WUSCHEL* and *AGAMOUS*. *Cell* **105**: 805–814.
- Lohmann, J.U., Hong, R.L., Hobe, M., Busch, M., Parcy, F., Simon, R., and Weigel, D. 2001. A molecular link between stem cell regulation and floral patterning in *Arabidopsis*. *Cell* **105**: 793–803.
- Long, J.A., Moan, E., Medford, J., and Barton M.K. 1996. A member of the *KNOTTED* class of homeodomain proteins encoded by the *STM* gene of *Arabidopsis*. *Nature* **379**: 66–69.
- Lynn, K., Fernandez, A., Aida, M., Sedbrook, J., Tasaka, M., Masson, P., and Barton, M.K. 1999. The *PINHEAD/ZWILLE* gene acts pleiotropically in *Arabidopsis* development and has overlapping functions with the *ARGONAUTE1* gene. *Development* **126**: 469–481.
- Nakajima, K., Sena, G., Nawy, T., and Benfey, P. 2001. Intercellular movement of the putative transcription factor SHR in root patterning. *Nature* **413**: 307–311.
- Nishimura, A., Tamaoki, M., Sato, Y., and Matsuoka, M. 1999. The expression of tobacco knotted1-type class1 homeobox genes corresponds to regions predicted by the cytohistological zonation model. *Plant J.* **18**: 337–347.
- Pysh, L., Wysocka-Diller, J.W., Camilleri, C., Bouchez, D., and Benfey, P. 1999. The *GRAS* gene family in *Arabidopsis*: Sequence characterization and basic expression analysis of the *SCARECROW-LIKE* genes. *Plant J.* **18**: 111–119.
- Reinhardt, D., Mandel, T., and Kuhlemeier, C. 2000. Auxin regulates the initiation and radial position of plant lateral organs. *Plant Cell* **12**: 507–518.
- Schoof, H., Lenhard, M., Haecker, A., Mayer, K.F., Jürgens, G., and Laux, T. 2000. The stem cell population of *Arabidopsis* shoot meristems is maintained by a regulatory loop between the *CLAVATA* and *WUSCHEL* genes. *Cell* **100**: 635–644.
- Sinha, N., Williams, R.E., and Hake S. 1993. Overexpression of the maize homeobox gene *KNOTTED-1* causes a switch from determinate to indeterminate cell fates. *Genes & Dev.* **7**: 787–795.
- Stewart, R.N. and Burk, L.G. 1970. Independence of tissues derived from apical layers in ontogeny of the tobacco leaf and ovary. *Amer. J. Bot.* **57**: 1010–1016.
- van den Berg, C., Willemsen, V., Hage, W., Weisbeek, P., and Scheres, B. 1995. Cell fate in the *Arabidopsis* root meristem is determined by directional signalling. *Nature* **378**: 62–65.
- van den Berg, C., Willemsen, V., Hendricks, G., Weisbeek, P., and Scheres, B. 1997. Short range control of cell differentiation in the *Arabidopsis* root meristem. *Nature* **378**: 62–65.
- Waites, R., Selvadurai, H.R.N., Oliver, I., and Hudson, A. 1998. The *PHANTASTICA* gene encodes a MYB transcription factor involved in growth and dorsoventrality of lateral organs in *Antirrhinum*. *Cell* **93**: 779–789.
- Weigel, D. and Jürgens, G. 2002. Stem cells that make stems. *Nature* **415**: 751–754.