## SUPPLEMENTARY FIGURES



Supplementary Figure 1. Domain-specific promoter activities, related to Figure 1.

(**a-f**) Immunolocalization of *pCLV3::HF-RPL18* (**a**), and expression of *pWUS>>GFP-ER* (**b**), *pUFO>>GFP-ER* (**c**), *pAS2>>GFP-ER* (**d**), *pFIL>>GFP-ER* (**e**) and *pATML1>>GFP-ER* (**f**) in longitudinal sections through vegetative shoot apex of 7 DAG wild-type like plants. Bars =  $10\mu$ m. (**g**) Pattern of GUS expression in a *pPTL>>GUS* plant displaying leaf marginal expression. Bars =  $500\mu$ m. A dotted line in (**a**) indicates the meristem.



## Supplementary Figure 2. Schematic diagram of TRAP-seq.

Shapes represent different kinds of protein, and lines show RNAs. Polysomes and anti-flag beads are indicated. The polysomes in salmon with a star represent the domain specifically expressed ones with HF-epitopes, which can be purified by anti-FLAG beads. Seedlings at 7 DAG (without roots) were ground and HF-tagged polysomes together with translating mRNA were purified by affinity to anti-FLAG beads. The mRNAs, expressed in specific domains, were then extracted and used for library construction and sequencing.



Supplementary Figure 3. Pearson correlation coefficients of translatome data.

Pearson correlation coefficients of translatome data from biological replicates for all profiled tissue domains. The three biological replicates are tightly related to each other and separate from other domains.



Supplementary Figure 4. Complete hierarchical clustering analysis.

Complete hierarchical clustering analysis of all expressed genes using expression value (Log<sub>10</sub>RPKM) from biological replicates for different domains, related to Fig. 2. The three biological replicates clustering together indicates the reproducibility of the translatome data.



**Supplementary Figure 5.** Expression patterns of known marker genes, related to Fig. 1b and Supplemental Data 1.



Supplementary Figure 6. Histograms of expression patterns.

Histograms of expression patterns of expressed transcripts represented by the *z*-score. Divergent distributions are displayed with domain-specific pattern, related to Fig. 2.



Supplementary Figure 7. Domain specifically expressed genes.

(a) Numbers of domain-specifically expressed genes identified by pair-wise comparison.
(b) Venn diagrams showing domain enriched genes identified by three methods.



**Supplementary Figure 8.** GO analysis of domain-enriched genes showing enrichment of selected GO categories.

Expression patterns of enriched genes in each category are depicted in heat maps. Colors represent the relative expression values indicated by *z*-score with low values in blue and high values in yellow.



Supplementary Figure 9. Enrichment analysis for different domains.

(a) Domain-specific enriched hormone-responsive genes. Red indicates enrichment and blue indicates depletion. Only significantly over-represented categories (FDR adjusted *P*-value < 0.05 by hypergeometric test) are colored. (b) Domain-specific enriched transcription factor families. Only significantly over-represented families (FDR adjusted *P*-value < 0.05 by hypergeometric test) are colored. (c) Domainspecific enriched of known *cis*-elements in specific domains. Only significantly overrepresented motifs (*E*-value < 10<sup>-4</sup>) are colored.



**Supplementary Figure 10.** Numbers of all genes and transcription factor-encoding genes identified in each GCN module.



**Supplementary Figure 11.** The gene structures and mutation sites of *RBE* (**a**) and *HAN* (**b**) genes.

Exons are shown in boxes and introns in lines. Boxes in dark blue represent coding sequences, in which key motifs are displayed in different colors. A T-DNA insertion site of *rbe-2* and two mutation sites of *han-2* and *han-30* are denoted with triangles.



Supplementary Figure 12. Expression pattern comparisons between epidermal domain and leaf primordial domain.

(a) Venn diagram of domain-enriched genes that exhibited significant (FC  $\geq$  two-fold with FDR adjusted *P*-value  $\leq$  0.01) up-regulation in epidermal domain (*ATML1*, red) or leaf primordial domain (*AS1*, blue). The number in the middle indicates expressed genes without domain specificity. (b) GO analysis of differentially expressed genes in epidermal domain identified significantly enriched gene categories in Cell Wall Organization, Wax Biosynthesis, and Response to Auxin. Aggregate size indicates significance levels for categories by a Yekutieli test with a FDR correction which was summarized by REVIGO. (c) Domain-specific enrichment of transcription factors families (FDR adjusted *P*-value  $\leq$  0.05) between the two domains. (d) Domain-specific enrichment of hormone activated and repressed genes (FDR adjusted *P*-value  $\leq$  0.05) between the two domains.



**Supplementary Figure 13.** Expression pattern comparisons between abaxial and adaxial leaf domains.

(a) Venn diagram of cell domain-enriched genes that exhibited significant (FC  $\geq$  twofold with FDR adjusted *P*-value  $\leq$  0.01) up-regulation in abaxial (*FIL*, red) or adaxial (*AS2*, blue) leaf domains. The number in the middle indicates expressed genes without domain specificity. (b) Domain-specific enrichment of hormone activated and repressed genes (FDR adjusted *P*-value  $\leq$  0.05) between the two domains. (c) Domain-specific enrichment of transcription factor families in the two domains (FDR adjusted *P*-value  $\leq$  0.05). (d) Domain-specific enriched known *cis*-elements in abaxial and adaxial leaf domains. Only significantly overrepresented (*E*-value < 10<sup>-4</sup>) classes were colored.



Supplementary Figure 14. Cutoffs and threshold (a) and dynamic tree cut (b and c) used for co-expression network construction by WGCNA.

## SUPPLEMENTARY TABLES

Domain	Library Siza	Unique Menned	Multi Mannad	Unique
Domain	Library Size	ibrary Size Unique Mapped Mu		Mapped Ratio
AS2-R1	20,407,777	19,103,058	1,304,719	93.61%
AS2-R2	20,948,752	19,550,404	1,398,348	93.32%
AS2-R3	21,461,115	20,106,508	1,354,607	93.69%
<i>ATML1</i> -R1	45,740,291	43,416,745	2,323,546	94.92%
ATML1-R2	19,578,140	18,306,700	1,271,440	93.51%
ATML1-R3	12,341,409	11,609,417	731,992	94.07%
<i>CLV3</i> -R1	13,089,870	12,208,916	880,954	93.27%
CLV3-R2	18,068,682	16,774,903	1,293,779	92.84%
CLV3-R3	21,013,986	19,786,117	1,227,869	94.16%
<i>FIL</i> -R1	20,480,184	19,470,951	1,009,233	95.07%
FIL-R2	21,343,660	20,334,273	1,009,387	95.27%
FIL-R3	23,568,618	22,415,861	1,152,757	95.11%
<i>PTL</i> -R1	20,676,628	19,655,125	1,021,503	95.06%
PTL-R2	19,327,737	18,332,593	995,144	94.85%
PTL-R3	22,624,393	21,468,061	1,156,332	94.89%
<i>UFO'</i> -R1	23,725,720	22,727,306	998,414	95.79%
<i>UFO'</i> -R2	24,694,401	23,705,747	988,654	96.00%
<i>UFO'</i> -R3	29,554,451	28,327,045	1,227,406	95.85%
WUS-R1	18,284,149	17,631,889	652,260	96.43%
WUS-R2	18,487,847	17,864,222	623,625	96.63%
WUS-R3	16,726,233	16,137,075	589,158	96.48%

Supplementary Table 1. Summary of reads mapping.

GO term	Ontology	Description	Number in	Number in	
Goterni		Description	input list	BG/Ref	TDK
GO:0044085	<b>Biological Process</b>	Cellular component biogenesis	66	433	1.54E-15
GO:0009058	<b>Biological Process</b>	Biosynthetic process	198	2933	2.66E-06
GO:0044249	<b>Biological Process</b>	Cellular Biosynthetic process	196	2783	2.00E-07
GO:0034645	<b>Biological Process</b>	Cellular macromolecule biosynthetic process	162	1920	1.61E-11
GO:0009853	<b>Biological Process</b>	Photorespiration	7	34	3.99E-02
GO:0009059	<b>Biological Process</b>	Macromolecule biosynthetic process	163	1937	1.61E-11
GO:0015979	<b>Biological Process</b>	Photosynthesis	21	107	1.16E-06
GO:0006091	<b>Biological Process</b>	Generation of precursor metabolites and energy	19	173	1.77E-02
GO:0019538	<b>Biological Process</b>	Protein metabolic process	132	2139	1.05E-02
GO:0019684	<b>Biological Process</b>	Photosynthesis, light reaction	12	63	1.71E-03
GO:0010467	<b>Biological Process</b>	Gene expression	162	2059	2.48E-09
GO:0022900	<b>Biological Process</b>	Electron transport chain	10	52	7.31E-03
GO:0045454	<b>Biological Process</b>	photosynthetic electron transport in photosystem I	6	13	1.06E-03
GO:0022613	<b>Biological Process</b>	Ribonucleprotein complex biogenesis	45	179	6.71E-19
GO:0042254	<b>Biological Process</b>	Ribosome biogenesis	45	173	2.25E-19
GO:0009834	<b>Biological Process</b>	Secondary cell wall biogenesis	5	14	1.77E-02
GO:0034622	<b>Biological Process</b>	Cellular macromolecular complex assembly	16	143	3.99E-02
GO:0006412	<b>Biological Process</b>	Translation	102	505	9.29E-34
GO:0044267	<b>Biological Process</b>	Cellular protein metabolic process	127	1853	2.06E-04
GO:0019725	<b>Biological Process</b>	Photosynthetic electron transport chain	7	27	1.08E-02

Supplementary Table 2. Gene Ontology terms of leaf adaxial enriched genes.

Lines	Sources	Background	Expression pattern
pAS2::LhG4	This work	Ler	Leaf adaxial
pATML1::LhG4	From Thomas Laux <sup>1</sup>	Ler	L1 layer
pCLV3::HF-RPL18	This work	Ler	Central zone
pFIL::LhG4	From Yuval Eshed <sup>2</sup>	Ler	Leaf abaxial
pOp::GFP-ER	From Yuval Eshed <sup>2</sup>	Ler	
pOp::HF-RPL18 35S::AP1-GR ap1-1 cal1	This lab <sup>3</sup>	Ler	
pPTL::LhG4	From David Smyth⁴	Col-0	Leaf margin
pUFO::LhG4	This work	Ler	Peripheral zone
pWUS::LhG4	From Thomas Laux <sup>5</sup>	Ler	Organization center
rbe-2	From Tengbo Huang <sup>6</sup>	Col-0	
han-2	From Elliot M. Meyerowitz <sup>7</sup>	Col-0	
han-30	From Hirokazu Tsukaya <sup>8</sup>	Col-0	

Supplementary Table 3. Transgenic and mutant lines used in this work.

Supplementary Table 4. Primers used in this work.

Primer name	Primer sequence	Usage
pAS2-F-Sall	gtcgacTAGGGCCCGAAATGTGGAAA	Construction of pAS2::LhG4
pAS2-R-Pstl	aactgcagTGTTGAAGAAGATGCCAT	Construction of pAS2::LhG4
pCLV3-FP-Sall	aaagtcgacCATATCCATAAAATTAATCGAATT	Construction of pCLV3::HF-RPL18
pCLV3-RP-Pstl	aaactgcagTTTTAGAGAGAAAGTGACTGAG	Construction of pCLV3::HF-RPL18
tCLV3-FP-Clal	aaaatcgatCCTAATCTCTTGTTGCTTTAAA	Construction of pCLV3::HF-RPL18
tCLV3-RP-BamHI	aaaggatccCTACAATGTTGATTACTTTGGTATA	Construction of pCLV3::HF-RPL18
pCLV3-iden-F	CTGTCCCCTTCTCATTTC	Sequencing
HF-RPL18-R	AACCTTGAATCCACGACTCTTC	Sequencing
pUFO-FP	atgtcgacCGACACGCAACGACTCCTATCG	Construction of pUFO::LhG4
pUFO-RP	gggtaccTTTAGCTGAAAAATGAAAAGATTT	Construction of pUFO::LhG4
SALK_037010-RP	CTTTATTTTGGCCGTTTAGGG	Genotyping of <i>rbe-2</i>
SALK_037010-LP	GGAGTTCACATGGAATACCAGAC	Genotyping of <i>rbe-2</i>
LB1.3	ATTTTGCCGATTTCGGAAC	Genotyping of <i>rbe-2</i>
han-30-FP	ATGCAGACTCCGTACACTAC	Genotyping of han-30
han-30-RP	TTGGTGTGAATCAAGTCCCAAA	Genotyping of han-30
han-2-FP	tttgcagTCCCTATGCAACGC	Genotyping of han-2
han-2-RP	CTATCCGCTACATTAAGCCTC	Genotyping of han-2

## **Supplementary References**

- Lenhard, M. & Laux, T. Stem cell homeostasis in the *Arabidopsis* shoot meristem is regulated by intercellular movement of CLAVATA3 and its sequestration by CLAVATA1. *Development* **130**, 3163-3173 (2003).
- Goldshmidt, A., Alvarez, J.P., Bowman, J.L. & Eshed, Y. Signals derived from YABBY gene activities in organ primordia regulate growth and partitioning of Arabidopsis shoot apical meristems. *Plant Cell* 20, 1217-1230 (2008).
- Jiao, Y. & Meyerowitz, E.M. Cell-type specific analysis of translating RNAs in developing flowers reveals new levels of control. *Mol Syst Biol* 6, 419 (2010).
- Brewer, P.B. et al. *PETAL LOSS*, a trihelix transcription factor gene, regulates perianth architecture in the *Arabidopsis* flower. *Development* **131**, 4035-4045 (2004).
- Gross-Hardt, R., Lenhard, M. & Laux, T. WUSCHEL signaling functions in interregional communication during *Arabidopsis* ovule development. *Genes & development* 16, 1129-1138 (2002).
- Huang, T., Lopez-Giraldez, F., Townsend, J.P. & Irish, V.F. RBE controls microRNA164 expression to effect floral organogenesis. *Development* 139, 2161-2169 (2012).
- Zhang, X. et al. Transcription repressor HANABA TARANU controls flower development by integrating the actions of multiple hormones, floral organ specification genes, and GATA3 family genes in *Arabidopsis*. *Plant Cell* 25, 83-101 (2013).
- Kanei, M., Horiguchi, G. & Tsukaya, H. Stable establishment of cotyledon identity during embryogenesis in *Arabidopsis* by *ANGUSTIFOLIA3* and *HANABA TARANU. Development* **139**, 2436-2446 (2012).
- Sessions, A., Weigel, D., & Yanofsky, M.F. The Arabidopsis thaliana MERISTEM LAYER 1 promoter specifies epidermal expression in meristems and young primordia. Plant J. 20, 259-263 (1999).
- 10. Abe, M., Katsumata, H., Komeda, Y., & Takahashi, T. Regulation of shoot

epidermal cell differentiation by a pair of homeodomain proteins in *Arabidopsis*. *Development* **130**, 635-643 (2003).

- Siegfried, K.R.*et al.* Members of the YABBY gene family specify abaxial cell fate in Arabidopsis. Development **126**, 4117-4128 (1999).
- Sawa, S.*et al. FILAMENTOUS FLOWER*, a meristem and organ identity gene of *Arabidopsis*, encodes a protein with a zinc finger and HMG-related domains. *Genes Dev.* 13, 1079-1088 (1999).
- Kerstetter, R.A., Bollman, K., Taylor, R.A., Bomblies, K., & Poethig, R.S. KANADI regulates organ polarity in *Arabidopsis*. *Nature* 411, 706-709 (2001).
- Palatnik, J.F. *et al.* Control of leaf morphogenesis by microRNAs. *Nature* 425, 257-263 (2003).
- 15. Iwakawa, H.*et al.* Expression of the ASYMMETRIC LEAVES2 gene in the adaxial domain of Arabidopsis leaves represses cell proliferation in this domain and is critical for the development of properly expanded leaves. *Plant J.* **51**, 173-184 (2007).
- Eshed, Y., Baum, S.F., Perea, J.V., & Bowman, J.L. Establishment of polarity in lateral organs of plants. *Curr. Biol.* **11**, 1251-1260 (2001).
- Greb, T. *et al.* Molecular analysis of the *LATERAL SUPPRESSOR* gene in *Arabidopsis* reveals a conserved control mechanism for axillary meristem formation. *Genes Dev.* **17**, 1175-1187 (2003).
- Vroemen, C.W., Mordhorst, A.P., Albrecht, C., Kwaaitaal, M.A., & de Vries, S.C. The CUP-SHAPED COTYLEDON3 gene is required for boundary and shoot meristem formation in Arabidopsis. Plant Cell 15, 1563-1577 (2003).
- 19. Takeda, S. *et al.* CUP-SHAPED COTYLEDON1 transcription factor activates the expression of *LSH4* and *LSH3*, two members of the ALOG gene family, in shoot organ boundary cells. *Plant J.* **66**, 1066-1077 (2011).
- 20. Belles-Boix, E. *et al. KNAT6*: an Arabidopsis homeobox gene involved in meristem activity and organ separation. *Plant Cell* **18**, 1900-1907 (2006).
- 21. Yadav, R.K., Tavakkoli, M., Xie, M., Girke, T., & Reddy, G.V. (2014). A highresolution gene expression map of the *Arabidopsis* shoot meristem stem cell

niche. Development 141, 2735-2744.

- Brand, U., Fletcher, J.C., Hobe, M., Meyerowitz, E.M., & Simon, R. Dependence of stem cell fate in *Arabidopsis* on a feedback loop regulated by *CLV3* activity. *Science* 289, 617-619 (2000).
- 23. Lee, I., Wolfe, D.S., Nilsson, O., & Weigel, D. A *LEAFY* co-regulator encoded by *UNUSUAL FLORAL ORGANS. Curr. Biol.* **7**, 95-104 (1997).
- 24. Mayer, K.F. *et al.* Role of *WUSCHEL* in regulating stem cell fate in the *Arabidopsis* shoot meristem. *Cell* **95**, 805-815 (1998).