SMALL ACIDIC PROTEIN1 Acts with RUB Modification Components, the COP9 Signalosome, and AXR1 to Regulate Growth and Development of Arabidopsis1[C][W][OA]

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Previously, a dysfunction of the SMALL ACIDIC PROTEIN1 (SMAP1) gene was identified as the cause of the anti-auxin resistant1 (aar1) mutant of Arabidopsis (Arabidopsis thaliana). SMAP1 is involved in the response pathway of synthetic auxin, 2,4-dichlorophenoxyacetic acid, and functions upstream of the auxin/indole-3-acetic acid protein degradation step in auxin signaling. However, the exact mechanism by which SMAP1 functions in auxin signaling remains unknown. Here, we demonstrate that SMAP1 is required for normal plant growth and development and the root response to indole-3-acetic acid or methyl jasmonate in the auxin resistant1 (aar1) mutation background. Deletion analysis and green fluorescent protein/glutathione S-transferase pull-down assays showed that SMAP1 physically interacts with the CONSTITUTIVE PHOTOMORPHOGENIC9 SIGNALOSOME (CSN) via the SMAP1 F/D region. The extremely dwarf phenotype of the aar1-1 csn5a-1 double mutant confirms the functional role of SMAP1 in plant growth and development under limiting CSN functionality. Our findings suggest that SMAP1 is involved in the auxin response and possibly in other cullin-RING ubiquitin ligase-regulated signaling processes via its interaction with components associated with RELATED TO UBIQUITIN modification.

The plant hormone auxin plays an indispensable role in regulating various morphogenic processes such as root growth, shoot branching, and flower bud formation (Davies, 2004). Understanding the mode of action of auxin has been a major issue in plant physiology ever since the plant hormone concept was developed (Davies, 2004). Recent studies revealed an elegant signaling model for auxin, which is centered on SCFTIR1/AFB ubiquitin E3 ligase (Woodward and Bartel, 2005). The SCF ubiquitin E3 ligase is a multisubunit complex that regulates the ubiquitin-dependent proteolysis of many proteins. This complex consists of S-PHASE KINASE-ASSOCIATED PROTEIN1 (SKP1; ASK1 for Arabidopsis SKP1), CULLIN1 (CUL1), RING H2 finger (RBX1), and substrate-recognition F-box proteins (Woodward and Bartel, 2005). In SCFTIR1/AFB, the F-box protein TRANSPORT INHIBITOR RESISTANT1 (TIR1) or its homologs, the AUXIN-SIGNALING F-BOX (AFB) proteins, function as auxin receptors (Dharmasiri et al., 2005a, 2005b; Kepinski and Leyser, 2005). Binding of auxin to the TIR1/AFB receptors facilitates ubiquitin-mediated degradation of auxin/indole-3-acetic acid (Aux/IAA) repressors, which interact with transcriptional factors, AUXIN RESPONSE FACTORS (ARFs), resulting in changes in the patterns of downstream gene expression (Kim et al., 1997; Ulmasov et al., 1997; Tiwari et al., 2001, 2004; Dharmasiri and Estelle, 2004). Another recently discovered F-box protein, SKP2A, directly binds auxin and promotes the degradation of cell cycle transcription factors (Jurado et al., 2010).

The RELATED TO UBIQUITIN (RUB, also known as NEDD8 in mammals) protein is a small conserved protein that covalently binds to several regulatory proteins. One such regulatory protein is CUL, a scaffold protein in CULLIN-RING UBIQUITIN E3 LIGASES (CRLs), including SCF E3 ubiquitin ligase.
(Dreher and Callis, 2007). This RUB/NEDD8 modification is processed through a series of ATP-dependent steps, similar to the ubiquitin conjugation cascade (Dreher and Callis, 2007). In Arabidopsis (Arabidopsis thaliana), a heterodimer of AUXIN RESISTANT1 (AXR1) and E1 C-TERMINAL-RELATED1 (ECR1) proteins functions as a RUB/NEDD8-activating enzyme (E1) and catalyzes the ATP-dependent formation of a thioester bond between ECR1 and RUB/NEDD8 (del Pozo and Estelle, 1999). Activated RUB/NEDD8 is transferred from ECR1 to RUB-CONJUGATING ENZYME1 (RCE1), which functions as a RUB E2 enzyme and directly binds to RBX1, which is thought to function as a RUB E3 ligase in the CRL complex (Dharmasiri et al., 2003). In the case of RUB modification of CUL, an e-amino group of a Lys residue in CUL interacts with the thioester of the RUB-E2 conjugate, resulting in the formation of an isopeptide bond between RUB and CUL (Hotton and Callis, 2008). Another protein family, DEFECTIVE IN CULLIN NEDDYLATION1 (DCN1), has also been reported as a RUB E3 ligase in human and yeast (Kurz et al., 2008). Although the biochemical activity of DCN1 proteins in plants is poorly understood, the loss of a DCN1-like gene causes a 2,4-dichlorophenoxyacetic acid (2,4-D)-resistant phenotype in Arabidopsis roots (Biswas et al., 2007). A large body of experimental evidence suggests that RUB modification mechanisms are significant for plant growth and development (Dreher and Callis, 2007).

An evolutionarily conserved complex, the CONSTITUTIVE PHOTOMORPHOGENIC9 (COP9) SIGNALOSOME (CSN), is also required for the activity of CRLs (Schwechheimer et al., 2001). One of the major activities of the CSN is to deconjugate RUB/NEDD8 from RUB/NEDD8-conjugated CULs via the metalloprotease activity of its CSN5 subunit. RUB/NEDD8-associated and -dissociated CRLs are active and inactive forms, respectively (Saha and Deshaies, 2008). A recent model of the RUB/NEDD8 modification cycle suggests that both RUB/NEDD8 conjugation and deconjugation of CUL proteins are required for optimal E3 activity (Cope and Deshaies, 2003; Hotton and Callis, 2008). The stability and efficiency of CRLs depend on the RUB/NEDD8 modification status of CULs, and this status subsequently affects their E3 activity (Schwechheimer and Isono, 2010). Indeed, mutations in either component (i.e. those promoting RUB/NEDD8 conjugation or deconjugation) cause auxin-resistant phenotypes in Arabidopsis (Gray et al., 2001; Schwechheimer et al., 2001; del Pozo et al., 2002).

To better understand auxin signaling mechanisms, we used an inhibitor of auxin action, p-chlorophenoxyisobutyric acid (PCIB), to screen Arabidopsis mutants. This resulted in the isolation of several anti-auxin resistant (aar) mutants (Oono et al., 2003; Biswas et al., 2007). Some aar mutations were located in previously known auxin-related loci such as tir1 and cul1, whereas other mutations were found in unknown loci (Biswas et al., 2007). One of the mutants, aar1-1, showed 2,4-D-specific resistance without any changes in 2,4-D transport or metabolism. The causal gene of aar1 was identified by map-based cloning and designated as SMALL ACIDIC PROTEIN1 (SMAP1) because it encodes a 62-amino acid protein (6.9 kD) with a pI of 3.4. SMAP1 confers the sensitivity of Arabidopsis roots to 2,4-D and PCIB (Rahman et al., 2006). The Arabidopsis genome has another copy of the SMAP gene, SMAP2, which is expressed only in siliques and anthers and potentially mediates the root response to 2,4-D, as SMAP2 overexpression restores the sensitivity of aar1 to 2,4-D (Nakasone et al., 2009). Physiological and genetic analyses of aar1 mutants and the SMAP1 gene suggested that the SMAP1 protein acts upstream of the degradation step of Aux/IAA proteins in auxin signaling (Rahman et al., 2006). Although the SMAP protein has no known functional motifs, there is a highly conserved Phe (F)- and Asp (D)-rich 18-amino acid sequence (F/D region) in the C-terminal region. This region is found in SMAP genes from a wide variety of plants and animals, implying that the SMAP genes are evolutionarily indispensable (Rahman et al., 2006; Nakasone et al., 2009). Although SMAP1 has been shown to regulate 2,4-D sensitivity and to work upstream of the ubiquitin proteasome pathway, the functional significance of this gene in auxin signaling remains unclear. In this study, we tried to elucidate SMAP1 function by examining the genetic relationships among known auxin-related mutants and attempted to identify SMAP1-interacting proteins. Our results suggested that SMAP1 physically interacts with CSN in Arabidopsis extracts and that its function is linked to the RUB modification components AXR1 and CSN.

RESULTS

Genetic Interaction of aar1 and Auxin-Related Mutants

To investigate the relationship between aar1 and other known auxin mutants, we crossed the aar1-1 mutant with various mutants in the auxin signaling pathway: tir1-1, a mutant of an auxin receptor (Dharmasiri et al., 2005a); ecrl-1, a mutant of a subunit of the RUB-activating enzyme E1 (Woodward et al., 2007); axr1-12, a mutant of another subunit of the RUB-activating enzyme E1 (Leyser et al., 1993); and aar3-2, a mutant of a DCN1-like gene (Biswas et al., 2007). Then we attempted to establish double mutants. aar1-1 tir1-1, aar1-1 ecrl-1, and aar1-1 aar3-2 double mutants were successfully obtained. However, the axr1-12 aar1-1 mutant showed severe morphological defects; nearly one-quarter of the offspring from the AXR1/axr1-12 aar1-1 parents did not germinate or died at the early seedling stage without developing roots (Figs. 1, A–F, and 2, A and B). Genotyping with PCR markers confirmed that the rootless siblings were axr1-12 aar1-1 double mutants (data not shown). In normal embryos, the hypophysis divides asymmetrically and forms a quiescent center in the root meristem. Formation of an auxin gradient is required for this process, which can
be readily visualized by the authentic auxin reporter

**Figure 1.** Genetic interaction between **SMAP1** and auxin-related mutants. A, Comparison of the early seedling phenotype in the wild type (WT), axr1-12, aar1-1, axr1-12 aar1-1 double mutant (axr1 aar1), and the double mutant containing the genomic SMAP transgene (axr1 aar1+ gSMAP1). The seedlings were grown for 6 d in GM and photographed. Note the rootless phenotype of axr1 aar1 double mutant and complementation of the phenotype by genomic SMAP1. B and C, Normal (left) and abnormal (right) early globular (B) and torpedo (C) stage embryos in siliques of AXR1/axr1-12 aar1-1 plants. D, Visualization of the auxin response pattern in normal (left) and abnormal (right) torpedo stage embryos in siliques of AXR1/axr1-12 aar1-1 plants containing DR5rev:GFP. GFP expression was absent in the hypophysis of abnormal embryos. E, Mature seeds with normal (left) and abnormal (right) embryos obtained from AXR1/axr1-12 aar1-1 plants. F, Venation patterns in cotyledons of 5-d-old wild type, axr1-12, aar1-1, and axr1 aar1. G, Response of single and double mutants to 2,4-D in a root growth assay. Seedlings were germinated and grown on GM containing 2,4-D. Elongation of roots was measured from day 4 to 7. Data are means ± SD (n = at least 10 seedlings). * The rootless seedlings (see Fig. 2Z) were excluded. DMSO, Dimethyl sulfoxide.

the abnormal phenotype in double mutants is not allele dependent.

The **aar1-1** mutation has an approximately 44-kb deletion in chromosome 4; this deleted region contains at least 10 open reading frames, including **SMAP1** (Rahman et al., 2006). Therefore, the abnormal phenotype could be caused by the lack of other genes rather than **SMAP1**. To eliminate this possibility, we crossed *axr1-12* with two independent transgenic lines harboring a 3.7-kb *BamH*I/Sac*I* genomic DNA fragment containing the **SMAP1** open reading frame (gSMAP1). As shown in Figures 1A and 2, G to I, the seedling population **AXR1/axr1-12 aar1-1 gSMAP1** lines showed no rootless offspring, in contrast to the line harboring the control genomic fragment (X/B). In
reverse, the homozygous offspring of the AXR1/axr1-12 heterozygous parental lines harboring the SMAP1 RNA interference (RNAi) construct were rootless (Fig. 2, J–N). Taken together, these results suggest that SMAP1 is necessary for root meristem formation and normal embryo development in the axr1 background.

We successfully obtained the double mutants aar1-1 tir1-1, aar1-1 ecr1-1, and aar1-1 aar3-2. Most of these double mutants did not show any severe morphological defects compared with their parental lines (data not shown). However, approximately 10% of seedlings from the aar1-1 ecr1-1 parent showed a rootless
phenotype similar to that of the axr1 aar1 double mutants (Fig. 2Z). The root elongation assay against 2,4-D revealed that the double mutants axr1-1 tir1-1, axr1-1 ecr1-1, and aar1-1 aar3-2 are more resistant compared with their corresponding single mutants, suggesting an additive relationship between the aar1-1 mutation and the auxin-related mutations tir1-1, ecr1-1, and aar3-2 for regulating the root auxin response (Fig. 1G).

Expression of SMAP1 under the Control of the Cauliflower Mosaic Virus 35S Promoter in the aar1 Background Partially Restored the Wild-Type Phenotype

SMAP1 was fused to GFP and expressed under the control of the cauliflower mosaic virus (CaMV) 35S promoter (35S:SMAP1-GFP or 35S:GFP-SMAP1) in aar1-1 plants. The expression of fusion proteins in the transgenic lines restored wild-type sensitivity to PCIB and 2,4-D in the aar1-1 roots, suggesting that the SMAP1-GFP and GFP-SMAP1 fusion proteins are functionally active (Supplemental Fig. S1). The mature 35S:SMAP1-GFP or 35S:GFP-SMAP1 (collectively designated as 35S:SMAP1~GFP) transgenic Arabidopsis plants were morphologically indistinguishable from nontransgenic plants in either the wild-type or aar1-1 background (data not shown). When 35S:SMAP1~GFP was introduced into the aar1-12 background (35S:SMAP1-GFP/aar1-12 or 35S:GFP-SMAP1/aar1-12), morphological phenotypes of aar1-12 such as dwarfism, reduced apical dominance, protruding pistils, and low fertility (Lincoln et al., 1990) were partially alleviated (Fig. 3A). In a root growth assay, the aar1-12 mutant is strongly resistant to auxin and other plant hormones such as jasmonic acid (JA; Lincoln et al., 1990; Tiryaki

Figure 3. Phenotypic changes resulting from the introduction of 35S:SMAP1~GFP fusion genes into the aar1-12 mutant. Independent transgenic lines of 35S:SMAP1-GFP (lines 1H and 2G) and 35S:GFP-SMAP1 (line 2B) were crossed with aar1-12, and homozygous lines were established for aar1-12, transgene, and wild-type (WT) AAR1 loci. A, Mature plants (27 d old; top row), inflorescences (middle row), and flowers (bottom row) are shown. Bars = 1 cm (top and middle rows) and 0.5 cm (bottom row). B, Seeds were germinated and grown on GM containing 2,4-D at the indicated concentrations. Increase in root length was measured from day 4 to 7. Data are means ± SD (n = at least 9 seedlings). DMSO, Dimethyl sulfoxide. C, Seeds were germinated and grown on GM for 4 d, and seedlings were then transferred onto GM containing IAA and grown for a further 3 d. Root elongation after transfer was measured and plotted as a relative value compared with that on medium without chemicals. Data are means ± SD (n = at least 21 seedlings). Mean values (cm) ± SD in the absence of chemicals controlling root elongation were as follows: 1.56 ± 0.25 (wild type), 2.23 ± 0.41 (aar1-12), 1.58 ± 0.37 (35S:SMAP1-GFP/aar1-12 1H), 1.76 ± 0.24 (35S:SMAP1-GFP/aar1-12 2G), and 1.64 ± 0.47 (35S:GFP-SMAP1/aar1-12 2B). D, Seeds were germinated and grown on GM containing 10 μM methyl jasmonate (JA-Me). Root elongation was measured from day 5 to 8 and plotted as a relative value compared with that on medium without chemicals. Data are means ± SD (n = at least 5 seedlings). Mean values (cm) ± SD in the absence of chemicals controlling root elongation were as follows: 2.05 ± 0.26 (wild type), 2.63 ± 0.19 (aar1-12), and 2.19 ± 0.24 (35S:SMAP1-GFP/aar1-12 1H).
and Staswick, 2002). The introduction of 35S:SMAP1-GFP also partially restored the 2,4-D sensitivity of axr1-12 roots (Fig. 3B). Furthermore, although the aar1 mutant is reported to be a 2,4-D-specific response mutant (Rahman et al., 2006), the sensitivity of 35S:SMAP1-GFP/axr1-12 to IAA (Fig. 3C) and methyl jasmonate (Fig. 3D) was intermediate between that of the wild type and axr1-12, suggesting that in the absence of functional AXR1, SMAP1 function is required for normal IAA and JA responses.

Unexpectedly, unlike the SMAP1 genomic fragment, 35S:SMAP1-GFP fusion constructs expressed under the control of the CaMV 35S promoter did not effectively recover the rootless phenotype of the axr1 aar1 double mutant (Fig. 2, O–W). Although we established homozygous axr1-12 aar1-1 35S:SMAP1-GFP (line 1H) and axr1-12 aar1-1 35S:GFP-SMAP1 (line 2B) lines by crossing, many seeds from these lines did not germinate, and most of those that did germinate were rootless (Fig. 2, Q and S). This may be due to the low expression level of SMAP1-GFP at the early stage of embryo development, which could not complement the loss of the SMAP1 gene from hypophyseal initial cells. GFP expression was hardly detected in the globular embryos of the two 35S:SMAP1-GFP lines (Supplemental Fig. S2). The transgene of SMAP1-GFP under the control of the SMAP1 promoter (SMAP:SMAP1-GFP line 1D) that was strongly expressed (as observed by GFP fluorescence) in the lower part of the globular embryo (Supplemental Fig. S2) effectively rescued the morphological defects in the axr1-12 aar1-1 double mutants (Fig. 2X). Another independent line of SMAP:SMAP1-GFP (line 1B) that showed low-level GFP expression in the globular embryo did not recover the root development in axr1-12 aar1-1 double mutant background (Fig. 2Y; Supplemental Fig. S2). This suggests that strong SMAP1 expression in the globular stage embryo is required to establish root initial organization in the axr1-12 background.

A Conserved F/D-Rich Region Is Significant for SMAP1 Function

To investigate the functional aspects of the SMAP1 protein, we constructed several versions of SMAP1 derivatives, including those with deleted versions of the conserved F/D region, fused to GFP, under the control of the SMAP1 promoter (approximately 5 kb; Fig. 4A). These constructs were transformed into the aar1-1 mutant, and their ability to complement the aar1 mutation was investigated (Fig. 4, B–D). The GFP fluorescence of the fusion proteins was detected in both nuclei and the cytosol (Supplemental Fig. S3).

The aar1 mutant is resistant to the anti-auxin PCIB and the synthetic auxin 2,4-D and exhibits long hypocotyl when grown in the light (Rahman et al., 2006). As shown in Figure 4, B to D, independent transgenic lines harboring SMAP1-GFP in the aar1 background recovered sensitivities to PCIB (Fig. 4B) and 2,4-D (Fig. 4C) and showed similar hypocotyl length to that of

Figure 4. Construction and analyses of GFP fusion constructs with SMAP1 and SMAP1 deletion series. A, Schematic diagram of constructs. SMAP1 and SMAP1 deletion series are shown by white and green boxes, respectively. The F/D-rich region in SMAP1 is shown by blue boxes. The SMAP1 promoter is shown by black arrows. The nos terminator is not shown. Dotted lines indicate deleted regions in the constructs. SMAP1-GFP contains full-length SMAP1. D1, D2, and D3 are N-terminal deleted constructs. Numbers on white boxes indicate the positions of amino acids within the N and C termini of modified SMAP1. B and C, Transgenic aar1-1 lines, the wild type (WT), and untransformed aar1-1 were planted on GM containing 20 μM PCIB (B) or 50 nM 2,4-D (C), and root length was measured after 10 d. Data are means ± SD (n = at least 13 seedlings). D, Hypocotyl length of 7-d-old seedlings grown on GM in the light. Data are means ± SD (n = at least 15 seedlings). [See online article for color version of this figure.]
wild type (Fig. 4D), suggesting that the SMAP1-GFP fusion protein is functional. The construct SMAP1ΔF/D-GFP, which contained a deleted version of the F/D-rich region, did not complement the aar1 mutation, although the fusion protein was expressed in the transgenic lines as confirmed by GFP fluorescence. This finding implies that the F/D region is necessary for SMAP1 function. The N-terminal deletion construct D1, in which one-third of the N-terminal amino acids were deleted, was still able to complement the aar1 phenotype. However, the D2 and D3 constructs, which consisted of 26 and 18 C-terminal amino acids, respectively, did not complement the aar1 phenotype, suggesting that the F/D-rich region alone is not sufficient for SMAP1 function.

SMAP1 Interacts with the COP9 Signalosome

Using the transgenic plants described above, we next attempted to identify SMAP1-interacting proteins in Arabidopsis extracts. Total proteins were extracted from SMAP1-GFP, SMAP1ΔF/D-GFP, and GFP lines using a nondenaturing extraction buffer and subjected to pull-down assays using anti-GFP microbeads. The purified GFP- and/or SMAP1-binding proteins were separated by SDS-PAGE and subjected to silver staining and western blotting with an anti-GFP antibody (Fig. 5A). Silver staining revealed that seven distinct bands are present in the SMAP1-GFP extract but absent in the SMAP1ΔF/D-GFP extract, suggesting that the proteins present in these bands are possibly SMAP1-interacting proteins (Fig. 5A). These proteins were digested by trypsin, eluted from the gel, and analyzed by liquid chromatography (LC)-tandem mass spectrometry (MS/MS). A Mascot database search of the resulting MS/MS spectra identified the proteins as CSN1, CSN2, CSN3, CSN4, CYTOSOLIC GLYCOLALDEHYDE-3-PHOSPHATE DEHYDROGENASE2 (GAPC2), CSN5A, and CSN8 (Fig. 5A; Table I). Of all the identified proteins, six out of seven were subunits of the large CSN complex (approximately 321.3 kD) that typically consists of eight subunits (Schwechheimer et al., 2001). Taken together, these results suggest that SMAP1 physically interacts with CSN in Arabidopsis extracts and that this interaction requires the presence of the F/D region.

To further elucidate the interaction between SMAP1 and CSN, we established an in vitro binding assay with glutathione S-transferase (GST)-tagged fusion proteins. Two transgenic lines, SMAP1-GFP/aar1-1 and SMAP1ΔF/D-GFP/aar1-1, were used in pull-down assays. Pulled down proteins were separated by SDS-PAGE and detected by western blotting (WB) using anti-GFP or by silver staining. Seven protein bands (marked by red circles) that bind to SMAP1-GFP but not to SMAP1ΔF/D-GFP were eluted and identified by LC-MS/MS (Table I). B. In vitro pull-down assay using GST-tagged protein constructs. The top panel shows a schematic diagram of constructs of GST-tagged proteins; GST control, GST-tagged full-length SMAP1 (GST-SMAP1), F/D region-deleted version of SMAP1 (GST-SMAP1ΔF/D), and the F/D region only (GST-F/D).
proteins expressed in *Escherichia coli* (Fig. 5B). The GST-tagged SMAP1 protein and its derivatives (GST-SMAP1, GST-SMAP1ΔF/D, GST-F/D, and GST) extracted from *E. coli* were immobilized to glutathione columns. Total proteins extracted from the Arabidopsis *aar1-1* mutant were added to these columns. Candidate interacting proteins as well as the GST fusion proteins themselves were eluted from the columns by SDS sample buffer, separated by SDS-PAGE, and subjected to Coomassie blue staining and western-blot analysis with an anti-CSN4 antibody (Fig. 5B). The results showed that GST-SMAP1 and GST-F/D but not GST-SMAP1ΔF/D or GFP pulled down CSN4 protein from the *aar1-1* extracts, suggesting that the F/D region of SMAP1 is necessary and sufficient for the interaction between SMAP1 and CSN.

**Genetic Interaction of SMAP1 and CSN**

CSN was originally identified through the biochemical characterization of the COP9 protein complex (Wei et al., 1994; Chamovitz et al., 1996), which plays a significant role in regulating photomorphogenesis and postembryonic development in Arabidopsis (Kwok et al., 1996). Furthermore, auxin responses are partially impaired in *csn* mutants (Dohmann et al., 2008). To investigate the interaction between SMAP1 and CSN at the genetic level, we crossed the *aar1-1* mutant with the *csn5a-1* homozygous homozygous and heterozygous *csn5a-1* (*aar1-1 CSN5A/csn5a-1*). In the seedling population from the *aar1-1 CSN5A/csn5a-1* parental line, both in light- and dark-grown conditions, we observed distinct segregation of dwarf seedlings readily distinguishable from either *aar1-1* or *csn5a-1* seedlings (70 dwarf seedlings out of 363 light-grown seedlings; χ² = 6.33, P > 0.01; Fig. 6, A and B; Supplemental Fig. S4B). Genotyping analyses confirmed that these dwarf plants are the *aar1-1 csn5a-1* double mutant (data not shown). The longer hypocotyl phenotype of *aar1* was suppressed by the *csn5a* mutation. In fact, the double mutant showed an extreme short hypocotyl phenotype. Similarly, a slower root growth phenotype was found in the double mutant, although the growth of the roots of *aar1-1* and *csn5a-1* is comparable to the wild type (Fig. 6A; Supplemental Fig. S4A). In addition, the double mutant showed 2,4-D resistance for root growth like its parental lines (Fig. 6C). The *aar1-1 csn5a-1* plants at the rosette stage also showed an extremely small seedling phenotype (Fig. 6D), and most of the plants died before producing seeds (data not shown).

When the 35S:SMAP1-GFP (line 1H) gene was introduced into the *aar1-1 csn5a-1* background (35S:SMAP1-GFP/aar1-1csn5a-1), the ratio of RUB-modified CUL1 in the double mutants and transgenic lines was measured by western blotting using anti-CUL1 antibody (Fig. 7). The results from two independent experimental approaches implied that SMAP1 interacts with the RUB modification-related factors AXR1 and CSN. Thus, we next examined the RUB modification status of the CUL1 protein, which is a core subunit of SCF E3 ligase and one of the most characterized RUB-modified proteins, in the double mutants and transgenic lines by western blotting using anti-CUL1 antibody (Fig. 7). As previously published, the ratio of RUB-modified and unmodified CUL1 is lower in *axr1-12* and higher in *csn5a-1* compared with that in the wild type (Gray et al., 2001; del Pozo et al., 2002; Gusmaroli et al., 2007). No significant difference was observed in RUB modification depending on the presence or absence of the SMAP1 gene, except that a slight increase in RUB-modified CUL1 ratio was detected in the flower extracts.

### Table 1. LC-MS/MS identification of pull-down proteins using SMAP1-GFP protein

| Band No. | Protein Name       | Theoretical Mass | Mascot Score | Matched Peptides | Sequence Coverage |
|----------|--------------------|------------------|--------------|------------------|-------------------|
| 1        | CSN1 (At3g61140.1) | 50.6             | 400          | 13               | 35                |
| 2        | CSN2 (At2g26990.1) | 51.1             | 142          | 8                | 25                |
| 3        | CSN4 (At5g42970.1) | 45.0             | 864          | 25               | 65                |
| 4        | CSN3 (At5g14250.1) | 47.7             | 370          | 15               | 40                |
| 5        | CSN5A (At1g22920.1)| 39.7             | 288          | 14               | 51                |
| 6        | GAC2 (At1g13440.1) | 36.9             | 91           | 6                | 24                |
| 7        | CSN8 (At4g14110.1) | 22.5             | 111          | 3                | 18                |

*The numbers are shown in Figure 5A.*

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of transgenics harboring 35S:SMAP1−GFP in the axr1-12 background (Fig. 7B).

DISCUSSION

The functional significance of the SMAP1 protein, which confers 2,4-D sensitivity in Arabidopsis, in the auxin signaling pathway remained obscure in spite of the genetic and physiological characterization of the axr1 mutants (Rahman et al., 2006; Nakasone et al., 2009). In this study, using a combinatorial approach of genetics and biochemistry, we demonstrated that the SMAP1 protein interacts with the RUB modification components, AXR1 and CSN, and plays an important role in regulating the growth and development of Arabidopsis seedlings under limiting AXR1 or CSN.

Figure 6. Genetic interaction of CSN5A and SMAP1. We could not obtain a sufficient number of seeds of the axr1-1 csn5a-1 double mutant; therefore, dwarf axr1-1 csn5a-1 seedlings were selected from a seed population from the axr1-1 CSN5A/ csn5a-1 parental line for analysis. A and B, Images (A) and root lengths (B) of 7-d-old seedlings grown on GM without growth regulators. Bar in A = 1 cm. Values in B are means ± sd (n = at least 13 seedlings). C, Seeds were germinated and grown on GM for 4 d and then transferred onto GM containing 2,4-D at the indicated concentrations and grown for an additional 3 d. Root elongation after transfer was measured and plotted as a relative value compared with that on medium without chemicals. Values are means ± sd (n = at least 17 seedlings). Mean values (cm) ± sd in the absence of chemicals controlling root elongation were as follows: 1.56 ± 0.18 (wild type [WT]), 1.64 ± 0.22 (axr1-1), 1.71 ± 0.30 (csn5a-1), and 1.21 ± 0.22 (axr1-1 csn5a-1). D, Photographs of plants at the rosette stage. Seeds were germinated on GM and grown for 9 d, and seedlings were then transferred to soil and grown for an additional 11 d under 16-h-light/8-h-dark conditions. White arrowheads indicate axr1-1 csn5a-1 double mutants. The photograph at bottom left is an enlarged image of a axr1-1 csn5a-1 double mutant. E, Photographs of adult plants. Seeds were germinated and grown on GM for 9 d, and seedlings were then transferred to soil and further grown for 35 d under 16-h-light/8-h-dark conditions. Bar = 5 cm.
function. Both AXR1 and CSN have important roles in CRL-mediated signaling processes, including the auxin response, via RUB modification. AXR1 facilitates the RUB modification and CSN functions in deconjugating RUB from RUB-modified proteins. The functional significance of these proteins in auxin signaling has been well demonstrated (Schwechheimer et al., 2001). The genetic and biochemical evidence presented here suggest that SMAP1 associates with the RUB modification cycle, which is consistent with the previous report, where we demonstrated that SMAP1 functions upstream of Aux/IAA protein degradation in Arabidopsis (Rahman et al., 2006).

Our initial characterization of SMAP1 demonstrated that plants with diminished SMAP1 function (aar1 mutants and SMAP1 RNAi lines) show altered responses to 2,4-D but not to IAA or other major plant hormones (Rahman et al., 2006). However, the results in this work suggest that under limited functionality of the RUB modification components AXR1 and CSN, SMAP1 potentially regulates IAA and JA signaling pathways along with 2,4-D. The root phenotypic difference observed between axr1 aar1 and axr1 tir1 double mutants is striking. The axr1 aar1 double mutant showed severe morphological defects with no root meristem formation, while the tir1 aar1 double mutant showed no apparent root developmental phenotype, although they showed increased 2,4-D resistance compared with their respective parental lines. The tir1 axr1 double mutant mimics the tir1 aar1 root phenotype, showing no apparent change in root development and increased auxin resistance (Ruegger et al., 1998). The inability of axr1 aar1 to form a root meristem suggests a strong functional interaction between AXR1 and SMAP1. This idea is further substantiated by the fact that in adult plants, the overexpression of SMAP1 restored a wild-type-like phenotype in the axr1 mutant background, suggesting that SMAP1 functionally cooperates with AXR1 in regulating various biological processes, including embryogenesis, morphology of mature plants, and hormonal responses. A similar genetic feature was reported for the AXR1 homolog gene AXR1-LIKE (AXL). The single mutant of axl did not show any remarkable phenotype, while the axr1 axl double mutant exhibited a rootless phenotype (Dharmasiri et al., 2007). Ectopic expression of AXL under the control of the CaMV 35S promoter in the axr1 background complemented the axr1 phenotype (Dharmasiri et al., 2007). Since both AXR1 and AXL encode a subunit of the RUB E1 enzyme and promote RUB modification to regulate CRL activity, one possible explanation for SMAP function could be its involvement in the RUB modification process.

In vivo GFP and in vitro GST pull-down assays revealed that SMAP1 interacts with CSN. The CSN genes were originally identified as causal genes of cop, deetiolated, or fus mutants, which show defects in photomorphogenesis or embryogenesis (Chamovitz et al., 1996). CSN influences numerous plant hormone signaling pathways, including the auxin, jasmonate, and strigolactone pathways, as well as light signaling, cell cycle progression, and circadian rhythm (Somers and Fujiwara, 2009; Schwechheimer and Isono, 2010). The best-defined function of CSN is to regulate the ubiquitin/proteasome-dependent protein degradation system by removing the covalently conjugated RUB protein from CULs and by stabilizing CRL components, such as TIR1, CUL1, and ASK1 (Stuttmann et al., 2009; Schwechheimer and Isono, 2010). The fact that the functionally significant F/D region is necessary and sufficient for SMAP1 to interact with CSN implies that SMAP1 is involved in the regulation of CRL activity together with CSN.

In our pull-down analysis of SMAP1-GFP, we detected six out of eight CSN subunits but did not detect CSN6 and CSN7 (Table I; Fig. 5A). This could be due to the similar molecular masses of CSN6, CSN7, and GFP-SMAP1, which results in an inadequate...
separation of these proteins on the gel. Alternatively, SMAP1 may interact with a CSN subcomplex that lacks CSN6 and CSN7. A structural analysis in mammalian cells demonstrated that several CSN subcomplexes exist, for example, CSN1/2/3/8, CSN4/5/6/7, or smaller versions including CSN1/3/8 and CSN4/6/7 (Sharon et al., 2009). Two symmetrical modules, CSN1/2/3/8 and CSN4/5/6/7, are connected by interaction between CSN1 and CSN6 to form an eight-subunit CSN complex. In the CSN1/2/3/8 and CSN4/5/6/7 subcomplexes, the most peripheral subunits are CSN2 and CSN5, respectively. Furthermore, subcomplex CSN4/5/6/7 was found to be very stable (Sharon et al., 2009). Therefore, it is difficult to predict that CSN subcomplexes lacking CSN6 or CSN7 are present in plant extracts, although the binding patterns of CSN6 and CSN7 are unknown in plant cells. Further biochemical experiments are required to clarify how SMAP1 physically interacts with CSN. CSN in mammalian cells is involved in various biological responses, including embryonic development, cell cycle progression, T-cell development, signal transduction, oocyte maturation, autophagy, and circadian rhythm (Seeger et al., 1998; Kato and Yoneda-Kato, 2009). Thus, a future challenge is to determine whether SMAP proteins in mammalian cells also interact with mammalian CSN and have a regulatory role in CRL-mediated biological processes.

The strong morphological defects of axr1 axr1 double mutants and the increase of RUB-modified CUL1 in 35S:SMAP1-GFP/axr1-12 transgenic plants suggest that SMAP1 acts as a positive regulator for RUB modification. Because CSN activates the dissociation of RUB from RUB-modified CULs, one possible explanation for SMAP1 function is that SMAP1 inhibits RUB dissociation activity of CSN by its binding to CSN. Given the marked morphological defects in axr1 and csn mutants, a small change of RUB modification status could be enough to cause the observed morphological changes in the presence or absence of SMAP1. However, we cannot eliminate the possibility that an increase of RUB-modified CUL1 in flower extracts of 35S:SMAP1-GFP/axr1-12 transgenic plants might be a secondary effect resulting from the morphological changes. Even if this is the case, the morphological defects of the double mutants and the physical interaction between SMAP1 and CSN imply that SMAP1 interacts with the RUB cycle-related regulation of CRL activity. In a recent model in fission yeast, CSN prevents the autocatalytic ubiquitin-mediated degradation of certain F-box proteins that are assembled in the CRL, resulting in maintenance of the CRL complex in the absence of the substrates (Schmidt et al., 2009). The stabilization of CRL components by CSN in Arabidopsis extracts has also been reported (Stuttmann et al., 2009). Thus, it may be possible that SMAP1 affects the stability of CRL components and CRL assembly by binding with CSN. The construction of more sensitive biochemical experimental systems for monitoring RUB modification status, CRL activity, and CRL stability in plants and in vitro would clarify how SMAP1 contributes to the RUB cycle-related regulation of CRL activity.

In conclusion, two independent experimental approaches indicate that the conserved SMAP1 protein interacts with RUB modification-related components. Although the precise mode of action of SMAP1 is still unclear, further biochemical research on the functions and the relationship of SMAP1 with AXR1 and CSN might reveal its biological role in the RUB cycle as well as in the CRL-regulated ubiquitin-proteasome system in plant cells.

MATERIALS AND METHODS

Plant Materials

The Arabidopsis (Arabidopsis italiana) lines axr1-1 and axr1-2 (Rahman et al., 2006), axr3-2 (Biswas et al., 2007), axr1-3 and axr1-12 (Leyser et al., 1993), tir1-1 (Ruegger et al., 1998), ccr1-1 (Woodward et al., 2007), csn5a-1 (Gusmaroli et al., 2007), and DR5rev:GFP (Friml et al., 2003) were described previously. The SMAP1 RNAi lines (520i), the transgenic lines transformed with the 3.7-kb SMAP1 genomic fragment (B/S; described as gSMAP1 in this report) in the axr1-1 background, and its control line (X/B) were described by Rahman et al. (2006). 35S:SMAP1-GFP/axr1-1, 35S:GFP/CSN1-1, 35S:CSN1-1, and 35S:CSN1-1/GFP/axr1-1, and their derivative lines were generated in this work as described below. All transgenic and mutant lines were derived from the Columbia accession of Arabidopsis except for axr1-2, which is in the Wassilewskijia background. The AART locus was genotyped by PCR using primers T6G15 9538F and T6G15 53634R to detect the DNA deletion in the axr1-1 genome, and its control line (X/B) were described by Rahman et al. (2006). 35S:SMAP1-GFP/axr1-1, 35S:GFP/CSN1-1, and 35S:CSN1-1/GFP/axr1-1, and their derivative lines were generated in this work as described below. All transgenic and mutant lines were derived from the Columbia accession of Arabidopsis except for axr1-2, which is in the Wassilewskijia background. The AART locus was genotyped by PCR using primers T6G15 93890 and T6G15 53634R to detect the DNA deletion in the axr1-1 genome and primers T6G15 24488F and AT4G1 3490F2 to detect the presence of the corresponding wild-type DNA. For axr1-2, the mutant locus was tracked by the kansamycin-resistant phenotype resulting from the NPTII gene in the transgene construct (Rahman et al., 2006). The axr1-2 mutation was genotyped using the primers KSK13 F38890 and KSK13 R39225 to detect the wild-type gene and primers KSK13 F38890 and LB1 to detect the T-DNA. The tir1-1 mutation was detected by PCR amplification using the primers TIR1gF547 and TIR1gR913, and the amplified product was digested with DprII, which cuts mutant but not wild-type products. The axr1-12 mutation was detected by PCR with the primers AXR1-12F and AXR1-12R, and the amplified product was digested with Drai, which cuts mutant but not wild-type products. For csn5a-1, the primers CSN5A1F and CSN5A1R2 were used to detect the wild-type locus and the primers CSN5A1F and CSN5A1R2 were used to detect the wild-type locus and the primers CSN5A1F and LB1 were used to detect the mutation. The primer sequences are listed in Supplemental Table S1. The ccr1-1 and axr1-3 mutations were detected by derived cleaved-amplified polymorphic sequence analysis as described by Woodward et al. (2007).

Growth Analyses

Seed sterilization and plant growth conditions were as described by Biswas et al. (2007). Briefly, seeds were plated on germination medium (GM; one-half-strength Murashige and Skoog salts, 1% [w/v] Suc, and 0.5 g L\(^{-1}\) MES, pH 5.8, containing 1× B5 vitamins and 0.8% [w/v] Bacto agar) in rectangular plates, unless otherwise mentioned. For synchronous germination, the seeds were kept in the dark for 2 d at 4°C, and then the plates were transferred to a growth chamber at 23°C. For the hypocotyl and root growth assays, the plants were grown vertically on square culture plates under continuous white light (20–30 µmol m\(^{-2}\) s\(^{-1}\)). For transfer assays, seedlings were first grown on GM without plant growth regulators and then transferred after 4 to 5 d to fresh GM containing plant growth regulators. For the IAA root inhibition assay, seedlings were transferred onto GM 1.2 (the same as GM except that the agar concentration is 1.2% to prevent roots from penetrating into the agar). The hypocotyl and root lengths were analyzed using the NIH ImageJ software package (National Institutes of Health) after the plants were photographed with a DP00 digital camera (Olympus). The percentage of root growth inhibition was calculated relative to root growth on medium without growth regulators. To observe the morphology of plants grown on soil, the seedlings were grown in a growth chamber (Biotron; LH-200-RDS, NK system) under a 16-h-light (100–130 µE m\(^{-2}\) s\(^{-1}\))/8-h-dark photoperiod at 23°C.
Plasmid Construction and Plant Transformation

All plasmids were constructed using standard recombinant DNA tech-
niques, and their authenticity was confirmed by DNA sequencing. The SMAP1
DNA fragment was amplified from Arabidopsis bacterial artificial chromo-
some clone TgG15. The SMAP1-GFP fusion construct and its deleted
versions were made by inserting appropriate DNA fragments into the binary
vector pEGAD containing GFP and the nos terminator (Culler et al., 2000). The
35S promoter was removed from pEGAD by digestion with ScaI and AgeI,
then an approximately 5-kb region of the SMAP1 promoter and the full or
deleted version of SMAP1 coding sequences were inserted into the restriction
enzyme sites. To generate 35S-SMAP1-GFP and 35S-GFP-SMAP1, SMAP1
coding sequences without or with the stop codon were amplified with the primer
sets 13520ATG-togo/13520-TAA and 13520ATG-togo/13520R-TAA
(Supplemental Table S1), respectively, and cloned into the pENTR/D-TOPO
vector (Invitrogen). The resulting plasmids, 13520-TAA ENTR or 13520+TAA
ENTR, were assembled into the Gateway binary vector pK7FW2 to generate
35S-SMAP1-GFP or into pK7FW2 to generate 35S-GFP-SMAP1, respectively,
by a site-specific recombination reaction between attL and attR sites (Karimi
et al., 2005). For the in vitro pull-down assay, GST-SMAP1 and its derivatives
were constructed with the pCEx4T-2 expression vector (GE Healthcare).
Full details of plasmid construction are available on request.

Observations of Embryo Morphology and GFP Fluorescence

For observations of embryo morphology, silique stigmas at different develop-
mental stages were harvested, dissected using a stereoscope, and cleared in a
7% Glc, mounted whole, and then observed with an Olympus BX60 microscope
for fluorescent signals in roots and

Pull-Down Assay

To detect SMAP1-GFP-binding proteins, we used a µMAGS GFP isolation
kit (Milleniyti Biots). Seven-day-old plants grown under continuous light were
grown in liquid nitrogen and then ground in ice-cold elution buffer (50 µs
Tris-HCl [pH 8.0], 50 mM NaCl, 10% glycerol, 1% Triton X-100, and Protease
Inhibitor Complete Mini [Roche Diagnostics]). The extract was centrifuged for
20 min at 13,000g, and the supernatant was filtered through a syringe-driven
filter unit with 0.45-µm pore size (Millipore). The solution containing 1 mg of
total protein was mixed with magnetic anti-GFP microbeads and then incu-
bated overnight to allow GFP to bind to anti-GFP microbeads. The mixture
was transferred to a prewashed column containing magnetic beads, and then
the column was washed four times with wash buffer 1 (50 µs Tris-HCl, pH
8.0, 150 mM NaCl, and 1% Nonidet P-40) and once with wash buffer 2 (20 µs
Tris-HCl, pH 8.0) to flush unbound proteins and excess salt. GFP and proteins
bound to it were eluted with 50 µL of SDS sample buffer. Eluted proteins were
heat denatured and separated by SDS-PAGE on a 12.5% gel. The protein spots
on the silver-stained gel were cut out and protein (380 amino acid residues of the N terminus
of CUL1) was inserted into the pET16b vector (Novagen) and expressed in the
E. coli strain BL21(DES) (Novagen). The insoluble protein fraction was solu-
bilized in His-binding buffer (20 mM Tris-HCl, pH 7.9, 0.5 M NaCl, 5 mM im-
idazole, and 6 M guanidine-HCl), and eluted with elution buffer (20 mM
Tris-HCl, pH 7.9, 0.5 M NaCl, 250 mM imidazole, and 6 M guanidine-HCl).
The eluted protein was precipitated by the addition of 80% cold acetone, and then
proteins were separated by SDS-PAGE and purified by electroelution from the
gel. The anti-CUL1 antibody was prepared from a rabbit injected with the
purified CUL1 protein. For RUB modification analysis of CUL1, Arabidopsis
seedlings or flowers were grown in lysis buffer (125 mM Tris-HCl, pH 8.5, 50
mM NaCl, 1% SDS, and 1% glycerol). The extracts were centrifuged at
13,000g for 10 min, and the protein concentration in the supernatant diluted 1/9
with water was determined by the Bradford assay (Bio-Rad Laboratories).
Protein samples were boiled in SDS-PAGE sample buffer, run on SDS-PAGE
gels, and blotted onto polyvinylidene difluoride membranes. For detection,
alkaline phosphatase-conjugated anti-rabbit IgG (A3687; Sigma-Aldrich) as
secondary antibody and nitroblue tetrazolium/3-bromo-4-chloro-3-indolyl
phosphate solution (Roche Diagnostics) were used. For densitometry analyses,
the digital images of the blots were acquired by scanning the blots, and the
peak intensity of the bands was determined by Quantity One software (Bio-
Rad Laboratories). The values of the ratio of RUB-modified CUL1 to total
CUL1 (RUB-modified plus unmodified CUL1) were analyzed by Fisher’s LSD
test in KaleidaGraph software (Synergy Software).

Supplemental Data

The following materials are available in the online version of this article.

Supplemental Figure S1. Complementation of the aart-1 phenotype with
SMAP1 and GFP fusion protein derived by 35S promoter.

Supplemental Figure S2. Expression of SMAP1-GFP in early embryonic
stage.

Supplemental Figure S3. Cellular localization of GFP fluorescence in root
tip epidermal cells of 7- to 8-old plants.

Supplemental Figure S4. Additional data for genetic interaction of the
CSN5A and SMAP1.

Supplemental Table S1. List of primers used in this study for genotyping of
mutant and transgenic lines.

For the GST pull-down assay, Escherichia coli lines harboring the constructs
were grown to an optical density at 595 nm of 0.5 to 0.6, and then the ex-
pression of fusion proteins was induced by isopropyl β-D-thiogalactopy-
oroside at a final concentration of 0.1 mM for 2 h. Harvest of fusion proteins and
their immobilization to the glutathione column were performed using a
GST-Pin Trap Purification Module kit (GE Healthcare) according to the
manufacturer’s protocol. The columns containing immobilized GST-SMAP1
fusion protein were washed with phosphate-buffered saline and then flushed
with ice-cold plant GST extraction buffer (50 mM Tris-HCl, pH 7.0, 10 mM
MgCl2, 150 mM NaCl, 10% glycerol, 0.01% Nonidet P-40, 0.5 mM phenyl-
methylsulfonyl fluoride, and Protease Inhibitor Complete Mini [Roche Diag-
nostics]). Total protein was prepared from 7-d-old aart-1 plants as described
above for the GFP pull-down assay, except that plant GST extraction buffer
was used, and applied to the columns. The columns were incubated at room
temperature for 10 min and washed five times with 600 µL of plant GST ex-
traction buffer. The GST and proteins bound to it were eluted with GST elu-
tion buffer (50 mM Tris-HCl, pH 7.0, and 10 mM glutathione), and 10 µg of
the eluted protein was separated by SDS-PAGE.

Antibodies and Western-Blot Analyses

We used antibodies against GFP (1181446001; Roche Diagnostics) and CSN4
(PW8360; Enzo Life Sciences). For the Arabidopsis CUL1 antibody, the N-
terminal region of CUL1 protein (380 amino acid residues of the N terminus
of CUL1) was inserted into the pET16b vector (Novagen) and expressed in the
E. coli strain BL21(DES) (Novagen). The insoluble protein fraction was solu-
bilized in His-binding buffer (20 mM Tris-HCl, pH 7.9, 0.5 M NaCl, 5 mM im-
idazole, and 6 M guanidine-HCl), and eluted with elution buffer (20 mM
Tris-HCl, pH 7.9, 0.5 M NaCl, 250 mM imidazole, and 6 M guanidine-HCl).
The eluted protein was precipitated by the addition of 80% cold acetone, and then
proteins were separated by SDS-PAGE and purified by electroelution from the
gel. The anti-CUL1 antibody was prepared from a rabbit injected with the
purified CUL1 protein. For RUB modification analysis of CUL1, Arabidopsis
seedlings or flowers were grown in lysis buffer (125 mM Tris-HCl, pH 8.5, 50
mM NaCl, 1% SDS, and 1% glycerol). The extracts were centrifuged at
13,000g for 10 min, and the protein concentration in the supernatant diluted 1/9
with water was determined by the Bradford assay (Bio-Rad Laboratories).
Protein samples were boiled in SDS-PAGE sample buffer, run on SDS-PAGE
gels, and blotted onto polyvinylidene difluoride membranes. For detection,
alcaline phosphatase-conjugated anti-rabbit IgG (A3687; Sigma-Aldrich) as
secondary antibody and nitroblue tetrazolium/3-bromo-4-chloro-3-indolyl
phosphate solution (Roche Diagnostics) were used. For densitometry analyses,
the digital images of the blots were acquired by scanning the blots, and the
peak intensity of the bands was determined by Quantity One software (Bio-
Rad Laboratories). The values of the ratio of RUB-modified CUL1 to total
CUL1 (RUB-modified plus unmodified CUL1) were analyzed by Fisher’s LSD
test in KaleidaGraph software (Synergy Software).
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