D14-SCF[D3]-dependent degradation of D53 regulates strigolactone signaling

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Abstract

Strigolactones (SLs) are a new class of carotenoid-derived phytohormones essential for developmental processes shaping plant architecture and interactions with parasitic weeds and symbiotic arbuscular mycorrhizal fungi. Despite the rapid progress in elucidating the SL biosynthetic pathway, the perception and signaling mechanisms of SL remain poorly understood. Here we show that DWARF53 (D53) acts as a repressor of SL signaling and SLs induce its degradation. We found that the rice *d53* mutant, which produces an exaggerated number of tillers compared to wild type plants, is caused by a gain-of-function mutation and is insensitive to exogenous SL treatment. The *D53* gene product shares predicted features with the class I Clp ATPase proteins and can form a complex with the α/β hydrolase protein DWARF14 (D14) and the F-box protein DWARF3 (D3), two previously identified signaling components potentially responsible for SL perception. We demonstrate that, in a D14- and D3-dependent manner, SLs induce D53 degradation by the proteasome and abrogate its activity in promoting axillary bud outgrowth. Our combined genetic and biochemical data reveal that D53 acts as a repressor of the SL signaling pathway, whose hormone-induced degradation represents a key molecular link between SL perception and responses.

Shoot branching (tillering in crops) is a major determinant of plant architecture and crop yield, which is under integrated control of hormonal, developmental and environmental factors[1–3]. Although the existence of a root-derived transmissible shoot-repressing signal was proposed more than 70 years ago[4], the identity of this signal(s) has remained elusive. Recent studies with branching mutants in several plants species have demonstrated that strigolactones (SLs), a specific group of terpenoid lactones, are the long-sought branching repressing hormone, whose function is highly conserved in both monocots and eudicots[5,6]. In addition to repressing shoot branching, SLs also play a role in regulating root growth, leaf senescence and flower development[7]. SLs also act as exogenous signals to promote the symbioses between land plants and arbuscular mycorrhizal fungi[8] and stimulate the germination of the parasitic weeds *Striga* and *Orobanche*, which are serious agricultural pests in many parts of the world[9].

Previous studies have shown that *DWARF3 (D3)*, *D10*, *D14*, *D17(HTD1)*, and *D27* in rice[10–14], *MORE AXILLARY GROWTH 1* (*MAX1*), *MAX2*, *MAX3* and *MAX4* in *Arabidopsis*[15–17], *RAMOSUS 1* (*RMS1*), *RMS4* and *RMS5* in pea[18], *DECREASED APICAL DOMINANCE 1* (*DAD1*), *DAD2* and *DAD3* in petunia[7,19] are involved in either the biosynthesis or signaling of SLs. Among these genes, *MAX3/RMS5/D17/DAD3*, *MAX4/RMS1/D10/DAD1*, *MAX1* in *Arabidopsis* and *D27* in rice encode the carotenoid cleavage dioxygenase (CCD) 7, CCD8, CYP711A1 (a cytochrome P450) and a novel β-carotene isomerase, respectively, and they are involved in the sequential cleavage of β-carotene and synthesis of SLs[20]. In contrast, *MAX2/RMS4/D3* and *D14/DAD2*, which encode an F-box protein and a protein of the α/β-hydrolase superfamily, respectively, likely play a role in SL signaling[12,19,21]. The structural similarity between MAX2/RMS4/D3 proteins with the auxin receptor TRANSPORT INHIBITOR RESPONSE 1 (TIR1)[22,23] and jasmonate receptor CORONATINE INSENSITIVE 1 (COI1)[24], and D14/DAD2 proteins with the gibberellin receptor GIBBERELLIN INSENSITIVE DWARF 1 (GID1)[25] has sparked the speculation that both MAX2/RMS4/D3 and D14/DAD2 could be candidates for the SL receptors[26] and that binding and hydrolysis of SLs by D14/DAD2 (refs 19, 27, 28) might be required for

triggering proteasome-mediated degradation of an unknown repressor by the SCF[MAX2] complex[29,30]. However, the identity of such a repressor and its regulatory mechanisms in SL signaling have remained unknown.

In this study, we identified a gain-of-function rice mutant *d53*, which displays a SL insensitive and increased tillering phenotype. Map-based cloning revealed that *D53* encodes a protein sharing predicted features with the class I Clp ATPase proteins and that it can form a complex with the α/β hydrolase protein D14 and the F-box protein D3. We show that SLs induce D53 degradation by the proteasome-ubiquitin pathway in a D14- and D3-dependent manner. Our studies establish D53 as a repressor of the SL signaling pathway, whose hormone-induced degradation is essential for SL signaling.

d53 is a rice SL insensitive mutant

Previous studies have identified several rice mutants defective in SL biosynthesis or signaling[10–14]. Because of their highly branched and dwarf phenotype, these mutants were termed "*d* mutants", such as *d3*, *d10*, *d14* (also known as *d88* or *htd2*), *d17* (*htd1*), and *d27*. The rice *d53* mutant[31] also displayed reduced height and increased tillering, as well as thinner stem and shorter crown root, compared to the wild-type strain (Fig. 1a, b and Extended Data Fig. 1a, b). Kinetic analysis showed that at the heading stage, the total tiller number of *d53* was about three times that of the wild type, resulting from an increase in both higher-order and high nodes tillers (Fig. 1c and Extended Data Fig. 1c, d). Histological analysis revealed that the sizes of vascular bundles and parenchyma cells in internodes were largely comparable between *d53* and wild-type plants, implying that the shortening and thinning of *d53* stem were mainly caused by a reduction in cell number (Extended Data Fig. 1e–h). The phenotypes of F_1 heterozygous plants were intermediate between the homozygous parental plants (Extended Data Fig. 2a–g). Genetic analyses of an F_2 population derived from a cross of *d53* and the wild-type parent (Norin 8) showed that the normal, intermediate and dwarf plants segregated as 1:2:1 (33:58:28, $\chi^2=0.09$, $P>0.05$), indicating that the *d53* mutation behaved in a semi-dominant manner (Extended Data Fig. 2h).

The phenotypic similarity between *d53* and the previously reported rice *d* mutants prompted us to examine whether *d53* is defective in SL-mediated inhibition of axillary bud outgrowth. Quantitative real-time RT-PCR analysis showed that expression of *D10 (CCD8)*, was similarly up-regulated in *d53* (Fig. 1d) as in other *d* mutants due to feedback regulation in the SL pathway[11]. In addition, expression of an inhibitor of axillary bud outgrowth, *FINE CULM1 (FC1)*[32], which is orthologous to the maize *TEOSINTE BRANCHED1 (TB1)*[33] and the Arabidopsis *BRANCHED1 (BRC1)*[34] was also similarly down-regulated in the *d53*, *d14* and *d27* mutants (Fig. 1d), suggesting that *D53* is likely involved in SL biosynthesis or signaling. Moreover, exogenous application of a SL analogue, GR24, effectively inhibited the outgrowth axillary buds of *d27*, but not *d14* or *d53* (Fig 1e and Extended Data Fig. 3a–c). Further, measurement of SLs produced in the root exudates showed that *d53* accumulated markedly higher levels of *2′-epi*-5-deoxystrigol, a native SL of rice, than the wild-type cultivar Norin 8 (Fig. 1f). These results indicate that *d53* is a SL-insensitive mutant.

D53 acts as a repressor of SL signaling

D53 was previously mapped to the terminal region of the short arm of rice chromosome 11 (ref. 35). To decipher the molecular defect in *d53*, we isolated *D53* by a map-based cloning approach. Using an F_2 population of ~ 12,000 plants generated from the cross between *Ketan Nangka* and the mutant, we further delimited the *D53* locus to a 34-kb DNA region on the BAC clone OSJNBa0032J07, which contains three putative genes (Fig. 2a). Sequence analysis revealed a single-nucleotide substitution and 15 nucleotides deletion in the third exon of LOC_Os11g01330 in *d53*, which resulted in an amino acid substitution (R812T) and deletion of five amino acids ($G_{813}KTGI_{817}$) (Fig. 2b). To verify that this mutation caused the tillering dwarf phenotype, we generated transgenic plants expressing the wild type or mutant *D53* gene under the control of rice *Actin1* promoter, in a wild-type background. Strikingly, all transgenic plants expressing the mutant *d53* gene showed a more exaggerated tillering phenotype than those expressing the wild-type *D53* gene. The severity of tillering phenotype in these transgenic plants was correlated with the expression level of the transgene. Notably, overexpression of the wild-type *D53* gene also caused a moderate increase in tillering, compared to the vector control plants (Fig. 2c–e). These observations suggested that the D53 protein acts as a repressor in the SL-mediated branching-inhibition pathway and that the dominant tillering phenotype of the *d53* mutant was most likely caused by a gain-of-function mutation in *d53*. To further confirm this, we generated *D53* knockdown transgenic plants using a RNA interference (RNAi) approach. As expected, reducing *D53* expression in *d53* background drastically reduced the tiller number (Extended Data Fig. 3d, e). Taken together, these data support the proposition that *d53* mutation enhanced D53 activity in repressing SL signaling.

D53 is predicted to encode a protein of 1131 amino acids. A BLAST search identified a closely related homolog of D53 (designated D53-like, LOC_Os12g01360) with 96.6% amino acid sequence identity in the rice genome. In addition, D53 homologs were found in other monocots and dicots, but not in lower plants, animals or microbes, indicating that D53-like proteins are specific in higher plants (Extended Data Fig. 4). Sequence analysis by the HHpred structure prediction server revealed that D53 shares a similar secondary structure composition, despite low primary sequence homology, to proteins of the class I Clp ATPases family, which are characterized by an N-terminal domain, a D1 ATPase domain, an M domain, and a D2 ATPase domain[36]. Notably, the D2 domain of D53 contains a highly conserved linear sequence, FDLNL, which closely matches the ETHYLENE RESPONSE FACTOR-associated amphiphilic repression (EAR) motif (Extended Data Fig. 5), which is known to interact with the TOPLESS family of proteins and involved in transcriptional repression[37].

Real-time PCR analysis revealed that *D53* was widely expressed in the examined rice tissues (Fig. 2f). *D53* promoter driven GUS (β-glucuronidase) reporter gene *(pD53::GUS)* assay showed that GUS staining was observed in vasculature in roots, shoots, leaves, leaf sheaths, nodes, internodes and young panicles, preferentially in the parenchyma cells surrounding the xylem (Extended Data Fig. 6a–h). Moreover, *D53* expression was up-regulated by GR24 treatment in wild-type plants, but down-regulated in six *d* mutants, suggesting that expression of *D53* is regulated by SLs signaling (Fig. 2g, h). The D53-GFP (green

fluorescent protein) fusion protein is exclusively localized to the nucleus in rice protoplasts and the *pActin::D53-GFP* transgenic root cells (Extended Data Fig. 6i–p).

SLs promote D53-D14 and D14-D3 interaction

Previous studies have identified the F-box protein D3 and the α/β hydrolase D14 as two key components of SL signaling in rice[10,12], of which D14 and its orthologues in Arabidopsis (AtD14) and Petunia (DAD2) have been proposed to directly participate in SL perception[19,27,28]. Yeast two-hybrid assay (Y2H) showed that both D53 and d53 could physically interact with D14 in the presence of GR24 (Fig. 3a). Domain deletion analysis indicated that the D1 domain of D53 was essential for the GR24-dependent D53-D14 interaction. Interestingly, its binding activity was inhibited by the M and D2 domains, although their negative effect can be overcome by the N domain (Extended Data Fig. 7). We verified the D53-D14 interaction in *N. benthamiana* leaf cell nucleus both in the presence or absence of exogenously applied GR24 using a bimolecular fluorescence complementation (BiFC) assay (Fig. 3b). The observed interaction between D53 and D14 in the absence of exogenously applied GR24 might be due to the effect of endogenous SLs present in the tobacco leaf cells. Consistent with the previously reported GR24-depedent interaction between DAD2 and PhMAX2A (an ortholog of D3 in Petunia) in yeast[19], our *in vitro* pull-down assay also revealed a direct physical interaction between D14 and D3 in a GR24-dependent manner (Fig. 3c). Furthermore, using recombinant GST-D3-OSK1 fusion protein as the bait, our *in vitro* pull-down assay showed that D14 could be more efficiently co-immunoprecipitated (co-IP) from *d3* plant extracts in the presence of exogenously applied GR24 (Fig. 3d). Together, these results suggest that SLs may act to promote complex formation among D14, D3 and D53, linking D53 to the hormone-perception components of the SL signaling pathway.

GR24 promotes D53 protein degradation

To investigate how SL regulates D53, we performed a set of additional experiments. Both western blot analysis and fluorescence microscopy examination showed that GR24 treatment induced rapid degradation of the D53 protein in wild-type cells, but not in *d3* and *d14* mutant cells (Fig. 4a–c). We further showed that D53 was degraded by the proteasome, as a proteasome inhibitor, MG132, but not other protease inhibitors, effectively blocked GR24-induced D53-GFP degradation (Fig. 4c, d and Extended Data Fig. 8a, b). Notably, unlike the wild-type D53-GFP fusion protein, the mutant d53-GFP fusion protein appeared to be stable in the presence of GR24 (Fig. 4c and Extended Data Fig. 8b). Interestingly, we noted that D53-GFP and D53-LUC were still degraded in the *d53* mutant cells, but not in *d3* or *d14* mutant cells (Fig. 4d and Extended Data Fig. 8c), indicating that the D53 degradation pathway was still operational in the *d53* mutant. Together, these results suggest that SL triggers proteasome-mediated degradation of D53 in a D14- and D3-dependent manner. Importantly, the insensitivity of d53 protein to SL-triggered turnover is consistent with the observed dominant gain-of-function mutant phenotype of *d53*.

To provide genetic support for the functional relationship between *D53*, *D3* and *D14*, we generated *d3 d53* and *d14 d53* double mutants. The *d3* mutant had more tillers and it was

shorter than the *d14* and *d53* single mutants (Extended Data Fig. 9a). The *d14 d53* double mutants exhibited a dwarf tillering phenotype resembling the *d14* and *d53* parental plants, whereas the *d3 d53* double mutant exhibited a dwarf tillering phenotype resembling *d3* (Fig. 4e, f and Extended Data Fig. 9a). The lack of obvious additive effects among these mutants suggests that *D3*, *D14* and *D53* act in the same signaling pathway. To further test their epistasis relationship, we knocked down *D53* gene expression in the *d3* and *d14* backgrounds. As shown in Fig. 4g and Extended Data Fig. 9b–d, the mutant phenotype of *d3* and *d14* was restored to nearly wild type levels, demonstrating that *D53* acts downstream of *D3* and *D14*, and that accumulation of D53 protein is responsible for blocking SL signaling and conferring the dwarf tillering phenotype in these mutants.

DISCUSSION

It has been speculated that perception of SLs triggers the degradation of putative repressors by the SCF[MAX2] ubiquitin ligase complex to suppress shoot branching[21,29,30]. In this study, we established that D53 acts as a repressor of SL signaling in rice. Consistent with the previous observation of GR24-dependent interaction between DAD2 and PhMAX2 (ref. 19), we found that GR24 also promotes the interaction between D14 with D53 and D3 (Fig. 3). Further, we showed that D53 is targeted for degradation by the proteasome in a D14- and D3-dependent manner (Fig. 4a–d and Extended Data Fig. 8). Together, these data collectively support the notion that SL perception by D14 acts to promote ubiquitination of D53 by the D14-SCF[D3] ubiquitin ligase, and subsequent degradation of D53 by the proteasome, leading to the propagation of SL signal and downstream physiological responses (Fig. 5). Our findings revealed a remarkable similarity between the hormonal perception and signaling mechanism of SL and several other classes of plant hormones, including auxin, jasmonate and gibberellin[25,38–40].

Interestingly, a recent study reported that a *D53* homologue in Arabidopsis, *SMAX1*, acts downstream of *MAX2* (orthologue of rice *D3*) in repressing the seed germination and seedling photomorphogenesis phenotypes of *max2*, but not the lateral root formation, axillary shoot growth, or senescence phenotypes of *max2* (ref. 41). Further, as observed for *D53*, three closest homologs of *D53* in Arabidopsis (designated *SMXL6*, *SMXL7* and *SMXL8*) were also induced by GR24 treatment[41], suggesting that *D53* and its homologues play a broad role in regulating different developmental processes and that the D3/D53 functional module is conserved between monocots and dicots. Consistent with this notion, the SL-analogous compounds karrikins also employ a MAX2 and KAI2 (D14-like)-dependent pathway to regulate seed germination and seedling growth[29,42]. The identification and characterization of *D53* in SL signaling now set the stage for further dissection of the mechanisms by which SLs regulate plant form and its complex interactions with parasitic weeds and symbiotic arbusular mycorrhizal fungi[3].

METHODS

Plant materials and growth conditions

The wild-type rice (*Oryza sativa* L. subspecies Japonica) varieties used in this study were Norin 8 (for *d53*, *d14* and *d27*), Akumuro (for *d3*), Shiokari (for *d10* and *d17*) and Kitaake

(for transformation analyses). The *d53*-carrying line, KL908 was initially identified from a mutant library of Norin 8 mutagenized by ^{60}Co-γ irradiation[31]. The *d27* and *htd2/d14* mutants were described previously[14,43]. The *d3* mutant used in this work carries a premature stop codon (unpublished data). The *d14 d53* and *d3 d53* double mutants were generated by crossing their respective parental lines and identified by genotyping from their respective F_2 or F_3 populations.

Rice plants were cultivated in the experimental field at the Nanjing Agriculture University in Nanjing in the natural growing seasons. For real-time RT-PCR, GR24 treatment, strigolactone analyses and transient expression assay, the seedlings of wild-type and mutants were grown in climate chambers (HP1500GS, Ruihua) at 70% humidity, under long-day conditions with daily cycles of 14 h of light at 30 °C and 10 h of dark at 25 °C. Light was provided by fluorescent white-light tubes (400–700 nm, 250 μmol $m^{-2}s^{-1}$).

Map-based cloning of *D53*

To map the *D53* locus, the *d53* mutant was crossed with a polymorphic *javanica* variety, *Ketan Nangka*. Rough mapping was performed with simple sequence repeat (SSR) markers W1 (5′-GGATGATGAGATCCTAATGTAGAA-3′ and 5′-CATCCTTGGAAAATAGTGGG-3′) and W2 (5′-GGCTTCATCTTTGGCGACC-3′ and 5′-CCGGATTCACGAGATAAACTCTC-3′) using 242 normal plants obtained in the F_2 population. Using 2,893 normal plants, *D53* was finally mapped to a region between the SSR marker Z7 (5′-CAGAGACCAAGCAACAGAG-3′ and 5′-GGAGACGTGTGAGCTACAAC-3′) and a derived cleaved amplified polymorphic sequence (dCAPS) marker Z3 (5′-GTTACCGCTCCTCCTCAGC-3′ and 5′-AGTAAAAATGTGGAGGGGCA-3′).

Vector construction and plant transformation

To generate the *pActin::D53-GFP* and *pActin::d53-GFP* construct, full-length cDNAs of *D53* and *d53* were amplified, and the PCR products were then cloned into the binary vector AHLG[44] using the In-Fusion™ Advantage PCR Cloning Kit (Cat: PT4065, Clontech) and sequenced (Biomed). The *pActin::GFP*, *pActin::D53-GFP* and *pActin::d53-GFP* constructs were introduced into the wild-type variety Kitaake. The *pActin::D53-GFP* transgene was introduced into *d3* and *d14* backgrounds by genetic crosses.

To knock down the *D53* gene, the *D53*-RNA interference (*D53*-RNAi) vector was constructed using a 475 bp cDNA fragment. The PCR fragment was cloned into the p*EASY*-Blunt cloning vector (TransGen), and the gene-specific fragment was prepared by double digestion with *Kpn*I-*Sac*I and *Pst*I-*BamH*I, respectively, and then the PCR products were ligated to both sides of the *FAD2* intron in the binary vector pCUbi1390- FAD2 in opposite orientation[45] and sequenced. The resultant construct *D53*-RNAi and the control vector pCUbi1390- FAD2 were introduced into *d53*, *d3* and *d14* mutants, respectively.

To analysis the expression pattern of *D53* gene, a 2.7-kb promoter fragment was cloned into the binary vector pCAMBIA-1381Z and sequenced to create the *pD53::GUS* reporter gene construct, which was then introduced into wild-type rice cultivar Norin 8. All transgenic rice

plants were generated using *Agrobacterium*-mediated transformation of rice calli, as previously described[46].

Rice hydroponic culture and LC/MS-MS Analysis of *epi*-5DS

Preparation of hydroponic culture solution and treatment with GR24 (http://www.chiralix.com/) were performed as described previously[6]. In brief, surface-sterilized rice seeds were incubated in sterile water at 30 °C in the dark for 2 days. The germinated seeds were transferred into hydroponic culture medium solidified with 0.5% agar and cultured at 30 °C for 5 days. The one-week-old seedlings were then transferred to a plastics pot containing hydroponic culture solution (500 ml) and grown for an additional 7 days (total two weeks) or 28 days (total five weeks). The hydroponic culture solution with or without 1 μM GR24 was renewed every week. For strigolactone analysis, the one-week-old seedlings were grown in hydroponic culture medium without Pi for an additional 14 days (total three weeks).

Strigolactone analysis in rice root exudates was performed according to a previously described method[14] with minor modifications. For each sample, the weight of fresh rice roots was recorded and 50 ml hydroponic culture medium was loaded onto a pre-equilibrated Oasis HLB 3cc cartridges (Waters) after adding 1 ng d_6-5DS as the internal standard. Subsequently, the columns were washed with de-ionized water. The strigolactones-containing fraction, eluted with acetone, was collected and dried up under nitrogen gas, then reconstituted in acetonitrile and subjected to UPLC-MS/MS analysis. The UPLC-MS/MS system consists of a triple quadruple tandem mass spectrometer (Quattro Premier XE; Waters MS Technologies) and an Acquity Ultra Performance Liquid Chromatograph (Acquity UPLC; Waters) equipped with a reverse phase column (BEH-C18, 2.1×100 mm, 1.7 μm; Waters). Mobile phase A (99.5% H_2O + 0.5% acetic acid) and mobile phase B (99.5% ACN + 0.5% acetic acid) were pumped at a rate of 0.4 ml/min. The gradient started with 50% A and increased B from 50% to 70% in 5 min. The column temperature was set to 25 °C. MS parameters were set to the following values: desolvation gas flow 800 L/h, capillary voltage 3.8 kV, cone voltage 24 V, desolvation temperature 350 °C, source temperature 110 °C, collision energy 14 V, using MRM 331>216 transition for the *epi*-5DS detection, and MRM 337>222 transition for the d_6-5DS detection. Data were analyzed with MassLynx software (V.4.1). The *epi*-5DS concentrations of rice exudates were calculated by comparing the MRM relative response of *epi*-5DS with the d_6-5DS ones then divided by the fresh rice roots weight.

Quantitative real-time reverse transcriptase (RT)-PCR

Total RNA was extracted using the RNeasy Plant Mini Kit (QIAGEN). Then, 20 μl cDNA was synthesized using 1 μg RNA with the QuantiTect® Reverse Transcription Kit (QIAGEN). Quantitative RT-PCR (20 μl reaction volume) was carried out using 0.5 μl cDNA, 0.2 μM each gene-specific primer and SYBR® Premix Ex Taq™ Kit (TaKaRa) in an ABI PRISM 7900HT (Applied Biosystems) according to the manufacturer's instructions. The rice *Ubiquitin* gene was used as the internal control. The gene specific primers used for qRT-PCR are: *Ubiquitin* gene (LOC_Os03g13170) (5′-AACCAGCTGAGGCCCAAGA-3′ and 5′-ACGATTGATTTAACCAGTCCATGA-3′); *D53* gene (5′-

CCAAGCAGTTTGAAGCGAC-3′ and 5′-CCGCAAGTTTATCAAAGTCAA-3′); *D10*
gene (5′-CGTGGCGATATCGATGGT-3′ and 5′-CGACCTCCTCGAACGTCTT-3′); *FC1*
gene (5′-CGACAGCGGCAGCTACTAC-3′ and 5′-GCGAATTGGCGTAGACGA-3′). Data
from three replicates were analyzed following the relative quantification method[47].

Histological analysis and GUS staining

Culm segments of rice were fixed in FAA (formalin/acetic acid/50% ethanol, 2:1:17, v/v/v)
overnight, followed by a series of dehydration and infiltration, and embedded in paraffin
(Paraplast Plus; Sigma-Aldrich). The tissues were sliced into 8–12 μm sections with a
microtome (Leica RM2265), and affixed to microscope slides. Paraffin was removed from
the sections using xylene, and the sections were dehydrated through a gradient ethanol
series, and stained with toluidine blue. Sections were viewed under a light microscope
(Leica DM5000B) and photographed using a Micro Color CCD camera (Leica DFC490).

GUS histochemical staining was performed according to a method described previously[48].
Various tissues or hand-cut sections of *pD53::GUS* T_1 generation transgenic plants were
incubated in a staining solution containing 100 mM $NaPO_4$ buffer pH 7.0, 2 mM X-Gluc,
0.5 mM $K_3Fe(CN)_6$, 0.5 mM $K_4Fe(CN)_6$, 0.1% Triton X-100 and 10 mM Na_2EDTA at 37
°C. Samples were vacuum-infiltrated briefly at the initiation of staining with X-Gluc
solution. After staining, the staining solution was removed and the sample was washed with
several changes of 50% ethanol until the tissue becomes clear. Images were taken directly or
under the stereomicroscope.

Yeast two-hybrid (Y2H) assay

Coding regions of the rice *D53*, *d53* and various domain deletion variants of *D53* were
cloned into the Y2H 'prey' vector pGADT7 (Clontech). Coding region of rice *D14* was
cloned into the Y2H 'bait' vector pGBKT7 (Clontech). Bait and prey constructs were co-
transformed into the yeast (*Saccharomyces cerevisae*) strain Y2HGold (Clontech) and the
resultant yeast stains were grown on SD–Leu/–Trp plates for 3 days at 30 °C. Interactions
between bait and prey were examined on the control media –LT (SD–Leu/–Trp) and
selective media –LTHA (SD–Leu/–Trp/–His/–Ade) or –LTHA+X-α-gal in the presence or
absence of 5 μM GR24. Plates were incubated for 4 days at 30 °C. Yeast strains containing
D53, *d53*, or *D14* in combination with pGADT7 or pGBKT7 were used as controls.

Antibody preparation

For detection of D53, a His-fused protein-specific polypeptide (amino acids 963–1112 of
D53 protein) was expressed in *BM*Rosetta (DE3) cells (Biomed), and then affinity purified.
Subsequently, the recombination protein was injected into rabbits to produce polyclonal
antibodies against D53 (Abmart). For detection of D14 protein, a His-fused polypeptide
(amino acids 153–302 of D14) was expressed in *BM*Rosetta (DE3) cells. Purified fusion
protein was injected into rats to produce polyclonal antibodies against D14 (Abmart). The
loading control used is anti-HSP82 antibody (Cat: AbM51099-31-PU, Beijing Protein
Innovation).

In vivo degradation assay of D53

One-week-old rice seedlings were treated with or without 5 μM GR24 and collected at different time points. To block proteasomal protein degradation, seedlings were pre-treated with 40 μM MG132 for 2 hours. Total protein was extracted and denatured in the SDS sample buffer containing 5% β-Mercaptoethanol (β-ME) at 95 °C for 10 minutes. Western blots were performed using the antiserum against D53 and visualized by an enhanced HRP-DAB substrate kit (Cat: PA110, TIANGEN).

The *pActin::D53-GFP* and *pActin::d53-GFP* transgenic plants were treated with 5 μM GR24 with or without the following inhibitors: MG132 (40 μM), AEBSF (500 μM), Pepstatin A (1 μM) or Leupeptin (20 μM). GFP fluorescence in nuclei of transgenic root cells was observed and imaged with a Zeiss LSM 700 laser scanning confocal microscope.

BiFC assay

The full-length *D14* and *D53* cDNAs were cloned into the binary vector pSPYNE173 and pSPYCE(M) to create the D14-NY and D53-CY vector, respectively. For transient expression, *A. tumefaciens* strain EH105 carrying the gene of interest (at an OD_{600} of 0.1) was co-infiltrated with p19 strain into 5-week-old *N. benthamiana* leaf as previously described[49]. After two days, infiltrated leaves were sprayed with (+) or without (−) 5 μM GR24 twenty-four hours prior to leaf excision. The eYFP and DAPI fluorescent signals of the infiltrated leaves were monitored sequentially using a laser confocal scanning microscope. The wavelengths for eYFP and DAPI were 514 and 405 nm for excitation, and 527 and 488 nm for detection, respectively.

Recombinant protein preparation

The full-length open reading frames of D3-OSK1 and D14 were cloned into the pGEX4T-1 vector, generating a fusion with the glutathione S-transferase (GST) protein. The full-length cDNA of D3 was cloned into the pMAL-c2x vector to generate a fusion with the maltose-binding protein (MBP). Expression of GST, GST-D3-OSK1, GST-D14, MBP and MBP-D3 in BL21 Rosetta cells was induced with 0.1 mM isopropyl-1-thio-D-galactopyranoside (IPTG) at 20 °C for 16 hours. Fusion proteins were purified using GST Bind™ Resin (Cat: 70541-3, Novagen) or amylose-affinity chromatography (Cat: E8021L, New England Biolabs) according to the manufacturer's protocols and quantified by the Bio-Rad protein assay reagent.

Pull-down assays

One-week-old rice seedlings were pre-treated with 40 μM MG132 for one hour and then treated with or without 5 μM GR24 for 30 minutes. Total proteins were subsequently extracted in a RIPA buffer (50 mM Tris-HCl pH 7.2, 150 mM NaCl, 10 mM $MgCl_2$, 1× complete protease inhibitor cocktail, Roche) containing 40 μM MG132. After centrifugation (16,000 g at 4 °C), the supernatant was collected. Total protein concentration was determined by the Bradford protein assay kit (Bio-Rad). Roughly equal amounts of purified GST and GST-D3-OSK1 fusion proteins (about 50 μg) were affixed to GST Bind™ Resin and then mixed with 300 μl of plant extract (containing 1.5 mg total proteins). After gently

shaken at 25 °C for 15 minutes, the resin was washed five times with RIPA buffer. During all the procedures, 5 μM GR24 was added to the assay mixture for GR24-treated seedlings and was not added to the protein samples from seedlings not treated with GR24. Proteins were detected with anti-D14 antibodies at 1:1000 dilution and visualized with enhanced chemiluminescence (ECL) reagent (GE Healthcare).

For *in vitro* pull-down, GST-D14 (5 μg) was incubated with MBP-D3 (6 μg) or MBP (2 μg) at 25 °C for 15 minutes in 300 μl of binding buffer (20 mM Tris-HCl, pH 7.6, 2.5 mM β-ME, and 0.1 M NaCl) with or without 10 μM GR24. After incubation, 20 μl of GST resin was added. After further incubation at 25 °C for 15 minutes, the resin was washed five times with the washing buffer (20 mM Tris-HCl, pH 7.5, 500 mM NaCl, and 0.5% Triton X-100). The washing buffer also contains 10 μM GR24 where appropriate. After washing, 40 μl of 1× SDS-PAGE sample buffer were added, the mixture was denatured, and the sample was loaded on a 8% SDS-PAGE gel, and proteins were detected by the horseradish peroxidase (HRP)-conjugated anti-MBP antibody (New England Biolabs) and visualized with enhanced chemiluminescence (ECL) reagent (GE Healthcare).

Transient expression assays

The *D53*, *d53*, *D3* and *D14* full-length cDNAs were cloned into the vector pA7-GFP or pGreen II 0800-LUC to generate D53-GFP or D53-LUC recombinant plasmids for transient expression assays, respectively. Plasmids DNA was prepared using the Plasmid Midi Kits (Cat: 12143, QIAGEN) according to the manufacturer's instructions. For preparation of rice protoplasts, the sterilized seeds were germinated and grown in climate chambers for 7–10 days. Green tissues from the stem and sheath of 40–60 rice seedlings were used. The protoplasts isolation and PEG-mediated transfections procedure were carried out as described previously[50]. GFP fluorescence from protoplasts was observed and imaged as described above. For measurement of the relative luciferase activity, protoplasts were incubated overnight and then treated with or without 5 μM GR24 for 4 hours. For MG132 treatment, 40 μM MG132 was added to the sample and incubated for one hour before addition of GR24. Activities of fLUC and rLUC were determined with the Dual-Glo® Luciferase Assay System (Cat: E2920, Promega). The data was represented as the ratio of firefly to the Renilla luciferase activity (fLuc/rLuc).

Acknowledgments

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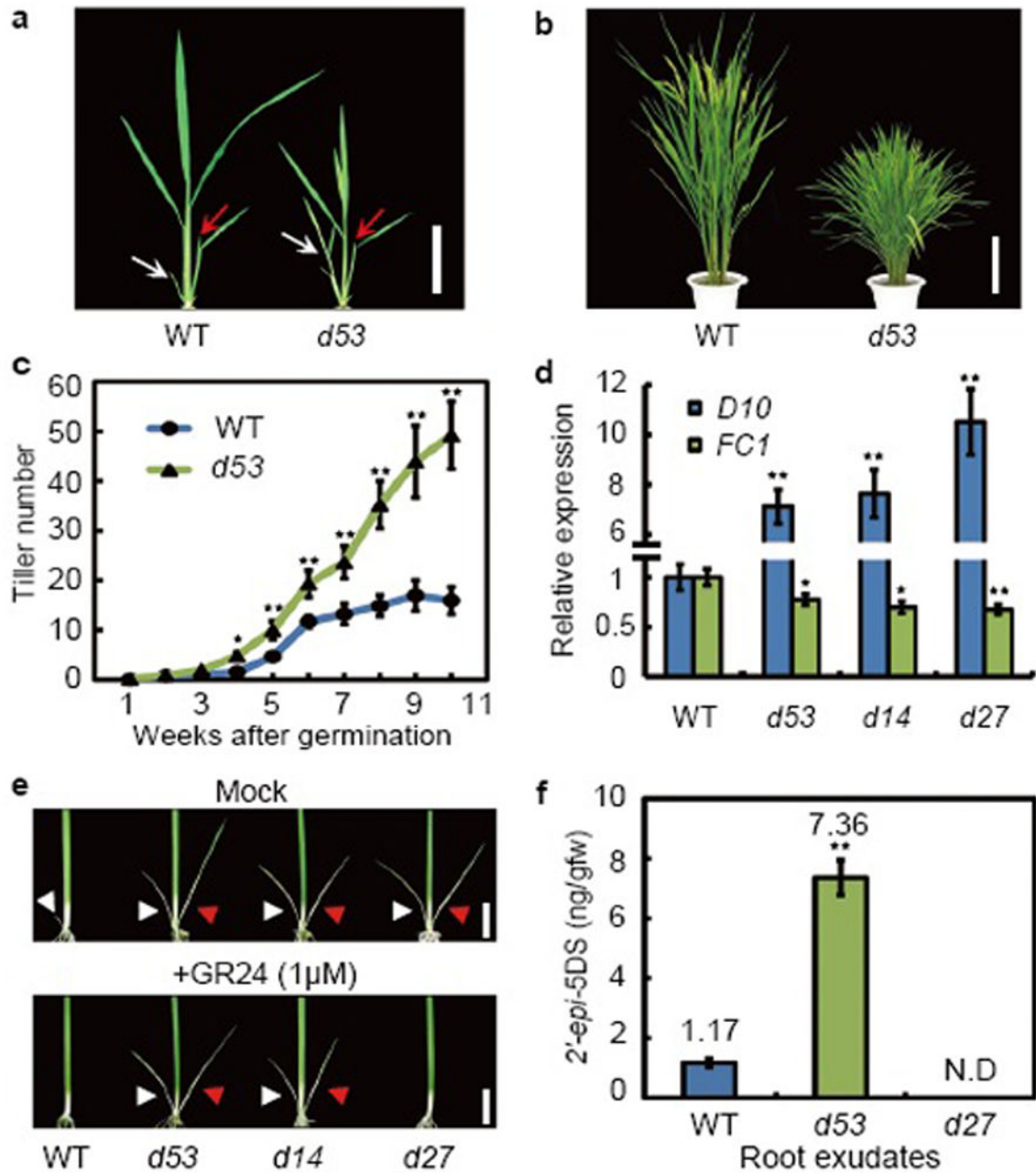
Figure 1. Phenotype of *d53* mutant

a, b, Phenotype of wild type (WT) and *d53* mutant at four-week-old seedling stage **(a)** or heading stage **(b)**. White arrows indicate the first tillers in *d53*, which is usually absent in WT, and red arrows show the second tillers. **c,** Comparison of tillering kinetics at different developmental stages. **d,** Real-time RT-PCR assay showing altered expression of *D10* and *FC1* in *d* mutants. **e,** Responses of rice seedlings to GR24 treatment. Red and white arrowheads indicate the first and second tillers, respectively. **f,** LC/MS-MS measurement of *epi*-5DS levels in root exudates. gfw, per gram fresh weight. Scale bars, 5 cm in **a,** 30 cm in **b** and 2 cm in **e**. Values are means ± s.d. (**c,** $n = 30$ plants; **d** and **f,** $n = 3$ replicates). The
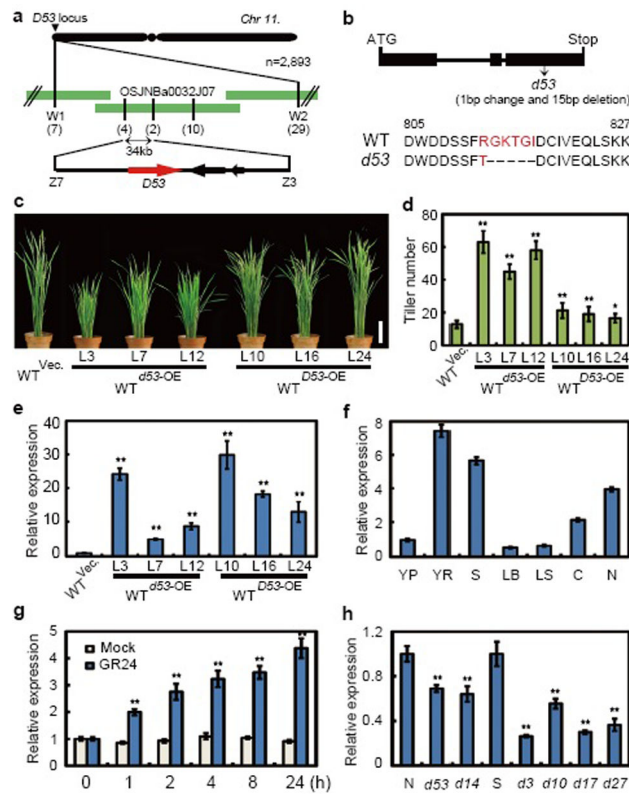
Student's *t*-test analysis indicates a significant difference (compared with WT, *P<0.05, **P<0.01).

Figure 2. Map-based cloning and characterization of *D53*

a, *D53* was fine-mapped on chromosome 11. The numbers of recombinants are shown in brackets. **b,** Molecular lesions in *d53* mutant. **c,** Phenotypic comparison of *proActin::D53-GFP* and *proActin::d53-GFP* transgenic plants. Vector, *proActin::GFP* control. Scale bar, 10 cm. **d, e,** Tiller number (**d**) and relative expression of *D53* (**e**) of transgenic plants in **c. f,** *D53* expression in various organs, including young panicles (YP), young roots (YR), shoots (S), leaf blades (LB), leaf sheaths (LS), culms (C) and nodes (N). **g,** GR24 treatment induces *D53* expression. **h,** Relative expression levels of *D53* in two wild-type varieties Norin 8 (N) and Shiokari (S), and six rice *d* mutants. Each value in **d**–**h** represents the mean ± s.d. (**d** and **f**–**h**, $n = 3$ replicates; **e**, $n = 20$ plants). The Student's *t*-test analysis indicates a significant difference (compared with control, *$P<0.05$, **$P<0.01$).
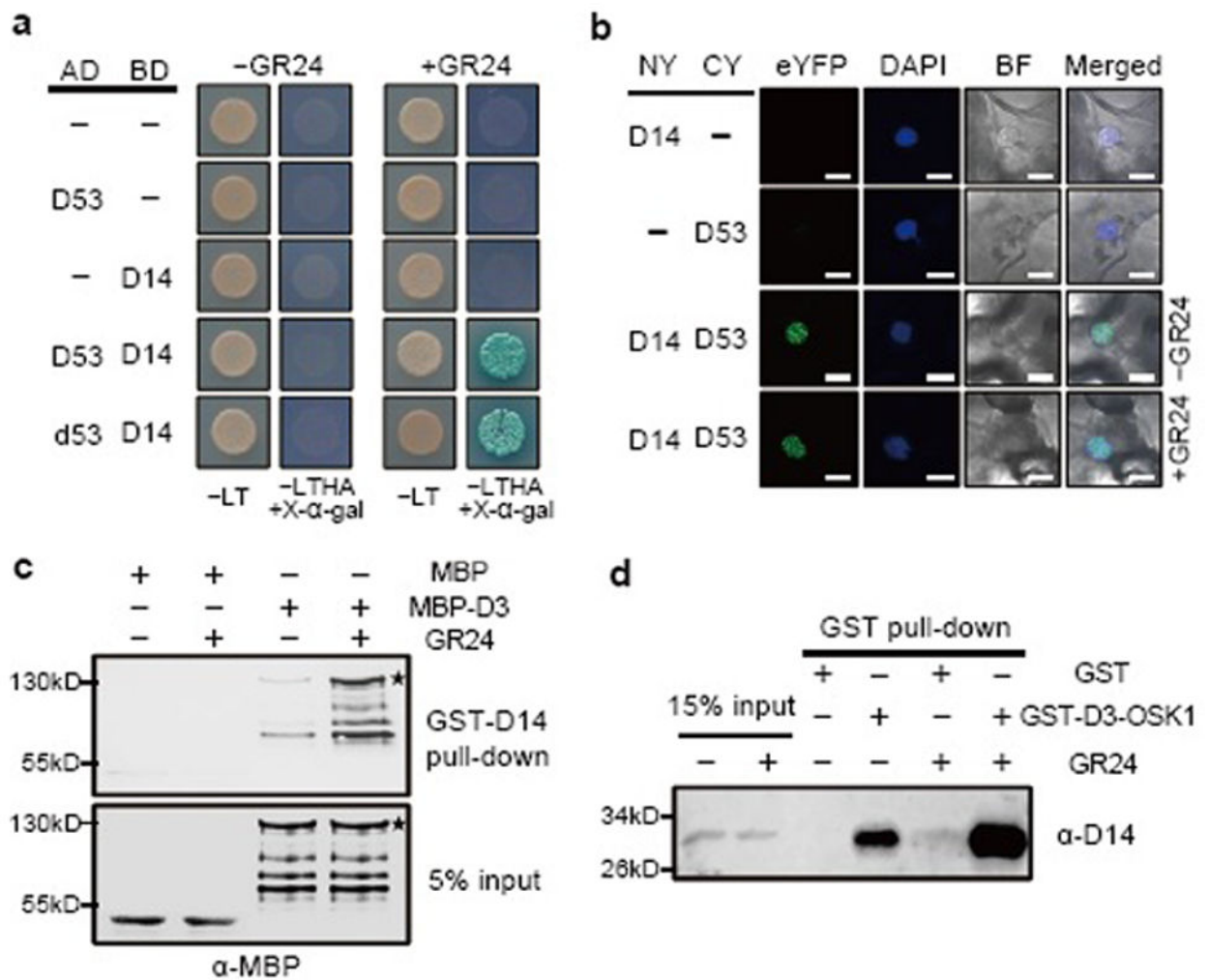
Figure 3. GR24 promotes the D53-D14 and D14-D3 interaction

a, Y2H showing that D53 and d53 interact with D14 in the presence of GR24. Yeast transformants were spotted on the control medium: –LT and selective medium: –LTHA + X-α-gal. **b,** BiFC analysis of D53 and D14. The positions of nuclei are indicated by DAPI staining. Scale bar, 10 μm. **c,** *In vitro* pull-down assay of recombinant MBP-D3 or MBP using resins containing GST-D14. Asterisks indicate the full-length MBP-D3 protein. **d,** Pull-down assay showing co-IP of D14 from the *d3* mutant plant extracts, using GST-D3-OSK1 as the bait. 'Input' shows that roughly equal amount of total plant proteins was used.
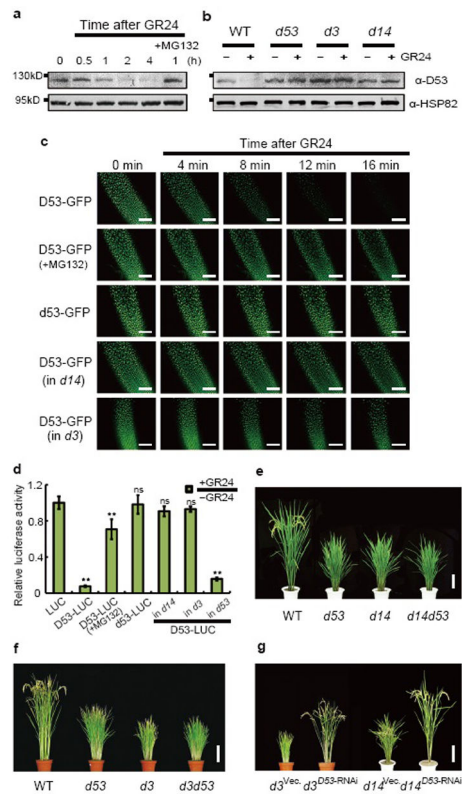
Figure 4. GR24 promotes D14- and D3-dependent proteasomal degradation of D53

a, b, Western blot analysis showing that GR24 promotes D53 protein degradation in wild-type **(a)**, but not in *d* mutants **(b)**. 10 μg of total protein were applied in each lane. **c,** Confocal scanning images showing different degradation patterns of D53-GFP and d53-GFP fusion proteins in wild-type, *d3* or *d14* backgrounds. **d,** Relative luciferase activity of D53-LUC or d53-LUC in wild-type, *d14*, *d3* or *d53* protoplasts. Values are means ± s.d. of three independent experiments. The double asterisks represent significance difference compared with control (LUC) determined by the Student's *t*-test at *P*<0.01. ns, not significant. **e, f,** Phenotype of *d14 d53* **(e)** and *d3 d53* **(f)** double mutants. **g,** Phenotypes of *D53*-RNAi transgenic plants in *d3* and *d14* backgrounds. Scale bars, 100 μm in **c** and 20 cm in **e**–**g**.
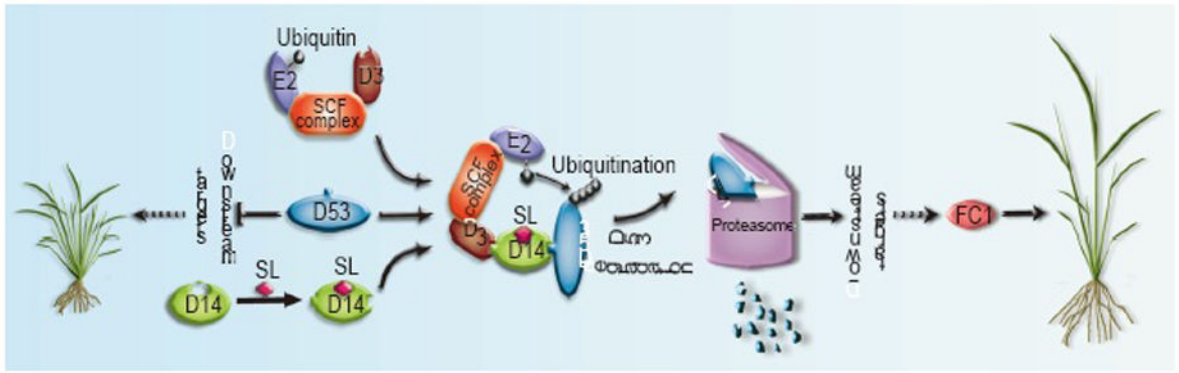
Figure 5.
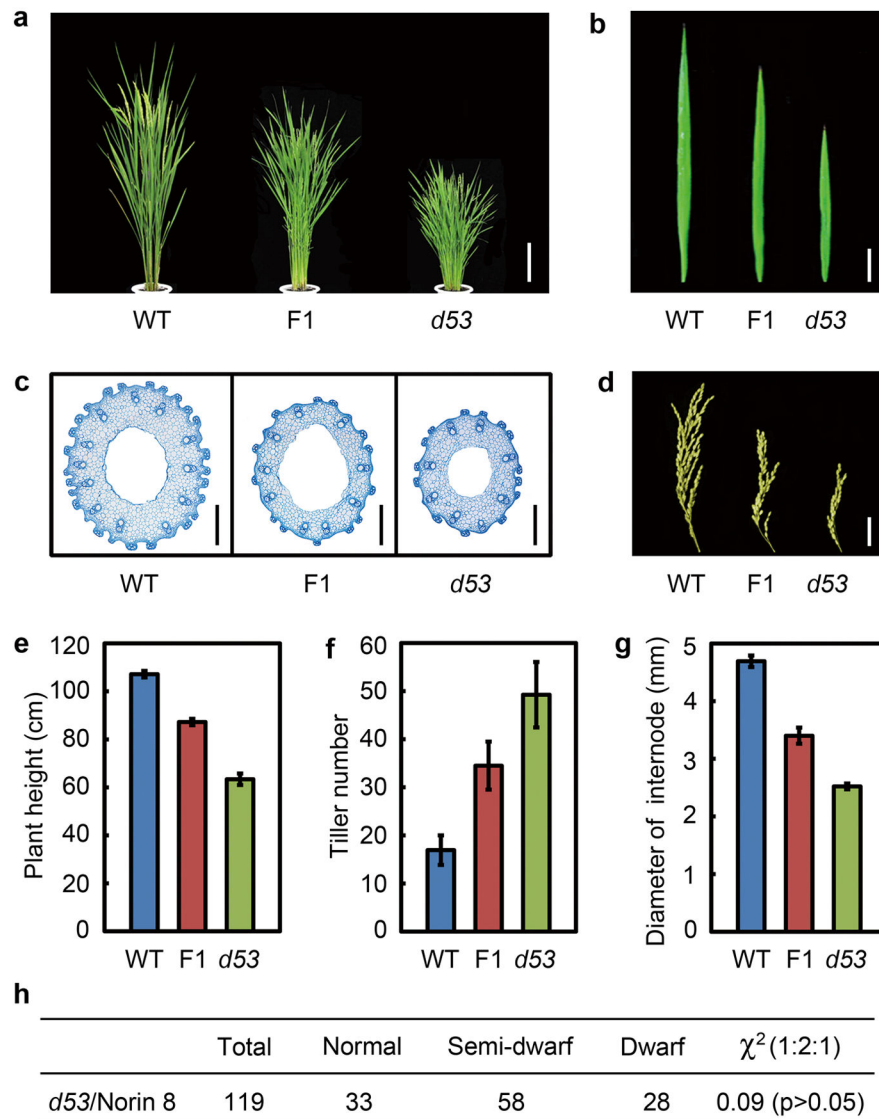A schematic model depicting SLs promote D14-SCF^{D3}-mediated degradation of D53.
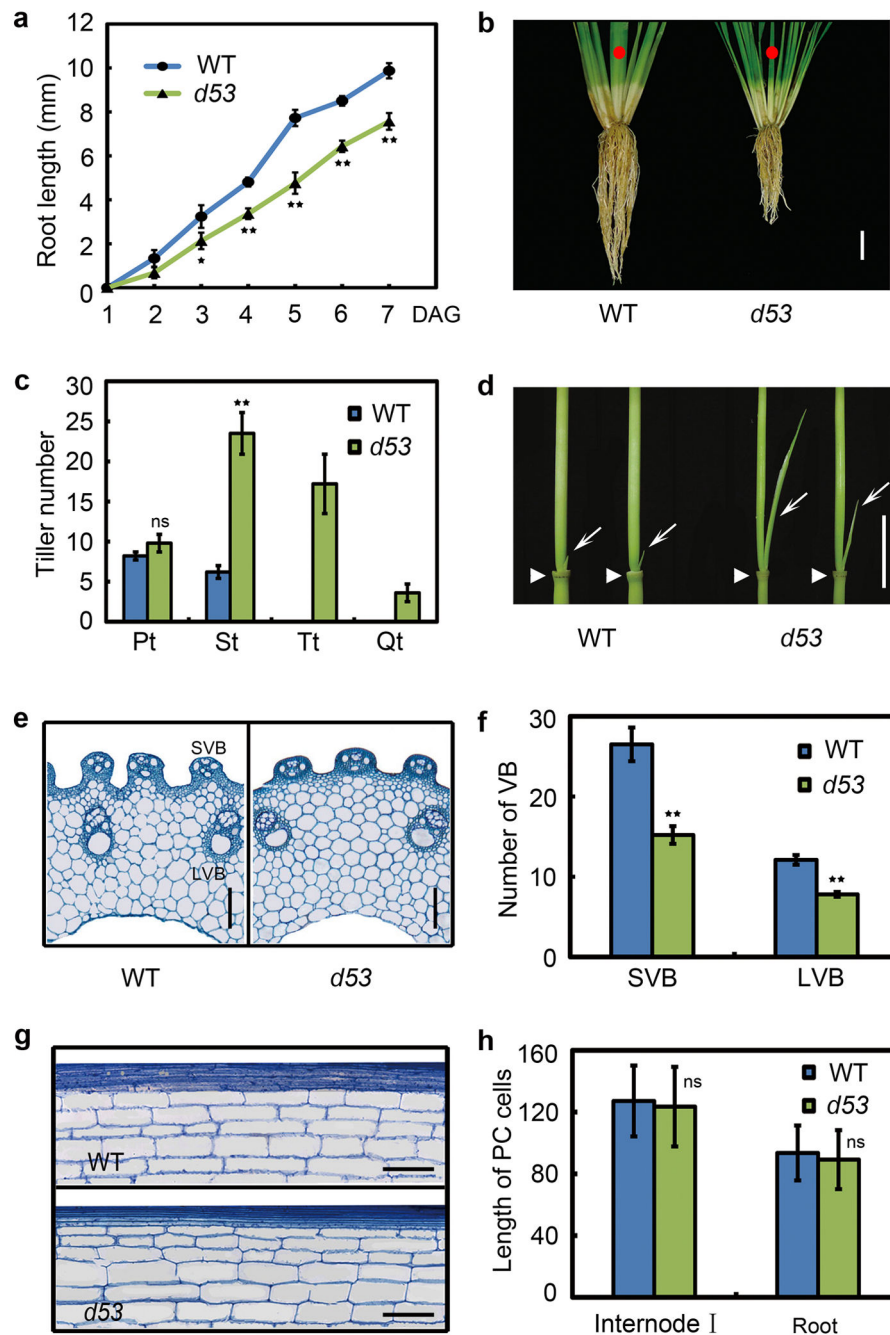
Extended Data Figure 1. Phenotypes of *d53* mutant

a, Comparison of crown root growth in wild type (WT) and *d53* mutant. DAG, day after germination. Each value represents the mean ± s.d. of 25 seedlings. **b,** Root phenotype of seven-week-old WT and *d53* at the tillering stage. Red dots indicate the main culms. **c,** Comparison of different types of tillers between WT and *d53* at the heading stage. Pt, primary tillers; St, secondary tillers; Tt, tertiary tillers; Qt, quaternary tillers. Each value represents the mean ± s.d. of 20 seedlings. **d,** Morphology comparison of tiller buds at the second node between WT and *d53*. White arrows and arrowheads indicate the tiller buds and

the second nodes, respectively. **e,** Transverse sections of the first internode of WT and *d53*. **f,** Number of vascular bundles (VB) calculated from transverse sections of the first internode of WT and *d53*. SVB, small vascular bundle; LVB, large vascular bundle. Data are means ± s.d. (n=10). **g,** Longitudinal sections of the first internode of WT and *d53*. **h,** Comparison of parenchyma (PC) cell length in first internode and root between WT and *d53*. Data are means ± s.d. (n=10). Differences with respect to the WT that were found to be significant in a *t*-test are indicated with asterisks (*P<0.05; ** P<0.01; ns, not significant). Scale bars, 10 cm in **b**, 2 cm in **d** and 100 μm in **e** and **g**.

|	Total	Normal	Semi-dwarf	Dwarf	χ^2 (1:2:1)
d53/Norin 8	119	33	58	28	0.09 (p>0.05)

Extended Data Figure 2. *d53* mutation behaves in a semi-dominant manner

Comparison of WT, heterozygous (F_1) and homozygous *d53* plants at the heading stage (**a**), flag leaf (**b**), cross section of the first internode (**c**), panicle (**d**), plant height (**e**), tiller number (**f**) and diameter of the third internode (**g**). Scale bars, 20 cm in **a**, 5 cm in **b** and **d** and 500 μm in **c**. For **e**–**g**, each value represents the mean ± s.d. (n=25). **h,** Segregation of F_2 progeny from a self-pollinated F_1 plant (*d53* × Norin 8).
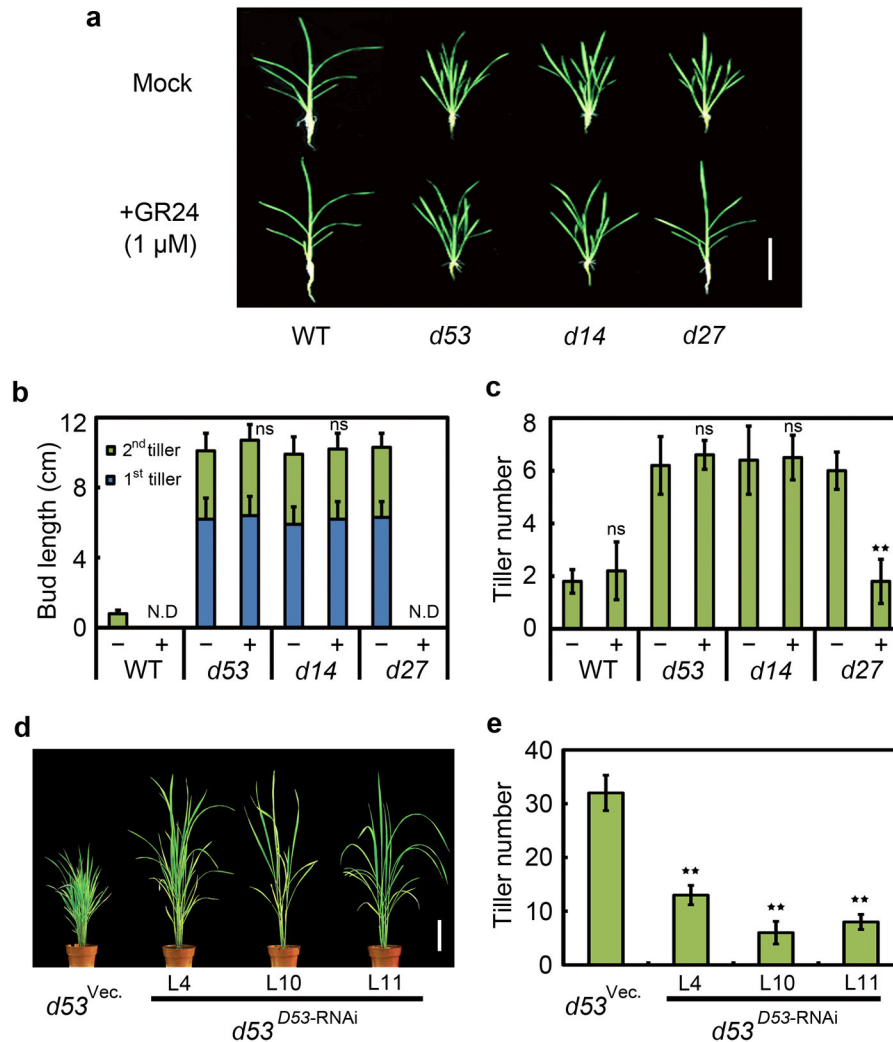
Extended Data Figure 3. *d53* **is insensitive to GR24 treatment and confers enhanced tillering promoting activity**

a, Response of five-week-old WT, *d53*, *d14* and *d27* to the application of 1 μM GR24. **b,** Tiller bud length of two-week-old WT, *d53*, *d14* and *d27* seedling treated with (+) or without (−) 1 μM GR24. Data are means + s.d. (n=10). **c,** Numbers of tillers showing outgrowth (>2 mm) for five-week-old WT, *d53*, *d14* and *d27* plants treated with (+) or without (−) 1 μM GR24. Data are means ± s.d. (n=10). Asterisks in **b** and **c** denote significant differences between treated and untreated samples within the same genotype (two tailed Mann-Whitney *U* test, *P*<0.01; ns, not significant). **d,** *D53* RNAi transgenic plants exhibit reduced tillering in the *d53* mutant background. Vector, *d53* transformed with the pCUbi1390- FAD2 control. **e,** Tiller number of RNAi transgenic lines in **d** at the tillering stage. Each value represents the mean ± s.d. of six plants (T$_1$ generation). L4, L10 and L11 represent three independent lines. The *t*-test analysis indicated a significant difference (compared with vector control, **P*<0.01). Scale bars, 20 cm in **a** and 10 cm in **d**.
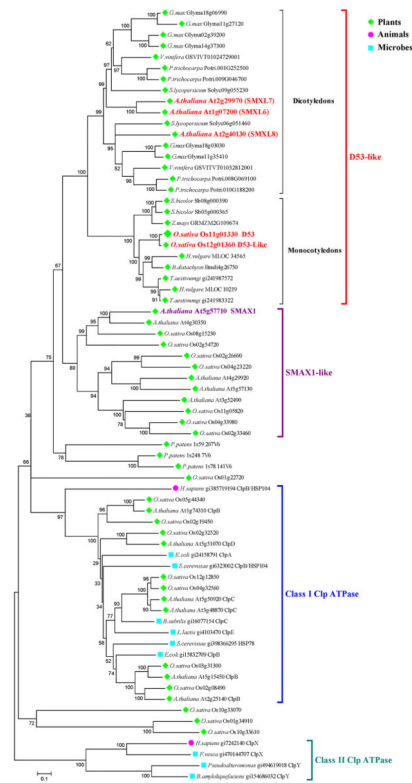
**Extended Data Figure 4. Phylogenetic analysis of D53 protein**
Using the D53 protein sequence as the query in tblastn searches, homologs were identified from different organisms with a permissive cutoff E value of 1E-3. The sequences chosen from representative genome were aligned and used to generate the neighbor-joining phylogenetic tree with 1,000 bootstrap replicates. The clade names were given based on known sequences in each clade, which is supported by a bootstrap value >85.
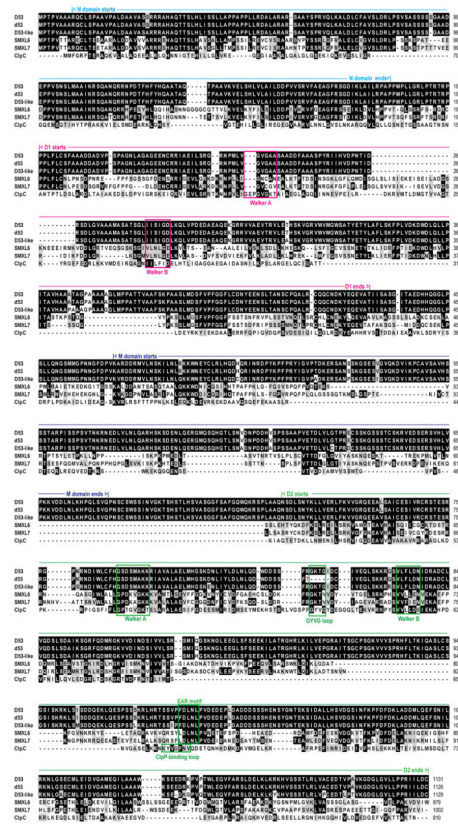
Extended Data Figure 5. Multiple sequence alignment of the deduced amino acid sequence of D53 with its homologs

D53 protein is predicted to contain an N-terminal domain, a D1 ATPase domain, an M domain, and a D2 ATPase domain (http://toolkit.tuebingen.mpg.de/hhpred). The beginning and ending sites of each domain are indicated above the sequences. The predicted Walker A (P-loop) and Walker B motifs are shown in red boxes in the D1 domain and green boxes in the D2 domain, respectively. Note that the deletion of five amino acids in the D2 domain of d53 protein overlaps with the GYVG loop in ClpC. The conserved putative EAR motif in D53 and ClpP-binding loop in ClpC are also shown. The sequences used for alignment are D53 (*Oryza sativa*, LOC_Os11g01330), D53-like (*Oryza sativa*, LOC_Os12g01360), SMXL6 (*Arabidopsis*, At1g07200), SMXL7 (*Arabidopsis*, At2g29970), and ClpC (*Bacillus subtilis*, GI: 16077154).
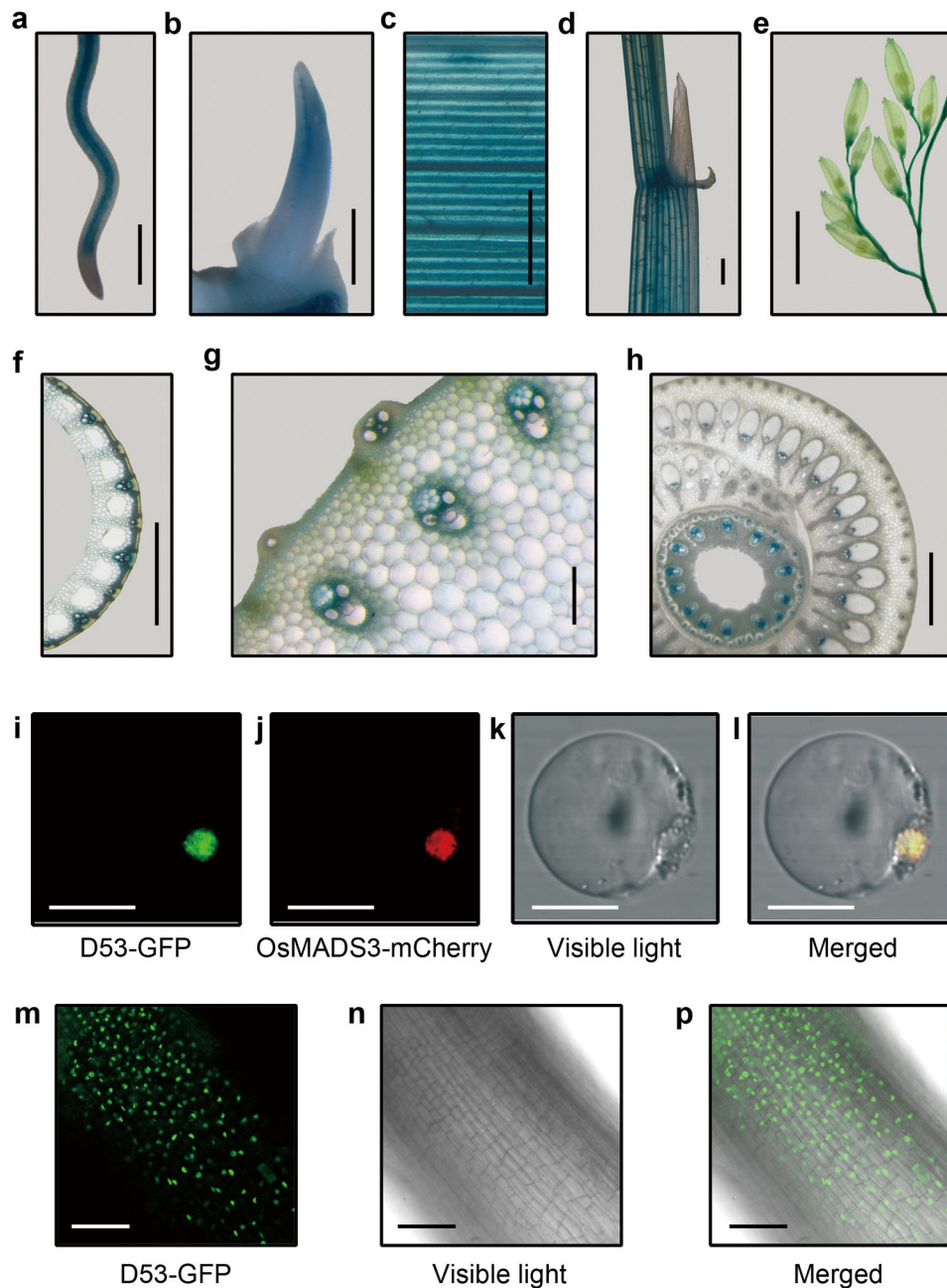
Extended Data Figure 6. Histochemical staining of the *pD53::GUS* reporter gene and subcellular localization of D53 protein

a–**h,** Histochemical staining of young root (**a**), shoot (**b**), leaf (**c**), leaf sheath (**d**), panicle (**e**), transverse section of the leaf sheath (**f**), stem (**g**) and node (**h**). Scale bars, 1 mm in **a**, **b**, **c**, **d**, **f** and **h**; 1 cm in **e** and 100 μm in **g. i**–**l,** Subcellular localization of D53-GFP fusion protein in rice protoplast cells. A nuclear marker protein, OsMADS3, fused with mCherry, was used as a positive control. Scale bars, 5 μm. **m**–**p,** Confocal scanning images showing nuclear localization of the D53-GFP fusion protein in transgenic root cells. Scale bars, 100 μm.
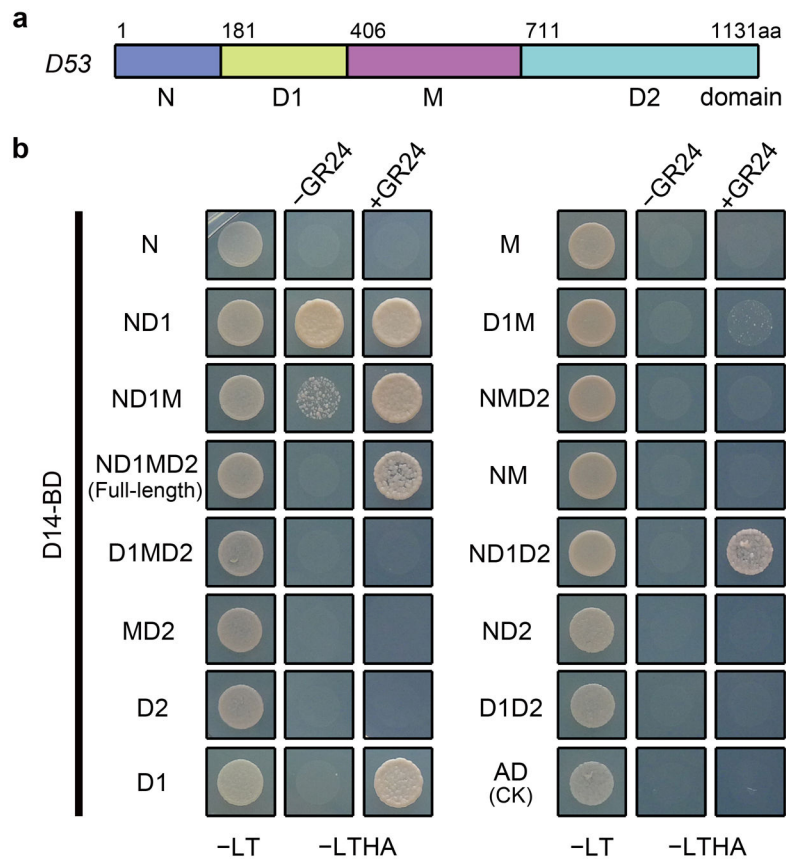
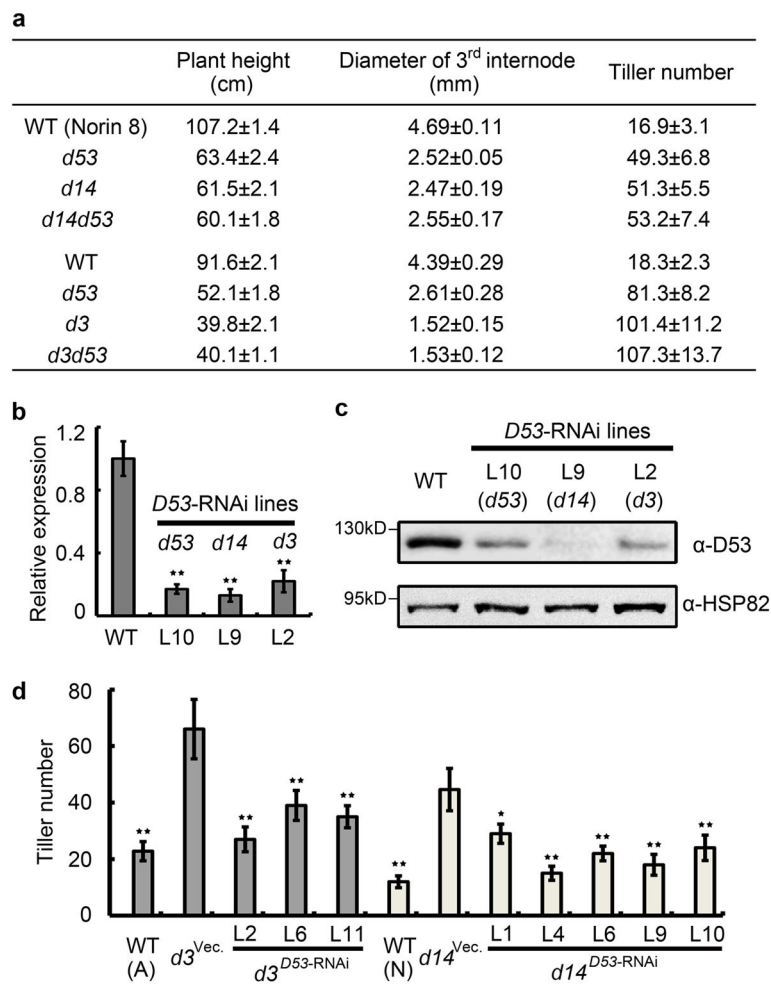**Extended Data Figure 7. Mapping of the D14-binding domain of D53**

a, Schematic structure of the D53 protein. Numbers indicate amino-acid (aa) residues. **b,** Y2H analysis showing interaction between full-length and various domain deletion variants of D53 with D14 in the presence or absence of 5 μM GR24. –LT, control medium (SD–Leu/–Trp); –LTHA, selective medium (SD–Leu/–Trp/–His/–Ade).
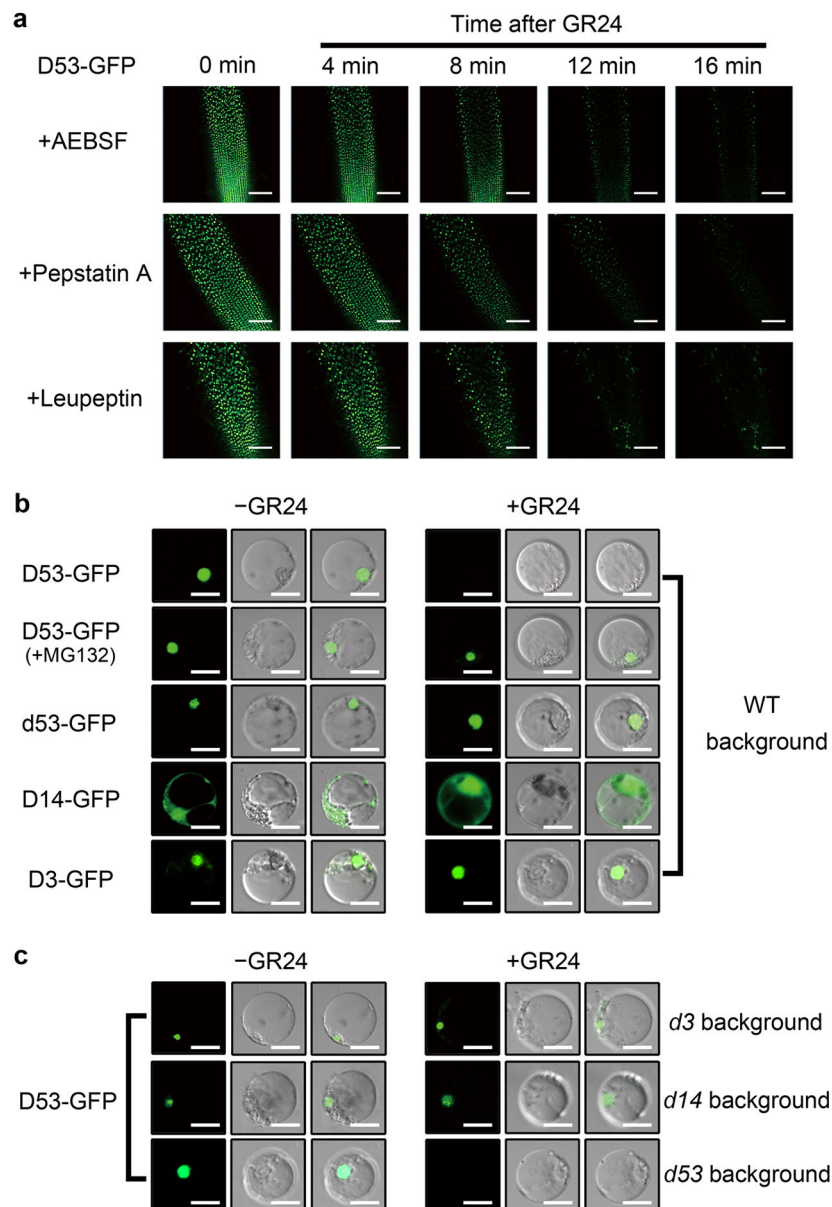
Extended Data Figure 8. GR24 promotes D53 protein degradation

a, Confocal scanning images showing that AEBSF, Pepstatin A and Leupetin are not effective in blocking D53-GFP fusion protein degradation in transgenic seedlings treated with 5 μM GR24. Scale bars, 100 μm. **b,** Degradation of D53-GFP fusion protein, but not D3-GFP and D14-GFP fusion proteins expressed in rice protoplasts, in the presence of 5 μM GR24. Pre-treatment with 40 μM MG132 for one hour before addition of GR24 effectively blocks D53-GFP degradation. **c.** D53-GFP is degraded in the *d53* mutant protoplasts in the presence of GR24, but not in *d3* or *d14* protoplasts. For **b** and **c**, each figure represents at least fifty cells observed. Scale bars, 10 μm.

a

	Plant height (cm)	Diameter of 3rd internode (mm)	Tiller number
WT (Norin 8)	107.2±1.4	4.69±0.11	16.9±3.1
d53	63.4±2.4	2.52±0.05	49.3±6.8
d14	61.5±2.1	2.47±0.19	51.3±5.5
d14d53	60.1±1.8	2.55±0.17	53.2±7.4
WT	91.6±2.1	4.39±0.29	18.3±2.3
d53	52.1±1.8	2.61±0.28	81.3±8.2
d3	39.8±2.1	1.52±0.15	101.4±11.2
d3d53	40.1±1.1	1.53±0.12	107.3±13.7

Extended Data Figure 9. *D53*-RNAi transgenic lines in *d3* and *d14* backgrounds

a, Comparison of plant height, diameter of the 3rd internode and tiller number between WT, *d53*, *d14*, *d3* and their double mutants. Values are mean ± s.d. (n=10). **b, c,** Real-time RT-PCR assay **(b)** and Western blot analysis **(c)** showing that the endogenous level of *D53* mRNAs and proteins are down-regulated in three representative *D53*-RNAi lines in *d53*, *d14* and *d3* mutant backgrounds, compared to WT control. Data are means ± s.d. (n=3). The double asterisks represent the significant difference determined by the *t*-test at $P<0.01$. Anti-HSP82 was used as a loading control. **d,** Tiller number of representative *D53*-RNAi transgenic lines in *d3* and *d14* mutant backgrounds at the heading stage. Each value represents the mean ± s.d. of six plants. L2, L6 and L11 represent three independent lines in *d3* background, and L1, L4, L6, L9 and L10 represent five independent lines in *d14* background. Akumuro (S) and Norin 8 (N) are the wild-type varieties correspond to *d3* and *d14* mutants, respectively. *d3*Vec. and *d14*Vec. transgenic lines were used as the controls. The single and double asterisks represent significance difference compared with vector control determined by the *t*-test at $P<0.05$ and $P<0.01$, respectively.