Ufd2p synthesizes branched ubiquitin chains to promote the degradation of substrates modified with atypical chains

Chao Liu1, Weixiao Liu1, Yihong Ye2 & Wei Li1

Ubiquitination of a subset of proteins by ubiquitin chain elongation factors (E4), represented by Ufd2p in *Saccharomyces cerevisiae*, is a pivotal regulator for many biological processes. However, the mechanism of Ufd2p-mediated ubiquitination is largely unclear. Here, we show that Ufd2p catalyses K48-linked multi-monoubiquitination on K29-linked ubiquitin chains assembled by the ubiquitin ligase (Ufd4p), resulting in branched ubiquitin chains. This reaction depends on the interaction of K29-linked ubiquitin chains with two N-terminal loops of Ufd2p. Only following the addition of K48-linked ubiquitin to substrates modified with K29-linked ubiquitin chains, can the substrates be escorted to the proteasome for degradation. We demonstrate that this ubiquitin chain linkage switching reaction is essential for ERAD, oleic acid and acid pH resistance in yeast. Thus, our results suggest that Ufd2p functions by switching ubiquitin chain linkages to allow the degradation of proteins modified with a ubiquitin linkage, which is normally not targeted to the proteasome.
Protein ubiquitination is an important post-translational modification in which ubiquitin, a 76-residue protein, is covalently attached to target proteins. Ubiquitination requires three classes of enzymes that act in concert: a ubiquitin-activating enzyme (E1) forms a thioester bond between itself and the C-terminus of ubiquitin, a ubiquitin-conjugating enzyme (E2) receives the activated ubiquitin from the E1 by transesterification, and a ubiquitin ligase (E3), including really interesting new gene (RING), homologous to E6-associated protein C-terminus (HECT) and RING-in-between-RING (RBR) type ligase, that transfers ubiquitin from the E2 to a lysine (K) or a serine/threonine residue in the target protein1−4. Ubiquitination regulates several cellular processes, including protein quality control, cell-cycle progression, endocytosis, cell signalling, and DNA damage responses1,2,5−7. The functional diversity of ubiquitination is achieved by the ability of ubiquitin to form different types of conjugates, each representing a distinct functional signal8,9. In addition to monoubiquitination and multi-monoubiquitination, polyubiquitination involves linking ubiquitin to one of the seven lysine residues or to the N-terminal methionine (Met1) of ubiquitin itself10−12. Ubiquitin can also be conjugated simultaneously to more than one lysine residue within a ubiquitin moiety to form branched ubiquitin chains13−15.

Although several general models have been proposed to explain the mechanisms underlying linkage-specific polyubiquitination, the precise mechanisms underlying many polyubiquitin chain assembly reactions remain unclear, particularly those synthesizing the recently discovered atypical ubiquitin chains or branched chains11,14,16−27. In addition to the three enzyme classes mentioned above, a class of chain elongation factor was also indentified more than ten years ago, termed the E4 or ubiquitin chain-binding protein Ufd3p (ref. 39) could efficiently pull down Ubc4p as the appropriate E2 for polyubiquitination by Ufd4p and Ufd2p (Supplementary Fig. 1). When purified Ufd2p was added to a reaction containing E1, Ubc4p and Ufd4p, it could efficiently enhance Ufd4p-mediated polyubiquitination of Ub-V-GFP (Fig. 1b), as previously demonstrated19,38.

To determine the linkage preference of the polyubiquitination system, we created a collection of Ub-V-GFP mutants that carry a single lysine in the ubiquitin moiety or have one of the seven lysine residues in ubiquitin mutated to arginine (K to R). In vitro ubiquitination using these Ub-V-GFP mutants showed that both K29 and K48 are required for Ufd4p and Ufd2p-mediated polyubiquitination of Ub-V-GFP because neither K48R, K29R nor K29, K48 single lysine Ub-V-GFP mutants could be efficiently polyubiquitinated when compared with wild-type (WT) Ub-V-GFP (Supplementary Fig. 2a,b).

To delineate the linkage specificity for each of the two enzymes, we first reconstituted the ubiquitination reaction using Ufd4p in the absence of Ufd2p. We found that in the Ufd4p-mediated polyubiquitination of Ub-V-GFP, K29 but not K48 on Ub-V-GFP was required (Supplementary Fig. 2c,d). This was because the ubiquitination pattern of Ub-V-GFP bearing only K29 and K48 was similar to that of the K29-only Ub-V-GFP mutant regardless of whether WT, K29R or K29-only ubiquitin mutants were used in the reaction (Fig. 1c). In addition, the K29-linked polyubiquitin chain-binding protein Ufd3p (ref. 39) could efficiently pull down Ufd4p synthesized Ubn-GFP (Supplementary Fig. 3a). Thus, we concluded that the K29 of ubiquitin is preferentially used by Ufd4p to build polyubiquitin chains on Ub-V-GFP, in line with previous findings31,40.

To examine the linkage specificity of Ufd2p-mediated ubiquitination, we used a previously established single-round ubiquitin turnover assay41. FLAG-tagged ubiquitin was charged onto Ubc4p. The charging reaction was quenched by addition of N-ethylmaleimide (NEM)/Ethylene diamine tetracetic acid...
K29/48-only Ub-V-GFP were tested for their ability to be ubiquitinated by the Ufd4p-Ufd2p ubiquitination system. See also Supplementary Figs 1–3.

on Ub-V-GFP are necessary and required for full ubiquitination by Ufd4p and Ufd2p. K48-only Ub-V-GFP, wild-type Ub-V-GFP, K29-only Ub-V-GFP, and K29-, K33-, K48-, or K63-only ubiquitin mutants (Lanes 2–9) was added in excess (10 fold of K29R ubiquitin) to the mixtures, together with Ufd2p, ubiquitin was charged onto the active cysteine of Ubc4p, and the reaction was quenched by adding NEM/EDTA. Wild-type ubiquitin or the K6-, K11-, K27-, K29-only ubiquitin. (Lanes 4–5). The reaction products were indistinguishable from the products of the K29-only Ub-V-GFP mutant (Lanes 1–3) in the presence of wild-type, K29R-or polyubiquitin chain assembly on Ub-V-GFP. Ufd4p-mediated polyubiquitination of a Ub-V-GFP mutant with only K29 and K48 in its ubiquitin region (Lanes 4–5). The reaction products were indistinguishable from the products of the K29-only Ub-V-GFP mutant (Lanes 1-3) in the presence of wild-type, K29R-or polyubiquitin chain assembly on Ub-V-GFP. Ufd4p-mediated polyubiquitination of a Ub-V-GFP mutant with only K29 and K48 in its ubiquitin region (Lanes 4–5). The reaction products were indistinguishable from the products of the K29-only Ub-V-GFP mutant (Lanes 1-3) in the presence of wild-type, K29R-or polyubiquitin chain assembly on Ub-V-GFP. Ufd4p-mediated polyubiquitination of a Ub-V-GFP mutant with only K29 and K48 in its ubiquitin region (Lanes 4–5).

Figure 1 | Ufd4p and Ufd2p synthesize K29- and K48-linked ubiquitin chains, respectively. (a) Coomassie blue-stained gels showing the expression and purification of relevant enzymes. (b) Reconstitution of the Ufd4p-Ufd2p ubiquitination system. Either Ufd2p (Lane 2), Ufd4p (Lane 3), or both (Lane 4) were added to the Ub-V-GFP ubiquitination system in vitro, which contains E1, E2, ubiquitin, ATP and Ub-V-GFP (Lane 1). (c) Ufd4p catalyses K29-linked polyubiquitin chain assembly on Ub-V-GFP. Ufd4p-mediated polyubiquitination of a Ub-V-GFP mutant with only K29 and K48 in its ubiquitin region (Lanes 4–5). The reaction products were indistinguishable from the products of the K29-only Ub-V-GFP mutant (Lanes 1-3) in the presence of wild-type, K29R- or K29-only ubiquitin. (d) The linkage specificity of Ufd2p-mediated ubiquitination was examined using a single-round ubiquitin turnover assay. K29R ubiquitin was charged onto the active cysteine of Ubc4p, and the reaction was quenched by adding NEM/EDTA. Wild-type ubiquitin or the K6-, K11-, K27-, K29-, K33-, K48-, or K63-only ubiquitin mutants (Lanes 2–9) was added in excess (10 fold of K29R ubiquitin) to the mixtures, together with Ufd2p, allowing transfer of the charged K29R ubiquitin to either wild-type ubiquitin or its variants to form free di-ubiquitin chains (Lanes 2 and 8). (e) K29 and K48 on Ub-V-GFP are necessary and required for full ubiquitination by Ufd4p and Ufd2p. K48-only Ub-V-GFP, wild-type Ub-V-GFP, K29-only Ub-V-GFP, and K29/48-only Ub-V-GFP were tested for their ability to be ubiquitinated by the Ufd4p-Ufd2p ubiquitination system. See also Supplementary Figs 1–3.

Ufd2p catalyses ubiquitination independent of Ufd4p. Because both Ufd4p and Ufd2p have E3 ligase activity, we asked how Ufd2p cooperates with Ufd4p to extend oligo-ubiquitin chains. Conceptually, the two enzymes can work either cooperatively or sequentially. In a cooperative model, the two proteins may form a complex together with the E2 enzyme Ubc4p. Alternatively, a substrate that has been ubiquitinated by Ufd4p may engage Ufd2p independently of Ufd4p, allowing addition of K48-linked ubiquitin conjugates to the substrate. To distinguish between these possibilities, we developed a stepwise ubiquitination assay (Fig. 2a–c). First, we incubated Ub-V-GFP with E1, Ubc4p, Ufd4p, ubiquitin and ATP to generate a short ubiquitin chain on Ub-V-GFP (Fig. 2b, Lane7). The reaction was then quenched by NEM/EDTA, which modifies the catalytic cysteine of Ufd4p and Ubc4p, inactivating and eliminating them from subsequent reactions. We then purified Ub-V-GFP and its modified forms using immunoprecipitation (IP) with an anti-GFP antibody. Immunoprecipitated materials were then incubated with E1, Ubc4p, Ufd2p, ubiquitin and ATP. The reaction was analysed by immunoblotting with an anti-GFP antibody. The results showed that under this experimental
condition, Ufd2p transferred one ubiquitin to Ub-V-GFP with a low-efficiency in the absence of Ufd4p (Fig. 2b, Lanes 3, 6). However, once Ub-V-GFP was modified with a ubiquitin moiety by Ufd4p, most mono-ubiquitinated Ub-V-GFP (Ub2-GFP) could be switched to Ub3-GFP by Ufd2p (Fig. 2b, Lane 9 and Supplementary Fig. 6a). Because immunoblotting showed that trace amounts of Ufd4p were still present in the second reaction, to exclude the possibility that inactive Ufd4p might influence Ufd2p activity by a dominant-negative mode, we added extra Ufd4p, which was inactivated by NEM/EDTA to the second

### Figure Legend

- **a**
  - **Step I:** Ufd4p + E1 + E2 + Ub → NEM/EDTA → IP + Ufd2p
  - **Step II:** Ufd4p + E1 + E2 + Ub + Ub-V-GFP

- **b**
  - IP: GFP
  - IB: GFP
  - IP: Xpress

- **c**
  - K29R Ub
  - K29R-Wt di-Ub
  - K29/48R Ub
  - K29/48R-Wt tri-Ub

- **d**
  - IP: GFP
  - IB: GFP

- **e**
  - IP: GFP
  - IB: GFP
reaction. We found that the presence of inactive Ufd4p did not affect the efficiency of Ufd2p (Supplementary Fig. 4a). Moreover, addition of inactivated Ufd4p to the Ufd2p-mediated Ub-V-GFP ubiquitination system affected neither Ub-V-GFP mono-ubiquitination nor free ubiquitin chain formation (Supplementary Fig. 4b). The efficiency of Ub3-GFP production was also not affected by the addition of fully active Ufd4p in the second reaction or when K29R ubiquitin was used in the second reaction (Supplementary Fig. 4c). To further demonstrate that the stepwise reaction faithfully recapitulates the physiological condition, we used Ufd4p together with FLAG-K29R Ub to synthesize Ub2-GFP. Ub2-GFP was then purified via immunoprecipitation with anti-FLAG M2 Affinity Gel, and eluted by FLAG peptides under the native conditions (Supplementary Fig. 4d). The purified Ub2-GFP was then used as the substrate to test the ubiquitin transfer efficiency of Ufd2p in the presence or absence of the Ufd4p condition. As shown in Supplementary Fig. 4e, the presence of Ufd4p in the second reaction could not enhance the production of Ub3-GFP, and the ubiquitin transfer efficiency of Ufd2p in the stepwise reaction was almost the same as the normal in vitro ubiquitination condition (Supplementary Fig. 4f). Moreover, further incubation of ubiquitinated Ub-V-GFP after the second reaction with Ufd4p, E1, E2, ubiquitin and ATP did not result in further addition of ubiquitin conjugates (Supplementary Fig. 4g, h). This suggests that Ufd4p cannot act on substrates modified by Ufd2p. Altogether, these results suggest that Ufd2p and Ufd4p function in sequence to assemble ubiquitin chains on Ub-V-GFP, with Ufd2p acting first to prime a substrate with a K29-linked ubiquitin moiety. Ufd2p subsequently works independently to add additional ubiquitin moieties.

Ufd2p and Ufd4p synthesize branched ubiquitin chains. To dissect the topology of the ubiquitin chains synthesized by Ufd4p and Ufd2p, we used the K29R ubiquitin mutant in step I (Fig. 2a). This allowed us to specifically assemble K29-linked Ub2-GFP (Fig. 2c, Lane1). After purification, this Ub2-GFP species was used as the substrate for Ufd2p in the second step. We found that Ufd2p could efficiently add one ubiquitin to Ub2-GFP to form Ub3-GFP (Fig. 2c, Lanes 2, 4, 5), regardless of whether WT ubiquitin, the K48R mutant (K48R Ub) or methylated ubiquitin (Met-Ub) was used in the second reaction. Since ubiquitin could be added to either the distal or proximal ubiquitin on Ub-V-GFP, to distinguish between these possibilities, we replaced the WT ubiquitin with the K29/48R ubiquitin mutant in step I. Because Ufd2p preferentially added ubiquitin to the K48 of ubiquitin, mutant ubiquitin should block chain extension by Ufd2p (if Ufd2p only extend the chain from the distal end). However, this substitution did not affect the addition of ubiquitin to Ub2-GFP by Ufd2p (Fig. 2c, Lanes 8–10 compared with Lanes 2, 4, 5), suggesting that Ufd2p does not extend the oligo-ubiquitin chain formed by Ufd4p from the distal end. Intriguingly, even when WT Ub was used in the first step, only a single ubiquitin moiety was transferred onto Ub2-GFP (Fig. 2c, Lanes 13–15). Thus, it appears that Ufd2p may preferentially add ubiquitin to the proximal ubiquitin of Ub2-GFP, resulting in the formation of branched ubiquitin chains (Supplementary Fig. 5a, b).

Next, we examined whether the proximal end preference is applicable to a longer ubiquitin chain on Ub-V-GFP. To this end, we synthesized and purified K29-linked di- or tri-ubiquitin chains (Supplementary Fig. 5) via Ufd4p-mediated polyubiquitination (see Methods)10. If di-ubiquitin carrying a Ub2-K9R mutant at the distal end was used in step I to synthesize Ub3-GFP (Fig. 2d, Lane 1), Ufd2p transferred only two ubiquitin moieties to Ub3-GFP regardless of whether WT Ub, K48R Ub or Met-Ub was used in step II (Fig. 2d, Lanes 3–5; Supplementary Fig. 5c). Since a di-ubiquitin bearing a K29R and K48R double mutant at the distal site yielded a similar result (Fig. 2d, Lanes 13–15), the distal ubiquitin is apparently dispensable for Ufd2p-mediated ubiquitination. In contrast, when K48 of the proximal ubiquitin was changed to arginine, only a single ubiquitin moiety could be transferred to Ub3-GFP regardless of whether the distal end ubiquitin included K48 (Fig. 2d, Lanes 8–10 and 18–20; Supplementary Fig. 5d). In this case, Ufd2p presumably only transfers ubiquitin to the middle ubiquitin moiety in Ub3-GFP. Likewise, when the K29R-WT-WT tri-ubiquitin was used in step I to assemble Ub4-GFP, Ufd2p transferred three ubiquitins onto Ub4-GFP, even when methylated ubiquitin was used (Fig. 2e, Lanes 13–15; Supplementary Fig. 5e). These results further support the conclusion that Ufd2p preferentially adds ubiquitin conjugates to proximal ubiquitins in a K29-linked ubiquitin chain, except for the distal one. We next expressed and purified all the other HECT domain E3s in yeast and tested their activities in the Ub-V-GFP ubiquitination reaction. We found that Hul4p could not catalyse the ubiquitination of Ub-V-GFP. However, Rsp5p, Tom1p CT (a C-terminal truncation of Tom1p) and Hul5p could synthesize Ub2-GFP (Supplementary Fig. 6a–d). Among them, Hul5p synthesized both K29 and K48-linked Ub2-GFP (Supplementary Fig. 6e). Strikingly, Ufd2p could only add one ubiquitin to the K29-linked Ub2-GFP generated by Hul5p, but ignored the K48-linked Ub2-GFP that was present in the same reaction (Supplementary Fig. 6e). This observation further supports our conclusion that Ufd2p preferentially initiates branched chains from special linkages such as K29-linked ubiquitin conjugates.

It is noteworthy that, when we extended the reaction time to 2 h, we detected a small amount of high-molecular weight species of Ub-n-GFP if WT Ub was used in step II. These high molecular weight species were reduced when either K48R Ub or Met-Ub was used in the second reaction (Fig. 2f, Lane 3 compared with Lanes 4 and 5). These results suggest that Ufd2p can...
form K48-linked ubiquitin chains, but only with low efficiency. By contrast, the efficiency of Ufd2p-mediated multi-monoubiquitination was more readily detected (Fig. 2f, Supplementary Fig. 7a). Therefore, although we cannot completely exclude the possibility that Ufd2p might assemble some K48-linked polyubiquitin chains in cells, we still favour a model in which Ufd2p mainly synthesizes multi-monoubiquitin chains on K29-linked polyubiquitin chains. Based on these observations, we propose that Ufd2p is a chain branching enzyme rather than a chain elongation factor; it forms branched chains by preferentially adding a ubiquitin moiety to K48 non-distal ubiquitin molecules in a K29-linked oligo-ubiquitin chain synthesized by Ufd4p (Fig. 2g).

In vivo branched ubiquitin chains assembly mediated by Ufd2p.

To determine whether Ufd2p and Ufd4p can form branched ubiquitin chains on Ub-V-GFP in vivo, we used a method developed by Meyer and Rapoport. Briefly, a Tobacco Etch Virus protease (TEV) cleavage site was introduced into ubiquitin after Gly53, or after Glu64 in combination with a FLAG-epitope (hereafter referred to as Ub-53TEV and Ub-64TEV/FLAG, respectively). Both ubiquitin variants could still be conjugated to Ub-V-GFP by Ufd4p and Ufd2p (Fig. 3b). After TEV digestion, a FLAG-epitope-containing peptide would remain attached to Ub-V-GFP if a single ubiquitin or a ubiquitin chain was conjugated to Ub-V-GFP. In contrast, if two ubiquitins were conjugated to Ub-V-GFP at the same time as in the case of branched chain formation, two FLAG peptide remnants would remain on Ub-V-GFP after TEV digestion (Fig. 3a). This method was first validated by Ufd4p and Ufd2p mediated in vitro ubiquitination: after TEV digestion, Ub-V-GFP containing one or two FLAG peptide remnants could be detected on Ub-V-GFP (Fig. 3b, Lanes 14, 16 and 18). If Ufd2p was omitted from the reaction, only cleaved Ub-V-GFP containing a single FLAG peptide was produced (Fig. 3b, Lanes 5, 7 and 9). These results suggested that this method is suitable for detecting branched ubiquitin chains on Ub-V-GFP.

We then co-expressed FLAG-UB-53TEV and Myc-Ub-64TEV/FLAG with His-tagged Ub-V-GFP in yeast cells. Ub-V-GFP modified by these engineered ubiquitin variants was significantly enriched after treatment with the proteasome inhibitor MG132, suggesting that Ub-V-GFP modified by these ubiquitin variants is still targeted to the proteasome for degradation (Fig. 3c, Lanes 1, 5, 9 versus Lanes 3, 7, 11). We then purified Ub-V-GFP and the modified species from these cells under denaturing conditions using Ni-NTA agarose. After TEV digestion, Ub-V-GFP conjugated with two FLAG-containing peptides could be detected using either anti-FLAG and anti GFP antibodies (Fig. 3c, Lanes 4, 8 and 12), suggesting the presence of branched ubiquitin chains on Ub-V-GFP in vivo. In a ufd2Δ strain, ubiquitinated Ub-V-GFP was reduced compared with a wild-type strain, and the level of Ub-V-GFP conjugated with two FLAG-containing peptides decreased significantly. This suggested that the formation of branched ubiquitin chains on Ub-V-GFP is dependent on Ufd2p in yeast cells (Fig. 3d). In addition, we found that the presence of branched ubiquitin chains on Ub-V-GFP was dependent on both K29 and K48 of ubiquitin in the Ub-V-GFP molecule because mutating either of these residues to arginine almost abolished the production of branched ubiquitin chains on Ub-V-GFP (Fig. 3e). When Ub-K29R-V-GFP was used, Ub-V-GFP bearing only a single FLAG peptide could be detected, and when Ub-K48R-V-GFP was used, a small amount of Ub-V-GFP bearing two FLAG peptides was detected, which may result from the conjugation of a ubiquitin to a lysine in Ub-V-GFP other than K29 or K48. Taken together, these results suggest that Ufd2p can synthesize branched ubiquitin chains on Ub-V-GFP in yeast cells.

Ufd2p binds K29-linked Ubn-GFP via its two N-terminal loops.

Although Ufd2p itself could transfer ubiquitin to both Ub-V-GFP and Ubn-GFP (Fig. 2), the ubiquitination efficiency was much lower with Ub-V-GFP than with Ub2-V-GFP as the substrate (Supplementary Fig. 7a). These results suggested that Ufd2p prefers K29-linked ubiquitin chains as substrate. This substrate preference might arise from a specific interaction with substrates bearing a K29-linked ubiquitin chain. To test that possibility, we labelled Ufd2p with biotin and used the labelled proteins to pull down Ub-V-GFP and Ubn-GFP. We found that Ufd2p interacted with both Ub-V-GFP and Ubn-GFP (Fig. 4a, Lanes 3 and 6), but it bound substrates bearing a K29-linked chain with much higher affinity (Fig. 4a,b). According to surface plasmon resonance experiments, Ufd2p bound the K29-linked Ub2-GFP with a KD of 0.29 μM, whereas the KD was 2.03 μM for Ub-V-GFP (Fig. 4f and Supplementary Table 1, Supplementary Fig. 7b and Supplementary Table 1). These results suggest an interaction between Ufd2p and the K29-linked ubiquitin chain.

Next, we defined the domain responsible for the higher affinity interaction of Ufd2p with substrates conjugated to K29-linked ubiquitin chains. To map this domain, we generated and purified a series of Ufd2p truncation mutants, as shown in Fig. 4c. GST pull-down experiments showed that deletion of the N-terminal region prevented Ufd2p from binding to Ubn-GFP (Fig. 4d, upper panel, Lanes 2–4; Supplementary Fig. 8a). Likewise, deleting either amino acids 55–87 or 75–87 of Ufd2p completely abolished its interaction with Ubn-GFP (Fig. 4d, upper panel, Lanes 11 and 12). In contrast, deleting the core region of Ufd2p (amino acids 111–879) did not affect the interaction with Ubn-GFP (Fig. 4d, upper panel, Lane 10). Thus, the substrate binding site is present in the N-terminal domain of Ufd2p.

The amino terminal region (amino acids 35–111) of Ufd2p contains three α-helices. Further mapping experiments showed that while deleting the middle α-helix, 55–75aa, did not influence the interaction of Ufd2p with Ubn-GFP (Fig. 4d, Lane 13), deleting the 35–55aa or 75–111aa helices did reduce the interaction (Fig. 4d, Lanes 8 and 9). These results suggest that the Ubn-GFP binding activity of Ufd2p is dependent on these two regions. Further mutagenesis analyses (Supplementary Fig. 8c–f) showed that two loops in these regions (51–55aa and 105–110aa) were critical for substrate binding (Supplementary Fig. 8g); when residues in these two loops were mutated to alanine (Ufd2p NLM), the interaction between Ufd2p and Ubn-GFP was almost completely abolished (Fig. 4e). Surface plasmon resonance experiments confirmed that the affinity of Ufd2p NLM to K29-linked Ub2-GFP was lower than for WT Ufd2p, but there was no significant difference between Ufd2p or Ufd2p NLM binding to Ub-V-GFP (Fig. 4f–h, Supplementary Fig. 7b–c and Supplementary Table 1), suggesting that the two N-terminal loops of Ufd2p are involved in recognition of K29-linked Ub2-GFP. Importantly, this mutant also failed to form branched ubiquitin chains on Ufd4p synthesized Ubn-GFP (Fig. 4h, Supplementary Fig. 9a), although it had similar ubiquitin ligase activity as wild-type Ufd2p (Supplementary Fig. 9b). The defects of the NLM mutant in branched chain formation could not be attributed to either protein misfolding or reduced stability because: (1) Circular Dichroism (CD) spectral analysis showed that the secondary structure of Ufd2p NLM was similar to that of WT Ufd2p (Supplementary Fig. 10a); and (2) a cycloheximide chase experiment indicated that the degradation rate of Ufd2p NLM was similar to that of Ufd2p or Ufd2p ΔU-box.
Figure 3 | Branched ubiquitin chains are detected on UFD substrates in cells. (a) Schematic representation of the method used to monitor the synthesis of branched ubiquitin chains on Ub-V-GFP. (b) In vitro validation of the method used to monitor branched ubiquitin chains on Ub-V-GFP. As illustrated in (a), Ub-V-GFP was ubiquitinated by Ufd4p or Ufd4p-Ufd2p. Reaction products were treated with the TEV enzyme and analysed by immunoblotting with the anti-GFP antibody. (c) Ub-V-GFP was modified with branched ubiquitin chains in yeast cells. Ub-V-GFP and its modified forms were expressed in a wild-type strain expressing FLAG-Ub53TEV and Myc-Ub64TEV/FLAG and purified under denaturing conditions in the presence or absence of the proteasome inhibitor MG132. The pull-down products were treated with the TEV enzyme and analysed by immunoblotting with anti-FLAG and anti-GFP antibodies. (d) The addition of branched ubiquitin chains to Ub-V-GFP is mainly dependent on UFD2 in vivo. Ub-V-GFP and its modified forms were purified from either a wild-type or ufd2Δ strain expressing both FLAG-Ub53TEV and Myc-Ub64TEV/FLAG and purified under denaturing conditions in the presence of MG132 under denaturing conditions. The pull-down products were treated with the TEV enzyme and analysed by immunoblotting with the anti-FLAG and anti-GFP antibodies. (e) K29 and K48 on Ub-V-GFP are required for the formation of branched ubiquitin chains in yeast cells. Wild-type, K29R, K48R or K29/48R Ub-V-GFP and their modified forms were purified from a wild-type strain expressing FLAG-Ub53TEV and Myc-Ub64TEV/FLAG. The pull-down products were treated with the TEV enzyme and analysed by immunoblotting with the anti-FLAG and anti-GFP antibodies.
Figure 4 | Ufd2p binds Ubn-GFP via two N-terminal loops. (a) Interactions between the substrate and Ufd2p. Biotinylated Ufd2p was immobilized on streptavidin-coated beads, which were then incubated with GFP, Ub-V-GFP or Ubn-GFP that was synthesized by Ufd4p. The precipitated proteins were analysed by immunoblotting with the indicated antibodies. (b) Quantification of the relative binding affinity between Ufd2p and Ubn-GFP. Relative amounts of Ub1–4-GFP were quantified using Odyssey software and compared with the input. Error bars denote the s.e.m. of three independent replicates. * indicated $P<0.05$, ** indicated $P<0.01$. (c) A summary of the Ufd2p variants used to map the Ubn-GFP binding motif, which include Ufd2p (amino acids 1–961), Ufd2p CF1 (amino acids 519–961), Ufd2p CF2 (amino acids 760–961), Ufd2p CF3 (amino acids 879–961), Ufd2p MFD1 (amino acids 35–55), Ufd2p MFD2 (amino acids 35–55), Ufd2p MFD3 (amino acids 75–111), Ufd2p MFD4 (amino acids 75–111), and Ufd2p NHD1 and Ufd2p NHD2. (d) Amino acids 35–55 and 75–111 of Ufd2p are necessary to bind to Ubn-GFP. GST-Ufd2p and its variants were purified and used to pull down Ubn-GFP, which was synthesized by Ufd4p; GST was used as a control. Asterisks in the bottom panel indicate GST products cleaved from the fused proteins. (e) Ufd2p binds to Ubn-GFP via its two N-terminal loops. GST-Ufd2p and GST-Ufd2p NLM (E51A K52A L53A D54A K55A E105A N106A M109A N110A) were purified and used to pull down Ubn-GFP, which was synthesized by Ufd4p; GST was used as a control. Asterisks in the bottom panel indicate GST products cleaved from the fused proteins. (f) Interactions between the substrate and Ufd2p. Biotinylated Ufd2p was immobilized on streptavidin-coated beads, which were then incubated with GFP, Ub-V-GFP or Ubn-GFP that was synthesized by Ufd4p. The precipitated proteins were analysed by immunoblotting with the indicated antibodies. (b) Quantification of the relative binding affinity between Ufd2p and Ubn-GFP. Relative amounts of Ub1–4-GFP were quantified using Odyssey software and compared with the input. Error bars denote the s.e.m. of three independent replicates. * indicated $P<0.05$, ** indicated $P<0.01$. (c) A summary of the Ufd2p variants used to map the Ubn-GFP binding motif, which include Ufd2p (amino acids 1–961), Ufd2p CF1 (amino acids 519–961), Ufd2p CF2 (amino acids 760–961), Ufd2p CF3 (amino acids 879–961), Ufd2p MFD1 (amino acids 35–55), Ufd2p MFD2 (amino acids 35–55), Ufd2p MFD3 (amino acids 75–111), Ufd2p MFD4 (amino acids 75–111), and Ufd2p NHD1 and Ufd2p NHD2. (d) Amino acids 35–55 and 75–111 of Ufd2p are necessary to bind to Ubn-GFP. GST-Ufd2p and its variants were purified and used to pull down Ubn-GFP, which was synthesized by Ufd4p; GST was used as a control. Asterisks in the bottom panel indicate GST products cleaved from the fused proteins. (e) Ufd2p binds to Ubn-GFP via its two N-terminal loops. GST-Ufd2p and GST-Ufd2p NLM (E51A K52A L53A D54A K55A E105A N106A M109A N110A) were purified and used to pull down Ubn-GFP, which was synthesized by Ufd4p; GST was used as a control. Asterisks in the bottom panel indicate GST products cleaved from the fused proteins. (f) SPR sensorgrams for the binding of GST-Ufd2p to K29-linked Ub2-GFP. A series of two-fold GST-Ufd2p dilutions was applied to the K29-linked Ub2-GFP surface (GFP served as a control). (g) SPR sensorgrams for the binding of GST-Ufd2p NLM to K29-linked Ub2-GFP. (h) The interaction between Ufd2p and Ubn-GFP is necessary for the E4 activity of Ufd2p. Ufd2p or Ufd2p NLM was incubated with E1, E2, Ufd4p, ATP, ubiquitin and Ub-V-GFP and detected using the anti-GFP antibody. See also Supplementary Figs 7–11.
Branched ubiquitin chains promote substrate degradation.

Next, we asked why Ufd2p is required for the degradation of a substrate modified by K29-linked ubiquitin chains. It is well established that K11- and K48-linked ubiquitin chains can target substrates to the proteasome for degradation\(^1\),\(^4\), and substrates conjugated with branched ubiquitin chains containing only K48 linkage can also be recognized and escorted to the proteasome\(^4\). It has been reported that Rad23p, Dsk2p, and Rpn10p can recognize K48- and K63-linked di-ubiquitin. The diversity of ubiquitin code information is achieved via distinct forms of ubiquitination, such as monoubiquitination, distal forms of polyubiquitination, chain-branching activity, and ubiquitin chain linkage switches\(^2\),\(^8\),\(^10\),\(^12\). However, these ubiquitin signals are not all recognized equally by the proteasome receptors.

We concluded that the branched ubiquitin chain forming activity of Ufd2p is essential for the UFD degradation pathway, acting as a ubiquitin chain linkage switcher to alter a non-optimized degradation signal to a signal preferred by the ubiquitin receptors of the proteasome.

Discussion

The function of branched chains in ERAD and OLE pathways.

Ufd2p is known to be involved in ERAD (refs 32,34) and in degradation of the short-lived HMG-CoA reductase, Hmg2p (refs 32,48). Indeed, we observed strong stabilization of Hmg2p in the ufd2Δ strain (Fig. 6a,c). Wild-type UFD2 fully restored the degradation of Hmg2p, but neither the UFD2\(^{NLM}\) mutant nor the UFD2\(^{NLM}\) mutant could do so (Fig. 6a,c). These results suggest that ubiquitin chain branching is essential for degradation of a physiological substrate in yeast.

Ufd2p also plays an important role in the OLE pathway in budding yeast by regulating the stability of Spt23p, a transcription factor that controls the synthesis of unsaturated fatty acids\(^22\). We therefore examined the degradation of Spt23p in a UFD2\(^{-}\)deletion strain and found that the Spt23 p90 protein was significantly stabilized. Wild-type UFD2, but not the UFD2\(^{NLM}\) mutant could fully restore Spt23p degradation in the ufd2Δ strain (Fig. 6b,d). Because one of the key target genes of Spt23p is OLE1, which regulates the synthesis of unsaturated fatty acids such as oleic acid, deregulation of this pathway is toxic to cells\(^49\),\(^50\). As a result, ufd2Δ mutants are sensitive to oleic acid\(^32\). Over-expression of wild-type UFD2 made the ufd2Δ mutant less sensitive to oleic acid, but both UFD2\(^{NLM}\) failed to do so (Fig. 6e). In addition, we found that these two mutants also failed to rescue the growth defect of the Alpha2 mutant under an acidic condition (Fig. 6e). Altogether, these results suggest that the N-terminal loops of Ufd2p are important in regulation of ERAD, the OLE pathway and cell growth under acidic conditions.

Discussion

The diversity of ubiquitin code information is achieved via distinct forms of ubiquitination, such as monoubiquitination, multi-monoubiquitination and polyubiquitination via different linkages\(^2\),\(^8\),\(^10\),\(^12\). However, these ubiquitin signals are not necessarily static and can undergo further modification to alter the fate of substrate. For example, ubiquitin chains attached to a substrate can be removed by a deubiquitinase and revert to the fate conferred by the original ubiquitin modification\(^51\).
Figure 5 | Substrates modified with branched ubiquitin chains are recognized by Rad23p, Dsk2p and Rpn10p. (a) Rad23p prefers to bind to branched ubiquitin chains on Ub-V-GFP over Ub-V-GFP with K29-linked ubiquitin chains or unmodified Ub-V-GFP. Ub-V-GFP, Ub-V-GFP with K29-linked ubiquitin chains (synthesized via Ufd4p) or Ub-V-GFP with branched ubiquitin chains (synthesized via Udf4p-Ufd2p) immobilized on protein A-sepharose beads were used to immunoprecipitate GST-Rad23p. The precipitated proteins were analysed by immunoblotting with the indicated antibodies. The asterisks indicate GST products cleaved from GST-Rad23p. (b) SPR sensorgrams for the binding of GST-Rad23p to Ub-V-GFP. A series of two-fold GST-Rad23p dilutions was applied to the Ub-V-GFP surface (GFP served as a control). (c) SPR sensorgrams for the binding of GST-Rad23p to K29-linked Ub2-GFP. (d) SPR sensorgrams for the binding of GST-Rad23p to branched Ub3-GFP. (e) Proteasome adaptor proteins such as Rad23p fail to recognize Ufd2p NLM-mediated ubiquitination products. Either wild-type Ufd2p-mediated ubiquitination products or Ufd2p NLM-mediated ubiquitination products immobilized on protein A-sepharose beads were used to immunoprecipitate GST-Rad23p. Ubn-GFP modified by Ufd2p NLM could not immunoprecipitate GST-Rad23p efficiently. Asterisks indicate nonspecific background signals from the IgG chains. (f) The ubiquitin chain linkage switch is necessary for substrate degradation. Protein expression in wild-type or ufd2Δ strains expressing Ub-V-GFP under the control of a galactose-induced promoter was stopped upon transfer to 2% glucose. The ufd2Δ strains harboured empty vector, UFD2 or UFD2ΔNLM under the control of a galactose-induced promoter. The degradation of Ub-V-GFP over time was analysed by immunoblotting. Pgtk1p served as a loading control. (g) Quantification of relative Ub-V-GFP levels in (f) using Odyssey software. Error bars denote the s.e.m. of three independent replicates. See also Supplementary Figs 12–14.
In addition, a ubiquitin conjugate can be re-modelled or ‘edited’ to execute a distinct function. Ubiquitin chain editing occurs when one type of chain is replaced by another\(^8\). A20, which contains an N-terminal deubiquitinating Ovarian Tumour (OTU) domain and a C-terminal RING domain, was proposed to be a chain-editing factor. A20 down-regulates NF-\(\kappa\)B signalling through its OTU domain-mediated removal of K63-linked ubiquitin chains from the receptor interacting protein (RIP),

**Table 2** | SPR parameters of GST-Rad23p binding to Ub-V-GFP, K29-linked Ub2-GFP and branched Ub3-GFP.

| Ka (1/Ms) | Kd (1/s) | KA (1/M) | KD (\(\mu\)M) |
|-----------|----------|----------|--------------|
| Rad23p::Ub-V-GFP | 4.88e3 | 4.48e-3 | 1.09e6 | 0.92 |
| Rad23p::Ub2-GFP | 3.80e3 | 3.36e-3 | 1.01e6 | 1.00 |
| Rad23p::Ub3-GFP | 255 | 6.06e-5 | 4.22e6 | 0.24 |

Figure 6 | The branched chain forming activity of Ufd2p is required for ERAD, the OLE pathway and acid resistance in yeast. (a) The Ub-GFP binding motif of Ufd2p is essential to the degradation of a subset of proteins during ERAD. Protein expression in wild-type and mutant strains (\(ufd2\)\(^D\) strain contains empty vector, \(UFD2\), \(UFD2\)\(^{\Delta\text{U-box}}\) or \(UFD2\)\(^{\text{NLM}}\) under the control of a galactose-induced promoter) expressing Myc-Hmg2p under the control of a galactose-induced promoter was stopped by transfer to 2% glucose and addition of cycloheximide. The degradation of Myc-Hmg2p over time was analysed by immunoblotting. Pgk1p served as a loading control. (b) The Ub-GFP binding motif of Ufd2p affects the stability of Spt23p in the OLE pathway. The degradation and detection of Myc-Spt23p expression under the control of a galactose-induced promoter were analysed as described above. (c) Quantification of Myc-Hmg2p turnover. The relative amounts of Myc-Hmg2p were quantified using Odyssey software; the high molecular weight smear representing the ubiquitylated species was considered when determining the turnover rate. Error bars denote the s.e.m. of three independent replicates. (d) Quantification of Myc-Spt23p p90 turnover. Error bars denote the s.e.m. of three independent replicates. (e) The Ub-GFP binding motif of Ufd2p is essential to the OLE pathway and acidic resistance: 10-fold serial dilutions of the indicated strains (the wild-type strain contains empty vector; \(ufd2\)\(^D\) strains contain empty vector, \(UFD2\), \(UFD2\)\(^{\Delta\text{U-box}}\) or \(UFD2\)\(^{\text{NLM}}\) under the control of the \(UFD2\) promoter) were plated on media supplemented with either 0.02% oleic acid or 100 mM MES (pH 1.0) and grown at 30\(^\circ\)C for 2 days.
Armadillo-like repeats with two pronounced helical hairpin for the chain branching activity of Ufd2p. The crystal structure of Ufd2p contribute to the substrate binding site, which is essential. Experiments show that the two loops in the N-terminal region of bearing a K29-linked ubiquitin chain. The nature of substrate Ufd4p act in sequence and Ufd2p is a chain branching factor synthesized K29-linked ubiquitin chains, resulting in branched ubiquitin conjugates to proximal ubiquitin moieties on Ufd4p-ubiquitinated substrates by the proteasome.

Ufd2p was initially proposed to be a chain elongating factor because it can add ubiquitin conjugates to a substrate primed by another ligase. However, the mechanism of Ufd2p-mediated ubiquitination had remained unclear. Previous studies have shown that Ufd2p can cooperate with an E2 enzyme in a manner similar to a RING ubiquitin ligase. Addition of additional ubiquitin conjugates in the form of branched ubiquitin chains by Ufd2p leads to more efficient degradation of the modified substrates by the proteasome.

Ufd2p was initially proposed to be a chain elongating factor because it can add ubiquitin conjugates to a substrate primed by another ligase. However, the mechanism of Ufd2p-mediated ubiquitination had remained unclear. Previous studies have shown that Ufd2p can cooperate with an E2 enzyme in a manner similar to a RING ubiquitin ligase, but the question remained as to how it collaborates with Ufd4p to assemble a productive degradation signal on diverse substrates. We previously proposed two possible models for Ufd2p- and Ufd4p-mediated ubiquitination: the sequential model and the cooperative model. Here, we established a stepwise ubiquitination assay to distinguish between these two possibilities. We found that Ufd2p can independently add ubiquitin moieties to a UFD substrate as long as it carries a K29-linked ubiquitin chain formed by Ufd4p (Fig. 2b). We further demonstrate that Ufd2p preferentially added ubiquitin conjugates to proximal ubiquitin moieties on Ufd4p-synthesized K29-linked ubiquitin chains, resulting in branched ubiquitin chains on the substrate (Fig. 2c–g). Thus, Ufd2p and Ufd4p act in sequence and Ufd2p is a chain branching factor rather than a chain elongator.

Mechanistically, we found that Ufd2p prefers substrates bearing a K29-linked ubiquitin chain. The nature of substrate preference lies in a high affinity interaction between Ufd2p and Ubn-GFP bearing a K29-linked ubiquitin chain. Our mapping experiments show that the two loops in the N-terminal region of Ufd2p contribute to the substrate binding site, which is essential for the chain branching activity of Ufd2p. The crystal structure of Ufd2p has an elongated shape resembling a running dog: the E2 binding U-box domain forms the head and multiple irregular Armadillo-like repeats with two pronounced helical hairpin protrusions form the trunk. The newly discovered substrate binding site is located in the tail, far from the U-box, raising the question of how Ufd2p transfers ubiquitin from a cognate E2 to the substrate. One possible solution is the use of a Ufd2p dimer in the chain branching reaction. In this case, the N-terminal motif recognizes K29-linked ubiquitin chains on a substrate and the U-box domain of a second Ufd2p catalyses the chain branching reaction. However, we have not found any evidence that supports oligomerization or dimerization of Ufd2p. Importantly, a mixture of the AU-box and Ufd2p NLM mutants could not restore Ufd2p activity (Supplementary Fig. 15), thus excluding the possibility that Ufd2p functions as a dimer. Alternatively, Ufd2p may catalyse ubiquitination in a manner analogous to the Skp1–Cul1–F-box-protein (SCF) ubiquitin E3 ligase complex, which shares a similar substrate and E2 binding configuration (Supplementary Fig. 16). The Skp1–Cul1–F-box-protein (SCF) ubiquitin E3 ligase complex contains a cullin subunit as a molecular scaffold, its C-terminus interacts with a RING-finger protein, and the N-terminus interacts with the crucial substrate binding adaptor Skp1 (S-phase-kinase-associated protein-1), which in turn binds the F-box-protein and substrate. Although the substrate-binding F-box protein is distant from the RING domain, ubiquitin charged on E2 can be efficiently transferred to the substrate. This is thought to be achieved through conformational rearrangements modulated by a post-translational modification of the complex. In this regard, Ufd2p may similarly undergo a conformational change during the reaction to allow efficient transfer of ubiquitin to substrate.

Although branched ubiquitin chains have been detected both in vivo and in vitro, the function of branched ubiquitin chains is still poorly understood. The anaphase-promoting complex (APC/C) can synthesize branched ubiquitin chains that contain both K11- and K48-linkages, and chain branching appears to strongly enhance substrate recognition by the proteasome, thereby promoting the degradation of several cell-cycle regulators during early mitosis. In this study, we found that branched ubiquitin chains containing both K29 and K48 linkages participate in ERAD, the OLE pathway and acid resistance. Recently, K29-linked heterotypic ubiquitination has also been reported in mammalian cells. Although K29-linked ubiquitin chains were found to direct the disassembly of certain proteins or to target proteins for lysosomal degradation, there is no evidence that it could target substrates to the proteasome for degradation. It appears that K29-linked ubiquitin chain does not form a robust degradation signal for the proteasome. However, after being modulated into branched chains containing a K48 linkage, the modified substrate is capable of being targeted to the proteasome for degradation. Whether it is the branched ubiquitin chain or simply the K48 linkage that improves substrate degradation probability remains to be elucidated. However, given that several proteasome adaptor proteins are known to selectively recognize K48 linked ubiquitin chains, we propose that Ufd2p works as a ubiquitin chain branching ‘switch’, that transforms a non-degradable ubiquitin signal to a type that is favored by the proteasome, a paradigm that may be applicable to other non-canonically linked ubiquitin linkages (K6, K27 and K33).

Figure 7 | The functional role of Ufd2p in ubiquitination. Model of how Ufd2p functions as a ubiquitin chain-linkage switcher to promote the degradation of proteins conjugated to non-canonically linked ubiquitin chains.
Methods
Antibodies and proteins. The FLAG (1:2,000, M20080L), GST (1:2,000, M20077L), Myc (1:1,000, M20002M) and mouse GFP (1:1,000, M20004M) antibodies were purchased from Abmart (Shanghai, China), and the Xpress (1:2,000, R910-25) antibody was purchased from Invitrogen (Carlsbad, CA). The GFP (1:1,000), Ufd2 (1:1,000), and Pgdk1(15:000) polyclonal antibodies were generated in rabbits using the corresponding recombinant proteins as antigens.

The E1 enzyme, FLAG ubiquitin, K48-linked di-ubiquitin, K63-linked di-ubiquitin and methylated ubiquitin were purchased from Boston Biochem (Cambridge, MA).

Strains and plasmids. All yeast strains and plasmids used in this study are described in Supplementary Tables 2 and 3.

Protein purification. Ufd2p and its variants, Ubc4p, Ub-V-GFP and its variants, Ufd3p, Otu1p, Rad23p, Dsk2p and Rpn10p, Hul4p, Hu5p, Rsp5p and C-terminal truncation of Tom1p were expressed as N-terminal hexahistidine-tagged or Glu- tathione S-transferase (GST)-tagged fusion proteins in PET28a or pGEX-4t-1 vectors. Briefly, the plasmid was introduced into Rosetta (DE3) cells and grown in Terrific Broth at 37 °C to an optical density of 0.8. The temperature of the culture was then shifted to 16 °C, and cells were induced with 0.25 mM isopropyl-β- thiogalactoside (IPTG) for 16 h. Cells were harvested by centrifugation and resuspended in lysis buffer (20 mM Tris, pH 7.4/300 mM NaCl/10 mM imidazole/10% glycerol for hexahistidine-tagged fusion protein; 50 mM Tris, pH 7.4/300 mM NaCl/2 mM MgCl2/5% glycerol for GST-tagged fusion proteins) with 1 mM PMSF. After lysis by sonication and high-speed centrifugation of the supernatant, the supernatant was incubated with Ni Sepharose 6 Fast Flow (GE Healthcare, Marlborough, MA) or Glutathione Sepharose 4B (GE Healthcare, Marlborough, MA) for 2 h at 4 °C. The resin was washed, and the protein was eluted using the lysis buffer supplemented with 250 mM imidazole or 10 mM glutathione. Ufd4p was expressed under the control of the galactose-induced promoter in the SF10 yeast strain and purified under native conditions using Ni Sepharose 6 Fast Flow (GE Healthcare, Marlborough, MA). Ub-V-GFP, Ufd4p, Rad23p, Dsk2p, Rpn10p, Ufd3p, Otu1p and Ufd2p and its variants were further fractionated using size exclusion chromatography on a Superdex 75 or Superdex 200 column (GE Healthcare, Marlborough, MA), Ub-V-GFP, Ufd4p, Rad23p, Dsk2p, Rpn10p, Ufd3p, Otu1p and Ufd2p were refolded in PBS and dialyzed in the aforementioned protein purification buffer.

In vitro ubiquitination assay. The ubiquitination experiments were conducted as follows: E1 (60 nM), Ubc4p (200 nM), Ufd4p (300 nM), Ufd2p (200 nM) and Ub-V-GFP (400 nM) were incubated with FLAG-Ub (10 nM) at 30 °C for 1 h. The reaction was treated with 1 mM DTT and 10 mM NEM for 15 min at room temperature. The reaction products were incubated with the GFP antibody (1:2,000, R910-25) antibody was purchased from Invitrogen (Carlsbad, CA). The GFP (1:1,000), Ufd2 (1:1,000), and Pgdk1(15:000) polyclonal antibodies were generated in rabbits using the corresponding recombinant proteins as antigens. The E1 enzyme, FLAG ubiquitin, K48-linked di-ubiquitin, K63-linked di-ubiquitin and methylated ubiquitin were purchased from Boston Biochem (Cambridge, MA).

Synthesis of K29-linked ubiquitin oligomers. Ubiquitin oligomers carrying FLAG-UbK29R at the distal end were synthesized in vitro as described below. Briefly, E1(60 nM), Ubc4p (200 nM), Ufd4p (300 nM), Ub(D77) (200 nM) and FLAG-UbK29R (8 mg ml−1 each) were incubated in a buffer containing 25 mM Tris–HCl pH 7.4, 2 mM magnesium chloride, 0.1 mM DTT and an ATP regenerating system at 30 °C for 1 h. At the end of the incubation, the reaction was treated with 1 mM DTT and 1 mM EDTA at 25 °C for 20 min. Yuh1 was added to the reaction at 20 μg ml−1 and the reaction was incubated at 30 °C for 2 h. The reaction was further in activated AtrX and treated with 50 mM EDTA and 10 mM NEM for 15 min at room temperature. The reaction was treated with 1 mM DTT and the reaction was incubated at 30 °C for 2 h. The beads were extensively washed three times with the high-salt buffer, and SDS loading buffer was added. The samples were analysed by immunoblotting with the anti-GFP antibody. This reaction yields mostly di-ubiquitin modifications, with FLAG-UbK48R at the distal end. Tri-ubiquitin was synthesized using a similar method, except that various di-ubiquitin chains were used in the reaction.

Branched chain detection. To detect branched linkages on Ub-V-GFP in vitro, the Ub-V-GFP ubiquitination experiment was conducted using Ufd4p and Ufd4p-Ufd2p as described above. FLAG-Ub, FLAG-UbK34STEV and FLAG-UbK48STEV/FLAG were used in these ubiquitination experiments. The TEV enzyme was added in excess, and the reactions were incubated for 16 h at 30 °C. Reactions were resolved via 12% Tricine SDS–polyacrylamide gel electrophoresis (SDS–PAGE).

GST pull-down experiments. GST-Ufd2p binding assays. Biotin binding assays were performed as described below. Biotinylated labelling was performed using EZ-link sulfo-NHS-Biotin (Thermo Scientific, Waltham, MA) according to the manufacturer’s instructions. The Ub-V-GFP and Ubn-GFP binding experiments were conducted at 4 °C in 300 μl of a low-salt buffer (50 mM Tris, pH 7.4, 0.5 mM magnesium chloride, 2 mM magnesium chloride, 0.1% Triton). GST-Ufd2p was incubated with 5 μg of biotinylated protein for 1 h, and the reaction was harvested as described above. The precipitated complexes were analysed by immunoblotting, and the data were quantified using Odyssey software.

GST pull-down experiments. GST-Ufd2p and its variants were incubated with Glutathione Sepharose 4B (GE Healthcare, Marlborough, MA) at 4 °C for 2 h. The GST protein was used as a negative control. The GST beads were centrifuged and washed with a high-salt buffer (50 mM Tris, pH 7.4, 50 mM sodium chloride, 2.5 mM magnesium chloride, 0.1% Triton) and the beads were centrifuged and washed three times with the high-salt buffer, and SDS loading buffer was added. The samples were analysed by immunoblotting with anti-GST and rabbit anti-GFP antibodies.

Ubiquitin binding assays. Either Ub-V-GFP or Ubn-GFP, stabilized by Ufd4p or Ufd4p-Ufd2p, respectively, was coupled to protein A–sepharose beads (GE Healthcare, Marlborough, MA) with the immobilized anti-GFP antibody in a high-salt buffer at 4 °C for 2 h. After washing, GST-Rad23p, Dsk2p or Rpn10p were added, and the samples were incubated at 4 °C for 2 h. The beads were centrifuged and washed three times with the high-salt buffer, and SDS loading buffer was added. The samples were analysed by immunoblotting with anti-GST and rabbit anti-GFP antibodies.

Surface plasmon resonance (SPR) interaction analysis. Surface plasmon resonance interaction experiments were carried out at 25 °C in a Biacore 3,000 instrument (GE Healthcare, Marlborough, MA). GFP, Uv-V-GFP, K29-linked Ub2-GFP and branched Ub3-GFP were covalently attached to a carboxymethyl dextran-coated gold surface (CM5 Chip; GE Healthcare, Marlborough, MA) using an amide couple kit following the manufacturer’s suggested protocol. The reactions were activated in a solution of 10 μM 3-(3-dimethylaminopropyl) carbodiimide (EDC) and N-hydroxysuccinimide (NHS). GFP, Uv-V-GFP, K29-linked Ub2-GFP and branched Ub3-GFP in a buffer
containing 10 mM sodium acetate (pH 4.5) were injected and immobilized. Any remaining reactive sites in the flow cell were blocked with ethanolamine. Analyses were performed in a buffer containing 10 mM HEPES (pH 7.4), 150 mM NaCl, 3 mM EDTA and 0.005% (vol/vol) surfactant P20. Regeneration was achieved by injecting a buffer containing 2.5 mM NaOH. The SPR signal was normalized using a reference flow cell containing GFP.

Circular dichroism spectroscopy. To determine the secondary structure of GST-Ufd2p and GST-Ufd2p NLM, CD spectra of the proteins were collected at 0.1 mg/ml in 50 mM Tris (pH 8.0), 300 mM NaCl, 2 mM MgCl2, 10% Glycerol using a Chirascan Plus CD spectrometer (Applied Photophysics, Surrey, UK) and a 1 mm quartz sample cell equilibrated at 20 °C. Scans between 200 and 260 nm were performed at a scan rate of 50 nm per min.

Expression shutoff experiments. For the Ub-V-GFP, Spt23p and Hmg2p degradation assays, yeast strains harbouring (1) the pESC-Ub-V-GFP-His6-FLAG plasmid and the pY2C-Ufd2 plasmid or its variants; (2) the pYe2s-9 x myc Hmg2 plasmid and the pESC-Ufd2 plasmid or its variants; or (3) the pYe2s-4 x myc Hmg2 plasmid and the pESC-Ufd2 plasmid or its variants were grown in synthetic complete glucose medium (s.d., containing 0.67% (wt/vol) yeast nitrogen base without amino acids and 0.3% (wt/vol) glucose) and leucine to an OD600 of 1.0. Yeast cells were harvested and washed three times with sterile water. The harvested cells were resuspended in synthetic complete galactose medium lacking uracil and leucine and incubated for 24 h. The expression of Ub-V-GFP, Myc-Spt23p or Myc-Hmg2p was shut off by changing the medium to medium lacking uracil and leucine and incubated for 24 h. Yeast growth sensitivity representing the ubiquitinated species was also considered to determine the protein turnover rate.

Yeast growth sensitivity. Yeast strains were grown in synthetic complete medium lacking histidine (SD-His) at 30 °C to an OD600 of 1.0. The cultures were serially diluted ten times, and each dilution was spotted onto SD-His plates containing either 0.02% oleic acid or 100 mM MES (pH 5.5). The plates were incubated at 30 °C for three days.

Statistical analysis. All data are presented as the mean ± s.e.m. The statistical significance of the differences between the mean values for the different genotypes was measured by Student’s t-test with a paired, two-tailed distribution. The data were considered significant when the P value was less than 0.05 (*) or 0.01 (**).

Data availability. The authors declare that all other data supporting the findings of this study are available within the article and its Supplementary Information files or from the corresponding author on reasonable request.

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Author contributions
C.L. planned and performed most of the experiments and helped to write the manuscript. W.L. and W.X.L. performed and analysed some biochemical experiments. W.L. and Y.Y. supervised the project, designed the experiments and wrote the article.

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