

# Mixed-Linkage Ubiquitin Chains Send Mixed Messages

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## SUMMARY

Research on ubiquitin (Ub) signaling has focused primarily on homogeneously linked polyUb. Although polyUb containing different linkages within the same chain exist, their structures and signaling properties are unknown. These mixed-linkage chains could be unbranched (i.e., no more than one lysine or methionine linkage per Ub) or branched. Here, we examined the structure, dynamics, receptor selectivity, and disassembly of branched and unbranched tri-Ub containing both K48 and K63 linkages. Each linkage was virtually indistinguishable from its counterpart in homogeneously linked polyUb. Linkage-selective receptors from hHR23A and Rap80 preferentially bound to the K48 or K63 linkages in the branched trimer. Linkage-selective deubiquitinases specifically cleaved their cognate Ub-Ub linkages in mixed-linkage chains, and the 26S proteasome recognized and processed branched tri-Ub. We conclude that mixed-linkage chains retain the distinctive signaling properties of their K48 and K63 components and that these multiple signals can be recognized by multiple linkage-specific receptors. Finally, we propose a new, comprehensive notation for Ub and Ub-like polymers.

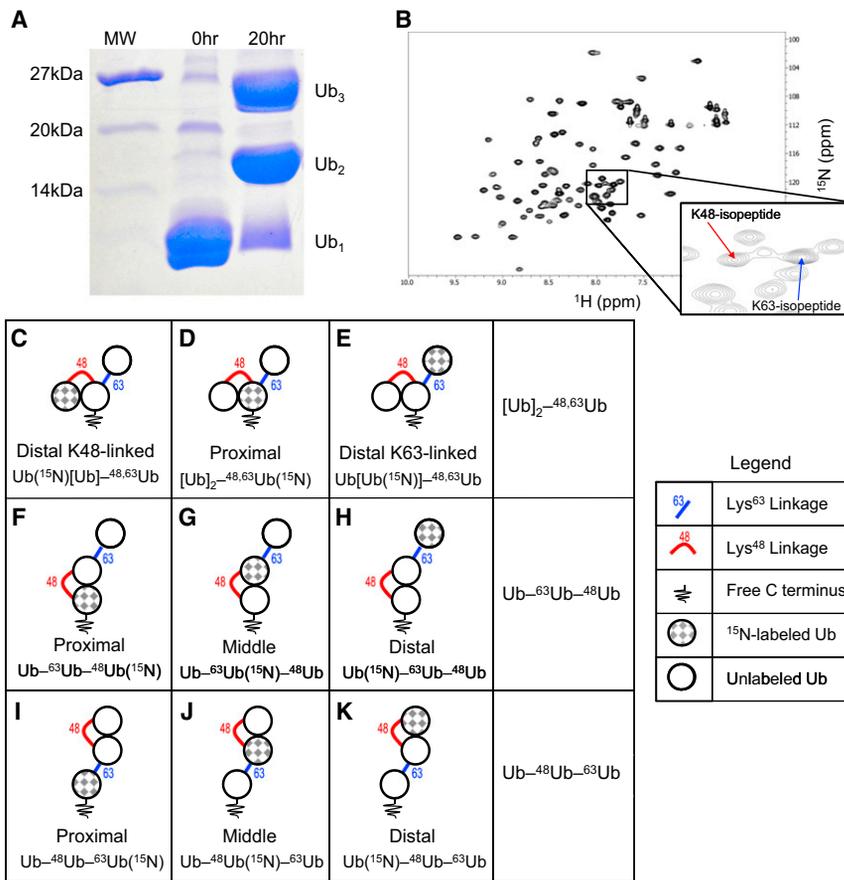
## INTRODUCTION

Critical eukaryotic cell functions such as DNA repair, cell-cycle control, protein turnover, and receptor-mediated endocytosis depend on the posttranslational modification of target proteins by covalent attachment of polyubiquitin (polyUb) (Fushman and Wilkinson, 2011; Hershko and Ciechanover, 1998). In ubiquitination, the C terminus of a “proximal” Ub typically is attached to an  $\epsilon$ NH<sub>2</sub> of a lysine on a substrate protein. Similarly, the C terminus of another Ub can be ligated to the N terminus (M1) or, more commonly, to an  $\epsilon$ NH<sub>2</sub> of K6, K11, K27, K29, K33, K48, or K63 of the previously ligated Ub (Xu et al., 2009). This process is tightly regulated in the cell, and, once a Ub is linked to a particular lysine (e.g., K48) on another Ub, it is often assumed that subsequent ubiquitination events will propagate this same linkage throughout the polyUb chain (Gregori et al., 1990). However, there is mounting evidence from both in vivo and in vitro studies

that more than one linkage type can be present in the same polyUb chain (Ben-Saadon et al., 2006; Crosas et al., 2006; Goto et al., 2010; Kim et al., 2007, 2009; Kirkpatrick et al., 2006; Kravtsova-Ivantsiv and Ciechanover, 2012; Newton et al., 2008). Currently, it is not known if linkage mixing or branching in polyUb confers signaling properties unavailable to homogeneous chains, or if the signaling properties of the individual linkages in mixed-linkage chains are preserved.

Recent mass spectrometry studies have confirmed that Ub-Ub linkages involving all seven lysines as well as M1 exist in vivo (Xu et al., 2009). Additionally, quantitative analyses have determined the relative abundances of each Ub linkage in whole-cell lysates and showed that K48 and K63 linkages occur most frequently by a large margin (Dammer et al., 2011; Ziv et al., 2011). However, these approaches generally fail to differentiate between homogeneous linkage, mixed linkage, and branched polyUb chains; therefore, it is unclear if linkages within chains are commonly mixed or branched in the cell. The use of linkage-specific anti-polyUb antibodies has revealed that polyUb is remodeled in several substrates (Dammer et al., 2011; Newton et al., 2008; Seyfried et al., 2010). Precedent for a functional mixed-linkage chain comes from Ring1b, an E3 Ub ligase that requires autoubiquitination by a mixed-linkage chain containing K6, K27, and K48 linkages (Ben-Saadon et al., 2006). In this example, our understanding of the structural and signaling properties is limited because the topology and sequence of Ub-Ub linkages in this polyUb chain are unknown. K11 and K63 mixed-linkage polyUb chains have also been suggested to serve a signaling function (Goto et al., 2010). During processing on the 26S proteasome, the substrate's polyUb signal can be remodeled by several deubiquitinases (DUBs), and, in yeast, the E3/E4 Ub ligase Hul5 has been shown to make linkages primarily through K63, but also K11 and K48 (Crosas et al., 2006). Although these and other studies indicate that mixed-linkage forms of polyUb occur in vivo, they do not provide clues about the structures of the chains or whether linkage mixing promotes specific interactions or processes. Thus, our understanding of the roles and signaling properties of mixed or branched forms of polyUb has been quite limited.

One challenge to studying mixed-linkage polyUb is that, depending on the number of individual Ub units, there could be an enormous number of unique chains (see Table S1 available online). Even discussions of the problem are confounded by the absence of a standardized nomenclature for mixed and branched polyUb. In this study, we have focused on branched and unbranched mixed-linkage chains containing K48 and K63 linkages as a logical starting point. The fact that, when in



**Figure 1. Assembly and Nomenclature of the Branched and Unbranched Mixed-Linkage Chains Used in This Study**

(A) 15% SDS-PAGE gel showing enzymatic synthesis of  $[\text{Ub}]_2^{-48,63}\text{Ub}$  in one step using E2-25K and Ubc13:Mms2 as the E2 ubiquitin-conjugating enzymes.

(B)  $^1\text{H}$ - $^{15}\text{N}$  SOFAST-HMQC spectrum of  $^{15}\text{N}$  labeled proximal Ub in  $[\text{Ub}]_2^{-48,63}\text{Ub}(^{15}\text{N})$ . The inset zooms on the region containing characteristic signals from  $\epsilon\text{NH}$  of K48 and K63 as a result of isopeptide bond formation.

(C–K) Chain schematics, nomenclature, and unit-specific  $^{15}\text{N}$ -enrichment for branched and unbranched mixed-linkage chains studied here. The rows depict isotope labeling schemes for each Ub unit in (C–E)  $[\text{Ub}]_2^{-48,63}\text{Ub}$ , (F–H)  $\text{Ub}^{-63}\text{Ub}^{-48}\text{Ub}$ , (I–K)  $\text{Ub}^{-48}\text{Ub}^{-63}\text{Ub}$ . The name in each box refers to the particular  $^{15}\text{N}$ -labeled Ub in the chain (checked gray); the formal notation indicating the  $^{15}\text{N}$  Ub is shown on the bottom of the box. See also Figures S1–S3.

homogeneous chains, the two linkages are essentially “orthogonal” with respect to their (1) location on the Ub surface, (2) conformations (i.e., compact versus extended; Varadan et al., 2002, 2004), and (3) signaling properties (i.e., proteolytic versus regulatory; Pickart and Fushman, 2004) suggests that combinations of K48 and K63 linkages could provide an extreme example of linkage mixing and branching. Additionally, their high relative abundances in the cell suggest that these linkages would predominate in randomly assembled mixed-linkage chains. Although a cellular process that requires both K48 and K63 linkages has yet to be identified, it has been reported that K48 and K63 linkages colocalize in the cell, both linkages have been detected on the same substrate, and at least one DUB, ataxin-3, preferentially cleaves mixed K48- and K63-linkage chains in vitro (Dammer et al., 2011; Newton et al., 2008; Winborn et al., 2008). These studies hint that polyUb chains containing both K48 and K63 linkages form in the cell, but whether they serve a specific function or are simply a mistake that is later “edited” remains to be seen.

For this study, we created the simplest possible model system for mixed-linkage polyUb by limiting chains to just three Ubs with combinations of K48 and K63 linkages. The resulting set of chains (all tri-Ubs) includes a single branched chain,  $[\text{Ub}]_2^{-48,63}\text{Ub}$ , and two unbranched mixed-linkage chains,  $\text{Ub}^{-63}\text{Ub}^{-48}\text{Ub}$  and  $\text{Ub}^{-48}\text{Ub}^{-63}\text{Ub}$  (Figure 1). We used established nuclear magnetic resonance (NMR) methods to examine how mixing of the linkages affects the polyUb structurally. Our

results show that linkage-specific inter-domain contacts observed with conventional, homogeneous chains are preserved in both the unbranched and branched mixed-linkage chains. To address the signaling properties of these chains, we demonstrate their recognition by K48- and K63-linkage-selective receptors, as well as by linkage-specific antibodies. We then show that linkage-selective DUBs can efficiently select and process the respective linkages in these mixed chains. Finally, we monitored the activity of purified yeast proteasomes and observed that  $[\text{Ub}]_2^{-48,63}\text{Ub}$  and homogeneous K48- and K63-linked trimers are processed similarly. We conclude that when the K48 and K63 linkages are contained in the same polyUb chain, the properties of each linkage are preserved. Hence, mixed-linkage Ub chains can send mixed messages.

## RESULTS

### Notation for Polymers of Ubiquitin or Ubiquitin-like Proteins

To facilitate discussion of different forms of polyUb conjugates, we propose a new, systematic notation. This scheme, which can be used to describe unambiguously polymers of Ub assembled from any combination of Ub–Ub linkages and also conjugates that include Ub-like proteins such as SUMO, is described in detail in the Supplemental Experimental Procedures. Fundamental to our notation is a convention whereby Ub units are connected by an *en dash* (–), and the polarity of a polyUb chain is indicated by placing the distal-end Ub unit(s) to the *left* and the proximal Ub (or the target molecule it is conjugated to) to the *right*. Specific residues (i.e., Ub Met1 or lysines that contribute an  $\epsilon$ -amine) in each linkage are indicated as superscripts. Multiple (poly)Ub moieties branching from a single Ub are indicated with brackets that, as needed, may be nested to specify the

structure of each branch. Modifications or variants of any Ub within the polymer can be indicated by parentheses that follow the Ub [e.g., Ub with a Lys63 to Arg63 substitution can be written as Ub(K63R), and completely Lys-to-Arg substituted Ub as Ub(K0)]. Thus, tri-Ub consisting of a distal-end Ub linked to K48 of a middle Ub that, in turn, is linked to K63 of a proximal Ub is written as Ub<sup>-48</sup>Ub<sup>-63</sup>Ub, whereas Ub[Ub]<sup>-48,63</sup>Ub or [Ub]<sub>2</sub><sup>-48,63</sup>Ub indicates two distal Ub units linked to K48 and K63 of a proximal Ub; the aforementioned branched tri-Ub would be written as Ub[Ub]<sup>-48,63</sup>Ub(<sup>15</sup>N) to indicate isotopic enrichment with <sup>15</sup>N specifically in the proximal Ub. If it is necessary to distinguish between the distal moieties in a branched chain, those should be listed in the same order (i.e., left to right) as the superscripts that correspond to the linkage sites. For example, the same branched tri-Ub chain with <sup>15</sup>N-enrichment of only the K48-linked or the K63-linked distal Ub would be written as Ub(<sup>15</sup>N)[Ub]<sup>-48,63</sup>Ub or Ub[Ub(<sup>15</sup>N)]<sup>-48,63</sup>Ub, respectively (see examples in Figure 1). As a final example, tetra-Ub made by conjugation of Ub(K0) to K11, K48, and K63 of a single Ub containing a C-terminal His6-tag would be [Ub(K0)]<sub>3</sub><sup>-11,48,63</sup>Ub(G76-H<sub>6</sub>). The proposed notation applies as well to homogeneous-linkage chains. While the existing notation for such polymers (e.g., K48-linked Ub<sub>2</sub>) is straightforward, for consistency we will use the new notation (i.e., Ub<sup>-48</sup>Ub) throughout this paper.

#### Assembly of Mixed-Linkage polyUb Chains

Trimeric mixed-linkage unbranched and branched chains, Ub<sup>-63</sup>Ub<sup>-48</sup>Ub, Ub<sup>-48</sup>Ub<sup>-63</sup>Ub, and [Ub]<sub>2</sub><sup>-48,63</sup>Ub, were produced using enzymatic chain assembly as detailed in Figure S1. Importantly, our method ensured that fully natural isopeptide linkages were formed with full control of the order of the linkages in each chain and, for NMR studies, which Ub unit is <sup>15</sup>N enriched. We found that the K48-linked dimer can accept Ub in a K63 linkage on either the distal or proximal end, and likewise the K63-linked dimer can accept Ub in a K48 linkage at either end. In addition, the K48,K63-branched trimer can be created from appropriate Ub monomers conjugated in one step with both K48- and K63-specific E2s (i.e., E2-25K and Ubc13:Mms2, respectively) in the reaction mixture (Figure 1A).

#### K48-Linked Ubiquitins in Unbranched or Branched Mixed-Linkage Chains Form the Classical K48 Hydrophobic Interface

We used NMR chemical-shift perturbation (CSP) mapping (e.g., Varadan et al., 2005a) to identify interactions between the Ub units in the mixed-linkage chains. In order to unravel monomer-specific contacts, each Ub unit in [Ub]<sub>2</sub><sup>-48,63</sup>Ub, Ub<sup>-63</sup>Ub<sup>-48</sup>Ub, and Ub<sup>-48</sup>Ub<sup>-63</sup>Ub (see Figure 1 for chain notations) was individually <sup>15</sup>N enriched (i.e., one <sup>15</sup>N-Ub per chain) resulting in nine distinct <sup>15</sup>N-labeled tri-Ub constructs. <sup>1</sup>H-<sup>15</sup>N NMR spectra of each chain (e.g., Figure 1B) were acquired under identical conditions and compared to those of monomeric Ub and of the corresponding Ub unit in Ub<sup>-48</sup>Ub and Ub<sup>-63</sup>Ub (see examples in Figures S2 and S3).

Consistent with the absence of detectable noncovalent contacts between Ub units in Ub<sup>-63</sup>Ub, the only CSPs (versus mono-Ub) observed in that chain (Varadan et al., 2004) were in the C-terminal residues of the distal Ub and those immediately

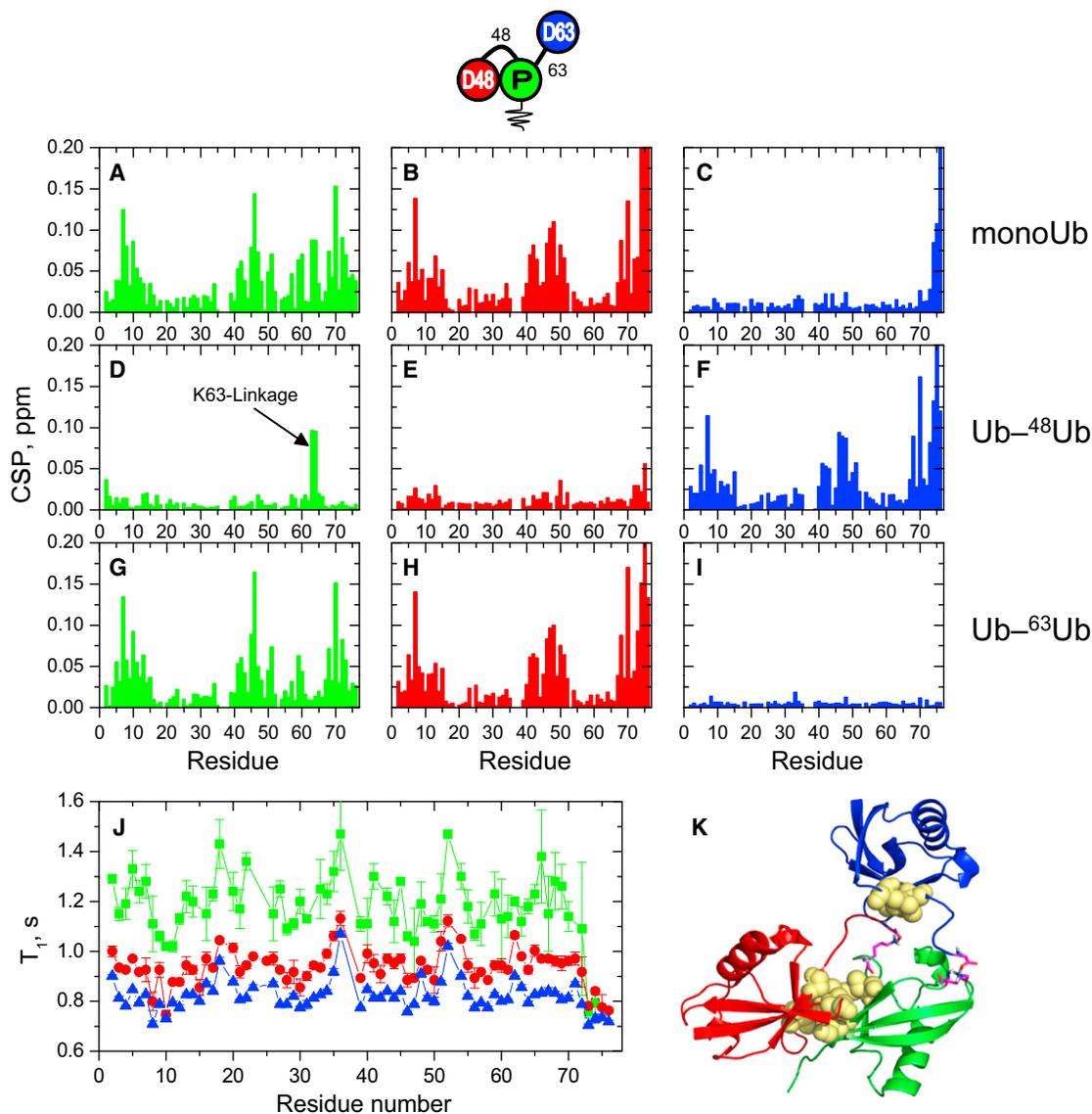
surrounding K63 of the proximal Ub, reflecting the residues involved in the isopeptide linkage between these two units. By contrast, CSP mapping of Ub<sup>-48</sup>Ub revealed (Varadan et al., 2002) that both the distal and the proximal Ubs exhibited highly specific spectral perturbations in and around the hydrophobic surface patch residues (L8, I44, V70). These CSPs, observed in addition to those in the vicinity of the isopeptide linkage between the C-terminal G76 of the distal Ub and K48 of the proximal Ub, are a clear indicator of the hydrophobic interface between the two Ubs in Ub<sup>-48</sup>Ub. These distinctive features of the NMR spectra of Ub<sup>-63</sup>Ub and Ub<sup>-48</sup>Ub serve as hallmarks of the corresponding linkages and the resulting Ub-Ub contacts.

NMR spectra of [Ub]<sub>2</sub><sup>-48,63</sup>Ub show that the amide resonances in the distal K63-linked Ub are almost identical to those in monomeric Ub with the exception of the C-terminal residues 74–76, where the observed CSPs are caused directly by ligation to K63 of the proximal Ub (Figure 2C). Consistent with this observation, there are virtually no spectral differences between distal K63-linked Ub and the distal Ub in Ub<sup>-63</sup>Ub, indicating their close structural similarity and the absence of noncovalent contacts with the proximal Ub in the corresponding chains (Figure 2I).

A similar comparison of the spectra of the distal K48-linked Ub in [Ub]<sub>2</sub><sup>-48,63</sup>Ub revealed a strikingly different picture. Here, we detected large site-specific CSPs between the distal K48-linked Ub and mono-Ub (Figure 2B) and almost negligible spectral differences between the distal K48-linked Ub and the distal Ub in Ub<sup>-48</sup>Ub (Figure 2E). These results show that the distal K48-linked Ub in [Ub]<sub>2</sub><sup>-48,63</sup>Ub makes essentially the same interdomain contacts (i.e., the hydrophobic interface with the proximal Ub and the isopeptide linkage through G76) as the distal Ub in Ub<sup>-48</sup>Ub. Predictably, these contacts result in large site-specific spectral differences between the distal K48-linked Ub and the distal Ub of Ub<sup>-63</sup>Ub (Figure 2H), which are also similar to the CSPs between the distal K48-linked Ub and monomeric Ub (Figure 2E).

Careful analysis of the spectra for the proximal Ub of [Ub]<sub>2</sub><sup>-48,63</sup>Ub revealed two signals originating from the isopeptide εNH groups of K48 and K63 (Figures 1B and S3). These signals are diagnostic for the isopeptide linkage through the ε-amino group of the corresponding lysine. That they are at the same resonance frequencies as in the respective isolated diubiquitins (K48- or K63-linked) proves the presence of both expected linkages in [Ub]<sub>2</sub><sup>-48,63</sup>Ub and also demonstrates that NMR can be used for linkage-diagnostic purposes. Note in this regard that for both unbranched mixed-linkage tri-Ub chains (see below) we also observed the isopeptide εNH signals that corresponded to each respective linkage.

Because the proximal Ub of [Ub]<sub>2</sub><sup>-48,63</sup>Ub is ubiquitinated at two lysines simultaneously, its NMR spectrum could differ from the proximal-Ub spectrum in the respective isolated di-Ubs. Nevertheless, a clear picture emerged from comparison of the corresponding spectra. For example, a comparison to mono-Ub indicates that the hydrophobic-patch residues of the proximal Ub form an interface with another Ub unit, while its C terminus is unligated (as in monomeric Ub) unlike those of the distal K48- and K63-linked Ubs (Figure 2A). The spectral differences between the proximal Ubs of [Ub]<sub>2</sub><sup>-48,63</sup>Ub and Ub<sup>-48</sup>Ub are minimal except for the region around K63 which is linked to



**Figure 2. NMR Characterization of [Ub]<sub>2</sub><sup>-48,63</sup>Ub**

Data for the proximal Ub are shown in green, for distal K48-linked Ub in red, and for distal K63-linked Ub in blue (see pictograph, top).

(A–I) Spectral differences (quantified as CSPs) between each Ub in [Ub]<sub>2</sub><sup>-48,63</sup>Ub and monomeric Ub (top row) or the corresponding Ub units in Ub<sup>-48</sup>Ub (middle row) or Ub<sup>-63</sup>Ub (bottom row). Left column: CSPs between the proximal Ub in [Ub]<sub>2</sub><sup>-48,63</sup>Ub and (A) Ub<sub>1</sub>, (D) proximal Ub in Ub<sup>-48</sup>Ub, and (G) proximal Ub in Ub<sup>-63</sup>Ub. Middle column: CSPs between distal K48-linked Ub in [Ub]<sub>2</sub><sup>-48,63</sup>Ub and (B) Ub<sub>1</sub>, (E) distal Ub in Ub<sup>-48</sup>Ub, and (H) distal Ub in Ub<sup>-63</sup>Ub. Right column: CSPs between the distal K63-linked Ub in [Ub]<sub>2</sub><sup>-48,63</sup>Ub and (C) Ub<sub>1</sub>, (F) distal Ub in Ub<sup>-48</sup>Ub, and (I) distal Ub in Ub<sup>-63</sup>Ub.

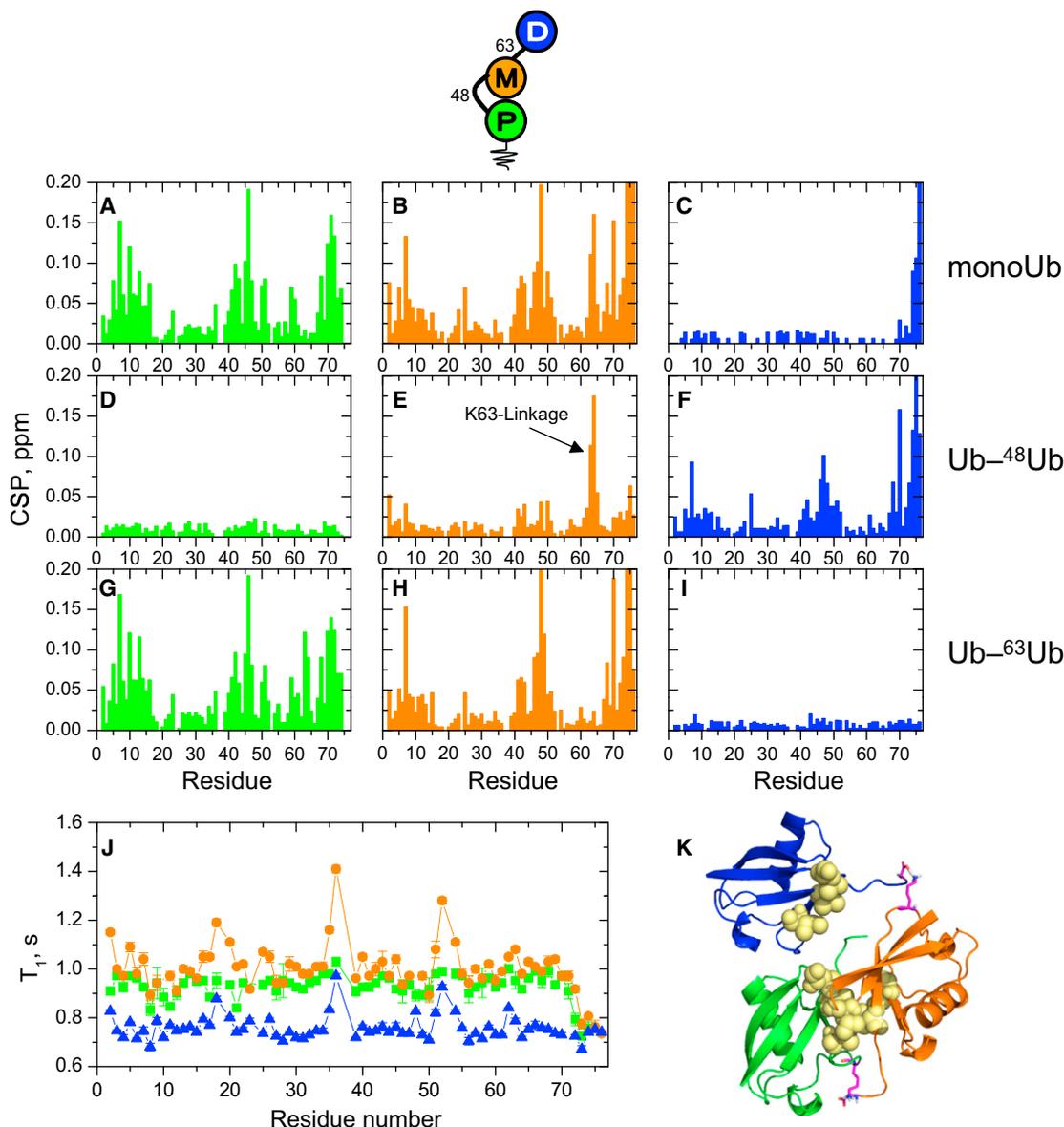
(J)  $^{15}\text{N}$   $T_1$  values as a function of the residue number for each Ub in [Ub]<sub>2</sub><sup>-48,63</sup>Ub. The error bars represent SDs.

(K) A representative structural model of [Ub]<sub>2</sub><sup>-48,63</sup>Ub from the lowest energy cluster. The coloring of the Ub units is the same as above, the isopeptide linkages are shown in magenta, and the hydrophobic-patch residues (L8, I44, V70) are shown as yellow spheres.

See also Figure S4.

another Ub only in the branched trimer (Figure 2D). As with the distal K48-linked Ub (see above), the strong site-specific CSPs with respect to the proximal Ub in Ub<sup>-63</sup>Ub reflect the hydrophobic-patch contacts between the two K48-linked Ubs (Figure 2G). Based on these and the abovementioned data, we conclude that the two K48-linked Ubs in [Ub]<sub>2</sub><sup>-48,63</sup>Ub form essentially the same hydrophobic-interface contact as in Ub<sup>-48</sup>Ub, whereas only the distal K63-linked Ub behaves as in Ub<sup>-63</sup>Ub, where no noncovalent contact was detected.

When we rearranged the linkages to form an unbranched mixed-linkage chain, Ub<sup>-63</sup>Ub<sup>-48</sup>Ub, the Ub units maintained their expected contacts and spectral properties. The proximal Ub of Ub<sup>-63</sup>Ub<sup>-48</sup>Ub has its K48 ligated, and we detected a hydrophobic interface by the large site-specific CSPs relative to mono-Ub and Ub<sup>-63</sup>Ub (Figures 3A and 3G). The spectral differences are minimal between the proximal Ub in Ub<sup>-48</sup>Ub and that in Ub<sup>-63</sup>Ub<sup>-48</sup>Ub, which indicates close similarity between the two units ligated via K48 (Figure 3D). The next (i.e., middle) Ub



**Figure 3. NMR Characterization of the Unbranched, Mixed-Linkage Ub<sup>63</sup>Ub<sup>48</sup>Ub Chain**

Data for the proximal Ub are shown in green, for the middle Ub in orange, and for the distal Ub in blue (see pictograph, top).

(A–I) Spectral differences (quantified as CSPs) between each Ub in Ub<sup>63</sup>Ub<sup>48</sup>Ub and monomeric Ub (top row) or the corresponding Ub units in Ub<sup>48</sup>Ub (middle row) or Ub<sup>63</sup>Ub (bottom row). Left column: CSPs between the proximal Ub in Ub<sup>63</sup>Ub<sup>48</sup>Ub and (A) Ub<sub>1</sub>, (D) proximal Ub in Ub<sup>48</sup>Ub, and (G) proximal Ub in Ub<sup>63</sup>Ub. Middle column: CSPs between the middle Ub in Ub<sup>63</sup>Ub<sup>48</sup>Ub and (B) Ub<sub>1</sub>, (E) distal Ub in Ub<sup>48</sup>Ub, and (H) distal Ub in Ub<sup>63</sup>Ub. Right column: CSPs between the distal Ub in Ub<sup>63</sup>Ub<sup>48</sup>Ub and (C) Ub<sub>1</sub>, (F) distal Ub in Ub<sup>48</sup>Ub, and (I) distal Ub in Ub<sup>63</sup>Ub.

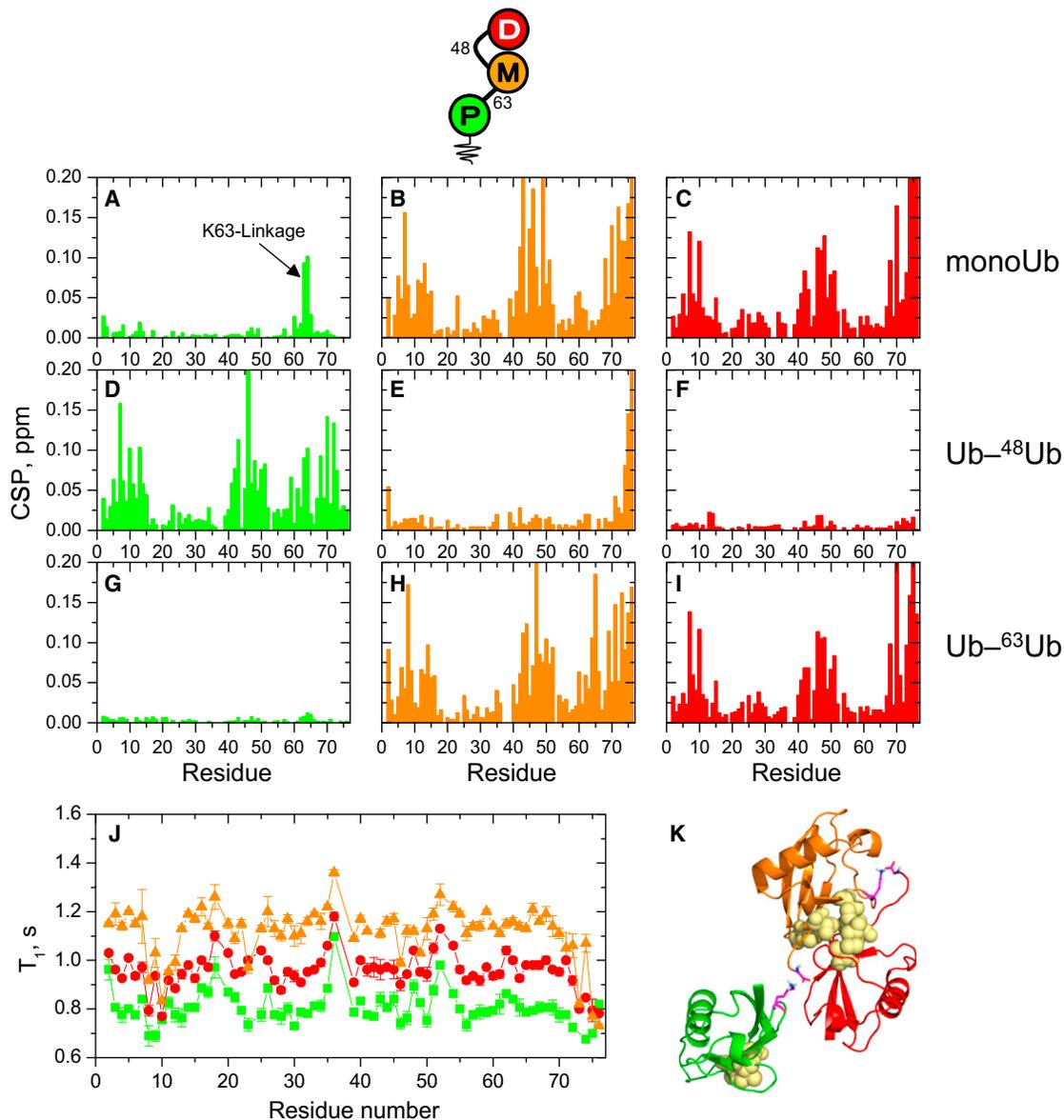
(J) <sup>15</sup>N T<sub>1</sub> values as a function of the residue number for each Ub in Ub<sup>63</sup>Ub<sup>48</sup>Ub. The error bars represent SDs.

(K) A representative structural model of Ub<sup>63</sup>Ub<sup>48</sup>Ub from the lowest-energy cluster. The coloring of the Ub units is the same as above, the isopeptide linkages are shown in magenta, and the hydrophobic-patch residues (L8, I44, V70) are shown as yellow spheres.

of Ub<sup>63</sup>Ub<sup>48</sup>Ub has features of both a proximal unit (linked through its K63) and a distal unit linked through its C-terminal G76. The large site-specific CSPs versus mono-Ub (Figure 3B) clearly indicate that the middle Ub participates in a hydrophobic interface. Based on the spectral similarity with the distal Ub of Ub<sup>48</sup>Ub (Figure 3E), we conclude that the K48-linked proximal and middle Ubs in Ub<sup>63</sup>Ub<sup>48</sup>Ub retain the hydrophobic interface characteristic of Ub<sup>48</sup>Ub. Note also that the large CSPs

observed around K63 in the middle Ub reflect its linkage to the distal Ub.

The NMR spectra of the distal Ub in Ub<sup>63</sup>Ub<sup>48</sup>Ub indicate that this unit is excluded from the hydrophobic interface formed between the middle and proximal Ubs. Indeed, the CSPs in distal versus monomeric Ub are localized to the (ligated) C-terminal region (Figure 3C), whereas the spectral differences between the distal Ubs of Ub<sup>63</sup>Ub<sup>48</sup>Ub and Ub<sup>63</sup>Ub are negligible



**Figure 4. NMR Characterization of the Unbranched, Mixed-Linkage Ub<sup>-48</sup>Ub<sup>-63</sup>Ub Chain**

Proximal Ub is shown in green, middle Ub in orange, and distal Ub in red (see pictograph, top).

(A–I) CSPs between each Ub in Ub<sup>-48</sup>Ub<sup>-63</sup>Ub versus monomeric Ub (left column), Ub<sup>-48</sup>Ub (middle column), or Ub<sup>-63</sup>Ub (right column). (A–I) Spectral differences (quantified as CSPs) between each Ub in Ub<sup>-48</sup>Ub<sup>-63</sup>Ub and monomeric Ub (first row) or the corresponding Ub units in Ub<sup>-48</sup>Ub (middle row) or Ub<sup>-63</sup>Ub (bottom row). Left column: CSPs between the proximal Ub in Ub<sup>-48</sup>Ub<sup>-63</sup>Ub and (A) Ub<sub>1</sub>, (D) proximal Ub in Ub<sup>-48</sup>Ub, and (G) proximal Ub in Ub<sup>-63</sup>Ub. Middle column: CSPs between the middle Ub in Ub<sup>-48</sup>Ub<sup>-63</sup>Ub and (B) Ub<sub>1</sub>, (E) distal Ub in Ub<sup>-48</sup>Ub, and (H) distal Ub in Ub<sup>-63</sup>Ub. Right column: CSPs between the distal Ub in Ub<sup>-48</sup>Ub<sup>-63</sup>Ub and (C) Ub<sub>1</sub>, (F) distal Ub in Ub<sup>-48</sup>Ub, and (I) distal Ub in Ub<sup>-63</sup>Ub.

(J) <sup>15</sup>N T<sub>1</sub> values as a function of the residue number for each Ub in Ub<sup>-48</sup>Ub<sup>-63</sup>Ub. The error bars represent SDs.

(K) A representative structural model of Ub<sup>-48</sup>Ub<sup>-63</sup>Ub from the lowest energy cluster. The coloring of the Ub units is the same as above, the isopeptide linkages are shown in magenta and the hydrophobic-patch residues (L8, I44, V70) are shown as yellow spheres.

(Figure 3I). This indicates strongly that the distal Ub of Ub<sup>-63</sup>Ub<sup>-48</sup>Ub has the same structure and contacts (or absence thereof) as the distal Ub in Ub<sup>-63</sup>Ub.

In Ub<sup>-48</sup>Ub<sup>-63</sup>Ub, the proximal Ub is expected to be the only unit that does not form a hydrophobic interface. Indeed, this is evident from the strong spectral similarity of that proximal Ub with the proximal Ub of Ub<sup>-63</sup>Ub (Figure 4G) as well as with

mono-Ub (Figure 4A). In contrast to the proximal Ub, which exhibits no noncovalent contacts, the middle and distal units of Ub<sup>-48</sup>Ub<sup>-63</sup>Ub form a characteristic K48-linked Ub–Ub interface. This is particularly apparent from the minimal CSPs (except for the ligated C terminus) between the middle Ub of Ub<sup>-48</sup>Ub<sup>-63</sup>Ub and the proximal Ub of Ub<sup>-48</sup>Ub (Figure 4E), and the virtual absence of CSPs between the distal Ubs

(Figure 4F). These and the other spectral comparisons in Figure 4 show that the structures and interfaces of the middle and distal units in Ub<sup>-48</sup>Ub<sup>-63</sup>Ub are nearly identical to those in Ub<sup>-48</sup>Ub.

### Structural Models of the Mixed-Linkage Tri-Ub Chains

Based on the above NMR data, we generated structural models of the three tri-Ub chains studied here using the biomolecular docking program HADDOCK (de Vries et al., 2010) and site-specific CSPs to guide the docking (see Supplemental Information). Although we observed slightly different overall structures in the top clusters for these chains, all featured a distinct arrangement of the interdomain contacts in which the two K48-linked Ubs formed a well-defined contact mediated by their hydrophobic-patch residues, whereas the K63-linked Ub was represented in essentially random positions. For example, the generated structures for [Ub]<sub>2</sub><sup>-48,63</sup>Ub (Figure 2K) clearly show that the proximal and distal K48-linked Ubs form the “canonical” L8, I44, V70 hydrophobic interface, whereas the distal K63-linked Ub samples several different orientations, depending on the HADDOCK cluster (Figure S4). This variation in the K63-linked Ub is not unexpected given that (1) the CSP data show no close noncovalent interactions involving this Ub unit and (2) the <sup>15</sup>N T<sub>1</sub> relaxation data (see below) indicate that this unit has greater mobility compared to the other two Ubs. Moreover, the two K48-linked Ubs superimpose quite well (rmsd = 1.72 Å) with the crystal structure (Protein Data Bank [PDB]: 1AAR) of Ub<sup>-48</sup>Ub (Figure S4).

### Interdomain Mobility Supports Distinct K48- and K63-Linkage Behavior

The NMR spectra above clearly indicate that the two K48-linked units in all three tri-Ub chains form a specific interface, while the third unit, linked via K63, shows no contacts except for being tethered (via its flexible C terminus or via K63) and therefore is expected to be more mobile than the other two Ubs. This distinction is further seen in the generated CSP-docked structures of these chains. To test this prediction, we measured and compared <sup>15</sup>N longitudinal relaxation times, T<sub>1</sub>, for each Ub unit in a chain. Generally, T<sub>1</sub> senses the overall tumbling of the chain (reflecting its size and shape), the relative intrachain mobility (on a ns timescale) of the Ub unit under observation (which could depend on its location and intrachain contacts), and the local polypeptide backbone dynamics within the Ub protomer (which are expected to be similar for all Ubs) (Ryabov and Fushman, 2007). Whereas the overall shape could differ between chains, the T<sub>1</sub> values within the same chain are expected to reflect the relative mobility of each Ub unit. For example, if a particular Ub forms a hydrophobic interdomain interface it becomes less mobile and this should result in an increase in T<sub>1</sub>. Furthermore, a Ub that is linked to two other Ubs should also show an increase in T<sub>1</sub> as its reorientation will partially depend on the movement of the other two Ubs. Therefore, one would expect the proximal Ub of [Ub]<sub>2</sub><sup>-48,63</sup>Ub and the middle Ubs of Ub<sup>-63</sup>Ub<sup>-48</sup>Ub and Ub<sup>-48</sup>Ub<sup>-63</sup>Ub to have the largest T<sub>1</sub> values.

Indeed, in [Ub]<sub>2</sub><sup>-48,63</sup>Ub the proximal Ub has the longest T<sub>1</sub>, whereas of the two distal (singly linked) Ubs, the distal K63-linked Ub shows the shortest T<sub>1</sub> (i.e., the least restricted mobility), and the distal K48-linked Ub has a longer T<sub>1</sub>, consistent

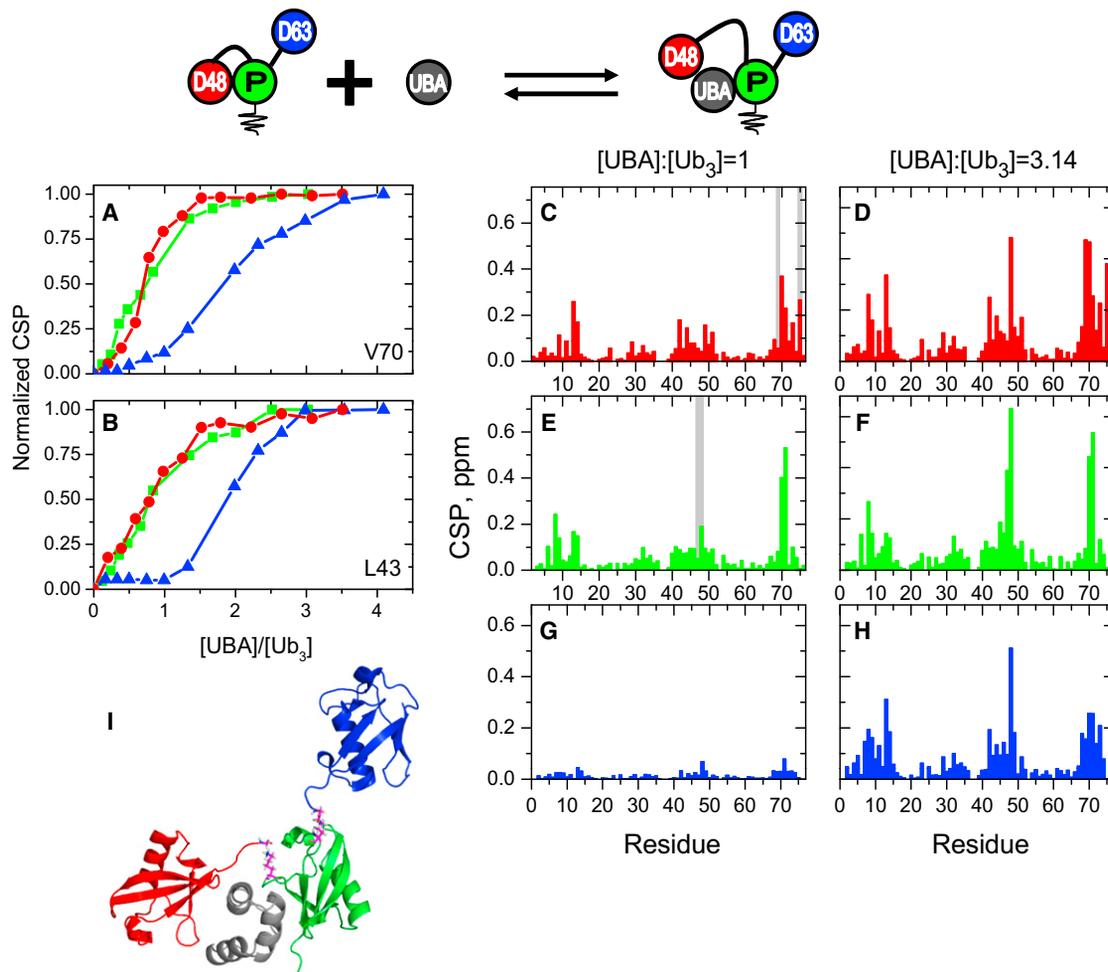
with its contact with the proximal Ub (Figure 2J). These results fully agree with the spectroscopic and structural data presented above. The same holds for the two unbranched chains, where the middle Ub shows the longest T<sub>1</sub> (most restricted mobility), while the K63-linked Ub is consistently the most mobile (i.e., shortest T<sub>1</sub>) regardless of whether it is at the distal or the proximal end of the chain (Figures 3J and 4J). K48-linked Ub shows intermediate T<sub>1</sub> values, as its tumbling is slowed by the hydrophobic interaction with the middle Ub. Note that in all of these chains the two K48-linked units have different T<sub>1</sub> values; thus, despite forming a hydrophobic contact, they are not rigidly locked. This is consistent with our previous findings for Ub<sup>-48</sup>Ub (Ryabov and Fushman, 2007; Varadan et al., 2002) and indicates dynamic opening and closing of the hydrophobic interface, which is critical for the ability of Ub-receptors to bind to this chain.

### pH Dependence of the Inter-Ub Contacts in [Ub]<sub>2</sub><sup>-48,63</sup>Ub Supports the Classical K48-Linked Interface

One hallmark of K48-linked di-Ub is that its conformation is pH dependent: lowering the pH shifts the equilibrium from the predominantly closed conformation at neutral pH to predominantly open, and this manifests itself as nearly complete disappearance of the CSPs at pH 4.5 (Varadan et al., 2002). Indeed, our NMR data (Figure S5) indicate that the pH-dependent behavior of [Ub]<sub>2</sub><sup>-48,63</sup>Ub is nearly identical to that reported for the corresponding Ubs in homogeneous K48- and K63-linked polyUb. This supports the conclusion that interdomain interactions in the branched tri-Ub are governed by the same forces as in the respective homogeneous-linkage chains.

### A K48-Selective Receptor, UBA(2) of hHR23a, Selectively Binds K48-Linked Units within the Branched [Ub]<sub>2</sub><sup>-48,63</sup>Ub Chain

Having established that the branched and unbranched mixed-linkage chains combine the structural properties characteristic of both linkages, we set out to examine whether these chains can be recognized by linkage-selective receptors. To determine if the K48 linkage in [Ub]<sub>2</sub><sup>-48,63</sup>Ub retains its receptor selectivity, we titrated [Ub]<sub>2</sub><sup>-48,63</sup>Ub (separately for each Ub unit <sup>15</sup>N-enriched; see Figure 1) with the K48-linkage selective receptor UBA(2) from the proteasomal shuttle protein hHR23A. UBA(2) binds Ub<sup>-48</sup>Ub in a “sandwich” mode and significantly more tightly than to Ub<sup>-63</sup>Ub or monomeric Ub (Varadan et al., 2005b). We observed strong residue-specific CSPs in both the proximal and the distal K48-linked Ubs (Figure 5); these perturbations center around the hydrophobic-patch residues and the isopeptide linkage, consistent with UBA(2) insertion into the hydrophobic pocket formed by the K48-linked Ubs (Varadan et al., 2005b). We can gauge the extent of binding from titration curves of residue-specific CSPs or by comparing the CSPs for all residues at various points in titration. The CSPs in the proximal and distal K48-linked Ubs of [Ub]<sub>2</sub><sup>-48,63</sup>Ub show little change between [UBA(2)]:[Ub]<sub>2</sub><sup>-48,63</sup>Ub = 1 and 3 (Figure 5), suggesting that UBA(2) binding to both Ubs reached saturation at approximately 1:1 molar ratio. Additional changes occurring in these Ubs above the 1:1 molar ratio could reflect binding of a second UBA(2) molecule, as observed for Ub<sup>-48</sup>Ub. Interestingly, the



**Figure 5. Binding of hHR23A UBA(2) to [Ub]<sub>2-48,63</sub>Ub**

All data are color coded as indicated in the pictograph on the top. Proximal Ub data are shown in green, distal K48-linked Ub in red, and distal K63-linked Ub in blue.

(A and B) Representative titration curves plotted as normalized CSP versus [UBA(2)]:[Ub]<sub>2-48,63</sub>Ub for residues V70 and L43 in all three Ub units.

(C–H) Residue-specific CSPs in individual Ub units at two points in titration with UBA(2): at the 1:1 molar ratio (left column) and at saturation (right column). Gray bars indicate residues with signals broadened beyond detection at the 1:1 molar ratio. Note that the distal K63-linked Ub exhibits virtually no binding until [UBA(2)]:[Ub]<sub>2-48,63</sub>Ub > 1.

(I) A model of the [Ub]<sub>2-48,63</sub>Ub/UBA(2) complex at the 1:1 molar ratio, based on the observed CSPs and the structure of the Ub<sup>-48</sup>Ub/UBA(2) complex (PDB: 1ZO6). The coloring of the Ub units is as depicted above; UBA(2) is colored gray.

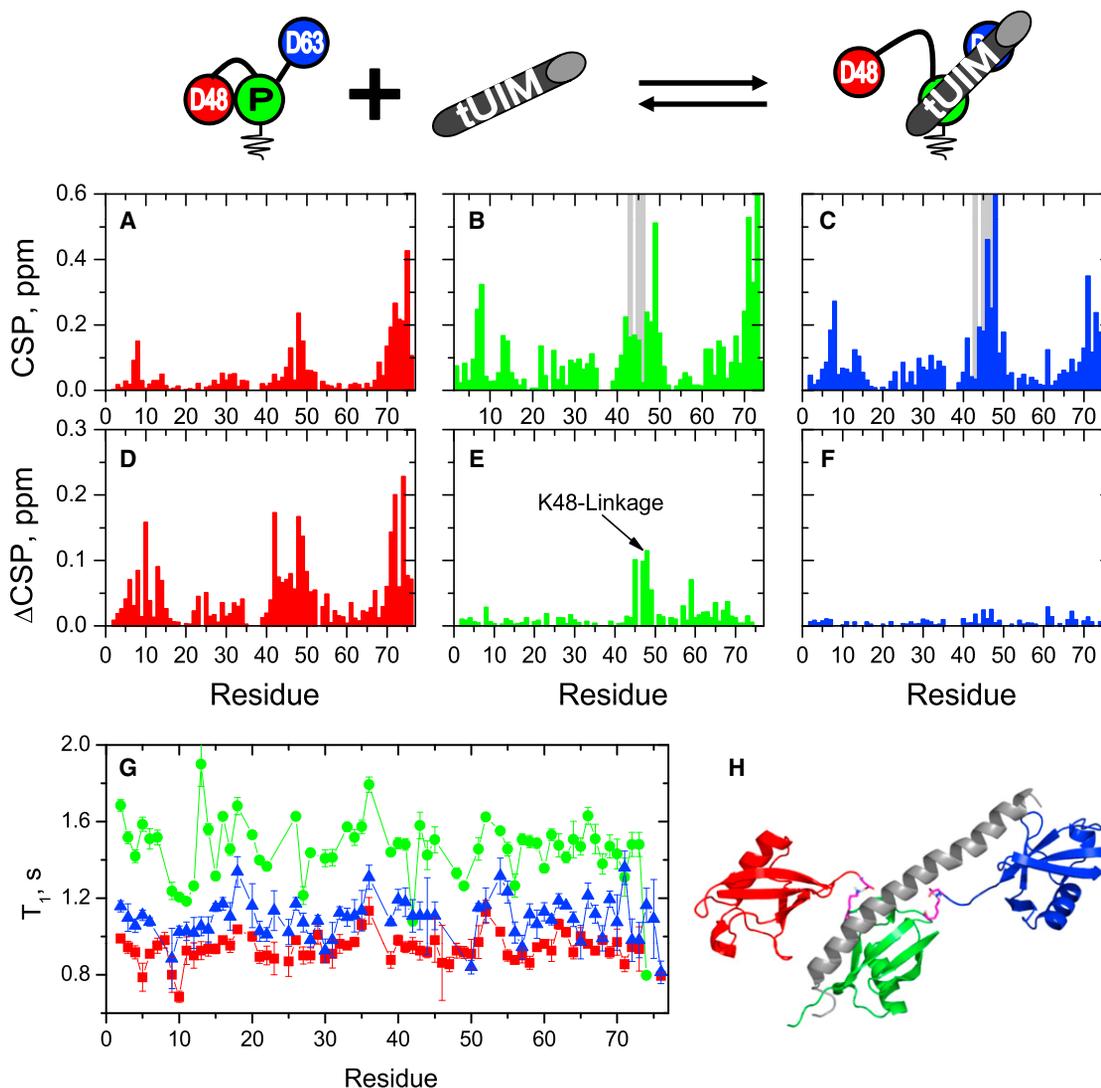
See also Figures S6 and S8.

distal K63-linked Ub exhibited an entirely different behavior and showed spectral perturbations only after the distal K48-linked and proximal Ubs were saturated with UBA(2) (Figure 5). Even at [UBA(2)]:[Ub]<sub>2-48,63</sub>Ub = 1, there are virtually no CSPs in the distal K63-linked Ub, and it was not until after this point that we finally observed UBA(2) binding to this Ub (Figures 5G and 5H). The titration curves for individual residues illustrate this trend from another angle by showing “standard” binding behavior for the distal K48-linked and proximal Ubs, but a “lag-phase” for the distal K63-linked Ub (Figures 5A, 5B, and 5G). This strong binding preference for K48-linked Ubs is also supported by our relaxation data that showed a larger increase in <sup>15</sup>N T<sub>1</sub> of the distal K48-linked Ub compared to distal K63-linked Ub at saturation (Figure S6). These results demonstrate

that a K48-linkage receptor can selectively recognize this linkage in a branched polyUb chain using the same binding mechanism as for homogeneous-linkage chains.

#### A K63-Selective Receptor, Rap80-tUIM, Avidly Binds across the K63 Linkage in the Branched [Ub]<sub>2-48,63</sub>Ub Chain

Rap80 contains a tandem Ubiquitin Interacting Motif (tUIM) that shows strong binding preference for Ub<sup>-63</sup>Ub versus Ub<sup>-48</sup>Ub (Sims and Cohen, 2009). The tUIM contains a flexible linker region between its two helical UIMs which adopts an  $\alpha$ -helical conformation upon binding to Ub<sup>-63</sup>Ub and perfectly aligns the two UIMs’ interacting surfaces for binding to adjacent Ubs in a K63 linkage (Sato et al., 2009; Sims and Cohen, 2009). To



**Figure 6. Binding of the Rap80 tUIM to  $[\text{Ub}]_2^{-48,63}\text{Ub}$  Chain**

All data are color coded as indicated in the pictograph on the top: proximal Ub data are shown in green, distal K48-linked Ub in red, and distal K63-linked Ub in blue.

(A–C) CSP plots for each Ub unit in  $[\text{Ub}]_2^{-48,63}\text{Ub}$  at saturation: (A) distal K48-linked Ub at  $[\text{tUIM}]:[\text{Ub}]_2^{-48,63}\text{Ub} = 2.43$ , (B) distal K63-linked Ub at  $[\text{tUIM}]:[\text{Ub}]_2^{-48,63}\text{Ub} = 1.52$ , and (C) proximal Ub at  $[\text{tUIM}]:[\text{Ub}]_2^{-48,63}\text{Ub} = 2.00$ . Residues that showed strong signal attenuation during the titration are represented with gray bars.

(D–F) Difference in the CSPs ( $\Delta\text{CSP}$ ) for each Ub of  $[\text{Ub}]_2^{-48,63}\text{Ub}$  and the corresponding Ub in the respective di-Ub controls at saturation with tUIM: (D) distal K48-linked Ub of  $[\text{Ub}]_2^{-48,63}\text{Ub}$  versus distal Ub of  $\text{Ub}^{-48}\text{Ub}$ , (E) proximal Ub of  $[\text{Ub}]_2^{-48,63}\text{Ub}$  versus proximal Ub of  $\text{Ub}^{-63}\text{Ub}$ , and (F) distal K63-linked of  $[\text{Ub}]_2^{-48,63}\text{Ub}$  versus distal Ub of  $\text{Ub}^{-63}\text{Ub}$ .

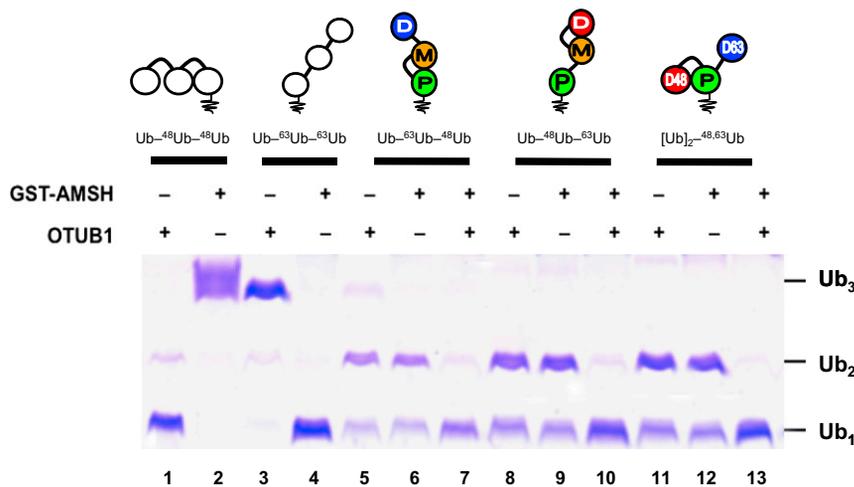
(G)  $^{15}\text{N}$   $T_1$  relaxation time for each amide in the proximal, distal K48-linked, and distal K63-linked Ubs in  $[\text{Ub}]_2^{-48,63}\text{Ub}$  at saturation with Rap80 tUIM. The error bars represent SDs.

(H) A model of the  $[\text{Ub}]_2^{-48,63}\text{Ub}/\text{tUIM}$  complex at 1:1 molar ratio, based on the observed CSPs and  $^{15}\text{N}$   $T_1$  values, and the  $\text{Ub}^{-63}\text{Ub}/\text{tUIM}$  complex structure (PDB: 3A1Q). The coloring of the Ub units is the same as above; tUIM is colored gray.

See also Figures S7 and S9.

examine how the tUIM interacts with branched chains, we titrated it into  $[\text{Ub}]_2^{-48,63}\text{Ub}$ , as well as  $\text{Ub}^{-48}\text{Ub}$  and  $\text{Ub}^{-63}\text{Ub}$  as controls, and used NMR to monitor the interactions separately for each  $^{15}\text{N}$ -enriched Ub unit. As shown in Figure 6, NMR signals of all three Ubs in  $[\text{Ub}]_2^{-48,63}\text{Ub}$  were altered by the tUIM. The observed perturbations were site specific and centered around the hydrophobic-patch residues as well as

the Ub C termini. The distal K48-linked Ub showed noticeably weaker CSPs at saturation (suggesting weaker binding) compared to the other two Ubs. Importantly, during these titrations some residues in the proximal and distal K63-linked Ubs, but not in the distal K48-linked Ub, showed strong attenuations or even disappearance of the NMR signals; this is caused by slow exchange (i.e., slow off-rates) on the NMR chemical-shift



**Figure 7. Linkage-Selective DUBs Cleave Their Cognate Ub-Ub Linkages in Homogeneous or Mixed-Linkage polyUb**

The indicated chains were incubated with OTUB1, GST-AMSH, or both and the products evaluated by SDS-PAGE and staining with Coomassie blue; see [Experimental Procedures](#) for details. Lanes 1–4 show reaction products of homogeneous Ub<sup>-48</sup>Ub<sup>-48</sup>Ub and Ub<sup>-63</sup>Ub<sup>-63</sup>Ub. Lanes 5 and 6, 8 and 9, and 11 and 12 show that alone each DUB can only process one linkage in the unbranched or branched mixed-linkage chains. Only with both DUBs present were the mixed-linkage chains hydrolyzed fully to monomeric Ub (lanes 7, 10, and 13).

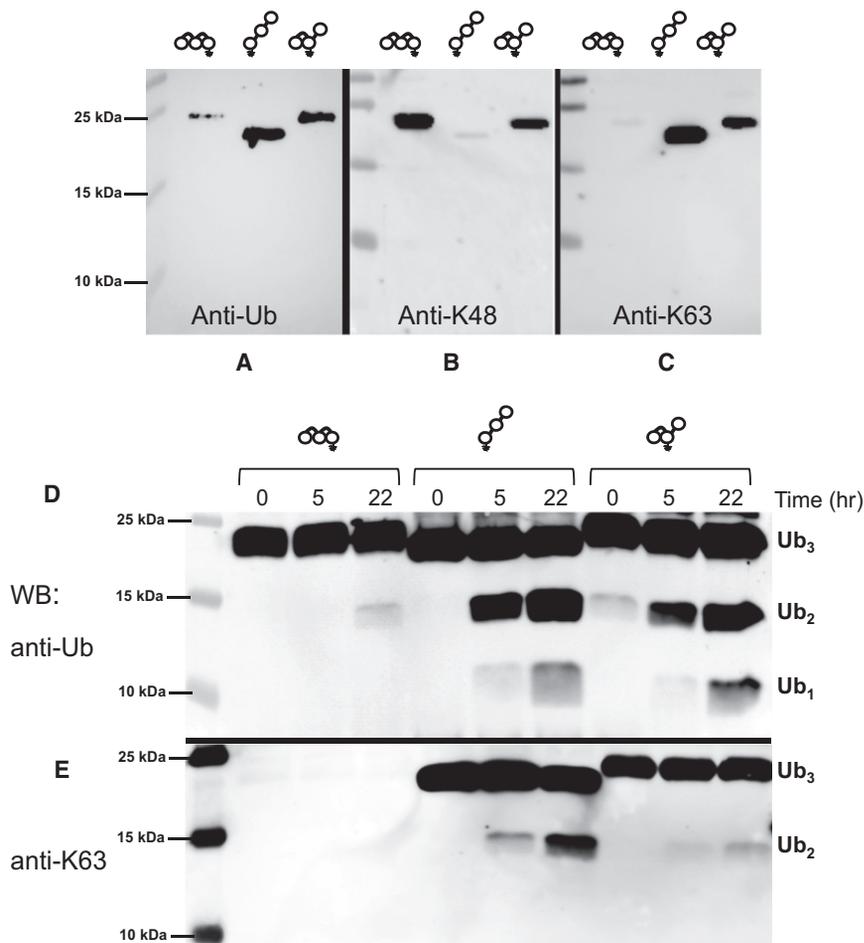
timescale and is indicative of tighter binding to the K63-linked Ubs. Comparison of the domain-specific CSPs in [Ub]<sub>2</sub><sup>-48,63</sup>Ub with those in the control di-Ub samples (Figures 6D–6F), as well as comparison of the signal trajectories upon titration (Figure S7), led us to the following conclusions. First, the distal K63-linked and proximal Ubs of [Ub]<sub>2</sub><sup>-48,63</sup>Ub bind the tUIM in the same mode as their respective units in Ub<sup>-63</sup>Ub (Figures 6D and 6E). This is supported by the fact that at saturation with the tUIM virtually all signals in the <sup>1</sup>H-<sup>15</sup>N NMR spectra of the distal K63-linked and proximal Ubs of [Ub]<sub>2</sub><sup>-48,63</sup>Ub overlay perfectly with the corresponding signals of Ub<sup>-63</sup>Ub. Also, the signal trajectories (in the course of the titration) for the same residue in the K63-linked components of the branched and control chains were nearly superimposable (Figure S7), consistent with identical binding interactions. Second, there is a stark difference between tUIM binding to the distal K48-linked Ub of [Ub]<sub>2</sub><sup>-48,63</sup>Ub and the Ubs of Ub<sup>-48</sup>Ub. Indeed, our data show spectral differences ( $\Delta$ CSP) at saturation between the distal K48-linked Ub and either Ub of Ub<sup>-48</sup>Ub (and even mono-Ub) that are much larger than the differences observed for the K63-linked components (Figure 6).

When the tUIM binds to Ub<sup>-48</sup>Ub, it has to disrupt the hydrophobic interface between the two Ubs in order to access the Ub hydrophobic surface. Even though avid tUIM binding to K48-linked di-Ub has not been observed, the reported interaction ( $K_d = 0.16$  mM) was still relatively strong (Sims and Cohen, 2009). In the branched chain, simultaneous tUIM binding to the proximal and distal K63-linked Ubs should displace the distal K48-linked Ub from its hydrophobic contact with the proximal Ub, thus freeing it and making it available for binding to excess tUIM molecules. To verify this model, we measured <sup>15</sup>N T<sub>1</sub> values for each Ub of [Ub]<sub>2</sub><sup>-48,63</sup>Ub at saturation with the tUIM (Figure 6G) and compared them with those of unbound [Ub]<sub>2</sub><sup>-48,63</sup>Ub (Figure 2J). The tUIM binding significantly increased the T<sub>1</sub> values (reflecting slower tumbling) for both the distal K63-linked and the proximal Ub but caused almost no change in the T<sub>1</sub> values of the distal K48-linked Ub, rendering it the lowest-T<sub>1</sub> unit in the chain (Figure 6G). This confirms that the Rap80 tUIM binds simultaneously to both the proximal and (formerly “free” and the most mobile) distal

K63-linked Ubs, thereby restricting their conformational freedom. At the same time, the tUIM binding displaced and freed the distal K48-linked Ub from its contact with the proximal Ub, making it the most mobile Ub in the chain. Thus, we conclude that the tUIM-induced perturbations detected in the distal K48-linked Ub are primarily due to this secondary effect rather than direct interaction with the tUIM. However, we cannot completely rule out that some small fraction of this distal Ub interacts with the excess Rap80 tUIM present at the saturation conditions.

#### Linkage-Specific Deubiquitinating Enzymes Retain Linkage Selectivity with Branched and Mixed-Linkage polyUb Substrates

Given that DUBs are critical to maintain the pool of free Ub and to regulate conjugate levels, we tested if linkage-selective DUBs would still retain their ability to recognize and cleave their cognate Ub-Ub linkage in the branched and unbranched mixed-linkage chains. For this purpose, we used OTUB1, which specifically cleaves K48 linkages (Wang et al., 2009), and associated molecule with the SH3-domain of STAM (AMSH), which is specific for K63 linkages (McCullough et al., 2004). As evident in Figure 7, OTUB1 readily reduced [Ub]<sub>2</sub><sup>-48,63</sup>Ub, Ub<sup>-63</sup>Ub<sup>-48</sup>Ub, and Ub<sup>-48</sup>Ub<sup>-63</sup>Ub to di- and mono-Ub. However, unlike the case with the homogeneously linked Ub<sup>-48</sup>Ub<sup>-48</sup>Ub substrate, OTUB1 alone could not further process the di-Ub species generated from any of the mixed-linkage chains (Figure 7). Only when both OTUB1 and AMSH were present was complete chain disassembly achieved, indicating that the uncleaved di-Ub was linked via K63. An essentially identical behavior was observed for AMSH, which readily cleaved the K63 linkage in all three tri-Ub chains while leaving the K48-linkage intact (Figure 7). Taken together, these results demonstrate that the K48 and K63 linkages in branched and unbranched mixed-linkage polyUb can each individually be processed by linkage-selective DUBs. Recently, it was shown that OTUB1 and AMSH can separately cleave their cognate linkages in a branched tri-Ub construct containing analogs of K48 and K63 linkages formed via a N $\epsilon$ -Gly-L-homothiaLys mimic (which is one C-C bond longer and contains a sulfur atom instead of carbon) of an isopeptide bond (Valkevich et al., 2012). Our study demonstrates this selectivity for fully natural isopeptide bonds.



**Figure 8. Disassembly of Homogeneous and Branched Chains by 26S Proteasomes and Their Recognition by Linkage-Specific Antibodies**

(A–C) Upper panel: 18% SDS-PAGE blots of  $Ub_{-48}Ub_{-48}Ub$ ,  $Ub_{-63}Ub_{-63}Ub$ , and  $[Ub]_2^{-48,63}Ub$  with (A) anti-Ub, (B) anti-K48, and (C) anti-K63 antibodies. All three tri-Ub species were detected with anti-Ub. Only the  $[Ub]_2^{-48,63}Ub$  was detected by both anti-K48 and anti-K63, confirming that both K48 and K63 linkages are present.

(D and E) Lower panel:  $Ub_{-48}Ub_{-48}Ub$ ,  $Ub_{-63}Ub_{-63}Ub$ , or  $[Ub]_2^{-48,63}Ub$  were incubated with purified yeast 26S proteasomes and the digests were monitored at 0, 5, and 22 hr by SDS-urea PAGE and blotting with (D) anti-Ub and (E) anti-K63 antibodies.

$[Ub]_2^{-48,63}Ub$  (Figures 8B and 8C). This result in itself clearly demonstrates that chain branching preserves both Ub-specific and linkage-specific epitopes for antibody recognition and also shows that linkage-specific antibodies can be used to probe if a particular polyUb chain contains multiple linkage types. Next, each chain was incubated with purified 26S proteasome, and reaction products were identified by blotting with anti-Ub antibodies. Processing of the homogeneous  $Ub_{-63}Ub_{-63}Ub$  chains was faster than that of  $Ub_{-48}Ub_{-48}Ub$  (Figure 8D); this result is consistent with previous reports (Cooper et al., 2009; Jacobson

### The Yeast 26S Proteasome Can Process Branched Chains

One of the most studied outcomes of Ub modification is targeting to the proteasome. In a poorly understood process, proteasome-associated DUBs mainly function to modify or remove the polyUb tag from conjugated substrates (e.g., [Glickman and Adir, 2004](#); [Lee et al., 2011](#)). The three major DUBs associated with the proteasomes are Rpn11 (Poh1 in humans), Ubp6 (Usp14 in humans), and Uch37 (in humans). Whether and how branched polyUb chains are processed by the proteasome is an open question.

A series of studies has shown that the so-called “forked” polyUb chains that are branched on two closely positioned lysine residues (i.e., K6 and K11, K27 and K29, or K29 and K33) are degraded slowly by purified proteasomes and inhibit proteasomal DUB activity and substrate degradation ([Kim and Goldberg, 2012](#); [Kim et al., 2007, 2009](#)). Whether this observation holds for other chains such as those containing the more abundant K48 and K63 linkages was not reported. To evaluate the efficiency of proteasome processing of  $Ub_{-48}Ub_{-48}Ub$ ,  $Ub_{-63}Ub_{-63}Ub$ , or  $[Ub]_2^{-48,63}Ub$  signals, we incubated these tri-Ubs with purified yeast 26S proteasomes and monitored the cleavage reaction by blotting with anti-Ub and linkage-selective antibodies.

We first established that both the K48- and K63-linkage-selective antibodies recognized their respective Ub–Ub linkages in

[et al., 2009](#)). Accordingly, the branched  $[Ub]_2^{-48,63}Ub$  chain was disassembled faster than  $Ub_{-48}Ub_{-48}Ub$ , and generation of di-Ub (by removal of the single K63-linked Ub) was as efficient as from  $Ub_{-63}Ub_{-63}Ub$  chains. These observations suggest that (1) the branching did not hinder proteasomal cleavage, and (2) the presence of a K63 linkage in this chain facilitates its conversion to di-Ub. The latter was verified using a K63-linkage-specific antibody, which revealed that only a minor fraction of the di-Ub products generated from  $[Ub]_2^{-48,63}Ub$  were K63-linked (Figure 8F). Taken together, these results demonstrate that the 26S proteasome can process branched polyUb chains.

### DISCUSSION

Ubiquitination machinery has been shown in some cases to be processive in generating homogeneously-linked chains. We demonstrated here that, when acting together simultaneously or sequentially, two linkage-specific E2 enzymes (K48-specific E2-25K and K63-specific Ubc13:Mms2) can form both branched and unbranched mixed-linkage forms of polyUb. This observation can be generalized further. As shown in [Figure S10](#), adding a K11-specific E2 (Ubc2s) to the abovementioned enzymes generates a triply-branched product,  $[Ub]_3^{-11,48,63}Ub$ , in which three distal Ubs are attached simultaneously to K11, K48, and K63 of the same proximal Ub. Altogether, our results indicate that

exposure of Ub (or polyUb) to more than one E2 enzyme can result in chains composed of more than one linkage type. A similar outcome could be expected in the case of a single linkage-promiscuous E2 (Kim et al., 2007). Depending on the factors involved (which are currently unclear), the resulting polyUb landscape would include chains of unbranched and branched topologies. This raises the question of why such chains have not been observed more often. Possible reasons are that intracellular DUBs rapidly edit mixed-linkage chains, or that assembly of mixed-linkage chains largely is prevented by regulation (e.g., via compartmentalization) of the various E2 and E3 enzymes in cells. However, because current methods are poorly suited for detection of mixed-linkage chains, such chains may in fact be quite abundant but remain unseen.

This study revealed that K48 and K63 linkages in mixed-linkage polyUb retain structural features of the corresponding homogeneously linked chains. Our NMR data show unambiguously that the hydrophobic interface characteristic of the K48 linkage is formed across the Ub–<sup>48</sup>Ub unit in both the branched and unbranched mixed-linkage trimers, and that this interface exhibits the same pH dependence as the “classical” interface in homogeneous K48-linked polyUb. Furthermore, the <sup>15</sup>N relaxation data revealed that in the mixed-linkage chains the K63-linked Ub is highly mobile compared to the other two Ubs, which are restricted by their K48-dimer interface contacts.

We found that highly selective receptors for each of the two linkage types bind the branched [Ub]<sub>2</sub>–<sup>48,63</sup>Ub chain in a linkage-specific manner; i.e., binding to the cognate Ub–Ub unit was essentially as if the other linkage was not present. It is noteworthy that the proximal Ub in [Ub]<sub>2</sub>–<sup>48,63</sup>Ub is adaptable and can bind in both K48- and K63-linkage-selective modes. Furthermore, chain branching also preserves both Ub-specific and linkage-specific epitopes for antibody recognition. Thus, the K48 and K63 linkages within the same polyUb chain can retain their characteristic signaling properties. This conclusion is further strengthened by the fact that linkage-selective DUBs can efficiently process their cognate linkages in both branched and unbranched mixed-linkage chains. Our finding that the 26S proteasome recognizes and cleaves the branched chain suggests that in vivo polyUb containing both K48 and K63 linkages can be disassembled by the proteasome essentially as a homogeneous chain. Taken together, these observations demonstrate that the K48 and K63 signaling properties can be encoded into the same polyUb via linkage mixing or branching, thus allowing the chain to carry two distinct signals. Just as there is little or no steric hindrance in assembling these chains, there appears to be no detectable hindrance in their reading and disassembly.

The reason for targeting conjugates to the proteasome is to degrade a substrate, not to disassemble the polyUb modification, yet inevitably the chain must be disassembled as well. Our observations using yeast 26S proteasomes together with published data for mammalian proteasomes (Cooper et al., 2009; Jacobson et al., 2009) demonstrate that homogeneous K63 linkages are disassembled faster than K48 linkages. This phenomenon holds even for a branched chain. A predicted outcome of the slower cleavage of K48 linkages is a longer residence on the proteasome's 19S regulatory complex, which may be important to allow sufficient time to prepare the substrate for degradation.

In principle, two or more (distal) Ub units attached to the same (proximal) Ub could be positioned to form a recognition surface or Ub–Ub interaction that is only available via branching (see Figure S11). However, in the case of the [Ub]<sub>2</sub>–<sup>48,63</sup>Ub system studied here, we detected no interaction between the two distal Ubs. Furthermore, the UBA(2) and Rap80 tUIM receptors selectively engaged Ub–Ub units linked via Ub K48 or K63, respectively, and bound in the same manner as with the di-Ub controls. Nevertheless, the potential exists for receptors, yet to be identified, to bind a branched chain in a completely different way, e.g., across the juxtaposed distal Ubs (in this case, distal K48- and distal K63-linked Ubs). That this was not observed suggests that, at least for the well-characterized receptors studied here, linkage specificity is maintained in the context of branched polyUb.

In order to explore whether a Ub–Ub interface could be formed between two distal Ubs attached to the same proximal Ub, we generated in silico structural models for all 28 possible branched tri-Ub chains (Table S1). We found that some combinations of linkages (e.g., K11 and K33, or K27 and K29) theoretically could promote close hydrophobic-patch contacts between the distal Ubs while simultaneously freeing the proximal Ub of any noncovalent interactions with the distal Ubs (Figure S11). Although we did not detect such contacts for [Ub]<sub>2</sub>–<sup>48,63</sup>Ub, the possibility of the distal units forming a Ub–Ub interface or novel binding interactions with receptors cannot be excluded for other branched chains. It should be emphasized that here we studied only the smallest possible branched chain. It is possible that an extension of a branched tri-Ub from one or both distal ends or from the proximal end could promote additional intrachain interactions.

Although this study is just a starting point for the structural investigation of unbranched and branched polyUb chains of mixed linkages, evidence of physiological relevance of such chains is mounting (Ben-Saadon et al., 2006; Dammer et al., 2011; Goto et al., 2010; Kravtsova-Ivantsiv and Ciechanover, 2012; Newton et al., 2008; Winborn et al., 2008). One cannot discount the possibility that the polyUb signal is constantly subject to editing or remodeling, and some branched or other mixed-linkage chains could simply be accidents that, in analogy to mismatched bases in DNA or misfolded proteins, are eventually corrected by cellular machinery. Nonetheless, as we have demonstrated here, linkage mixing or branching could enhance the signaling capability of polyUb. That mixed-linkage chains can carry multiple recognition signals is an exciting concept opening a new perspective on ubiquitin signaling.

## EXPERIMENTAL PROCEDURES

### Assembly and Purification of [Ub]<sub>2</sub>–<sup>48,63</sup>Ub, Ub–<sup>63</sup>Ub–<sup>48</sup>Ub, and Ub–<sup>48</sup>Ub–<sup>63</sup>Ub

The tri-Ub chains were assembled using enzyme-catalyzed reactions with full control of chain length, linkage composition, and the location of the isotope-enriched Ub unit, as detailed in Supplemental Information.

### Preparation of Linkage-Selective Receptors

Rap80-tUIM was prepared as described (Sims and Cohen, 2009). C121 on the C-terminal end of the second UIM was mutated to a tyrosine to prevent disulfide bond formation and to facilitate quantification by absorbance measurement at 280 nm. UBA(2) of hHR23A was expressed as a GST-fusion in *E. coli* as described (Varadan et al., 2004).

**26S Proteasome Deubiquitination Assay**

To follow 26S proteasome deubiquitination activity, tri-Ub molecules were incubated with purified *Saccharomyces cerevisiae* 26S proteasomes in an ATP regenerating system buffer at 30°C in ~1:10 proteasome/substrate ratio. Samples were taken at indicated time points and denatured in an SDS and 6 M urea loading buffer. Following low voltage (30 V for 2 hr) transfer to nitrocellulose membranes, the blots were blocked with 5% low-fat milk powder, and incubated either with anti-Ub (Dako, diluted 1:1,000), anti-K63Ub, or anti-K48Ub (Genentech Inc, diluted 1:500) and analyzed by enhanced chemiluminescence reaction.

**Linkage-Selective DUB Assay**

Recombinant human otubain-1 (OTUB1) and GST-AMSH were expressed in *E. coli* and purified essentially as described (McCullough et al., 2004; Wang et al., 2009). Deubiquitination reactions contained 1 µg/µl of polyUb substrate incubated at 37°C with OTUB1 (0.04 µg/µl) or GST-AMSH (~0.1 µg/µl) in a pH 7.5 buffer (50 mM NaHEPES, 0.05% BSA, and 2.5 mM DTT). After 2.5 hr, reactions were stopped with SDS-PAGE sample buffer and analyzed by SDS-PAGE and Coomassie staining.

**NMR Experiments**

All NMR experiments were performed at 23°C on a Bruker Avance III 600 spectrometer equipped with a cryogenic TCI probe. All experiments at pH 6.8 used protein samples in 20 mM sodium phosphate buffer (pH 6.8), containing 0.02% (w/v) Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub> and 5% D<sub>2</sub>O. For experiments at pH 4.5, proteins were exchanged into buffer containing 20 mM sodium acetate, 0.02% (w/v) Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub>, and 5% D<sub>2</sub>O, adjusted to pH 4.5 with acetic acid. See Supplemental Information for details on the NMR experiments.

**Chemical-Shift Perturbations**

Differences between <sup>1</sup>H-<sup>15</sup>N NMR spectra of two species (A and B) were quantified as chemical-shift perturbations, defined as follows: CSP =  $\{(\delta_{HA} - \delta_{HB})^2 + [(\delta_{NA} - \delta_{NB})/5]^2\}^{1/2}$ , where  $\delta_H$  and  $\delta_N$  are chemical shifts of <sup>1</sup>H and <sup>15</sup>N, respectively, for a given backbone N-H group. The same equation was used to quantify spectral perturbations upon titration; in this case, A refers to the unbound species, and B corresponds to various steps in the titration.

**SUPPLEMENTAL INFORMATION**

Supplemental Information includes Supplemental Experimental Procedures, one table, and 11 figures and can be found with this article online at <http://dx.doi.org/10.1016/j.str.2013.02.019>.

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