Selective Proteasomal Degradation of the B’β Subunit of Protein Phosphatase 2A by the E3 Ubiquitin Ligase Adaptor Kelch-like 15*

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Background: Proteasomal degradation of PP2A regulatory subunits has been described, but responsible E3 ubiquitin ligases have remained elusive.

Results: KLHL15 is an E3 ubiquitin ligase adaptor targeting the B’/B56β regulatory subunit for proteasomal degradation, promoting formation of alternative PP2A holoenzymes.

Conclusion: KLHL15 contributes to brain-specific expression of B’β and modifies PP2A holoenzyme composition.

Significance: E3 ligase-mediated B subunit degradation is a novel mechanism to remodel the PP2A heterotrimer.

Protein phosphatase 2A (PP2A), a ubiquitous and pleiotropic regulator of intracellular signaling, is composed of a core dimer (AC) bound to a variable (B) regulatory subunit. PP2A is an enzyme family of dozens of heterotrimerers with different subcellular locations and cellular substrates dictated by the B subunit. B’β is a brain-specific PP2A regulatory subunit that mediates dephosphorylation of Ca2+/calmodulin-dependent protein kinase II and tyrosine hydroxylase. Unbiased proteomic screens for B’β interactors identified Cullin3 (Cul3), a scaffolding component of E3 ubiquitin ligase complexes, and the previously uncharacterized Kelch-like 15 (KLHL15). KLHL15 is one of ~40 Kelch-like proteins, many of which have been identified as adaptors for the recruitment of substrates to Cul3-based E3 ubiquitin ligases. Here, we report that KLHL15-Cul3 specifically targets B’β to promote turnover of the PP2A subunit by ubiquitylation and proteasomal degradation. Comparison of KLHL15 and B’β tissue expression profiles suggests that the E3 ligase adaptor contributes to selective expression of the PP2A/Bβ holoenzyme in the brain. We mapped KLHL15 residues critical for homodimerization as well as interaction with Cul3 and B’β. Explaining PP2A subunit selectivity, the divergent N terminus of B’β was found necessary and sufficient for KLHL15-mediated degradation, with Tyr-52 having an obligatory role. Although KLHL15 can interact with the PP2A/B’β heterotrimer, it only degrades B’β, thus promoting exchange with other regulatory subunits. E3 ligase adaptor-mediated control of PP2A holoenzyme composition thereby adds another layer of regulation to cellular dephosphorylation events.

PP2A3 is one of four major serine/threonine phosphatases (in addition to PP1, PP2B, and PP2C) (1) and comprises up to 1% of total protein in mammalian cells (2). Predominantly found as a heterotrimer containing a core dimer of catalytic (C) and scaffolding (A) subunits, PP2A is a widely distributed and critical mediator of many signaling pathways. A third, variable B subunit defines the substrates and localization of the holoenzyme in the cell (3–5). PP2A B regulatory subunits are subdivided into three gene families, B (PR55/PPP2R2), B’ (B56/PR61/PPP2R5), and B” (PR72/PPP2R3), with little sequence or structural similarity between them. In mammals each gene family is comprised of four to five highly similar genes. Additional complexity arises through alternative splicing. The five mammalian B’ gene products (α/β/γ/δ/ε) feature a conserved, tandem α-helical central domain that binds to the AC dimer (6, 7). The variable N and C termini contain subcellular localization sequences and phosphorylation sites for isoform-specific regulation (8). Upon down-regulation of the A or C subunit, PP2A B and B’ subunits are rapidly degraded by the proteasome (9–11). However, whether proteasomal degradation constitutes a physiological PP2A regulatory mechanism involving dedicated E3 ligases has remained unclear.

We previously identified PP2A/B’β as a brain-specific PP2A holoenzyme that catalyzes the dephosphorylation and inactivation of tyrosine hydroxylase, the rate-limiting enzyme in dopamine synthesis (12), and uncovered B’β residues that mediate substrate specificity (7). Because PP2A/B’β could be a target for intervention into Parkinson disease, a hallmark of which is loss of dopamine in the striatum, we set out to discover novel interactors of this regulatory subunit by proteomic approaches.

These screens identified the E3 ubiquitin ligase scaffold Culin3 (Cul3) and an uncharacterized protein, KLHL15. KLHL15 belongs to a large family of proteins with a characteristic domain architecture consisting of an N-terminal BTB (broad-complex, tram track, and bric-a-brac) domain, a C-terminal

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3 The abbreviations used are: PP2A, protein phosphatase 2A; Cul3, Culin3; KLHL15, Kelch-like 15; BTB, broad-complex, tram track and bric-a-brac; CHX, cycloheximide.
Kelch β-propeller, and a middle BACK (between BTB and C-terminal Kelch) domain (13, 14). Several KLHL proteins have been shown to bridge Cul3-containing E3 ligases to specific substrates (15–21). Here we report on the characterization of the KLHL15-Cul3 complex as a B subunit-specific regulator of PP2A. We propose E3 ligase-mediated exchange of regulatory subunits as a novel mechanism to control PP2A holoenzyme composition and downstream dephosphorylation events.

**EXPERIMENTAL PROCEDURES**

**Cell Culture and Transfection—**HEK293 cells were cultured (37 °C and 5% CO2) in Dulbecco’s modified Eagle’s medium (Invitrogen) with 10% (v/v) fetal bovine serum. Cells were grown to 60% confluency on collagen-coated plates and transfected using Lipofectamine 2000 (Invitrogen) following the manufacturer’s protocol for transient transfection of adhered cells. 24 – 48 h post-transfection, cells were washed in phosphate-buffered saline and harvested. For experiments including MG132 (Sigma), cells were incubated with 50 μM concentrations of the drug (from a 50 mM stock in dimethyl sulfoxide, DMSO) for the final 12 h before lysis.

**Plasmids—**The green fluorescent protein (GFP)-tagged KLHL1 and KLHL24 plasmids were kindly provided by Dr. Michael Koob (University of Minnesota, Minneapolis, MN) and Dr. Fernando Laezza (Washington University, St. Louis, MO), respectively. The HA-Cul2, HA-Cul3, and HA-Cul3Δroc constructs were kind gifts of Dr. Mark Hannink (University of Missouri, Colombia, MO). The coding sequence of KLHL15 was amplified from HEK293 RNA isolated with TRIzol reagent (Invitrogen) using the one-step RT-PCR kit from Qiagen. Both constructs were verified by sequencing. All constructs were expressed shRNA and RNAi-resistant GFP-KLHL15. All mutations were introduced into the gene replacement plasmid expressing shRNA and RNAi-resistant GFP-KLHL15. All constructs were verified by sequencing.

**Antibodies—**Antibodies from commercial sources include: HA epitope (mouse monoclonal, Santa Cruz Biotechnology), GFP (rabbit polyclonal, Abcam), FLAG (rabbit polyclonal, Cell Signaling; mouse monoclonal M2, Sigma), EE epitope (mouse monoclonal, Covance), PP2A/A subunit (rabbit polyclonal, Cell Signaling), PP2A/C subunit (mouse monoclonal, BD Biosciences). The rabbit polyclonal antibody against B’β was described previously (12). To generate antisera specific to KLHL15, a GST fusion of the Kelch domain (amino acids 255–604) was expressed in *Escherichia coli* and purified on glutathione-agarose (Pierce/Thermo Scientific). Rabbit polyclonal antisera were raised at the Iowa State Hybridoma Facility (Ames, IA) and affinity-purified over an antigen column. To this end, GST-KLHL15255–604 was coupled to diaminodipropylamine (DADPA) resin using 1-ethyl-3-[3-dimethylaminopropyl]carbodiimide hydrochloride (EDC) cross-linking agent (Pierce/Thermo Scientific) according to the manufacturer’s instructions. Antisera diluted 1:1 in PBS were passed over the antigen column, and antibody was eluted with 20 mM glycine, pH 2.5. The eluate was neutralized with Tris base and concentrated using centrifugal filter columns (Amicon/Millipore).

**AFFINITY PURIFICATION COUPLED TO MASS SPECTROMETRY—**pcDNA5-FLAG-B’β (PP2R5B, BC045619) was stably transfected in T-Rex Flp-In 293 cells (Invitrogen) and submitted to affinity purification with FLAG antibody coupled to magnetic beads as described previously (27). pcDNA3-FLAG-B’β was stably transfected into HEK293 cells and submitted to affinity purification with FLAG antibodies coupled to aagarose beads (M2, Sigma) as described previously (28). Two biological replicates of the samples from the T-Rex Flp-In 293 cells were analyzed on an LTQ mass spectrometer, and one HEK293 sample was analyzed on an LTQ-Orbitrap mass spectrometer. In parallel several negative control runs (cells expressing the tag alone and/or the tag fused to GFP) were analyzed on the respective mass spectrometers. All acquisition conditions have been described previously (29). A database search was performed using the Mascot search engine against the human and adenovirus complements of the RefSeq library (v45), allowing for deamidation (NQ) and oxidation (M) with trypsin fixed as an enzyme, and one missed cleavage allowed. For LTQ data, the MS tolerance was fixed at 3 Da, and the MS/MS tolerance was fixed at 0.6 Da. For LTQ-Orbitrap data, the MS tolerance was fixed at 12 ppm, whereas the MS/MS tolerance was fixed at 0.6 Da. Hits identified in any of the negative control runs were removed from the list of putative interactors, and only those hits detected in all three of the biological replicates with at least 2 unique peptides were kept. Furthermore, proteins identified in ≥4% of 275 AP-MS analyses performed under the same conditions were deemed potential frequent fliers and removed from further analysis. With the exception of the core PP2A enzymes, only two proteins were identified, CUL3 and KLHL15.

**Immunoprecipitation and Immunoblot Analyses—**For immunoprecipitation, cells were lysed in buffer containing 20 mM Tris, pH 7.5, 150 mM NaCl, 1 mM EDTA, 1 mM EGTA, 1 mM phenylmethylsulfonlfur fluoride, 1 μg/ml leupeptin, 1 mM benzamidine, and 1% (v/v) Triton X-100. Lysates were allowed to rotate at 4 °C for 30 min to complete solubilization. Samples were then centrifuged at 13,000 × g for 15 min to pellet debris. A portion of the supernatant was added directly to SDS-PAGE sample buffer to provide an input sample. HA-tagged and FLAG-tagged proteins were immunoprecipitated as indicated for each experiment from HEK293 cells overexpressing the
indicated proteins via HA or FLAG antibody EZ-view Red Agarose gel (Sigma) for 4 h rotating at 4 °C. EE-tagged proteins were immunoprecipitated with protein A/G-Sepharose (Santa Cruz Biotechnology) preincubated with 4 µg of EE antibody. V5-tagged proteins were immunoprecipitated with goat anti-V5-agarose (Abcam). Beads were washed 4 times by centrifugation in lysis buffer and eluted by boiling in SDS-PAGE sample buffer. Samples were separated on 10% (w/v) SDS-PAGE gels, electrotransferred to nitrocellulose, and immunoblotted as indicated. Proteins were visualized using species-specific secondary antibodies conjugated to IRDye 680/800 (LI-COR) and a LI-COR Odyssey infrared scanner. Signals were quantified by densitometry using the gel analysis plugin of ImageJ (National Institutes of Health).

**RESULTS**

**KLHL15 Specifically Associates with B′β—**Mass spectrometry was carried out to identify proteins that associate with B′β. To this end, FLAG epitope-tagged B′β was stably expressed in HEK293 cells followed by FLAG immunoprecipitation and elution of precipitated proteins with ammonium carbonate. Triptipic peptides were then subjected to liquid chromatography and tandem mass spectroscopy. B′β-interacting proteins reliably identified in three independent experiments were the PP2A A and C subunits and Cul3 as well as KLHL15 (Table 1). KLHL15 is a member of the Kelch-like or BTB-Kelch protein family that is encoded by 39 genes in humans. As first shown for Keap1/ KLHL19 (16, 17, 30), these proteins function as substrate adaptors in Cul3-based E3 ubiquitin ligases (13, 31, 32).

To assess the specificity of the interaction between B′β and KLHL15, the human KLHL15 cDNA was isolated by RT-PCR, tagged at the N terminus with GFP and V5 epitopes, and transiently coexpressed with the five HA-tagged B′ family subunits (α–ε). Remarkable considering the high sequence similarity between B′ isoforms (60–80%), KLHL15 selectively coimmunoprecipitated B′β (Fig. 1A). Further analysis confirmed that members of the distantly related B and B′ subunits of PP2A do not interact with KLHL15 (data not shown). To test whether B′β discriminates among KLHL-family members, FLAG-B′β was coexpressed with GFP-tagged KLHL1, KLHL15, or KLHL24. Only KLHL15 coimmunoprecipitated robustly with B′β (Fig. 1B), suggesting that KLHL15 functions as a highly selective E3 ubiquitin ligase adaptor for B′β. Interrogating the fate of ubiquitylated B′β, we immunoisolated the FLAG-tagged protein from HEK293 cells also expressing HA-tagged ubiquitin after incubation with chloroquine or MG132 (50 µM, 12 h) to inhibit lysosomal or proteasomal degradation, respectively. Proteasome, but not lysosome inhibition, resulted in the accumulation of heterogeneous, high molecular weight species positive for both FLAG and HA epitopes (Fig. 1C).

**KLHL15 Promotes B′β Ubiquitylation and Degradation—**Because other KLHL proteins act as substrate recognition modules for an E3 ligase core complex composed of Cul3 and Roc1 (13), we sought to determine whether KLHL15 regulates B′β levels via ubiquitylation and proteasomal degradation. First, we generated KLHL15-directed polyclonal antibodies and effective shRNAs to confirm expression of endogenous KLHL15 in HEK293 cells (Fig. 2A). To assess whether B′β is ubiquitylated in a KLHL15-dependent manner, FLAG-B′β was immunoprecipitated from HEK293 cells also transfected with GFP-KLHL15, KLHL15-directed shRNAs, or control scrambled shRNAs. Proteasome inhibition by MG132 before cell lysis

| Gene  | Mascot score | Spectral counts | Unique peptides | Coverage % |
|-------|--------------|-----------------|----------------|------------|
| PPP2R5B | 12/19/933 | 8/6/45          | 18/13/13       | 54/36/35 |
| KLHL15 | 258/197/269 | 12/7/5          | 5/7/5          | 12/9/10  |
| CUL3   | 375/38/395  | 8/9/8           | 5/7/7          | 11/9/12  |
resulted in the appearance of multiple ubiquitin- and FLAG-immunoreactive species larger than unmodified B′β. Compared with control shRNA-transfected cells, KLHL15 knockdown decreased the abundance of higher molecular weight forms of B′β, whereas GFP-KLHL15 overexpression enhanced B′β ubiquitylation (Fig. 2B). GFP-KLHL15 recovered in FLAG-B′β immunoprecipitates migrated close to its predicted molecular weight, as well as a smear near the stacker/running gel interface. This suggests that KLHL15 undergoes autoubiquitylation, as was reported for other KLHL proteins (15, 33–36). To test whether KLHL15-mediated ubiquitylation promotes B′β protein turnover, we assessed B′β levels after inhibition of protein synthesis by cycloheximide (CHX) for up to 8 h (Fig. 2, C and D). As a loading control, B′β signals were normalized to extracellular signal-regulated kinase (ERK1/2) signals in the same lane, which were stable under these conditions (as compared with Ponceau S-stained total protein). Compared with cotransfection with a non-targeting shRNA, overexpression of KLHL15 lowered steady-state levels of B′β from ∼8 to ∼2.5 h. Conversely, silencing of endogenous KLHL15 increased B′β levels at steady state and stabilized the protein such that ∼90% of the protein remained after 8 h in CHX (Fig. 2, C and D). Comparison of the area under the curve (AUC) of the B′β degradation time courses as well as the 5- and 8-h time points revealed significant differences between KLHL15 overexpression, silencing, and control (Fig. 2D). Together these data indicate that KLHL15 recruits a Cul3-based E3 ligase to promote polyubiquitylation and proteasomal degradation of specifically the B′β subunit of PP2A. Near complete stabilization of B′β in the absence of KLHL15 further demonstrates that B′β degradation is principally mediated by ubiquitylation via KLHL15, at least in HEK293 cells.

KLHL15 May Contribute to Brain-specific Expression of B′β—We previously showed that B′β protein is abundantly expressed in the brain but undetectable in other tissues even after microcystin affinity purification of PP2A holoenzymes (12). We, therefore, wondered whether posttranslational regulation by KLHL15 may contribute to selective expression of B′β in the nervous system. Quantitative RT-PCR analysis showed that B′β mRNA is fairly uniformly expressed in a panel of nine rat tissues (Fig. 3A). KLHL15 mRNA expression is also ubiquitous, with the highest levels in lung, muscle, and spleen (Fig. 3A). Transcript profiles were similar regardless of whether β-actin, phosphoglycerate kinase 1 (Pgk1) or hypoxanthine phosphoribosyltransferase 1 mRNA was used for normalization (Fig. 3A and not shown). Plotting the ratio of B′β to KLHL15 mRNA levels revealed that forebrain, brainstem, and cerebellum contain more B′β than KLHL15 mRNA, whereas the opposite holds for all non-neuronal tissues examined (Fig. 3B). Because
transcripts are ubiquitous, brain-specific expression of the PP2A/B protein (12) may, therefore, at least in part result from relatively low levels of KLHL15 in the brain.

The BTB Domain Mediates KLHL15 Dimerization and Cul3-dependent B’/β Degradation—The mass spectrometry-based identification of both KLHL15 and Cul3 in B’/β immunoprecipitates is in line with previous evidence that BTB-Kelch proteins tether Cul3-containing E3 ubiquitin ligases to their targets (13, 31, 32). To confirm an interaction between Cul3 and KLHL15, overexpression promotes the appearance of large, ubiquitylated B’/β species (FLAG and ubiquitin-positive, indicated by brackets), whereas KLHL15 shRNA decreases B’/β ubiquitylation. C and D, FLAG-B’/β was coexpressed with GFP-KLHL15, non-targeting, control shRNA, or KLHL15-directed shRNA-expressing plasmids (1:1 mass ratio) followed by inhibition of protein synthesis by CHX (100 μg/ml) for 0–8 h. Total cell lysates were probed for the indicated proteins (representative blots in C), and B’/β degradation was quantified by dividing FLAG signals by ERK1/2 signals in the same lane and normalizing to the zero time point (D). Both individual time points and area under the curve (normalized to control) are shown as the means ± S.E. of four independent experiments. *, p < 0.05; **, p < 0.01 by Student’s t test. Molecular mass marker positions are shown in kDa. Asterisks on blots indicate nonspecific bands. Arrowheads in C point to transfected (xfect.), GFP-tagged, and endogenous (endo.) KLHL15.

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Cul3 ΔRoc, a dominant-negative C-terminal deletion mutant unable to bind the E3 core component Roc1. GFP-KLHL15 also preferentially associated with Cul3 over Cul2, underscoring Cul3 preference as a feature shared among KLHL family proteins.

To explore the role of the KLHL15 BTB domain in regulation of PP2A/B’/β, conserved amino acids were targeted by site-directed mutagenesis. Guided by a threading model of the BTB domain dimer (Fig. 4B), we mutated two buried residues near the predicted dimerization interface (D32A, H45L) as well as three adjacent exposed residues (Ile-72, Leu-73, Lys-74; ILK72AAA). To avoid confounds due to dimerization of endogenous with transfected, mutant KLHL15, we used a single plas-
mid, RNAi-based gene replacement strategy (23). To this end, H1 promoter-driven shRNA targeting KLHL15 (Fig. 2A) was expressed from the same plasmid as the mutant GFP-KLHL15 cDNA, which was rendered RNAi-resistant by silent mutagenesis of the RNAi target site. HEK293 cells coexpressing FLAG-tagged KLHL15 (Fig. 4) were obtained with a second set of KLHL15 primers (not shown).

FIGURE 3. Brain-specific expression of PP2A/B′β may arise from low KLHL15 mRNA levels in the brain. A and B, shown is quantitative RT-PCR analysis of KLHL15 (gray) and B′β (black) mRNA levels in the indicated rat tissues. In A, KLHL15 and B′β mRNA levels were normalized to phosphoglycerate kinase 1 (Pgk1) mRNA levels. In B, B′β is plotted relative to KLHL15 mRNA (means ± S.E. of 4–8 independent experiments). Essentially identical results were obtained with a second set of KLHL15 primers (not shown).

To confirm that KLHL15 and B′β interact directly via the Kelch domain and the divergent N terminus, respectively, we expressed and purified from *E. coli* His6-KLHL15255–604 and (ILK72AAA) or dimerization (D32A) of KLHL15 significantly slowed turnover of B′β (t1/2 ≈ 8 h, Fig. 4, E and F). These results indicate that KLHL15 mediates proteasomal degradation of B′β via Ile-72, Leu-73, Lys-74-dependent scaffolding of the Cul3-Roc1 E3 core complex, with Asp-32-dependent homodimerization also contributing to KLHL15 activity.

The Top of the KLHL15 B′Propeller Mediates B′β Association—According to high confidence structure modeling templated on the Kelch repeats of Keap1/KLHL19 (38, 39), the C terminus of KLHL15 folds into a cone-shaped β-propeller with six propeller blades composed of four antiparallel β-strands each (Fig. 5A). Keap1/KLHL19 was previously shown to interact with a peptide derived from its substrate Nrf2 via multiple, blade-spanning residues that form the top cavity of the Kelch β-propeller (39). The corresponding residues are perfectly conserved among KLHL15 orthologs in vertebrates but are distinct in other KLHL family members (Fig. 5B) and thus may function as substrate specificity determinants. A subset of these putative substrate binding residues mapping to Kelch blades 1 and 2 (asterisks in Fig. 5B) was assessed for effects on B′β interaction and down-regulation. As with BTB domain mutants, Kelch domain mutant GFP-KLHL15 replaced endogenous KLHL15 via shRNA expressed from the same plasmid. R318E, E335R/L337A, and E371R substitution strongly impaired association of GFP-KLHL15 with FLAG-B′β by communoprecipitation (Fig. 5C) and KLHL15-mediated degradation of B′β (t1/2 > 8 h, Fig. 5, D and E). On the other hand, mutation of a residue of similar evolutionary conservation but predicted localization at the bottom of the KLHL15 β-propeller (E518R) had no interaction phenotype (Fig. 5C). Thus, conserved residues unique to KLHL15 form a substrate binding pocket predicted to reside on top of the Kelch β-propeller.

The B′β N Terminus Interacts Directly with KLHL15—B′β subunits share a highly conserved central portion composed of tandem α-helical repeats that interacts with the PP2A A and C subunits (6). Because KLHL15 associates with B′β, but not other B′ isoforms (Fig. 1A), we surmised that instead, the divergent N- or C-terminal extensions of B′β contain the KLHL15 binding domain. In support, initial communoprecipitation experiments with truncated B′β cDNAs mapped critical residues to the N terminus (amino acids 31–65, Fig. 6A). Interestingly, truncation of the first 30 residues consistently enhanced the interaction of B′β with KLHL15, suggesting perhaps that the extreme N terminus folds back to occlude an adjacent KLHL15 binding domain.

Sequence alignments pointed to B′β Tyr-52 as a probable binding determinant, as this residue is perfectly conserved among B′β orthologs from fish to man but is divergent within the B′ family (Fig. 6A). Indeed, replacement of Tyr-52 with the corresponding residue in B′ε (Y52S), the isoform most similar to B′β, eliminated KLHL15 association (Fig. 6A). B′β degradation assays corroborated these findings, demonstrating enhanced stability of B′β Δ1–65 and Y52S compared with wild-type B′β or the Δ1–30 truncation (~80% versus ~40% remaining after 10 h in CHX; Fig. 6, B and C).

To confirm that KLHL15 and B′β interact directly via the Kelch domain and the divergent N terminus, respectively, we expressed and purified from *E. coli* His6-KLHL15255–604 and
GST-B′β1–82 (GST-B′β1–68 as a negative control). In glutathione-agarose pulldown experiments, GST-B′β1–82 precipitated wild-type KLHL15255–604 effectively, whereas GST-B′β1–68 displayed no detectable interaction. Paralleling association studies in intact cells (Fig. 5C), the E371R substitution weakened the association between the KLHL15-propeller and the B′β N terminus (Fig. 6D).

Because the first 65 residues are dispensable for incorporation of B′β into the PP2A heterotrimer (7), we next asked whether KLHL15 discriminates between free and holoenzyme-associated B′β. To this end, charge reversal substitutions were incorporated into B′β at tandem α-helical repeat 1 (KR103DE) and 4 (RK232ED) to block A and C subunit association (Fig. 6F and data not shown). Highly conserved across species and B′ subunit isoforms, but not contacting A or C subunits directly (40, 41), these residues instead appear to be necessary for proper folding of the two tandem α-helical repeats. Both monomeric B′β mutants efficiently associated with KLHL15; furthermore, the endogenous PP2A C subunit was detected in KLHL15 immunoprecipitations when wild-type B′β was coexpressed (Fig. 6F). Therefore, KLHL15 targets B′β regardless of whether or not the regulatory subunit is incorporated into the PP2A heterotrimer.

The B′β N terminus Is Sufficient for KLHL15-mediated Degradation—We next inquired whether the B′β N terminus contains a degron, i.e. sequence determinants sufficient for deg-
radiation. To this end we assessed turnover of the B′/β N terminus (amino acids 1–82) fused to GFP after overexpression or knockdown of KLHL15 (Fig. 7A). About 40% of GFP-B′/β1–82 degraded when protein synthesis was inhibited by CHX for 8 h. In contrast, GFP fusions of the B′/β N terminus (amino acids 1–68) or the B′/β C terminus (amino acids 476–497) were stable over this time course (Fig. 7, A–D). KLHL15 silencing prevented degradation of GFP-B′/β1–82, whereas KLHL15 overexpression accelerated degradation. Neither silencing nor forced expression of KLHL15 had an impact on the stability of GFP-B′/β476–497 or GFP-B′/β1–68. Therefore, the N terminus contains both E3 ligase docking and ubiquitylation sites sufficient for degradation of B′/β.

**KLHL15 Does Not Degrade the PP2A Holoenzyme but Influences Its Composition**—Because KLHL15 can associate with the B′/β-containing PP2A holoenzyme (Fig. 6F), we examined whether KLHL15 mediates wholesale degradation of the PP2A/B′/β heterotrimer. In cells transfected with B′/β, neither endogenous nor epitope-tagged A and C subunit levels were modulated by KLHL15 overexpression or knockdown (data not shown). However, this negative result could be explained by B′/β competing with endogenous regulatory subunits for association with A and C subunits such that only a small pool of A and C subunits becomes subject to proteasomal degradation via KLHL15. To provide B′/β with privileged access to the A subunit, we employed an A subunit point mutant (DW139RR) that interacts with B′/β but not B or B′/β subunits (22). EE-tagged A/β DW139RR, FLAG-B′/β1–82, and either GFP-KLHL15, control shRNA, or KLHL15-directed shRNA were coexpressed in HEK293 cells. Immunoblotting for the EE epitope revealed little to no effect of KLHL15 levels on A subunit stability in the presence of CHX (Fig. 8, A and B), whereas an inverse relation—
ship between KLHL15 and B\'/H11032/H9252 levels was confirmed by blotting for FLAG (Fig. 8, A and C). Thus, even though KLHL15 can bind to the PP2A/B\'/H11032/H9252 holoenzyme, the E3 adaptor only ubiquitylates and degrades the regulatory subunit.

To provide direct evidence that KLHL15 can impact PP2A holoenzyme composition, we assessed competition of B\'/H11032/H9252 and B\'/H11032/H9254 for incorporation into the PP2A heterotrimer as a function of KLHL15 levels. HA-tagged B\'/H11032 isoforms were coexpressed with wild-type EE-tagged A subunit, and cell lysates were subjected to EE-tag immunoprecipitation and immunoblotting (Fig. 8D). Association of both endogenous and HA-tagged B\'/H11032/H9252 with the A subunit increased when KLHL15 was silenced (1.4–2.2-fold, \( n = 2 \)), whereas PP2A holoenzyme incorporation of B\'/H11032/H9254 increased when KLHL15 was overexpressed (1.6–2.0-fold, \( n = 2 \)). Therefore, by virtue of degrading B\'/H11032, KLHL15 creates a window of opportunity for other regulatory subunits with distinct subcellular localization and substrate specificity to incorporate into the PP2A heterotrimer.
Site-directed mutagenesis guided by structure prediction supports the consensus view of KLHL proteins as modular adaptors, with the N-terminal BTB domain mediating homodimerization of KLHL15 and Cul3 recruitment via distinct interaction surfaces and the top cavity of the C-terminal Kelch β-propeller docking to the substrate, B′β. Predictably, interfering with either B′β or Cul3 interaction abrogated KLHL15-mediated down-regulation of B′β. Unexpectedly, a point mutation in the BTB domain that specifically blocked KLHL15 self-association (D32A) had the same effect (Fig. 4, E and F). Consistent with this finding, missense mutations in KLHL10 linked to male sterility were recently shown to interfere with homodimerization of the protein (43). Perhaps substrate ubiquitylation by KLHL proteins is an obligatory trans reaction, with one subunit of the dimer directing Cul3-Roc1 for ubiquitin transfer to a substrate bound by the other subunit.

This report also defines a novel regulatory mechanism for PP2A, one of the major Ser/Thr phosphatases. With at least 12 genes encoding regulatory subunit in mammals, PP2A comprises a family of signaling enzymes with unique substrate specificity, localization, and regulation by second messengers (3–5, 7, 8). We found that KLHL15 displays exquisite specificity for binding and down-regulation of B′β, one of five members of the B′/B56 family of regulatory subunits. PP2A/B′β is enriched in the nervous system, where it was shown to dephosphorylate and inactivate two important enzymes, 1) calcium/calmodulin-dependent protein kinase II, a key mediator of long term potentiation and spatial memory (44), and 2) tyrosine hydroxylase, the rate-limiting enzyme in dopamine synthesis (12). According to qRT-PCR, B′β mRNA is relatively uniformly expressed among across tissues (Fig. 3A), whereas B′β protein is only detectable in the brain (12). Because brain is the only tissue with higher B′β than KLHL15 mRNA levels, we propose that the brain-specific expression of the B′β protein is at least in part a result of efficient KLHL15-mediated proteasomal degradation in other tissues.

KLHL15 likely not only determines cell and tissue specificity of B′β expression but may also regulate abundance of this PP2A regulatory subunit dynamically and in response to signaling events. This could for instance occur through B′β phosphorylation at Tyr-52, an essential KLHL15 binding determinant (Fig. 6, B–D), or at nearby Ser-44, Ser-46, and Ser-47, Ser residues that have been identified in unbiased phosphoproteomic screens (PhosphoSitePlus).

The β-propeller of KLHL15 directly engages the divergent N terminus of B′β, a region of the regulatory subunit necessary and sufficient for proteasomal degradation (Figs. 6, B and E, and 7, A and B). Accordingly, KLHL15 targets both monomeric and holoenzyme-incorporated B′β (Fig. 6F). However, we found no evidence that KLHL15 degrades the A and C subunit within the PP2A/B′β heterotrimer. Instead, the E3 ligase adaptor appears to abstract B′β from the AC dimer, allowing other regulatory subunits to take the place of B′β (Fig. 8D). Selective ubiquitylation and proteasomal degradation of B′β by KLHL15 may, therefore, be a mechanism for dynamic regulation of PP2A holoenzyme composition and specific substrate dephosphorylation.

On a final note, our results raise the intriguing possibility that each of the dozen or more PP2A regulatory subunit is paired with a dedicated E3 ubiquitin ligase. Indeed, substrates for the
majority of the 39 human KLHL family proteins remain to be identified. KLHL protein-catalyzed exchange of PP2A regulatory subunits could thus shape dephosphorylation profiles in every cell.

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![FIGURE 8. KLHL15 modulates PP2A holoenzyme composition. A and B, FLAG-B′β and an EE-tagged PP2A/A subunit that only associates with B′ regulatory subunits (Ax DW139RR (22)) were coexpressed with GFF-KLHL15, control shRNA, or KLHL15-directed shRNA and assessed for stability in CHX (0–8 h). Representative blots are shown in A, whereas quantification of Ax and B′β levels are shown in B and C, respectively (means ± S.E., n = 3 experiments, see Fig. 2 legend). Although B′β stability is modulated by KLHL15, the stability of the scaffolding A subunit is not. AUC, area under the curve. D, HEK293 cells coexpressing HA-tagged B′β and B′β, EE-tagged Ax, and either GFF-KLHL15, control shRNA, or KLHL15-directed shRNA were subjected to immunoprecipitation (IP) with EE-tag antibodies to isolate PP2A holoenzymes containing the transfected A subunit. KLHL15 silencing promotes B′β, whereas KLHL15 overexpression promotes B′β incorporation into the PP2A heterotrimer. Molecular mass marker positions are shown in kDa. The asterisks indicates the IgG heavy chain. *, p < 0.05; **, p < 0.01 by Student’s t test.](image-url)
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