The autism-linked UBE3A T485A mutant E3 ubiquitin ligase activates the Wnt/β-catenin pathway by inhibiting the proteasome

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UBE3A is a HECT domain E3 ubiquitin ligase whose dysfunction is linked to autism, Angelman syndrome, and cancer. Recently, we characterized a de novo autism-linked UBE3A mutant (UBE3A T485A) that disrupts phosphorylation control of UBE3A activity. Through quantitative proteomics and reporter assays, we found that the UBE3A T485A protein ubiquitinates multiple proteasome subunits, reduces proteasome subunit abundance and activity, stabilizes nuclear β-catenin, and stimulates canonical Wnt signaling more effectively than wild-type UBE3A. We also found that UBE3A T485A activates Wnt signaling to a greater extent in cells with low levels of ongoing Wnt signaling, suggesting that cells with low basal Wnt activity are particularly vulnerable to UBE3A T485A muta-tion. Ligase-dead UBE3A did not stimulate Wnt pathway activation. Overexpression of several proteasome subunits reversed the effect of UBE3A T485A on Wnt signaling. We also observed that subunits that interact with UBE3A and affect Wnt signaling are located along one side of the 19S regulatory particle, indicating a previously unrecognized spatial organization to the proteasome. Altogether, our findings indicate that UBE3A regulates Wnt signaling in a cell context-dependent manner and that an autism-linked mutation exacerbates these signaling effects. Our study has broad implications for human disorders associated with UBE3A gain or loss of function and suggests that dysfunctional UBE3A might affect additional proteins and pathways that are sensitive to proteasome activity.

Loss of UBE3A causes a debilitating neurodevelopmental disorder called Angelman syndrome (1–3), whereas excess UBE3A, through gene duplication or gain-of-function mutation, increases the risk for autism (4–7). Moreover, ectopic activation of UBE3A (also known as E6AP), via the human papillomavirus E6 oncoprotein, contributes to cervical cancers (8). UBE3A is an E3 ubiquitin ligase that targets several substrate proteins, including itself, for proteasomal degradation, increasing the risk for autism (4–7). Moreover, ectopic activation of UBE3A (also known as E6AP), via the human papillomavirus E6 oncoprotein, contributes to cervical cancers (8).

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Figure 1. The canonical Wnt signaling pathway. Drugs used in epistasis experiments are indicated (red). C59 is a porcine inhibitor that suppresses Wnt secretion. XAV939 is a tankyrase inhibitor that stabilizes the destruction complex, LiCl and CT99021 are inhibitors of GSK3β, valproic acid (VPA) is an activator of β-catenin-dependent transcription, and iCRT14 inhibits β-catenin-dependent transcription. TCF/LEF, T cell factor/lymphoid enhancer factor; LRP, low-density lipoprotein receptor-related protein.

of a genome-wide small interfering RNA screen, we found that Wnt pathway activation is highly sensitive to knockdown of specific proteasomal subunits (Ref. 39 and see new analyses of these data below). Moreover, others found that wild-type (WT) UBE3A can stimulate Wnt pathway activation and protect β-catenin from proteasomal degradation with these activities dependent on the ubiquitin ligase activity of UBE3A (40 – 42).

Recently, we characterized a de novo autism-linked UBE3A T485A mutation that disrupts phosphorylation control of UBE3A and enhances UBE3A ubiquitin ligase activity (6). This UBE3A T485A mutation, along with an engineered UBE3A T485E mutation that inhibits UBE3A by mimicking phosphorylation, gave us new molecular tools to probe the link between UBE3A activity and Wnt signaling. Through extensive proteomic and functional experiments, we show that UBE3A and Wnt signaling converge at the proteasome with UBE3A impacting overall protein homeostasis, including β-catenin turnover, by ubiquitinating multiple proteasome subunits. Intriguingly, subunits that interact with UBE3A and affect Wnt signaling are located along one side of the 19S regulatory particle, suggesting functional organization of the proteasome. The UBE3A T485A mutant activated Wnt signaling more effectively than WT UBE3A, and ligase-dead UBE3A failed to activate Wnt signaling, raising the novel possibility that abnormal Wnt signaling contributes to neurodevelopmental disorders involving UBE3A loss or gain of function.

Results

UBE3A T485A enhances Wnt signaling in a cell-context dependent manner

WT UBE3A, but not ligase-dead (LD) UBE3A, was found previously to stimulate Wnt reporter gene expression in HEK293T cells and to do so independently of Wnt ligand (40, 42). Given that a significant number of autism-linked de novo mutations are found in genes associated with the Wnt pathway (14, 17), we sought to determine whether the autism-linked UBE3A T485A mutation, which disables phosphorylation control and hyperactivates ubiquitin ligase activity (6), had an equal or greater effect on Wnt pathway activation. To test this possibility, we transfected HEK293T cells with the β-catenin-activated luciferase reporter (BAR) (43) along with the following UBE3A expression constructs: WT UBE3A, UBE3A-LD, UBE3A T485A, and UBE3A T485E (phosphomimetic mutant; reduces UBE3A activity to near UBE3A-LD levels). We previously characterized the protein level and ubiquitin ligase activity of each construct in HEK293T cells (6). Cells were then acutely (12–16 h) treated with control (L-cell) or Wnt3α-conditioned medium (CM) prior to quantifying luciferase activity. We found that WT UBE3A and the UBE3A T485A mutant strongly stimulated Wnt pathway activation in the absence (Fig. 2A) or in the presence of Wnt ligand (Fig. 2B). The autism-linked UBE3A T485A construct activated Wnt signaling to a greater extent than WT UBE3A, particularly as the amount of plasmid transfected was increased (Fig. 2C). The phosphomimetic UBE3A T485E mutant, which impairs UBE3A ubiquitin ligase activity, was much less effective at stimulating Wnt pathway activation. Consistent with a previous study (40), UBE3A-LD did not activate the pathway (Fig. 2A and B), indicating that UBE3A ubiquitin ligase activity is required to stimulate Wnt pathway activation. Lastly, we found that UBE3A T485A stimulated Wnt pathway activation more than 11-fold (relative to T485E) in cells with low baseline Wnt pathway activation (L-cell control medium) and was relatively less effective in cells treated acutely (12–16 h; 2.6-fold relative to T485E) or chronically (one passage prior to transfection through to lysis; 1.8-fold relative to T485E) with Wnt ligand (Fig. 2D). These data suggest that UBE3A stimulates Wnt signaling in a cell-context dependent manner and is relatively more effective in cells with low levels of ongoing Wnt signaling, consistent with findings by Sherman and co-workers (42).

UBE3A and the Wnt pathway converge at the proteasome

Epistasis experiments suggested that UBE3A stimulated the Wnt pathway independently of GSK3β or adenomatous polyposis coli protein (41). We performed additional epistasis experiments, probing different parts of the pathway (Fig. 1), with the O-acetyltransferase inhibitor CS9 (inhibits Wnt palmitoylation and secretion), tankyrase inhibitor XAV939 (stabilizes the destruction complex), GSK3β inhibitors CT99021 and LiCl, the β-catenin transcriptional activator valproic acid, and the β-catenin transcriptional inhibitor iCRT14 (supplemental Fig. S1). These studies suggested that UBE3A acts at or downstream of the destruction complex and influences the transcriptional activity of β-catenin, consistent with work by Kuslansky et al. (42).

We next sought to identify a molecular link between UBE3A and the Wnt/β-catenin signaling pathway. Given that UBE3A is an E3 ubiquitin ligase that targets proteins for destruction, we hypothesized that any proteins that 1) interact with UBE3A, 2) are ubiquitinated by UBE3A, and 3) activate the Wnt pathway...
when knocked down could represent *bona fide* UBE3A sub-
strates and serve as core components of this signaling pathway. Published data sets addressed two of these criteria. Martínez-
Noël *et al.* (24) identified a comprehensive list of proteins that
selectively interact with UBE3A in HEK293T cells (relative to
NEDD4, another HECT domain E3 ubiquitin ligase). Of these
proteins, 63 interact with all three UBE3A isoforms (WT and
LD; Fig. 3A and supplemental Table S1). We recently identified
a comprehensive list of positive and negative regulators of Wnt
signaling in HEK293T cells from a near-saturation genome-
wide siRNA screen (39). Of these regulators, 1,958 proteins
significantly increased Wnt pathway activation when knocked
down (Fig. 3A and supplemental Table S2). Database for Anno-
tation, Visualization and Integrated Discovery (DAVID) analy-

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Figure 2. The autism-linked UBE3A<sup>T485A</sup> mutant exacerbates Wnt signaling. A and B, HEK293T cells were transfected with the BAR reporter, CMV-<i>Renilla</i>, and empty vector or the indicated UBE3A plasmid. Cells were serum-deprived and treated with control (L-cell) CM (A) or Wnt3a CM (B). Mean values are shown for firefly:<i>Renilla</i> ratios (n = 12). Error bars represent S.D. Statistical analysis was performed using one-way analysis of variance with Bonferroni post hoc correction, ***, p < 0.0005. C, HEK293T cells were transfected with increasing amounts of plasmid encoding WT UBE3A or UBE3A<sup>T485A</sup>. Mean values are shown for firefly:<i>Renilla</i> ratios (n = 3). Error bars represent S.D. Statistical analysis was performed using two-way analysis of variance with Bonferroni post hoc correction, ***, p < 0.0005. D, transfected HEK293T cells were grown for 12–16 h in L-cell CM or Wnt3a CM (acute) or in Wnt3a CM for one passage prior to transfection through lysis (chronic). Values are shown as box and whisker plots for firefly:<i>Renilla</i> ratios (n = 6). Whiskers represent the range of maximum and minimum values obtained in our experiments. Statistical analysis was performed using a two-sample t test (two-tailed). *, p < 0.05; ***, p < 0.0005.

UBE3A physically associates with the proteasome (24–26), and this association can be reversibly altered by certain stimuli (23). To evaluate whether Thr-485 phosphorylation affects the association of UBE3A with the proteasome, we isolated UBE3A protein complexes from HEK293T cells overexpressing LD versions of UBE3A<sup>T485A</sup>, which cannot be phosphorylated, and UBE3A<sup>T485E</sup>, a mutant that mimics phosphorylation. Using mass spectrometry and spectral counts to quantify interactions, we found that the UBE3A<sup>T485A</sup>–proteasome subunit interactions (supplemental Table S3 and supplemental Fig. S2A). To further evaluate whether phosphorylation at Thr-485 affects UBE3A binding to the proteasome, we biochemically purified proteasomes from HEK293T cells expressing LD versions of UBE3A<sup>T485A</sup> and UBE3A<sup>T485E</sup> mutants. UBE3A abundance was measured in proteasome-bound and unbound fractions by Western blot analysis. We found no difference between UBE3A<sup>T485A</sup> and UBE3A<sup>T485E</sup> mutants, suggesting that phosphorylation does not strongly influence UBE3A–proteasome interaction (supplemental Fig. S2, B and C).
Multiple proteasome subunits are ubiquitinated by UBE3A

We next sought to comprehensively identify proteins that are ubiquitinated by UBE3A in HEK293T cells. We combined quantitative stable isotope labeling of amino acids in cell culture (SILAC) with ubiquitin remnant immunoaffinity profiling (45, 46). This approach allowed us to isolate ubiquitinated peptides containing the characteristic diglycine motif exposed after tryptic digestion and to perform ratiometric analysis in cells expressing active UBE3AT485A relative to inactive UBE3AT485E (supplemental Table S4). We found that UBE3A ubiquitinated multiple components of the proteasome, including eight core proteasome subunits that physically interact with UBE3A and that negatively regulate Wnt signaling (Fig. 3B, shown in bold).

Of these eight subunits, we examined endogenous protein abundance of five (PSMA2, PSMB1, PSMD2, PSMD4, and PSMD11) by Western blot analysis in cells transfected with UBE3AT485A or UBE3AT485E expression constructs. Protein levels of most of these proteasome subunits were reduced in UBE3AT485A-expressing cells relative to T485E controls in the absence or presence of Wnt ligand (Fig. 4, A–C). We also examined endogenous levels of these five subunits in immortalized human lymphocytes derived from the UBE3AT485A proband and the unaffected (WT UBE3A) parents (47). We previously found that UBE3A ubiquitinated multiple components of the proteasome, including eight core proteasome subunits that physically interact with UBE3A and that negatively regulate Wnt signaling (Fig. 3B, shown in bold).

Of these eight subunits, we examined endogenous protein abundance of five (PSMA2, PSMB1, PSMD2, PSMD4, and PSMD11) by Western blot analysis in cells transfected with UBE3AT485A or UBE3AT485E expression constructs. Protein levels of most of these proteasome subunits were reduced in UBE3AT485A-expressing cells relative to T485E controls in the absence or presence of Wnt ligand (Fig. 4, A–C). We also examined endogenous levels of these five subunits in immortalized human lymphocytes derived from the UBE3AT485A proband and the unaffected (WT UBE3A) parents (47). We previously found that UBE3A ubiquitinated multiple components of the proteasome, including eight core proteasome subunits that physically interact with UBE3A and that negatively regulate Wnt signaling (Fig. 3B, shown in bold).

Our experiments suggested that PSMD2 (also known as RPN1) was a novel UBE3A substrate. Unlike other 19S components, PSMD2 is located at the edge of the regulatory complex (Fig. 5A, green) and anchors several accessory proteins to the proteasome (48). These accessory proteins include multiple UBE3A substrates, RAD23A and RAD23B (28, 49), ADRM1, PSMC3, UCHL5 (25), PSMD4 (6), and MCM7 (50), as well as proteins that associate with UBE3A, including the E2 enzyme UBE2L3 (51).

In titration experiments with increasing amounts of PSMD2 plasmid, we found that PSMD2 reduced UBE3AT485A-mediated Wnt pathway activation in the absence or presence of Wnt ligand (Fig. 5, B and C). These findings suggest that overexpression of PSMD2 can functionally rescue or compensate for the loss of endogenous PSMD2 in UBE3AT485A-expressing cells. Moreover, high-molecular-weight ubiquitin-conjugated forms of PSMD2 were detected in HEK293T cells expressing UBE3AT485A, PSMD2, and HA-tagged ubiquitin after treatment with the proteasome inhibitor MG-132 (30 μM; 4 h) (Fig. 5D). In contrast, polyubiquitinated PSMD2 was not detected in cells expressing the inactive UBE3AT485E mutant (Fig. 5D). We also performed in vitro ubiquitination reactions with recombi-
nant ubiquitin, E1, E2, and UBE3A enzymes along with PSMD2 purified from HEK293T cells. A single distinct ubiquitin-conjugated PSMD2 species was detected in reactions containing UBE3A (Fig. 5E) that differed from what was observed in cells (Fig. 5D), suggesting that additional cellular factors promote polyubiquitin chain formation on PSMD2. Collectively, these data indicate that PSMD2 is a substrate of UBE3A and that overexpression of PSMD2 dampens Wnt pathway activation in UBE3A*T485A-expressing cells.

UBE3A regulates Wnt signaling through multiple proteasome subunits

We next evaluated the extent to which additional proteasomal subunits rescue Wnt pathway activation in UBE3A*T485A-expressing cells. Plasmids encoding subunits that interact with UBE3A were transfected into HEK293T cells together with UBE3A*T485A at 1:1 and 1:2 UBE3A:subunit ratios or alone to confirm overexpression (supplemental Fig. S4). We were particularly interested in PSMD4 as PSMD4 is a well characterized UBE3A substrate (6, 25, 26, 29). However, PSMD4 failed to reduce Wnt pathway activation in UBE3A*T485A-expressing cells at the 1:1 or 1:2 transfection ratio (Fig. 6; ranked results shown in supplemental Table S5). Instead, PSMD4 augmented Wnt signaling by nearly 200% at the 1:2 transfection ratio. Of the remaining subunits tested, PSMB1, PSMC2, PSMD2, and PSMD7 consistently reduced Wnt pathway activation in UBE3A*T485A-expressing cells, whereas PSMC6 augmented the Wnt response (at 1:1 and 1:2 ratios; Fig. 6 and supplemental Table S5). PSMD2 was one of the most effective subunits to rescue UBE3A-dependent Wnt activation in HEK293T cells (supplemental Table S6). PSMD1, PSMD3, PSMD6, and PSMD11 had no effect or inconsistent effects across the four conditions. Our experiments collectively suggest that UBE3A stimulates Wnt pathway activation by interacting with, ubiquitinating, and reducing the levels of multiple (PSMB1, PSMC2, PSMD2, and PSMD7) proteasome subunits.

UBE3A inhibits proteasome function and stabilizes β-catenin

Several groups found that WT UBE3A inhibits the proteasome in cells (25, 34). To determine whether UBE3A*T485A similarly affected proteasome activity, we transfected HEK293T cells with UBE3A*T485A or UBE3A*T485E mutants along with a
destabilized G76V-ubiquitin (UbG76V)-GFP fusion reporter protein and mCherry as the normalization control. This UbG76V-GFP construct was used previously to quantify proteasome-dependent activity in cells (52). We found that the levels of this UbG76V-GFP reporter were higher in cells transfected with UBE3AT485A relative to the inactive T485E mutant, suggesting that elevated UBE3AT485A activity inhibits proteasome function in cells (Fig. 7, A and B). This difference in UbG76V-GFP reporter levels was abolished in cells treated with MG-132 (10 μM; 4 h), further suggesting a proteasome-dependent mechanism.

β-Catenin is ubiquitinated and then degraded by the proteasome, which limits entry to the nucleus and transcriptional activity (Fig. 1). Given that UBE3AT485A reduces proteasome activity, we hypothesized that nuclear β-catenin levels might be elevated in UBE3AT485A-expressing cells. Indeed, we found that nuclear β-catenin levels were slightly elevated in cells expressing the UBE3AT485A mutant relative to cells expressing UBE3AT485E in the absence of Wnt ligand (Fig. 7, C and D). Additionally, UBE3AT485A less effectively stimulated Wnt signaling (relative to UBE3AT485E) in cells overexpressing WT β-catenin and did not augment Wnt signaling (relative to UBE3AT485E) in cells overexpressing a constitutively active β-catenin construct (Fig. 7E). These data further demonstrate that UBE3AT485A most effectively stimulates Wnt signaling in cells with subsaturating levels of active β-catenin, that is when
baseline Wnt pathway activation is low. Altogether, our results suggest a mechanism whereby UBE3A ubiquitinates multiple subunits of the proteasome, leading to impairment of proteasome-mediated degradation of β-catenin and increased β-catenin/Wnt signaling (Fig. 7F).

**Discussion**

Despite the well known association between excess UBE3A and autism risk (4, 5, 53), including autism-linked mutations that elevate UBE3A alone (6, 7), and extensive research on UBE3A (10), it is currently unclear how excess UBE3A affects mechanisms linked to autism and neurodevelopment. UBE3A is thought to be a promiscuous enzyme (54), and this seeming promiscuity has confounded attempts to identify the physiological substrates of UBE3A. Recently, we found that an autism-linked point mutation in UBE3A (UBE3AT485A) disables a phosphorylation control switch and elevates UBE3A ubiquitin ligase activity (6). Here, we found that UBE3AT485A exacerbates Wnt signaling by inhibiting the proteasome and stabilizing nuclear β-catenin. Numerous autism-linked genes are implicated in Wnt signaling (14, 15, 17); valproic acid, an environmental risk for autism, activates Wnt signaling (supplemental Fig. S1E) (55, 56); and Wnt signaling regulates proliferation and differentiation of neuronal progenitors (57, 58). Our study and the work of others (40–42) firmly place UBE3A within the Wnt signaling pathway and provide fundamental new insights into how excess UBE3A could impair brain development and increase risk for autism.

Our biochemical and proteomics findings suggest that UBE3A stimulates Wnt signaling by ubiquitinating multiple proteasome subunits and reducing proteasome activity, which ultimately stabilize β-catenin. UBE3A T485A-mediated stimulation of Wnt signaling was reduced in cells overexpressing different proteasome subunits, suggesting that proteasome subunits are functionally relevant UBE3A substrates. Multiple laboratories have found that UBE3A can ubiquitinate proteasome subunits and inhibit proteasome function (25, 34), although one group found that UBE3A stimulated proteasome proteolytic activity (27). Discrepancies could relate to differences in cell context or experimental approach. Jacobson et al. (25) found that UBE3A inhibited the proteasome by *in situ* ubiquitination provided the levels of polyubiquitinated proteins in the cell were low. Cellular stresses that increase polyubiquitinated protein levels (oxidative stress and serum starvation) blocked UBE3A from ubiquitinating the proteasome, leading to increased proteasome activity (25). In a simplistic sense, the proteasome can “sense” whether global polyubiquitinated protein levels are low or high and, via UBE3A, adjust
proteasomal activity accordingly. Thus, differences in cell culture conditions (such as differences in serum composition, oxygen tension, and plating density) have the potential to influence polyubiquitinated protein levels and alter proteasome activity via UBE3A.

We found that knockdown of different proteasome subunits or overexpression of UBE3A*T485A or UBE3A*T485E led to activation of Wnt signaling in HEK293T cells, a cell line with low baseline Wnt pathway activation. In contrast, UBE3A*T485E-expressing cells are shown (n = 6). Error bars indicate S.D. *, p < 0.05, two-sample t test (two-tailed). E, the destabilized UbG76V-GFP reporter protein, and mCherry as a transfection and loading control. The proteasome was inhibited with MG-132 (10 μM; 12 h). Mean percent values relative to UBE3A*T458E-expressing cells are shown (n = 5). Error bars indicate S.D. *, p < 0.05, two-sample t test (two-tailed).

Figure 7. UBE3A inhibits the proteasome and increases nuclear β-catenin accumulation. A and B, representative Western blot (A) and quantification (B) of protein lysates from HEK293T cells expressing UBE3A*T485A or UBE3A*T485E, the destabilized UbG76V-GFP reporter protein, and mCherry as a transfection and loading control. The proteasome was inhibited with MG-132 (10 μM; 12 h). Mean percent values relative to UBE3A*T485E-expressing cells are shown (n = 5). Error bars indicate S.D. *, p < 0.05, two-sample t test (two-tailed). C and D, representative Western blot (C) and quantification (D) of nuclear lysates from HEK293T cells expressing UBE3A*T458A or UBE3A*T485T mutants. Cells were grown in L-cell CM or Wnt3a CM and analyzed for β-catenin abundance. Mean percent values relative to UBE3A*T485E-expressing cells are shown (n = 6). Error bars indicate S.D. *, p < 0.05, two-sample t test (two-tailed). E, β-catenin degradation is required for UBE3A-dependent Wnt pathway activation. BAR assays were performed with HEK293T cells expressing UBE3A*T485A or UBE3A*T485E and co-transfected with plasmids encoding GFP, WT β-catenin, or a stabilized β-catenin mutant. Values are shown as box and whisker plots for percent firefly:Renilla ratios relative to cells transfected with GFP alone (n = 9). Whiskers represent the range of maximum and minimum values obtained in our experiments. ***p < 0.0005, two-sample t test (two-tailed).

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verified, we speculate that such an arrangement could spatially restrict docking or shuttling of Wnt pathway components to the proteasome. The destruction complex scaffolding protein AXIN1 can directly bind to β-catenin and PSMD1, whereas β-catenin stabilization is dependent on its sequestration away from the proteasome (64, 65). Post-translational modifications of proteasome subunits, such as ubiquitination, could block the association of β-catenin with the proteasome, which would stabilize β-catenin and promote Wnt signaling. Thus, cells might physically segregate Wnt signaling from other ubiquitin-dependent signaling pathways to permit independent control of cellular processes that depend on proteasomal proteolysis.

Numerous pathways require protein degradation, but it has long remained unknown how the proteasome simultaneously processes signaling components to maintain homeostasis. Our study raises the possibility that distinct pathways are compartmentalized at the proteasome, and this provides the necessary molecular platforms to execute multiple parallel functions.

Lastly, our study has broader implications and suggests that UBE3A could affect additional proteins, signaling pathways, and cellular functions that are particularly sensitive to ubiquitination and proteasome degradation. Our findings hint that the apparent promiscuity of UBE3A may relate to its inhibitory effect on the proteasome as proteasome inhibition would elevate the levels of many ubiquitinated proteins, including those that are not direct UBE3A substrates.

**Experimental procedures**

**Molecular biology**

All UBE3A constructs used in this study were derived from human UBE3A isoform II (NCBI accession number NP_000453.2). Myc epitope tags were placed on the N terminus of UBE3A by polymerase chain reaction and cloned into pcIG2 using SacI and Xmal or into pEGFP-N1 using BamHI and NotI. All constructs were verified by sequencing. HA-tagged ubiquitin, pGL3-BAR, TK-**Renilla**, pHAGE β-catenin WT, and pHAGE β-catenin mutant constructs were described previously (39, 66, 67). Ub^{G76V,G79F} was obtained from Addgene (23969), pCMV6 M yc-D DK-tagged constructs for PSMB1 (RC201798), PSM C2 (RC200945), PSMC5 (RC201251), PSMC6 (RC202809), PSM D1 (RC210486), PSMD2 (RC203204), PSMD3 (RC202307), PSM D6 (RC202292), PSM D7 (RC203133), PSM D11 (RC201201), and untagged PSMD4 (SC111678) were all purchased from Origene.

**Antibodies and reagents**

Primary antibodies were used anti-Myc (1:1,000; EMD Millipore, 05-724), mouse anti-HA (1:1,000; EMD Millipore, 05-904), rabbit anti-GAPDH (1:1,000; Genetex, GTX100118), mouse anti-FLAG M2 (1:1,000; Sigma, PA5-17294), rabbit anti-PSMA2 (1:1,000; Thermo Fisher, PA5-49648), rabbit anti-PSMB1 (1:1,000; Thermo Fisher, PA5-27663), goat anti-PSMD4 (1:1,000; Boston Biochem, AF5540), rabbit anti-PSMD11 (1:1,000; Thermo Fisher, PA5-27447); mouse anti-GFP (1:5,000; Clontech, 632281), rabbit anti-mCherry (1:2,500; Abcam, ab167453), rabbit anti-β-catenin (1:1,000; Cell Signaling Technology, 9562), and mouse anti-HDAC1 (1:1,000; Cell Signaling Technology, 5356). LiCl, XAV939, iCRT14, and valproic acid were purchased from Sigma; CT99021 was purchased from Axon Med Chem; and C59 was purchased from Cellagen Technology. MG-132 was purchased from Calbiochem. Wnt3a medium was made according to the ATCC protocol (68).

**Tissue culture**

HEK293T cells (ATCC) were maintained in a 5% CO₂ humidified incubator in DMEM (Thermo Fisher) containing 4.5 g/liter glucose, 584 mg/liter glutamine, and 110 mg/liter sodium pyruvate supplemented with 10% (v/v) fetal bovine serum (Hyclone) and 1× antibiotic-antimycotic (penicillin, streptomycin, amphotericin B; Life Technologies). For biochemical analyses, transfections were performed in 6-well dishes using Lipofectamine 2000 (Life Technologies) according to the manufacturer’s instructions. HEK293T cells were lysed in NuPAGE lithium dodecyl sulfate sample buffer (Life Technologies) supplemented with 1% β-mercaptoethanol and 1× Complete Mini protease inhibitor mixture (Roche Applied Science). Lysates were boiled, and proteins were resolved by 4–20% SDS-PAGE and transferred to nitrocellulose membranes (Bio-Rad). Membranes were blocked in Odyssey blocking buffer (LI-COR Biosciences) and probed with the appropriate primary antibodies overnight. Protein bands were visualized using the Odyssey CLx infrared imaging system (LI-COR Biosciences) with the following secondary antibodies: donkey anti-rabbit 800CW (926-32213), donkey anti-rabbit 680RD (925-68073), and donkey anti-mouse 680RD (926-68072; all from LI-COR Biosciences and used at 1:10,000). Nuclear fractionation was performed using the NE-PER kit (Thermo Fisher) according to the manufacturer’s instructions.

Immortalized human lymphocytes (Simons Simplex Collection) were maintained in a 5% CO₂ humidified incubator in RPMI 1640 medium supplemented with 15% (v/v) fetal bovine serum (HyClone), 1× GlutaMAX (Gibco), and 1× antibiotic-antimycotic. The following cell lines were used for our study: 13873.Fa, 13873.Mo, and 13873.P1. For protein analysis, 10 million lymphocytes were collected by centrifugation and lysed for 20 min on ice with periodic rocking in a lysis buffer consisting of 20 mM HEPES, pH 7.4, 1% Nonidet P-40, and 50 mM KCl supplemented with protease and phosphatase inhibitors. Samples were cleared at 1,000 × g for 5 min, and the supernatant was collected and subjected to a protein assay (Bio-Rad) to determine concentration. Samples were prepared by diluting in sample buffer supplemented with 1% β-mercaptoethanol, boiled, and resolved by SDS-PAGE.

**Luciferase assays**

BAR assays were performed in 96-well plates for all experiments except those shown in supplemental Fig. S1, B–F (see below). HEK293T cells were plated at a density of 10,000/well. Cells were transiently transfected with 10 ng of pRLL-TK-**Renilla**, 30 ng of BAR-pGL3, and 60 or 80 ng of the indicated constructs using the TransIT®-2020 transfection reagent (Mirus) or TransIT-L1 transfection reagent (Mirus). After 2 h, medium containing transfection reagents was replaced with fresh medium. The following day, cells were serum-deprived
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(0.2% FBS) for 6 h and stimulated with Wnt3a-conditioned medium or control L-cell medium for 12–16 h. Reporter gene expression was assessed using the Dual-Luciferase reporter assay system (Promega) and measured on an Enspire plate reader from PerkinElmer Life Sciences. For supplemental Fig. S1, B–F, HEK293T cells were plated at a density of 5,000/well in 384-well plates, and drugs were added 4 h postplating. 24 h after plating, cells were transfected with 5 ng of pRL-TK-\textit{Renilla}, 15 ng of BAR-pGL3, and 30 ng of the indicated UBE3A expression constructs using FuGENE HD (Promega). Reporter gene expression was assessed 24 h post-transfection using the Dual-Glo luciferase assay system (Promega). For all experiments, the luciferase activity was normalized against \textit{Renilla} activity.

Ubiquitin remnant immunoaffinity profiling

HEK293T cells were grown in high-glucose DMEM (Caisson Laboratories) supplemented with 10% dialyzed FBS (Gibco), 3.7 g/liter sodium bicarbonate, 1× GlutaMAX, and 1× antibiotic-antimycotic. For heavy medium, 50 μg/ml l-lysine (Lys8) and 40 μg/ml l-arginine (Arg10) were added (Cambridge Isotope Laboratories). 50 μg/ml standard l-lysine and 40 μg/ml standard l-arginine (Thermo Fisher) were added for light medium. For complete labeling, cells were passaged for at least seven generations in the above medium. Cells were lyzed in lysis buffer (8 M urea in 50 mM Tris-HCl, pH 7.5, 150 mM NaCl, 1 mM EDTA, 2 μg/ml aprotinin, 10 μg/ml leupeptin, 1 mM PMSF, 50 μM PR-619, and 1 mM iodoacetamide) and sonicated to clarify the lysate. The extract was centrifuged at 20,000 rpm at 4 °C for 15 min, and protein was estimated using the method of Bradford (69). Both heavy and light protein samples were mixed in a 1:1 ratio (mg:mg) and dialyzed in 50 mM Tris-HCl, pH 7.5, to reduce the urea concentration to 2 M. The total protein was digested overnight with sequencing grade trypsin (Promega). The trypsin:protein ratio was maintained at 1:100. Total peptides were purified on a Sep-Pak column (Waters) and eluted with a 40-ml step gradient of 10–50% acetonitrile and 0.1% trifluoroacetic acid gradient (10, 15, 20, 25, 30, 35, 40 and 50%; 5 ml each). The eluent fractions were lyophilized overnight until dry powder flakes were obtained.

A diglycine antibody kit was purchased from Cell Signaling Technology (PTMScan® Ubiquitin Remnant Motif (K-e-GG) kit 5562). Dried total cell peptide samples were solubilized in 1× immunoaffinity purification buffer, and peptide immuno-precipitations were performed at 4 °C for 4–6 h. The bead slurry was washed three times with PBS and two times with water. K-e-GG peptides were eluted with two washes of 50 μl of 0.15% trifluoroacetic acid (TFA). Eluted peptides were further purified on Pierce C_{18} spin columns (Cat 89870) using the manufacturer’s protocol. Peptides were eluted using 70% acetonitrile and 0.1% TFA solution in 50-μl volumes twice and dried in a SpeedVac at room temperature.

Mass spectrometry

Mass spectrometry analysis was performed at the Michael Hooker Proteomics Center at the University of North Carolina. Dried peptide samples were reconstituted in 1% acetonitrile and 0.1% formic acid. Each sample was analyzed by LC/MS/MS using an Easy nLC 1000 coupled to a QExactive HF mass spectrometer (Thermo Scientific). Samples were injected onto an Easy Spray PepMap C_{18} column (75-μm inner diameter × 25 cm; 2-μm particle size) (Thermo Scientific) and separated over a 2-h method. The gradient for separation consisted of 5–35% mobile phase B at a 250 nl/min flow rate where mobile phase A was 0.1% formic acid in water and mobile phase B consisted of 0.1% formic acid in acetonitrile. The QExactive HF was operated in data-dependent mode where the 15 most intense precursors were selected for subsequent fragmentation. Resolution for the precursor scan (m/z 400–1600) was set to 120,000 with a target value of 3 × 10^{6} ions. MS/MS scan resolution was set to 30,000 with a target value of 1 × 10^{5} ions. The normalized collision energy was set to 27% for higher-energy collisional dissociation, and the isolation window was set to 1.6 m/z. Peptide match was set to preferred, and precursors with unknown charge or charge states of 1 and ≤7 were excluded.

Raw data files were processed using Proteome Discoverer version 2.1 (Thermo Scientific). Peak lists were searched against a reviewed human UniProt database (containing 20,203 entries), appended with a contaminant database, using Sequest. The following parameters were used to identify tryptic peptides for protein identification: 10-ppm precursor ion mass tolerance; 0.02-Da product ion mass tolerance; up to two missed trypsin cleavage sites; carbamidomethylation of Cys was set as a fixed modification; and oxidation of Met, Gly-Gly of Lys, phosphorylation of Ser/Thr/Tyr, $^{13}$C\textsubscript{6}\textsuperscript{Arg}, $^{13}$C\textsubscript{6}\textsuperscript{Lys}, and acetylation of the N terminus were set as variable modifications. Percolator was used to estimate the number of false-positive identifications, and a q-value (false discovery rate) was assigned, a q-value of <0.05 (5% false discovery rate), which was used to filter all results. The ptmRS node was used to assign the post-translational modification site localization probability. Peptide quantitation was enabled to determine SILAC ratios for ubiquitin-modified peptides.

Ubiquitination assays

In vitro ubiquitination assays were performed using immunopurified PSMD2 in combination with a commercially available recombinant UBE3A ubiquitin ligase kit (Boston Biochem, K-230) according to the manufacturer’s instructions. In brief, HEK293T cells grown in 25-cm dishes were transfected with a plasmid encoding Myc-DDK-PSMD2 and allowed to grow for 48 h. Cells were lysed on ice for 20 min with gentle agitation using an immunoprecipitation buffer containing 20 mM HEPES, pH 7.4, 1% Nonidet P-40, 50 mM KCl and 100 mM NaCl. Lysates were clear by centrifugation, and PSMD2 was immunoprecipitated using an anti-Myc affinity gel (Sigma, A7470) at 4 °C for 2 h. The resulting beads were washed two times each (eight washes total) in immunoprecipitation buffer containing 100, 250, and 500 mM NaCl as well as two final washes in PBS. PSMD2 was eluted from the beads in PBS containing 10 μg/ml Myc peptide (Sigma).

For cell-based assays, HEK293T cells were transfected with constructs encoding UBE3A, Myc-DDK-PSMD2, and HA-ubiquitin. Cells were allowed to grow for 48 h and treated with 30 μM MG-132 for 4 h. The cells were then lysed in radioimmune precipitation assay buffer containing 1% SDS and 30 μM MG-132. Cell lysates were boiled for 20 min and clarified by
centrifugation at 15,000 × g for 10 min. The resulting supernatant was diluted 1:10 (v/v) in an immunoprecipitation buffer (20 mM HEPES, pH 7.4, 50 mM KCl, and 1% Triton X-100), and PSMD2 was immunoprecipitated using the EZ View anti-FLAG affinity gel (Sigma, F2426) at 4 °C for 1 h. The final complex was washed three times with immunoprecipitation buffer containing 125 mM NaCl, resuspended in sample buffer, and processed for SDS-PAGE and immunoblot analysis.

Statistical analysis

Statistical analyses were performed using GraphPad Prism software. Statistical treatments for each experiment are indicated in the figure legends. p < 0.05 was considered significant.

Author contributions—J. Y. conceived, designed, and performed experiments and wrote the manuscript. S. R. P. performed luciferase experiments. M. P. W. and J. M. W. performed epistasis experiments. R. C. performed ubiquitin remnant immunoaffinity profiling. G. F. performed quantitative PCR. M. J. E., M. B. M., and M. J. Z. conceived and designed experiments and wrote the manuscript.

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Note added in proof—In the version of the article that was published as a Paper in Press on May 30, 2017, the PSMA2 and PSMB1 immunoblots shown in Fig. 4D were derived from the same physical membrane. As these two proteins are the same size, the two images appeared to be similar. To avoid any confusion, the PSMB1 immunoblot was replaced using an image from a replicate immunoblot.

References


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