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Self-oligomerization regulates stability of survival motor neuron protein isoforms by sequestering an SCF^{S1mb} degron

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ABSTRACT Spinal muscular atrophy (SMA) is caused by homozygous mutations in human *SMN1*. Expression of a duplicate gene (*SMN2*) primarily results in skipping of exon 7 and production of an unstable protein isoform, *SMNΔ7*. Although *SMN2* exon skipping is the principal contributor to SMA severity, mechanisms governing stability of survival motor neuron (SMN) isoforms are poorly understood. We used a *Drosophila* model system and label-free proteomics to identify the SCF^{S1mb} ubiquitin E3 ligase complex as a novel SMN binding partner. SCF^{S1mb} interacts with a phosphor degron embedded within the human and fruitfly SMN YG-box oligomerization domains. Substitution of a conserved serine (S270A) interferes with SCF^{S1mb} binding and stabilizes *SMNΔ7*. SMA-causing missense mutations that block multimerization of full-length SMN are also stabilized in the degron mutant background. Overexpression of *SMNΔ7*^{S270A}, but not wild-type (WT) *SMNΔ7*, provides a protective effect in SMA model mice and human motor neuron cell culture systems. Our findings support a model wherein the degron is exposed when SMN is monomeric and sequestered when SMN forms higher-order multimers.

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INTRODUCTION

Spinal muscular atrophy (SMA) is a common neuromuscular disorder, recognized as the most prevalent genetic cause of early childhood mortality (Pearn, 1980). Patients with the most severe form of the disease, which is also the most common, become symptomatic

in the first 6 mo of life and rarely live past 2 yr (Prior, 2010; Wee et al., 2010). Because the onset of symptoms and their severity can vary, SMA has historically been classified into three subtypes (Ogino and Wilson, 2004). More recently, clinicians have recognized that SMA is better characterized as a continuous spectrum disorder, ranging from acute (prenatal onset) to nearly asymptomatic (Tiziano et al., 2013). Clinically, SMA patients experience degeneration of motor neurons in the anterior horn of the lower spinal cord (Crawford and Pardo, 1996). This leads to progressive atrophy of proximal muscle groups, ultimately resulting in loss of motor function and symmetrical paralysis. The cause of death is often restrictive respiratory failure (Kolb and Kissell, 2015).

SMA typically results from homozygous deletion of the *survival motor neuron 1 (SMN1)* gene (Lefebvre et al., 1995). A small fraction

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Abbreviations used:

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of SMA patients have lost one copy of *SMN1* and the remaining copy contains a point mutation (Burghes and Beattie, 2009). Humans have two *SMN* paralogues, named *SMN1* and *SMN2*, both of which contribute to total cellular levels of survival motor neuron (SMN) protein. *SMN2* exon 7 contains a silent base change that alters splicing to primarily produce a truncated, unstable protein product called SMN Δ 7 (Lorson *et al.*, 1999; Monani *et al.*, 1999; Lorson and Androphy, 2000). The last 16 amino acids of SMN are replaced in SMN Δ 7 by four amino acids, EMLA, encoded by exon 8. Current estimates suggest that *SMN2* produces 10–15% of the level of full-length protein produced by *SMN1* (Lorson *et al.*, 2010). Complete loss of SMN is lethal in all organisms investigated to date (O'Hearn *et al.*, 2016). Although the amount of full-length protein produced by *SMN2* is not enough to compensate for loss of *SMN1*, *SMN2* is sufficient to rescue embryonic lethality (Monani *et al.*, 2000). SMA is therefore a disease that arises due to a hypomorphic reduction in SMN levels (Lefebvre *et al.*, 1995). Furthermore, relative levels of the SMN protein correlate with the phenotypic severity of SMA (Coovert *et al.*, 1997; Lefebvre *et al.*, 1997).

Whereas a causative link between *SMN1* and SMA was established in the early 1990s, the molecular role of SMN in disease etiology remains unclear. SMN is the central component of a multimeric protein assemblage known as the SMN complex (Li *et al.*, 2014; Matera and Wang, 2014). The best-characterized function of this complex, which is found in all tissues of metazoan organisms, is in the cytoplasmic assembly of small nuclear ribonucleoproteins (snRNPs), core components of the spliceosome (Fischer *et al.*, 1997; Meister *et al.*, 2001; Pellizzoni *et al.*, 2002).

Although it is ubiquitously expressed, SMN has also been implicated in a number of tissue-specific processes related to neurons and muscles. These functions include actin dynamics (Oprea *et al.*, 2008; Ackermann *et al.*, 2013), axonal pathfinding (Fan and Simard, 2002; McWhorter *et al.*, 2003; Sharma *et al.*, 2005), axonal transport of β -actin mRNP (Rossoll *et al.*, 2003), phosphatase and tensin homolog-mediated (PTEN-mediated) protein synthesis pathways (Ning *et al.*, 2010), translational regulation (Sanchez *et al.*, 2013), neuromuscular junction formation and function (Chan *et al.*, 2003; Kariya *et al.*, 2008; Kong *et al.*, 2009; Voigt *et al.*, 2010), myoblast fusion (Shafey *et al.*, 2005), and maintenance of muscle architecture (Rajendra *et al.*, 2007; Walker *et al.*, 2008; Bowerman *et al.*, 2009).

Ubiquitylation pathways have been shown to regulate the stability and degradation of SMN (Chang *et al.*, 2004; Burnett *et al.*, 2009; Hsu *et al.*, 2010) as well as axonal and synaptic stability (Korhonen and Lindholm, 2004). In the ubiquitin proteasome system (UPS), proteins destined for degradation are tagged by linkage to ubiquitin through the action of three factors (Petroski, 2008). E1 proteins activate ubiquitin and transfer it to the E2 enzyme. E2 proteins conjugate ubiquitin to their substrates. E3 proteins recognize the substrate and assist in the transfer of ubiquitin from the E2. Because E3 ligases confer substrate specificity, they are typically considered as candidates for targeted inhibition of protein degradation. Ubiquitin homeostasis is thought to be particularly important for neuromuscular pathology in SMA (Groen and Gillingwater, 2015). Indeed, mouse models of SMA display widespread perturbations in UBA1 (ubiquitin-like modifier activating enzyme 1) levels (Wishart *et al.*, 2014). Furthermore, mutations in UBA1 are known to cause X-linked infantile SMA (Ramser *et al.*, 2008; Schmutzler *et al.*, 2008).

Given the importance of these processes to normal development as well as neurodegenerative disease, we set out to identify and characterize novel SMN binding partners. Previously, we developed *Drosophila melanogaster* as a model system wherein the endogenous *Smn* gene is replaced with a *Flag-Smn* transgene (Praveen

et al., 2012). Although it is highly similar to human *SMN1* and *SMN2*, the entire open reading frame of fruitfly *Smn* is contained within a single exon, and so only full-length SMN protein is expressed in *Drosophila* (Rajendra *et al.*, 2007). When modeled in the fly, SMA-causing point mutations recapitulate the full range of phenotypic severity seen in humans (Praveen *et al.*, 2014; Garcia *et al.*, 2016). Using this system, we carried out proteomic profiling of Flag-purified embryonic lysates and identified the SCF^{Smb} E3 ubiquitin ligase complex as a novel SMN interactor. Importantly, this interaction is conserved from flies to humans. We show that SCF^{Smb} binding requires a phosphodegron motif located within the SMN self-oligomerization domain, mutation of which stabilizes SMN Δ 7 and, to a lesser extent, full-length SMN. Additional studies in flies, mice, and human cells elucidate a disease-relevant mechanism whereby SMN protein stability is regulated by self-oligomerization. Other E3 ligases have been reported to target SMN for degradation in cultured human cells (Hsu *et al.*, 2010; Kwon *et al.*, 2013; Han *et al.*, 2016). Given our findings in fruit-fly embryos, SMN is likely targeted by multiple E3 ubiquitin ligases.

RESULTS

Flag-SMN interacts with ubiquitin proteasome system proteins

We previously generated transgenic flies that express Flag-tagged SMN proteins in an otherwise null *Smn* background (Praveen *et al.*, 2012). To preserve endogenous expression patterns, the constructs are driven by the native promoter and flanking sequences. As described under *Materials and Methods*, we intercrossed hemizygous *Flag-Smn*^{WT}, *Smn*^{X7}/*Smn*^D animals to establish a stock wherein all of the SMN protein, including the maternal contribution, is epitope tagged. After breeding them for >100 generations, essentially all of the animals are homozygous for the *Flag-Smn*^{WT} transgene, but second-site recessive mutations are minimized due to the use of two different *Smn* null alleles. Adults from this stock display no apparent defects and have an eclosion frequency (~90%) similar to that of wild-type (Oregon-R) animals.

We collected (0–12 h) embryos from *Flag-Smn*^{WT/WT}, *Smn*^{X7/D} (SMN) and Oregon-R (Ctrl) animals and analyzed Flag-purified lysates by “label-free” mass spectrometry. In addition to Flag-SMN, we identified SMN complex components Gemin2 and Gemin3, along with all seven of the canonical Sm-core snRNP proteins (Figure 1A). We also identified the U7-specific Sm-like heterodimer Lsm10/11 (Pillai *et al.*, 2003) and the Gemin5 orthologue, Rigor mortis (Gates *et al.*, 2004). Previous studies of Schneider2 (S2) cells transfected with epitope-tagged *Smn* had identified most of the proteins listed above as SMN binding partners in *Drosophila* (Kroiss *et al.*, 2008). However, despite bioinformatic and cell biological data indicating that Rigor mortis is part of the fruit-fly SMN complex, this protein failed to copurify with SMN in S2 cells (Kroiss *et al.*, 2008; Cauchi *et al.*, 2010; Guruharsha *et al.*, 2011). On the basis of our purification data, we conclude that the conditions are effective and that Rigor mortis/Gemin5 is an integral member of the SMN complex in flies.

A detailed proteomic analysis of these flies will be presented elsewhere. As shown in Figure 1B, our preliminary analysis identified 396 proteins, 114 of which were detected only in the Flag-SMN sample and not in the control. An additional 279 proteins were detected in both the Flag purification and control samples. In addition to SMN complex members, we copurified numerous factors that are part of the ubiquitin proteasome system (UPS; Figure 1C). Among these UPS proteins, we identified Cullin 1 (Cul1), Skp1-related A (SkpA), and supernumerary limbs (Smb), as being highly enriched

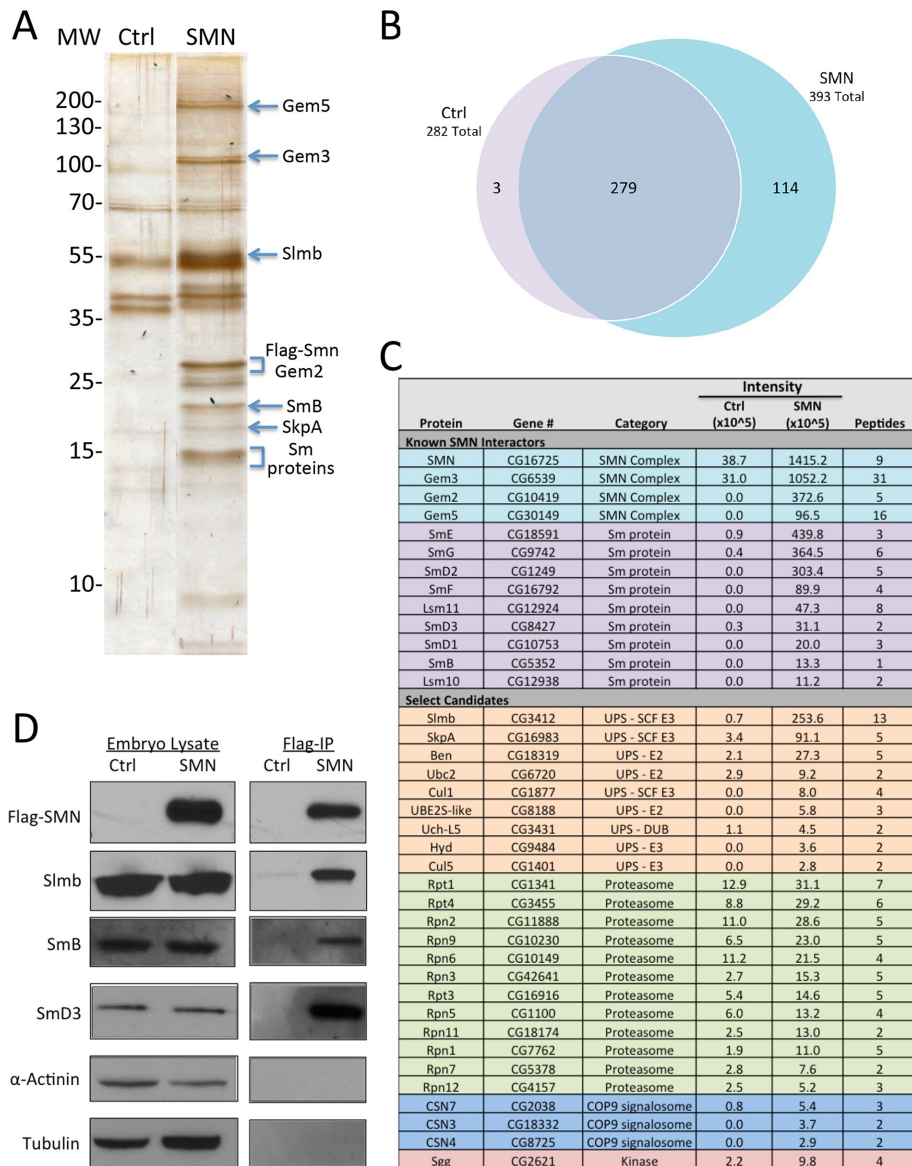


FIGURE 1: Flag-SMN immunopurified lysates contain known protein interaction partners and ubiquitin proteasome system (UPS) proteins. (A) Lysates from Oregon-R control (Ctrl) *Drosophila* embryos and embryos expressing only transgenic Flag-SMN (SMN) were Flag-immunopurified, and protein eluates were separated by gel electrophoresis and silver stained. Band identities predicted by size using information from panels C and D. (B) Direct mass spectrometric analysis of the eluates (which were not gel purified) identified a total of 396 proteins, 114 of which were detected only in SMN sample and 279 of which were detected in both SMN and Ctrl samples. (C) Flag-purified eluates were analyzed by “label-free” mass spectrometry. Numerous proteins that copurify with Flag-SMN are part of the ubiquitin proteasome system (UPS). Of these UPS proteins, Cullin 1 (Cul 1), SkpA, and supernumerary limbs (Slmb) were highly enriched (at least 10-fold) in the SMN sample as compared with Ctrl. (D) Western blot analysis of Flag-purified eluates was used to further validate the presence or absence of SMN interaction partners. Flag-SMN was successfully purified from SMN embryos but was undetectable in the control. As positive controls for known protein interaction partners of SMN, SmB and SmD3 were also easily detectable by Western blotting using anti-Sm antibodies. The presence of Slmb was verified using anti-Slmb. α -Actinin and tubulin were not enriched in our purification and are used as negative controls to demonstrate specificity.

(>10-fold) in Flag-SMN samples as compared with the control. Together, these proteins comprise the SCF^{Slmb} E3 ubiquitin ligase. Cul1 forms the major structural scaffold of this horseshoe-shaped, multisubunit complex (Zheng et al., 2002). Slmb is an F-box protein and is the substrate recognition component (Jiang and Struhl,

1998). SkpA is a bridging protein essential for interaction of Cul1 with the F-box protein (Patton et al., 1998a,b). Because of its role in substrate recognition, Slmb is likely to be the direct interacting partner of SMN within the SCF^{Slmb} complex. For this reason, we focused on Slmb for the initial validation. As shown, Slmb was easily detectable in Flag-purified eluates from embryos expressing Flag-SMN and nearly undetectable in those from control embryos (Figure 1D). SmB and SmD3 were also easily detectable by Western blot in Flag-purified embryonic lysates and were used as positive controls for known protein interaction partners of SMN. Tubulin and α -actinin were not detected as interacting with SMN in our purification and demonstrate the specificity of the detected SMN interactions.

SCF^{Slmb} is a bona fide SMN interaction partner that ubiquitylates SMN

As an E3 ubiquitin ligase, the SCF^{Slmb} complex is a substrate recognition component of the ubiquitin proteasome system. As outlined in Figure 2A, E3 ligases work with E1 and E2 proteins to ubiquitylate their targets. The interaction of SCF^{Slmb} with SMN was verified in a reciprocal coimmunoprecipitation, demonstrating that Flag-tagged SCF components form complexes with endogenous SMN (Figure 2B) in S2 cells.

SCF complexes are highly conserved from flies to humans: SkpA is 77% identical to human Skp1, Cul1 is 63% identical, and Slmb is 80% identical to its human homologues, B-TrCP1 and B-TrCP2. Slmb/B-TrCP is the SCF component that directly contacts substrates of the E3 ligase. We therefore tested the interaction of recombinant human SMN in complex with (SMN•Gem2) Gupta et al., 2015) with GST-tagged B-TrCP1 and -SMN proteins in an in vitro binding assay. As shown in Figure 2C, SMN•Gem2 did not interact with GST alone but was detected at high levels following pull down with either GST-SMN (positive control) or GST-B-TrCP1. We also tested the interaction of Flag-tagged *Drosophila* SCF components with endogenous human SMN in HEK 293T cells (Figure 2D). Accordingly, human SMN was coprecipitated with Flag-Cul1 and Flag-Slmb and at lower levels following Flag-SkpA immunoprecipitation. Flag-B-TrCP1 and Flag-B-TrCP2, the two human homologues of Slmb, also copurified with endogenous human SMN in HEK 293T

cells (Figure 2E). Altogether, these data demonstrate a conserved interaction between SMN and the SCF^{Slmb/B-TrCP} E3 ubiquitin ligase complex.

To test the functional consequences of this conserved interaction between SMN and SCF^{Slmb/B-TrCP}, a cell-based ubiquitylation assay

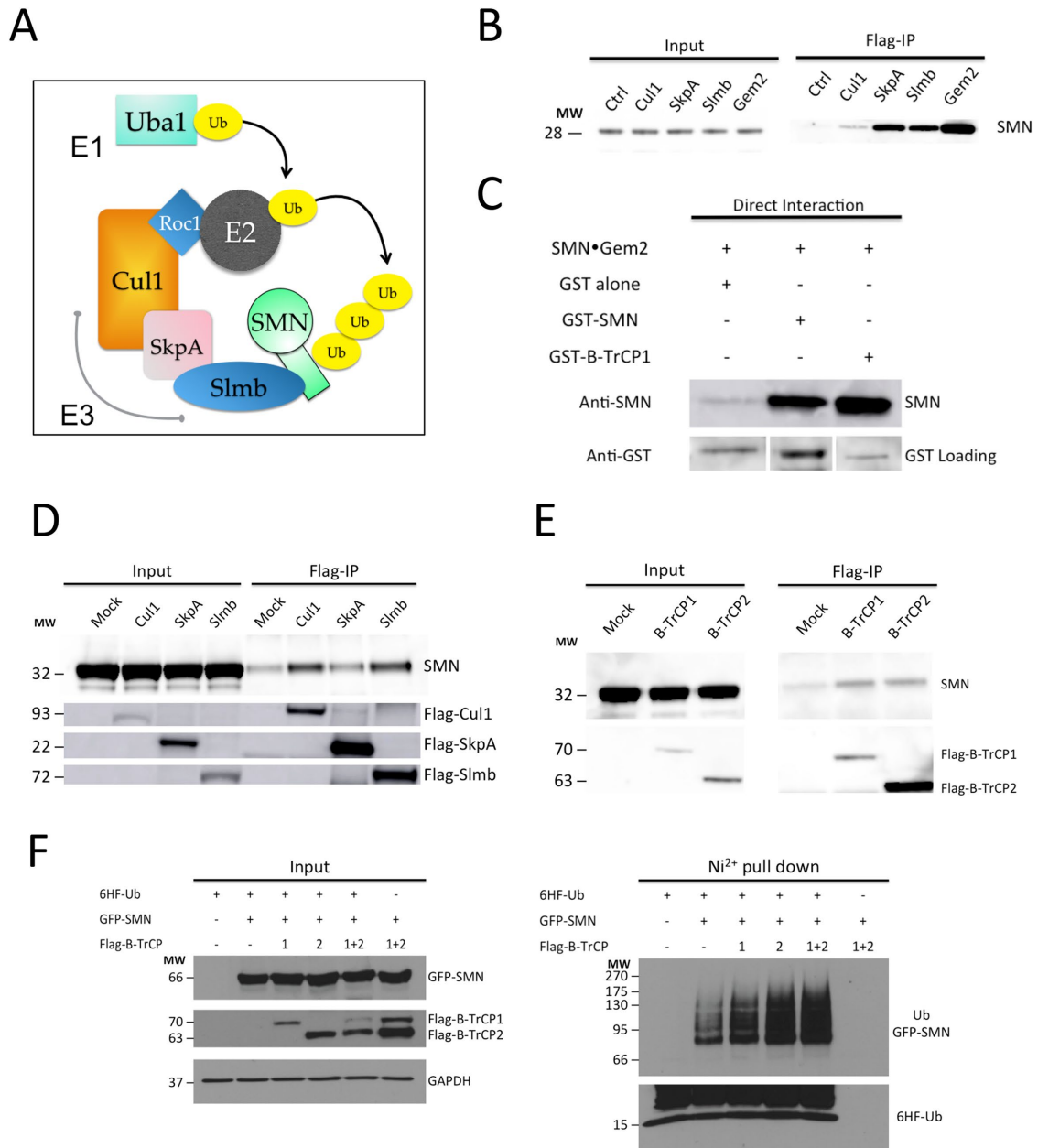


FIGURE 2: Conserved interaction between SMN and the SCF^{Slmb/B-TrCP} E3 ubiquitin ligase results in ubiquitylation of SMN. (A) E3 ligases work with E1 and E2 proteins to ubiquitylate their targets. The SCF^{Slmb/B-TrCP} E3 ubiquitin ligase is made up of three proteins: Cul1, SkpA, and Slmb. The E3 ubiquitin ligase is the substrate recognition component of the ubiquitin proteasome system. (B) Following Cul1-Flag, SkpA-Flag, Flag-Slmb, and Flag-Gem2 immunoprecipitation from *Drosophila* S2 cell lysates, Western analysis using anti-SMN antibody for endogenous SMN was carried out. Coprecipitation of each of the SCF components with endogenous SMN was detected. (C) An in vitro binding assay tested direct interaction between human SMN Δ 5-Gemin2 (SMN•Gem2) (Martin *et al.*, 2012; Gupta *et al.*, 2015) and purified GST-tagged proteins. SMN•Gem2 did not interact with GST protein alone but bound to GST tagged *Drosophila* SMN (GST-SMN) and GST tagged human B-TrCP1 (GST-B-TrCP1). Levels of GST alone, GST-SMN, and GST-B-TrCP1 were detected using anti-GST antibody. (D) The interaction of Flag-tagged *Drosophila* SCF components with endogenous human SMN was tested in HEK 293T cells. Human SMN was detected at high levels following immunoprecipitation of *Drosophila* Flag-Cul1 and Flag-Slmb and detected at a lower level following *Drosophila* Flag-SkpA immunoprecipitation. (E) Flag-tagged versions of the human homologues of Slmb, Flag-B-TrCP1, and Flag-B-TrCP2, interact with endogenous human SMN in HEK 293T cells demonstrated by Flag-immunoprecipitation followed by immunodetection of SMN. (F) Protein lysate from HEK 293T cells transfected with 6xHis-Flag-ubiquitin (6HF-Ub) and GFP-SMN was purified using a Ni²⁺ pull down for the tagged ubiquitin. Baseline levels of ubiquitylated GFP-SMN were detected using anti-GFP antibody. Following transfection of Flag-B-TrCP1 or Flag-B-TrCP2, the levels of ubiquitylated SMN markedly increased. Ubiquitylation levels were further increased following addition of both proteins together. In the input, GFP-SMN was detected using anti-GFP antibody, Flag-B-TrCP1 and Flag-B-TrCP2 were detected using anti-Flag antibody, and GAPDH was detected by anti-GAPDH antibody. In the Ni²⁺ pull down, ubiquitylated GFP-SMN was detected using anti-GFP antibody and 6HF-Ub was detected using anti-Flag antibody to verify successful pull down of tagged ubiquitin.

was performed (Figure 2F). Protein lysate from HEK 293T cells transfected with 6xHis-Flag-ubiquitin and GFP-SMN was purified using a Ni²⁺ pull down for the tagged ubiquitin. Baseline levels of ubiquitylated GFP-SMN were detected using anti-GFP antibody. Following transfection of Flag-B-TrCP1 or Flag-B-TrCP2, the levels of ubiquitylated SMN markedly increased (Figure 2F). Ubiquitylation levels were further increased following addition of both proteins together.

These experiments demonstrate that SCF^{Slmb/B-TrCP} can ubiquitylate SMN *in vivo*.

Depletion of Slmb/B-TrCP results in a modest increase in SMN levels

Given that one of the primary functions of protein ubiquitylation is to target proteins to the proteasome, we examined whether depletion

of Slmb by RNA interference (RNAi) using dsRNA in S2 cells would increase SMN levels (Figure 3A). Following Slmb RNAi, endogenous SMN levels were modestly increased as compared with cells treated with control dsRNA. We obtained similar results using an siRNA that targets both B-TrCP1 and B-TrCP2 in HeLa cells. As shown in Figure 3B, we detected a modest increase in levels of full-length SMN following B-TrCP RNAi but not control RNAi. Next, we treated S2 cells with cycloheximide (CHX), in the presence or absence of dsRNA targeting Slmb, to determine whether differences in SMN levels would be exacerbated when production of new proteins was prevented (Figure 3C). SMN protein levels were also specifically targeted using dsRNA against *Smn* as a positive control for the RNAi treatment. At 6 h post-CHX treatment, there was a modest increase in full-length SMN levels following Slmb RNAi as compared with the initial timepoint (0 h) or the negative control (Ctrl) RNAi (Figure 3C). Together, these data indicate that Slmb/B-TrCP participates in the regulation of SMN protein levels.

Identification and characterization of a Slmb/B-TrCP degradation signal in SMN

Studies of numerous UPS substrates in a variety of species have revealed the presence of degradation signals (called degrons) that are required for proper E3 target recognition and binding. Slmb/B-TrCP canonically recognizes a consensus DpSGXXpS/T degron, where p indicates a phosphoryl group (Fuchs *et al.*, 2004; Jin *et al.*, 2005; Frescas and Pagano, 2008). There are also several known variants of this motif, for example, DDGFVD, SSGYFS, and TSGCSS (Kim *et al.*, 2015). As shown in Figure 4A, we identified a putative Slmb/B-TrCP degron (²⁶⁹MSGYHT²⁷⁴) in the highly conserved self-oligomerization domain (YG Box) of human SMN. Interestingly, this sequence was previously identified as part of a larger degron motif (²⁶⁸YMSGYHT-GYYMEMLA²⁸²) that was thought to be created in SMNΔ7 by SMN2 alternative splicing (Cho and Dreyfuss, 2010). In particular, mutation of S270 (S201 in flies) to alanine was shown to dramatically stabilize SMNΔ7 constructs in human cells, and overexpression of SMNΔ7^{S270A} in SMN-deficient chicken DT40 cells rescued their viability

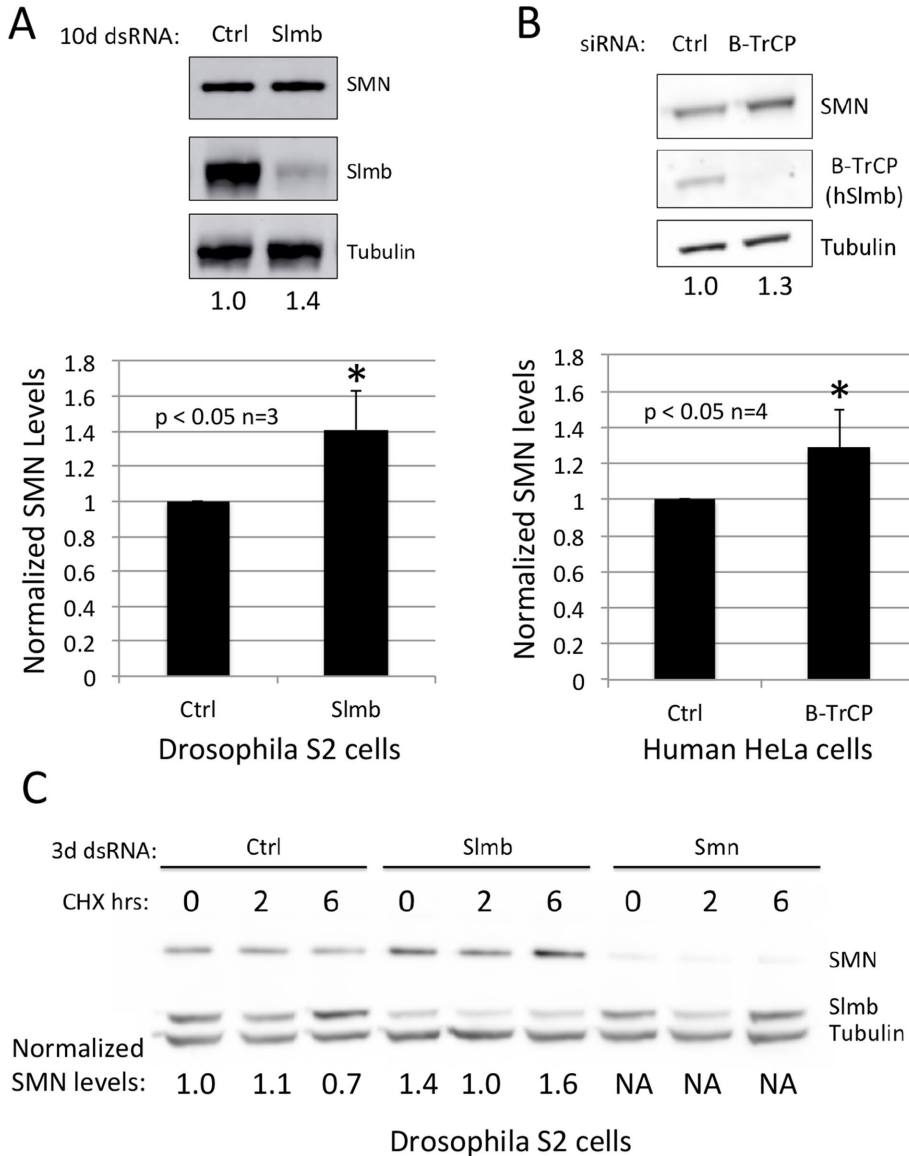


FIGURE 3: Depletion of Slmb/B-TrCP results in an increase of SMN levels. (A) Depletion of Slmb using 10-d (10d) treatment with dsRNA in *Drosophila* S2 cells resulted in modestly increased SMN levels. Following Slmb RNAi, full-length SMN levels were increased as compared with cells treated with control dsRNA against *Gaussia* Luciferase, which is not expressed in S2 cells. (B) The effect of B-TrCP depletion on SMN levels in human cells was tested using siRNA that targets both B-TrCP1 and B-TrCP2 in HeLa cells. We detected a modest increase in levels of full-length endogenous SMN after B-TrCP RNAi but not control (scramble) RNAi. (C) *Drosophila* S2 cells were treated with cycloheximide (CHX), an inhibitor of protein synthesis, following Slmb depletion following a 3-d dsRNA treatment to test whether differences in protein levels would be exacerbated when the production of new protein was prevented. SMN protein levels were also directly targeted using dsRNA against *Smn* as a positive control for the RNAi treatment. As a negative control (Ctrl), dsRNA against *oskar*, which is not expressed in S2 cells, was used. Protein was collected at 0, 2, and 6 h post-CHX treatment. At 6 h post-CHX treatment, there is a modest increase in full-length SMN levels following *Slmb* RNAi as compared with the initial time point (0 h) and as compared with control RNAi treatment.

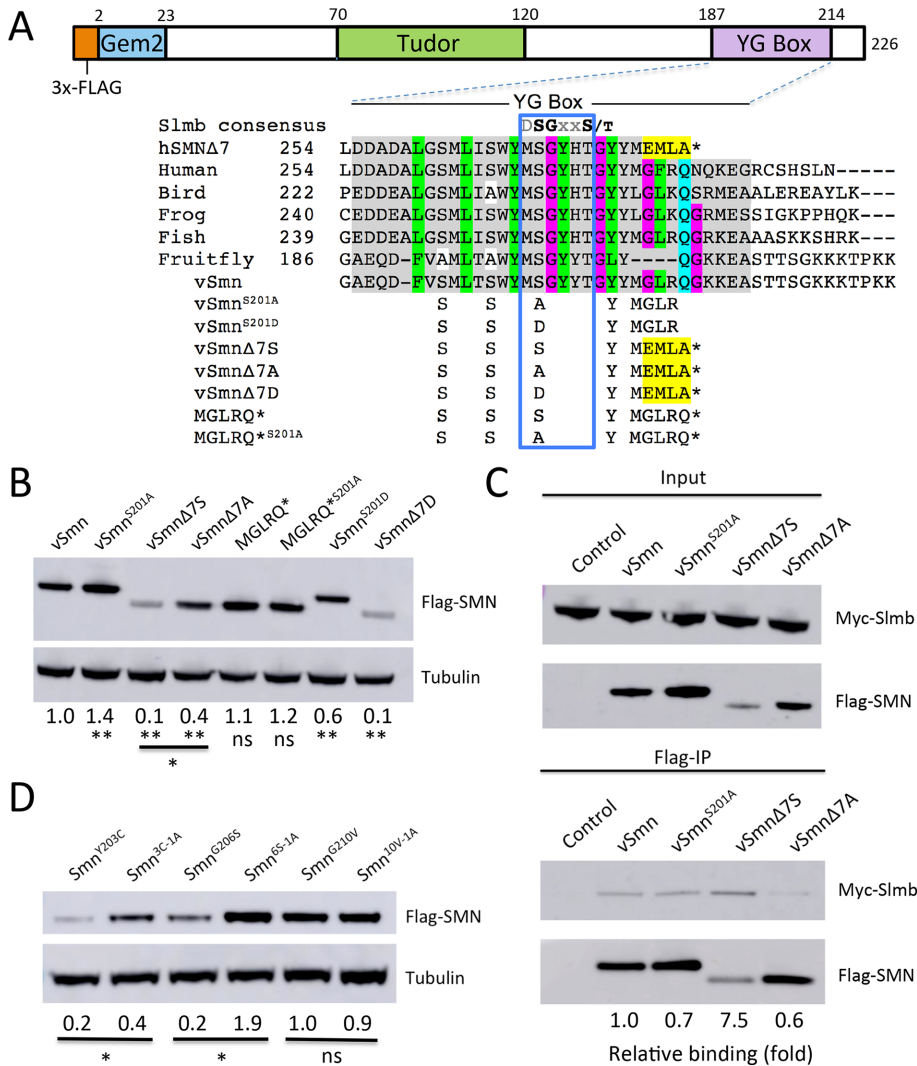


FIGURE 4: Identification and mutation of a putative Slmb/B-TrCP phosphodegron (A) Identification of a conserved putative Slmb phosphodegron (DpSGXXS/T motif variant) in the C-terminal self-oligomerization domain (YG Box) of SMN. The amino acid sequence of SMN from a variety of vertebrates is shown to illustrate conservation of this motif and rationale for the amino acid changes. Full-length human SMN is labeled as "Human," and the truncated isoform is labeled "hSMNΔ7." Endogenous *D. melanogaster* SMN is labeled "Fruitfly." To generate a more vertebrate-like SMN, key amino acids in *Drosophila* SMN were changed to amino acids conserved in vertebrates. Using this SMN backbone, a serine-to-alanine mutation was made in the putative degron in both full-length (vSMN^{S201A}) and truncated SMNΔ7 (vSMNΔ7A). An additional SMN construct that is the same length as SMNΔ7, but has the amino acid sequence GLRQ (the next amino acids in the sequence) rather than EMLA (the amino acids introduced by mis-splicing of SMN2), was generated. The same serine to alanine mutation was made in this construct as well (MGLRQ* and MGLRQ*^{S201A}). Finally, to mimic a phosphorylated serine, a full-length vSmn^{S201D} and truncated vSmnΔ7D were also employed. (B) Western blotting was used to determine protein levels of each of these SMN constructs, with expression driven by the endogenous promoter, in *Drosophila* S2 cells. Both the vSMN and vSMNΔ7S proteins show increased levels when the serine is mutated to an alanine, indicating disruption of the normal degradation of SMN. Additionally, MGLRQ* protein is present at higher levels than is vSMNΔ7S and protein levels do not change when the serine is mutated to an alanine. Normalized fold change as compared with vSmn levels is indicated at the bottom. **p* < 0.05, ***p* < 0.01, *n* = 3. (C) Flag-tagged SMN constructs were cotransfected with Myc-Slmb in *Drosophila* S2 cells. Protein lysates were Flag-immunoprecipitated and probed with anti-Myc antibody to detect SMN-Slmb interaction. In both full-length SMN (vSMN) and truncated SMN (vSMNΔ7), serine-to-alanine mutation decreased interaction of Slmb with SMN. Truncated SMN (vSMNΔ7) showed a dramatically increased interaction with Slmb as compared with full-length SMN (vSMN), despite the fact it is present at lower levels. (D) Full-length SMN constructs containing point mutations known to decrease self-oligomerization (Smn^{Y203C} and Smn^{G206S}) and a mutation that does not disrupt self-oligomerization in the fly (Smn^{G210V}) with and without the

(Cho and Dreyfuss, 2010). However, factors responsible for specifically mediating SMNΔ7 degradation have not been identified.

To develop a more disease-relevant *Drosophila* system to investigate SMN YG box function, we generated a "vertebrate-like" SMN construct, called vSmn (Figure 4A). Transgenic flies expressing Flag-vSmn and Flag-vSmn^{S201A} in the background of an *Smn*^{X7} null mutation are fully viable (Supplemental Figure S1). In fact, the eclosion frequencies of these animals are consistently higher than those that express Flag-Smn^{WT} (Supplemental Figure S1). Additional *Smn* mutant constructs were generated using the vSmn backbone, including both the full-length (e.g., vSmn^{S201A}) and truncated (e.g., vSmnΔ7A) versions of the protein (Figure 4A). To test the effects of overall protein length and distance of the putative degron from the C-terminus, we also generated vSmn constructs that are the same length as SMNΔ7, replacing the MEMLA* motif (the amino acids introduced by human SMN2 splicing) with MGLRQ*; see Figure 4A. The S201A mutation was created in this construct as well (MGLRQ*^{S201A}). To mimic a constitutively phosphorylated state, we also introduced serine to aspartate mutations, vSmn^{S201D} and vSmnΔ7D. We transfected each of these constructs, Flag-tagged and driven by the native *Smn* promoter, into S2 cells and measured protein levels by Western blotting (Figure 4B). We note that these constructs are expressed at levels far below those of endogenous SMN protein in S2 cells; moreover, they do not affect levels of endogenous SMN (Supplemental Figure S2). As shown, the vSmn^{S201A} and vSMNΔ7A constructs exhibited increased protein levels compared with their serine containing counterparts, whereas levels of the S201D mutants were reduced, suggesting that the phosphodegron motif identified in human SMNΔ7 (Cho and Dreyfuss, 2010) is also conserved in the fly protein. In addition to examining protein levels of each of these constructs in cell culture, transgenic flies expressing vSmn, vSmn^{S201A}, vSmnΔ7S, and vSmnΔ7A were created. Here again, we observed that the S201A mutation increased protein levels of both full-length SMN and SMNΔ7 (Supplemental Figure S3).

serine-to-alanine mutation were transfected in *Drosophila* S2 cells. The constructs containing the serine to alanine mutation are as follows: Smn^{Y203C}→Smn^{3C-1A}, Smn^{G206S}→Smn^{65-1A}, Smn^{G210V}→Smn^{10V-1A}. The serine to alanine mutation has a stabilizing effect on SMN mutants with poor self-oligomerization capability. **p* < 0.05, *n* = 3.

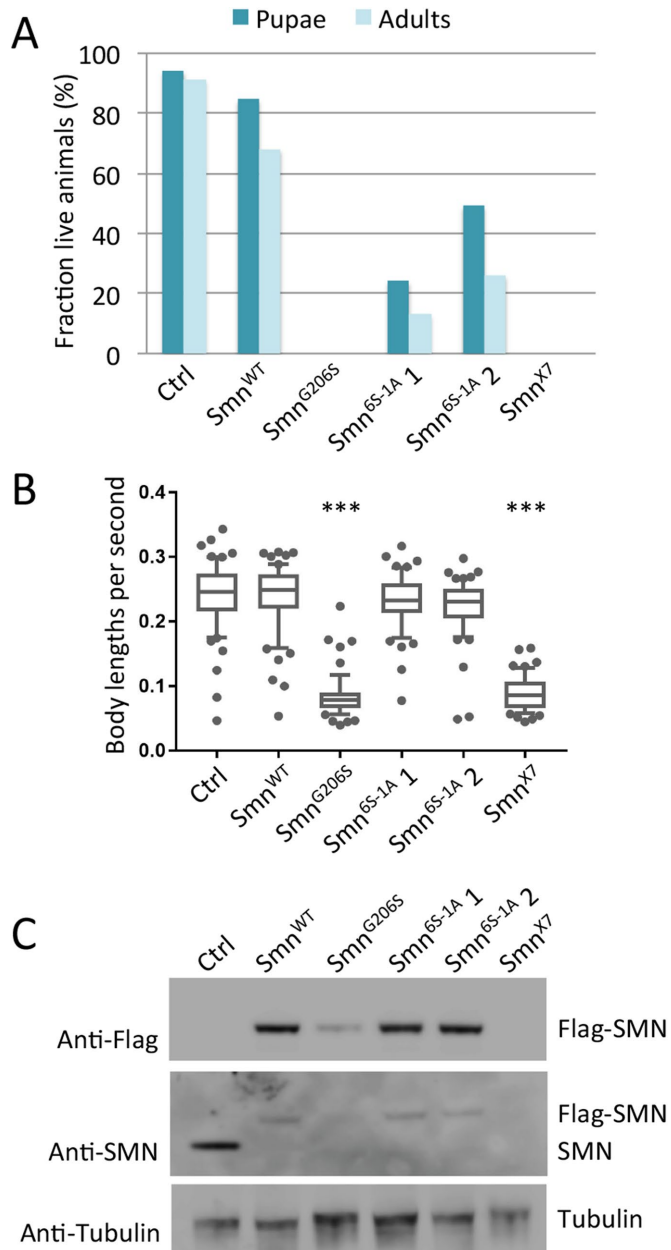


FIGURE 5: Mutation of the Slmb degon rescues defects in SMA model flies. (A) Viability analysis of an SMA point mutation (G206S) in the presence and absence of the degon mutation, S201A. Flies with the following genotypes were analyzed in this experiment: Oregon-R (Ctrl), *Flag-Smn^{WT}, Smn^{X7}/Smn^{X7}* (*Smn^{WT}*), *Flag-Smn^{G206S}, Smn^{X7}/Smn^{X7}* (*Smn^{G206S}*), *Flag-Smn^{G206S, S201A}, Smn^{X7}/Smn^{X7}* (*Smn^{65-1A}*), or *Smn^{X7}/Smn^{X7}* (*Smn^{X7}*). The data for each genotype are expressed as a fraction of pupae or adults over the total number of starting larvae, $n = 200$. Expression of the WT transgene (*Smn^{WT}*) shows robust rescue of the null (*Smn^{X7}*) phenotype (~68% adults). *Smn^{G206S}* is a larval lethal mutation. In two independent recombinant lines of *Smn^{65-1A}* (*Smn^{65-1A 1}* and *Smn^{65-1A 2}*) a fraction of the larvae complete development to become adults. (B) Locomotor ability of early third-instar larvae was determined by tracking their movement for 1 min and then calculating the velocity. To account for potential differences in larval size, speed is expressed as average body lengths per second moved. Genotypes are as in panel A. *Smn^{G206S}* larvae move similarly to null animals. The motility of *Smn^{65-1A 1}* and *Smn^{65-1A 2}* larvae is not different from Ctrl or *Smn^{WT}* larvae. $***p < 0.001$, $n = 50-60$ larvae. (C) Larval protein levels were examined by Western blotting; genotypes as in panel A. Lysates from hemizygous mutant lines were probed with anti-Flag or

The MGLRQ* construct is present at levels that are similar to wild type (vSmn) and much higher than vSmn Δ 7S. Based on the crystal structures of the SMN YG box (Martin *et al.*, 2012; Gupta *et al.*, 2015), the presence of the MGLR insertion in *Drosophila* SMN is predicted to promote self-oligomerization (unpublished data), thus stabilizing the protein within the SMN complex (Burnett *et al.*, 2009). By the same logic, the relative inability of vSmn Δ 7S to self-interact would be predicted to lead to its destruction. To determine whether the observed increase in SMN protein levels correlated with its ability to interact with Slmb, we cotransfected the appropriate Flag-Smn constructs with Myc-Slmb in S2 cells. Protein lysates were then Flag-immunoprecipitated and probed with anti-Myc antibody (Figure 4C). The S201A mutation decreased binding of Slmb to both the full-length and the truncated SMN isoforms (Figure 4C). However, the vSmn Δ 7S construct coprecipitated the greatest amount of Slmb protein, despite the fact that it is present at much lower levels in the input lysate (Figure 4C). Because SMN Δ 7 is defective in self-interaction, this result suggests that the degon is more accessible to Slmb when SMN is monomeric and cannot efficiently oligomerize.

SMN self-oligomerization regulates access to the Slmb degon

To examine the connection between SMN self-oligomerization and degon accessibility more closely, we took advantage of two SMA patient-derived point mutations (Y203C and G206S) that are known to destabilize the full-length protein and to decrease its self-oligomerization capacity (Praveen *et al.*, 2014). As a control, we also employed an SMA-causing mutation (G210V) that does not disrupt SMN self-oligomerization (Praveen *et al.*, 2014; Gupta *et al.*, 2015). Next, we introduced the S201A degon mutation into all three of these full-length SMN constructs, transfected them into S2 cells and carried out Western blotting (Figure 4D and Supplemental Figure S2). The S201A degon mutation has a clear stabilizing effect on the G206S and Y203C constructs, as compared with the effect of S201A paired with G210V. Hence, we conclude that the Slmb degon is exposed when SMN is present predominantly as a monomer, whereas it is less accessible when the protein is able to form higher-order multimers.

Mutation of the Slmb degon rescues viability and locomotion defects in SMA model flies

Next, we examined the effect of mutating the Slmb degon in the context of the full-length protein in vivo. We characterized adult viability, larval locomotion, and SMN protein expression phenotypes of the G206S mutants in the presence or absence of the degon mutation, S201A (Figure 5, A–C). As described previously (Praveen *et al.*, 2014), *Smn^{G206S}* animals express very low levels of SMN and fail to develop beyond larval stages. In contrast, flies bearing the S201A degon mutation in addition to G206S (*Smn^{65-1A}*) express markedly increased levels of SMN protein (Figure 5C), and a good fraction of these animals complete development (Figure 5A). Moreover, *Smn^{65-1A}* larvae display significantly improved locomotor activity as compared with *Smn^{G206S}* or *Smn^{X7}* null mutants (Figure 5B).

anti-SMN antibodies as indicated. The slower-migrating bands represent the Flag-tagged transgenic proteins and the faster migrating band corresponds to endogenous SMN, which is present only in the Ctrl (note Oregon-R has two copies *Smn*, whereas the transgenics have only one). *Smn^{G206S}* has very low levels of SMN protein. Flies bearing the S201A degon mutation in addition to G206S (*Smn^{65-1A}*) express markedly increased levels of SMN protein.

These results strongly suggest that both the structure of the G206S mutant protein as well as its instability contribute to the organismal phenotype.

GFP-SMN Δ 7 overexpression stabilizes endogenous SMN and SMN Δ 7 in cultured human cells

Increased SMN2 copy number correlates with a milder clinical phenotype in SMA patients (Oskoui *et al.*, 2016). This phenomenon was successfully modeled in mice in the early 2000s (Hsieh-Li *et al.*, 2000; Monani *et al.*, 2000), showing that high-copy-number SMN2 transgenes fully rescue the null phenotype, whereas low-copy transgenes do not. Moreover, transgenic expression of a human SMN Δ 7 cDNA construct in a low-copy SMN2 background improves survival of this severe SMA mouse model from P5 (postnatal day 5) to P13 (Le *et al.*, 2005). Although the truncated SMN likely retains partial functionality, the protective effect of SMN Δ 7 overexpression may not entirely be intrinsic to the protein. That is, SMN Δ 7 could also act as a “soak-off” factor, titrating the ubiquitylation machinery and stabilizing endogenous SMN. In such a scenario, the prediction would be that SMN Δ 7A is less protective than SMN Δ 7S because it is not a very good substrate for SCF^{Slmb}.

We therefore compared the stabilizing effects of overexpressing GFP-tagged SMN Δ 7^{S270A} (SMN Δ 7A) and SMN Δ 7 (SMN Δ 7S) on endogenous human SMN and SMN Δ 7. HEK 293T cells were transfected with equivalent amounts of GFP-SMN Δ 7A or -SMN Δ 7S. The following day, cells were harvested after treatment with cycloheximide (CHX) for zero to 10 h. As shown in Figure 6A, Western blotting with anti-SMN showed that the SMN Δ 7S construct exhibits a clear advantage over SMN Δ 7A in its ability to stabilize endogenous SMN and SMN Δ 7. By comparing band intensities within a given lane, we generated average intensity ratios for each time point using replicate blots (Figure 6A, table). We then calculated a “stabilization factor” by taking a ratio of these two ratios. As shown (Figure 6A, graph), the protective benefit of overexpressing Δ 7S versus Δ 7A at $t = 0$ h was roughly 3.0 \times for endogenous SMN Δ 7 and 1.75 \times for full-length SMN. Thus, as predicted above, the GFP-SMN Δ 7A construct was much less effective at stabilizing endogenous SMN isoforms. Because SMN Δ 7 is a relatively good SCF^{Slmb} substrate, overexpression of this isoform protects full-length SMN from degradation.

As mentioned above, experiments in an SMN-deficient chicken DT40 cell line showed that expression of SMN Δ 7A, but not SMN Δ 7S, rescued cellular proliferation (Cho and Dreyfuss, 2010). These results suggest that, when stable, SMN Δ 7 is intrinsically functional. To examine SMN Δ 7A functionality in a more disease-relevant cell type, control and SMA-induced pluripotent stem cell (iPSC) motor neuron cultures were transduced with lentiviral vectors expressing an mCherry control protein or SMN Δ 7A (Figure 6B). At 4 wk postdifferentiation, no statistical difference was observed between control and SMA motor neurons; however, by 6 wk, SMA motor neuron numbers had decreased significantly to ~7% of the total cell population (Figure 6B). In contrast, expression of SMN Δ 7A maintained motor neuron numbers to approximately the same level as the controls and nearly twofold greater than untreated cells (Figure 6B). Thus expression of SMN Δ 7A improves survival of human iPSCs when differentiated into motor neuron lineages.

SMN Δ 7A is a protective modifier of intermediate SMA mouse phenotypes

To examine the importance of the Slmb degron in a mammalian organismal system, two previously developed SMA mouse models were utilized. As mentioned above, the “Delta7” mouse

(*Smn*^{-/-};*SMN2*;*SMN* Δ 7) is a model of severe SMA (Le *et al.*, 2005), and affected pups usually die between P10 and P18 (avg. = P15). The “2B/-” mouse (*Smn*^{2B/-}) is a model of intermediate SMA (Bowerman *et al.*, 2012; Rindt *et al.*, 2015) and these animals survive much longer, typically between P25 and P45 (avg. = P32). Adeno-associated virus serotype 9 (AAV9) was selected to deliver the SMN cDNA isoforms to these SMA mice, as this vector has previously been shown to enter and express in SMA-relevant tissues and can dramatically rescue the SMA phenotype when expressing the wild-type SMN cDNA (Foust *et al.*, 2010; Passini *et al.*, 2010; Valori *et al.*, 2010; Dominguez *et al.*, 2011; Glascock *et al.*, 2012).

Delivery of AAV9-SMN Δ 7A at P1 significantly extended survival in the intermediate 2B/- animals, resulting in 100% of the treated pups living beyond 100 d, similar to the results obtained with the full-length AAV9-SMN construct (Figure 7A). In contrast, untreated 2B/- animals lived, on average, only 30 d. Mice treated with AAV9-SMN Δ 7S survived an average of 45 d (Figure 7A). Mice treated with AAV9-SMN Δ 7D, a phosphomimetic of the wild-type serine 270 residue, have an average life span that is equivalent or slightly shorter than that of untreated 2B/- mice (Figure 7A). These results not only highlight the specificity of the S270A mutation in conferring efficacy to SMN Δ 7 but also illustrate that AAV9-mediated delivery of protein alone does not improve the phenotype.

We also analyzed the effects of SMN Δ 7A expression in the severe Delta7 mouse model (Le *et al.*, 2005). Treatment with AAV9-SMN Δ 7A had only a very modest effect on Delta7 mice, as none of the animals (treated or untreated) survived weaning (Supplemental Figure S4). These findings are similar to the results in *Drosophila*. Transgenic expression of SMN Δ 7A in the *Smn* null background is not sufficient to rescue larval lethality (Supplemental Figure S3). Thus expression of SMN Δ 7A provides a clear protective benefit to the viability of intermediate mice but not to severe SMA models.

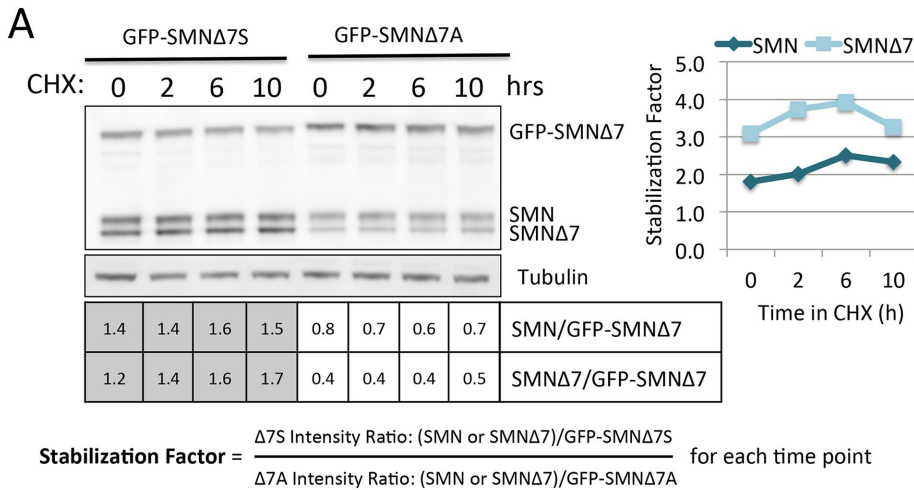
Consistent with the life-span data, AAV9-SMN Δ 7A treated 2B/- mice gained significantly more weight than either untreated or AAV-SMN Δ 7S-treated animals, nearly achieving the same weight as pups treated with full-length AAV-SMN (Figure 7B). Treatment with full-length SMN cDNA resulted in animals that were clearly stronger and more mobile, consistent with the weight data (Figure 7C). Although they did not perform as well as mice treated with full-length SMN cDNA, the SMN Δ 7A-treated animals retained strength and gross motor function at late time points (e.g., P100), as measured by their ability to splay their legs and maintain a hanging position using a modified tube test, (Figure 7C). Animals treated with AAV9-SMN Δ 7D and -SMN Δ 7S did not survive long enough for testing.

SCF^{Slmb} primarily targets unstable SMN monomers

As indicated in Figure 8, our findings suggest a model whereby SMN and SMN Δ 7 degradation is in part mediated by SCF^{Slmb}, a multicomponent E3 ubiquitin ligase composed of Slmb, SkpA, Cul1, and Roc1 (Jiang and Struhl, 1998; Patton *et al.*, 1998a,b; Zheng *et al.*, 2002). Our work demonstrates that B-TrCP/Slmb binds directly to SMN (Figure 2) and is one of a growing number of E3 ligases in the cell that can target SMN protein (Kwon *et al.*, 2013; Han *et al.*, 2016). SMN monomers, such as those created in SMN Δ 7, are the primary targets for degradation. As shown in the model, partially active SMN•SMN Δ 7 dimers and active SMN oligomers are also substrates but to a lesser extent.

DISCUSSION

Factors that recognize the putative SMN Δ 7-specific degron have not been identified, and the molecular mechanisms governing proteasomal access to SMN and SMN Δ 7 remain unclear. In this study, we



ligase complex as a novel SMN binding partner whose interaction is conserved in human. Depletion of Slmb or B-TrCP by RNAi resulted in an increase in steady-state SMN levels in *Drosophila* and human cells, respectively. We also showed that ectopic expression of SMNΔ7^{S270A}, but not SMNΔ7 or SMNΔ7^{S270D}, a phosphomimetic, is a protective modifier of SMA phenotypes in animal models and human iPSC cultures.

The SCF^{Slmb} degron is exposed by SMN2 exon skipping

A previous study posited that a phosphodegron was specifically created by exon 7 skipping and that this event represented a key aspect of the SMA disease mechanism (Cho and Dreyfuss, 2010). Our identification of a putative Slmb binding site located in the C-terminal self-oligomerization domain of *Drosophila* and human SMN has allowed us to explore the molecular details of this hypothesis. The mutation of a conserved serine within the Slmb degron not only disrupted the interaction between SMN and Slmb but also stabilized full-length SMN and SMNΔ7. Notably, the degron mutation has a greater effect on SMN levels (both full-length and Δ7) when made in the context of a protein that does not efficiently self-oligomerize. These and other findings strongly suggest that the Slmb degron is uncovered when SMN is monomeric, whereas it is less accessible when SMN forms higher-order multimers. On the basis of these results, we conclude that SMN2 exon skipping does not create a potent protein degradation signal; rather, it exposes an existing one.

SMN targeting by multiple E2 and E3 systems

SMN degradation via the UPS is well established (Chang *et al.*, 2004; Burnett *et al.*, 2009; Kwon *et al.*, 2011). Using candidate approaches, investigators have studied other E3 ligases that have been reported to target SMN for degradation in cultured human cells (Hsu *et al.*, 2010; Kwon *et al.*, 2013; Han *et al.*, 2016). Given our findings, it is therefore likely that SMN is targeted by multiple E3 ubiquitin ligases, as this regulatory paradigm has been demonstrated for a number of proteins (e.g., p53; Jain and Barton, 2010). Targeting of a single protein by multiple E3 ligases is thought to provide regulatory specificity by expressing the

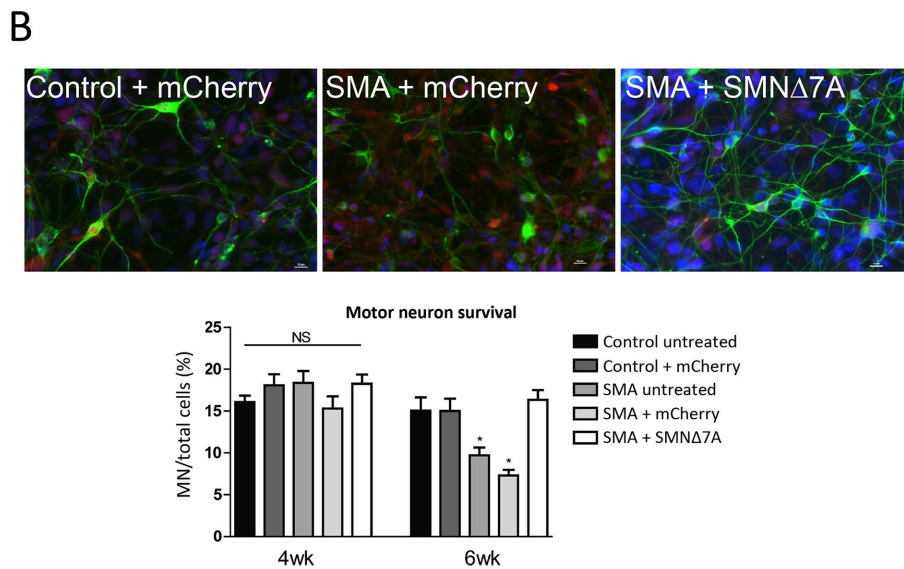


FIGURE 6: Stabilization of endogenous SMN and SMNΔ7 in cultured human cells. (A) HEK 293T cells were transfected with equivalent amounts of GFP-SMNΔ7A or -SMNΔ7S. The following day, cells were harvested after treatment with cycloheximide (CHX) for zero to 10 h. Western blotting with anti-SMN showed that SMNΔ7S stabilizes endogenous SMN and SMNΔ7 to a greater extent than SMNΔ7A. By comparing band intensities within a given lane, we generated average intensity ratios for each time point using replicate blots. We then calculated a “stabilization factor” by taking a ratio of these two ratios. The protective benefit of overexpressing Δ7S vs. Δ7A at t = 0 h was roughly 3.0× for endogenous SMNΔ7 and 1.75× for full-length SMN. (B) SMNΔ7A (S270A) expression protects SMA iPSC-derived motor neurons. Control motor neurons were left untreated or transduced with a lentiviral vector expressing an mCherry control. SMA motor neurons were left untreated or transduced with a lentiviral vector expressing an mCherry control or a lentiviral vector expressing SMNΔ7A (S270A). At 4 wk of differentiation, there was no difference in motor neuron survival between control and SMA iPSC motor neuron cultures in any of the treatment groups. However, at 6 wk, SMI-32-positive motor neurons showed selective loss in SMA iPSC motor neuron cultures in the untreated and lenti-mCherry groups compared with control iPSC motor neuron cultures. In contrast, lenti-SMNΔ7A expression fully protects SMA iPSC-derived motor neurons. Representative images of control and SMA iPSC-derived motor neurons labeled with SMI-32 (green) and mCherry (red). Nuclei are stained with DAPI and shown in blue. *p < 0.05 by ANOVA. NS = not significant. n = 3

isolated factors that copurify with SMN from *Drosophila* embryos that exclusively express Flag-SMN. This approach reduces potential bias towards SMN partner proteins that may be more abundant in a given tissue or cell line (Charroux *et al.*, 1999; Meister *et al.*, 2001; Pellizzoni *et al.*, 2002; Kroiss *et al.*, 2008; Trinkle-Mulcahy *et al.*, 2008; Guruharsha *et al.*, 2011). Here we identify the SCF^{Slmb} E3 ubiquitin

appropriate degradation complexes only within certain tissues, sub-cellular compartments, or developmental time frames. Moreover, ubiquitylation does not always result in immediate destruction of the target; differential use of ubiquitin lysine linkages or chain length can alter a protein's fate (Mukhopadhyay and Riezman, 2007; Ikeda and Dikic, 2008; Liu and Walters, 2010).

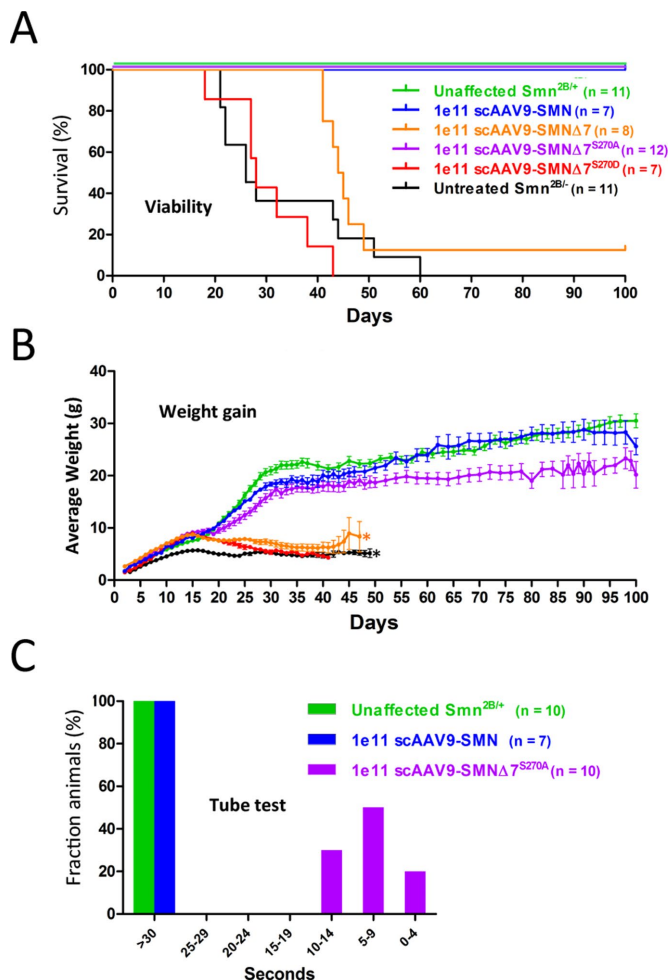


FIGURE 7: SMN Δ 7A is a protective modifier of intermediate SMA phenotypes in mice. (A) Mouse genotypes include control unaffected *Smn*^{2B/+} mice, which have a wild-type *Smn* allele, *Smn*^{2B/-} (2B/-) mice treated with scAAV9 expressing different versions of SMN, and untreated 2B/- mice, which are an intermediate mouse model of SMA. 1e11 denotes the viral dose. scAAV9-SMN expresses full-length SMN, scAAV9-SMN Δ 7 expresses truncated SMN, scAAV9-SMN Δ 7^{S270A} expresses truncated SMN with the S-to-A change in the degron, and scAAV9-SMN Δ 7^{S270D} expresses truncated SMN with a phosphomimic in the degron. Delivery of AAV9-SMN Δ 7A at P1 significantly extended survival in the intermediate 2B/- animals, resulting in 100% of the treated pups living beyond 100 d, similar to the results obtained with the full-length AAV9-SMN construct. Untreated 2B/- animals lived, on average, only 30 d. Mice treated with AAV9-SMN Δ 7S survived an average of 45 d. Mice treated with AAV9 expressing SMN Δ 7D had an average life span equivalent or slightly worse than that of untreated 2B/- mice. (B) Average weight (measured over time) of the animals used in panel A. AAV9-SMN Δ 7A treated mice also gained significantly more weight than either untreated or AAV-SMN Δ 7S-treated animals, nearly achieving the same weight as 2B/- pups treated with full-length SMN cDNA. (C) Mouse genotypes include control unaffected *Smn*^{2B/+} mice, which carry a wild-type *Smn* allele, and 2B/- mice treated with scAAV9 expressing different versions of SMN. scAAV9-SMN expresses full-length SMN and scAAV9-SMN Δ 7^{S270A} expresses truncated SMN with the S-to-A change in the degron. AAV-SMN Δ 7A-treated animals retained their improved strength and gross motor functions at late time points (P100), as measured by their ability to splay their legs and maintain a hanging position using a modified tube test.

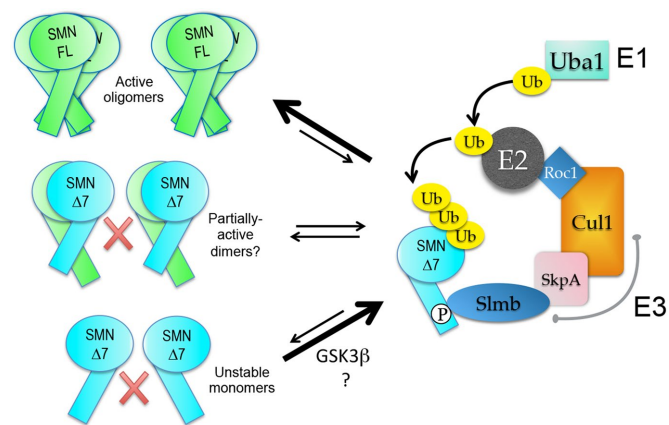


FIGURE 8: Proposed model of SMN as a substrate of SCF^{Slmb} E3 ubiquitin ligase. Unstable SMN monomers, such as those created in SMN Δ 7, are the primary substrates for degradation. Active oligomers of full-length SMN (SMN-FL) and partially active SMN-FL•SMN Δ 7 dimers (Praveen *et al.*, 2014; Gupta *et al.*, 2015) would be targeted to a lesser extent. SCF^{Slmb} is a multicomponent E3 ubiquitin ligase composed of Slmb, SkpA, Cul1, and Roc1 (see the text for details). This E3 ligase complex functions together with E1 and E2 proteins in the ubiquitin proteasome system (UPS) to tag proteins for degradation by linkage to ubiquitin (Ub). Phosphorylation (P) by GSK3 β and/or another kinase (see the text) is predicted to trigger ubiquitination.

Avenues of future exploration include determination of the E2 proteins that partner with SCF^{Slmb} as well as the types of ubiquitin lysine chain linkages they add to SMN. These two questions are interconnected, as ubiquitin linkage specificity is determined by the E2 (Ye and Rape, 2009). Lysine 48 (K48) linked chains typically result in degradation of the targeted protein by the 26S proteasome, whereas lysine 63 (K63) linkage is more commonly associated with lysosomal degradation and nonproteolytic functions such as endocytosis (Tan *et al.*, 2007; Kirkin *et al.*, 2009; Lim and Lim, 2010). Interestingly, recent work has implicated defects in endocytosis in SMA (Custer and Androphy, 2014; Dimitriadi *et al.*, 2016; Hosseinbar-kooie *et al.*, 2016). It remains to be determined how the ubiquitylation status of SMN might intersect with endocytic functions.

Does SMN function as a signaling hub?

In the Flag-SMN pull down, we identified three E2 proteins as potential SMN interacting partners (Figure 1C). Among them, Bendless (Ben) is particularly interesting. Ben physically interacts with TRAF6, an E3 ligase that functions together with Ube2N/Ubc13/Ben in human cells (Kim and Choi, 2017). TRAF6 is an activator of NF- κ B signaling, and its interaction with SMN is thought to inhibit this activity (Kim and Choi, 2017). Notably, Ube2N/Ben heterodimerizes with Uev1a to form K63 ubiquitin linkages on target proteins (Ye and Rape, 2009; van Wijk and Timmers, 2010; Komander and Rape, 2012; Marblestone *et al.*, 2013; Zhang *et al.*, 2013). Furthermore, Ben-Uev1a is involved in upstream activation of both JNK and IMD signaling in *Drosophila* (Zhou *et al.*, 2005; Paquette *et al.*, 2010). Previously, we and others have shown that JNK signaling is dysregulated in animal models of SMA (Garcia *et al.*, 2013, 2016; Genabai *et al.*, 2015; Ahmad *et al.*, 2016). Moreover, mutations in all three components of SCF^{Slmb} lead to constitutive expression of antimicrobial peptides, which are also downstream of the IMD pathway (Khush *et al.*, 2002). Together, these findings suggest the interesting possibility of SMN functioning as a signaling hub that links the UPS

to the JNK and IMD pathways, all of which have been shown to be disrupted in SMA.

Phosphorylation of the Slmb degnon within SMN

As Slmb is known to recognize phospho-degrons, one of the first questions raised by our study concerns the identity of the kinase(s) responsible for phosphorylating the degnon in SMN. A prime candidate is GSK3 β (Figure 8), as this kinase recognizes a motif (SxxxS/T; Liu *et al.*, 2007; Lee *et al.*, 2013) that includes the degnon and extends N-terminally (²⁶²SxxxSxxxSxxxT²⁷⁴, numbering as per human SMN). In support of this hypothesis, we identified the *Drosophila* GSK3 β orthologue, Shaggy (Sgg), in our SMN pull downs (Figure 1C). Moreover, GSK3 β inhibitors as well as siRNA-mediated knock-down of GSK3 β were shown to increase SMN levels, primarily by stabilizing the protein (Makhortova *et al.*, 2011; Chen *et al.*, 2012). Finally, GSK3 β is also responsible for phosphorylation of a degnon in β -catenin, a well-characterized SCF^{Slmb} substrate (Liu *et al.*, 2002). SMA mice have low levels of UBA1 (E1), ultimately leading to accumulation of β -catenin (Wishart *et al.*, 2014). Pharmacological inhibition of β -catenin improved neuromuscular pathology in *Drosophila*, zebrafish, and mouse SMA models. β -Catenin had previously been shown to regulate motor neuron differentiation and stability by affecting synaptic structure and function (Murase *et al.*, 2002; Li *et al.*, 2008; Ojeda *et al.*, 2011). β -Catenin also regulates motor neuron differentiation by retrograde signaling from skeletal muscle (Li *et al.*, 2008). The connections of UBA1 and multiple SCF^{Slmb} substrates to motor neuron health thus places the UPS at the center of SMA research interest.

Concluding remarks

In summary, this study identifies conserved factors that regulate SMN stability. To our knowledge, this work represents the first time that SMN complexes have been purified in the context of an intact developing organism. Using this approach, we have demonstrated that the SCF^{Slmb} E3 ligase complex interacts with a degnon embedded within the self-oligomerization domain of SMN. Our findings establish plausible connections to disease-relevant cellular processes and signaling pathways. Further, they elucidate a model (Figure 8) whereby accessibility of the SMN phosphodegnon is regulated by self-multimerization, providing an elegant mechanism for balancing functional activity with degradation.

MATERIALS AND METHODS

Fly stocks and transgenes

Oregon-R was used as the wild-type control. The Smn^{X7} microdeletion allele (Chang *et al.*, 2008) was a gift from S. Artavanis-Tsakonis (Harvard University, Cambridge, MA). This deficiency removes the promoter and the entire SMN coding region, leaving only the final 44bp of the 3' UTR. All stocks were cultured on molasses and agar at room temperature (24 \pm 1°C) in half-pint bottles. The Smn transgenic constructs were injected into embryos by BestGene (Chino Hills, CA) as described in Praveen *et al.*, 2014. In short, a ~3 kb fragment containing the entire Smn coding region was cloned from the *Drosophila* genome into the pAttB vector (Bischof *et al.*, 2007). A 3X FLAG tag was inserted immediately downstream of the start codon of dSMN. Point mutations were introduced into this construct using Q5 (NEB) site-directed mutagenesis according to manufacturer's instructions. The basal Smn construct used, vSmn, contained three single-amino-acid changes, and the addition of the MGLR motif to make fruit fly Smn more similar to the evolutionarily conserved vertebrate Smn. Subsequently generated constructs used vSmn as a template and consist of the amino acid changes detailed in

Figure 4. Y203C, G206S, and G210V were previously published in Praveen *et al.*, 2014.

Drosophila embryo protein lysate and mass spectrometry

Drosophila embryos (0–12 h) were collected from Oregon-R control and Flag-SMN flies, dechorionated, flash frozen, and stored at –80°C. Embryos (approximately 1g) were then homogenized on ice with a Potter tissue grinder in 5 ml of lysis buffer containing 100 mM potassium acetate, 30 mM HEPES–KOH at pH 7.4, 2 mM magnesium acetate, 5 mM dithiothreitol (DTT), and protease inhibitor cocktail. Lysates were centrifuged twice at 20,000 rpm for 20 min at 4°C and dialyzed for 5 h at 4°C in Buffer D (HEPES 20 mM, pH 7.9, 100 mM KCl, 2.5 mM MgCl₂, 20% glycerol, 0.5 mM DTT, PMSF (0.2 mM)). Lysates were clarified again by centrifugation at 20,000 rpm for 20 min at 4°C. Lysates were flash frozen using liquid nitrogen and stored at –80°C before use. Lysates were then thawed on ice, centrifuged at 20,000 rpm for 20 min at 4°C and incubated with rotation with 100 μ l of EZview Red Anti-FLAG M2 affinity gel (Sigma) for 2 h at 4°C. Beads were washed a total of six times using buffer with KCl concentrations ranging from 100 to 250 mM with rotation for 1 min at 4°C in between each wash. Finally, Flag proteins were eluted 3 consecutive times with one bed volume of elution buffer (Tris 20 mM, pH 8, 100 mM KCl, 10% glycerol, 0.5 mM DTT, PMSF 0.2 mM) containing 250 μ g/ml 3XFLAG peptide (sigma). The entire eluate was used for mass spectrometry analysis on an Orbitrap Velos instrument, fitted with a Thermo Easy-spray 50-cm column.

Tissue culture and transfections

S2 cell lines were obtained from the *Drosophila* Genome Resource Center (Bloomington, IL). S2 cells were maintained in SF900 SFM (Life Technologies) supplemented with 1% penicillin/streptomycin and filter sterilized. Cells were removed from the flask using a cell scraper and passaged to maintain a density of ~10⁶–10⁷ cells/ml. S2 cells were transferred to filter sterilized SF900 SFM (Life Technologies) without antibiotics prior to transfection with Cellfectin II (Invitrogen). Transfections were performed according to Cellfectin II protocol in a final volume of 4 ml in a T-25 flask containing 10⁷ cells that were plated 1 h before transfection. The total amount of DNA used in transfections was 2.5 μ g. Human embryonic kidney HEK-293T and HeLa cells were maintained at 37°C with 5% CO₂ in DMEM (Life Technologies) supplemented with 10% fetal bovine serum (FBS) and 1% penicillin/streptomycin (Life Technologies). Cells (1 \times 10⁶–2 \times 10⁶) were plated in T-25 flasks and transiently transfected with 1–2 μ g of plasmid DNA per flask using Lipofectamine (Invitrogen) or FuGENE HD transfection reagent (Roche Applied Science, Indianapolis, IN) according to the manufacturer's protocol. Cells were harvested 24–72 h posttransfection.

For siRNA transfections, HeLa cells were plated subconfluently in T-25 flasks and transfected with 10 nm of siRNA (gift from Mike Emanuele lab) and 17 μ l Lipofectamine RNAi MAX (Invitrogen) in 5 ml total media according to manufacturer's instructions. After 48 h of transfection, cells were harvested. For RNAi in S2 cells using dsRNA, 10⁷ cells were plated in each well of a six-well plate in 1 ml of media. Cells were treated approximately every 24 h with 10 μ g/ml dsRNA targeted against Slmb, Oskar, or Gaussia Luciferase (as controls) as described in Rogers and Rogers, 2008.

In vitro binding assay

GST and GST-SMN were purified from *Escherichia coli*. In brief, cells transformed with BL21*GST-SMN were grown at 37°C overnight and then induced using 1 mM IPTG. Recombinant protein was extracted and purified using Glutathione sepharose 4B beads. GST-B-TrCP1

was purchased from Novus Biologicals (cat# H00008945). SMN•Gem2 complexes were coexpressed in *E. coli* using SMNΔ5 and Gemin2(12–280) constructs, as described in Gupta et al. (2015). Glutathione sepharose 4B beads were washed 3× with phosphate-buffered saline (PBS). GST alone, GST-SMN, or GST-B-TrCP1 were attached to beads during 4-h-overnight incubation at 4°C in PBS [AQ 13] with rotation. Beads were then washed 3× with modified RIPA buffer (50 mM Tris-HCl, pH 7.5, 250 mM NaCl, 1 mM EDTA, 1% NP-40). Beads (20 μl) with ~2 μg attached GST-tagged protein (as determined by Coomassie stain with BSA standard) were added to 200 μl [AQ 14] modified RIPA buffer with 100 μg/ml BSA block. SMN•Gem2 (2 μg) was added, and the mixture was rotated end over end at 4°C overnight. Beads were then washed 3× with modified RIPA buffer, and 10 μl SDS loading buffer was added followed by boiling for 5 min.

In vivo ubiquitylation assay

The in vivo ubiquitylation assay was performed as described previously (Choudhury et al., 2016). Briefly, HEK-293T cells were transfected as indicated in 10-cm dishes using Lipofectamine2000 (Thermo Fisher Scientific). The day after, cells were treated with 20 μM MG132 for 4 h and then harvested in PBS. Of the cell suspension, 80% was lysed in 6 M guanidine-HCl-containing buffer and used to pull down His-Ubiquitinated proteins on Ni²⁺-NTA beads, while the remaining 20% was used to prepare inputs. Ni²⁺ pull-down eluates and inputs were separated through SDS-PAGE and analyzed by Western blot.

Cycloheximide treatment

Following RNAi treatment, S2 cells were pooled, centrifuged, and resuspended in fresh media. One-third of these cells were frozen and taken as the 0 h time point. The remainders of the cells were replated in six-well plates. Cycloheximide (CHX; 100 μg/ml) was added to each sample, and cells were harvested at 2 and 6 h following treatment.



Immunoprecipitation

[AQ 15] Clarified cell lysates were precleared with immunoglobulin G agarose beads for 1 h at 4°C and again precleared overnight at 4°C. The precleared lysates were then incubated with anti-FLAG antibody cross-linked to agarose beads (EZview Red Anti-FLAG M2 affinity gel, Sigma) for 2 h at 4°C with rotation. The beads were washed with lysis buffer or modified lysis buffer six times and boiled in SDS gel-loading buffer. Eluted proteins were run on an SDS-PAGE for Western blotting.

Antibodies and Western blotting

Larval and adult lysates were prepared by crushing the animals in lysis buffer (50 mM Tris-HCl, pH 7.5, 150 mM NaCl, 1 mM EDTA, 1% NP-40) with 1× (adults) or 10× (larvae) protease inhibitor cocktail (Invitrogen) and clearing the lysate by centrifugation at 13,000 rpm for 10 min at 4°C. S2 cell lysates were prepared by suspending cells in lysis buffer (50 mM Tris-HCl, pH 7.5, 150 mM NaCl, 1 mM EDTA, 1% NP-40) with 10% glycerol and 1× protease inhibitor cocktail (Invitrogen) and disrupting cell membranes by pulling the suspension through a 25-gauge needle (Becton Dickinson). The lysate was then cleared by centrifugation at 13,000 rpm for 10 min at 4°C. Human cells (293T and HeLa) were first gently washed in ice-cold 1× PBS and then collected in ice-cold 1× PBS by scraping. Cells were pelleted by spinning at 1000 rpm for 5 min. The supernatant was removed, and cells were resuspended in ice-cold lysis buffer (50 mM Tris-HCl, pH 7.5, 150 mM NaCl, 1 mM EDTA, 1% NP-40) and allowed to lyse on ice for 30 min. After lysing, the lysate was cleared by

centrifuging the cells for 10 min at 13,000 rpm at 4°C. Western blotting [AQ 16] on lysates was performed using standard protocols. Rabbit anti-dSMN serum was generated by injecting rabbits with purified full-length dSMN protein (Pacific Immunology Corp, CA) and was subsequently affinity purified. For Western blotting, dilutions of 1 in 2500 for the affinity purified anti-dSMN, 1 in 20,000 (fly) or 1 in 5000 (human) for anti-α-tubulin (Sigma), 1 in 10,000 for monoclonal anti-Flag (Sigma), 1 in 1000 for anti-Slmb (gift from Greg Rogers), 1 in [AQ 17] 2500 for anti-human SMN (BD Biosciences), 1 in 1000 for anti-B-TrCP (gift from MB Major lab), 1 in 10,000 for polyclonal anti-Myc (Santa Cruz), and 1 in 2000 for anti-GST (Abcam) were used.

Larval locomotion

Smn control and mutant larvae (73–77 h post-egg laying) were placed on a 1.5% agarose molasses tray five at a time. The tray was then placed in a box with a camera, and the larvae were recorded moving freely for 60 s. Each set of larvae was recorded three times, and one video was chosen for analysis based on video quality. The videos were then converted to AVI files and analyzed using the wrMTrck plug-in of the Fiji software. The “Body Lengths per Second” was calculated in wrMTrck by dividing the track length by half the perimeter and time (seconds). *p* Values were generated using a multiple comparison analysis of variance (ANOVA).

SMA mouse models

Two previously developed SMA mouse models were utilized. The “Delta7” mouse (*Smn*^{-/-}; *SMN2*; *SMNΔ7*) is a model of severe SMA (Le et al., 2005). The “2B/-” mouse (*Smn*^{2B/-}) is a model of intermediate SMA (Bowerman et al., 2012; Rindt et al., 2015). Adeno-associated virus serotype 9 (AAV9) delivered SMN cDNA isoforms to these SMA mice, as previously described (Foust et al., 2010; Passini et al., 2010; Valori et al., 2010; Dominguez et al., 2011; Glascock et al., 2012). Gross motor function was measured using a modified tube test which tests the ability of mice to splay their legs and maintain a hanging position.

Human iPSC cell culture

Human iPSCs from two independent unaffected control and two SMA patient lines were grown as pluripotent colonies on Matrigel substrate (Corning) in Nutristem medium (Stemgent). Colonies were then lifted using 1 mg/ml Dispase (Life Technologies) and maintained as floating spheres of neural progenitor cells in the neural progenitor growth medium Stemline (Sigma) supplemented with 100 ng/ml human basic fibroblast growth factor (FGF-2; Miltenyi), 100 ng/ml epidermal growth factor (EGF; Miltenyi), and 5 μg/ml heparin (Sigma-Aldrich) in ultra-low attachment flasks. Aggregates were passaged using a manual chopping technique as previously described (Svendsen et al., 1998; Ebert et al., 2013). To induce motor neuron differentiation, neural progenitor cells were cultured in neural induction medium (1:1 DMEM/F12 [Life Technologies], 1× N2 Supplement [Life Technologies], 5 μg/ml Heparin [Sigma], 1× Non-Essential Amino Acids [Life Technologies], and 1× Antibiotic-Antimycotic [Life Technologies]) plus 0.1 μM all-trans retinoic acid (RA) for 2 wk; 1 μM Purmorphamine (PMN; Stemgent) was added during the second week. Spheres were then dissociated with TrypLE Express (Life Technologies) and plated onto Matrigel-coated 12-mm coverslips in *NIM* plus 1 μM RA, 1 μM PMN, 1× B27 Supplement [AQ 18] (Life Technologies), 200 ng/ml ascorbic acid (Sigma), 1 μM cAMP (Sigma), 10 ng/ml *BDNF* (Peprotech), and 10 ng/ml *GDNF* (Peprot- [AQ 19] ech). One week postplating, cells were infected with lentiviral vectors (*MOI* = 5) expressing mCherry alone or SMN S270A-IRES- [AQ 20] mCherry. Transgenes in both viruses were under the control of the

EF1 α promoter. Uninfected cells served as controls. Cells were analyzed at 1 and 3 wk postinfection, which was 4 and 6 wk of total differentiation (Ebert *et al.*, 2009; Sareen *et al.*, 2013).

Immunocytochemistry

Coverslips were fixed in 4% paraformaldehyde (Electron Microscopy Sciences) for 20 min at room temperature and rinsed with PBS. Cells were blocked with 5% normal donkey serum (Millipore) and permeabilized in 0.2% Triton X-100 (Sigma) for 30 min at room temperature. Cells were then incubated in primary antibody solution for 1 h, rinsed with PBS and incubated in secondary antibody solution for 1 h at room temperature. Finally, nuclei were labeled with Hoechst nuclear stain (Sigma) to label DNA and mounted onto glass slides using FluoroMount medium (SouthernBiotech). Primary antibodies used were mouse anti-SMI-32 (Covance SMI-32R; 1:1000) and rabbit anti-mCherry (ThermoFisher; 1:1000). Secondary antibodies used were donkey anti-rabbit Cy3 (Jackson ImmunoResearch 711-165-152) and donkey anti-mouse AF488 (Invitrogen A21202).

Immunocytochemical analysis

Images were acquired from five random fields per coverslip using an inverted fluorescent microscope (Nikon) and NIS Elements software. Images were blinded and manually analyzed for antigen specificity with NIS Elements software.

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ETOC:

SMN protein levels inversely correlate with the severity of spinal muscular atrophy. The SCF^{Slmb}E3 ligase complex interacts with a degron embedded within the C-terminal self-oligomerization domain of SMN. The findings elucidate a model whereby accessibility of the SMN degron is regulated by self-multimerization.

Author Queries

- [AQ 1]: Please provide abbreviations list with definitions.
- [AQ 2]: Please define GST at first occurrence.
- [AQ 3]: Please define dsRNA at first occurrence.
- [AQ 4]: Matera and Van Duyne: if applicable please update and add to ref list. Otherwise unpublished per journal style.
- [AQ 5]: Please spell out DAPI.
- [AQ 6]: There is a Tan *et al.*, 2008 in list; change year here?
- [AQ 7]: Please define JNK at first occurrence.
- [AQ 8]: Please spell out UTR.
- [AQ 9]: Does gr mean grem (g)?
- [AQ 10]: Please define PMSF at first occurrence.
- [AQ 11]: Please define SFM at first occurrence.
- [AQ 12]: Please spell out IPTG.
- [AQ 13]: Please define RIPA at first occurrence.
- [AQ 14]: Please define BSA at first occurrence.
- [AQ 15]: Use of "immunoglobulin" instead of "immune-globulin" OK?
- [AQ 16]: rpm or $\times g$?
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- [AQ 18]: Please spell out NIM.
- [AQ 19]: Please spell out BDNF and GDNF.
- [AQ 20]: Please spell out MOI.
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- [AQ 25]: Monani 2005 not cited in text; delete from list?
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- [AQ 28]: added "a" and "b" labels Please check citations (a,b always cited together, OK?)
- [AQ 29]: Pearn 1982: Please provide volume number.
- [AQ 30]: Schrank B *et al.*, 1997 not cited in text; delete from list?
- [AQ 31]: Swoboda *et al.*, 2015 not cited in text; delete from list?
- [AQ 32]: please note that Tan *et al.*, 2007 was cited in text; add to list?
- [AQ 33]: Wirth 2000 not cited in text; delete from list?