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Intra-axonal translation of *Khsrp* mRNA slows axon regeneration by destabilizing localized mRNAs

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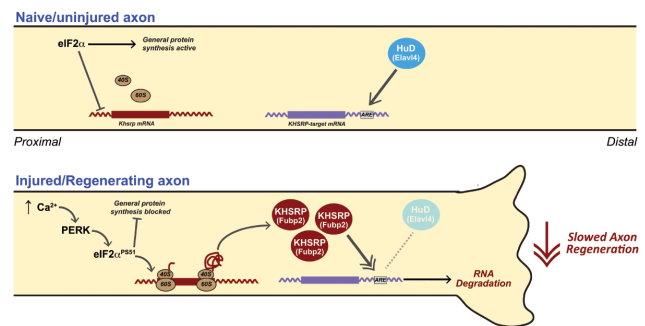
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ABSTRACT

Axonally synthesized proteins support nerve regeneration through retrograde signaling and local growth mechanisms. RNA binding proteins (RBP) are needed for this and other aspects of post-transcriptional regulation of neuronal mRNAs, but only a limited number of axonal RBPs are known. We used targeted proteomics to profile RBPs in peripheral nerve axons. We detected 76 proteins with reported RNA binding activity in axoplasm, and levels of several change with axon injury and regeneration. RBPs with altered levels include KHSRP that decreases neurite outgrowth in developing CNS neurons. Axonal KHSRP levels rapidly increase after injury remaining elevated up to 28 days post axotomy. *Khsrp* mRNA localizes into axons and the rapid increase in axonal KHSRP is through local translation of *Khsrp* mRNA in axons. KHSRP can bind to mRNAs with 3’UTR AU-rich elements and targets those transcripts to the cytoplasmic exosome for degradation. KHSRP knockout mice show increased axonal levels of KHSRP target mRNAs, *Gap43*, *Snap25*, and *Fubp1*, following sciatic nerve injury and these mice show accelerated nerve regeneration in vivo. Together, our data indicate that axonal translation of the RNA binding protein *Khsrp* mRNA following

nerve injury serves to promote decay of other axonal mRNAs and slow axon regeneration.

GRAPHICAL ABSTRACT



INTRODUCTION

Subcellular localization of mRNAs provides polarized cells with means to rapidly respond to environmental stimuli within different domains of those cells. Neurons are highly polarized cells with cytoplasmic processes, axons and dendrites, that extend great distances from the cell body or soma. Proteins synthesized in developing axons drive responses to some axon guidance cues and populations of proteins synthesized in axons changes during synaptogenesis pointing to dynamic post-transcriptional regulation of mRNAs in axons (1–3). In rodents, peripheral and some central nervous system (PNS and CNS, respectively) neu-

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rons can extend centimeters from their soma and intra-axonal protein synthesis can bring autonomy from the soma but it also must be tightly regulated (4). Several lines of evidence indicate that translation of mRNAs in mature axons of the peripheral nervous system (PNS) contributes to axon regeneration after injury (5). Since one mRNA can be translated many times over to generate multiple copies of a protein, the survival of an mRNA within an axon can substantially affect the spatial and temporal regulation of that axon's proteome (4). Much has been learned about how mRNAs are transported into and translated within axons over recent years (6–10), and it is clear that the axonal transcriptome is quite extensive in terms of numbers of different mRNAs and dynamic in terms of changes in mRNA populations with different physiological states (11). RNA binding proteins (RBP) and mRNAs assemble into ribonucleoprotein particles (RNP) for transport into axons, and those or other RBPs can also subsequently regulate the translation of interacting mRNAs in the axons or provide a storage depot to sequester the mRNAs until needed (4). However, we know of relatively few axonal RBPs that contribute to these mechanisms. There is also some intrinsic capacity for locally depleting specific mRNAs from axons in addition to regulating their transport, storage, and translation, since both nonsense-mediated decay (NMD) and microRNA (miRNA)-stimulated RNA degradation have been shown to occur in distal axons (12–16). But the extent to which localized mRNAs are subjected to regulation of their decay is not known.

RBP interactions have also been linked to mRNA stability, including axonally localized mRNAs. The neuronal protein HuD (also called ELAVL4) has been known to stabilize mRNAs by binding to AU-rich elements (AREs) in 3'UTRs of target mRNAs (17). HuD protein localizes into distal neurites and has been implicated in transport and translation of some mRNAs (18–20). For *Gap43* mRNA, HuD interaction is needed for axonal localization of the mRNA, but it also increases survival of the transcript (20). We previously showed that the KH splicing regulatory protein (KHSRP; also known as KSRP, MARTA1, ZBP2 and FUBP2) decreases neurite growth in cultures of embryonic cortical neurons and competes with HuD for binding to target mRNAs (21). In contrast to HuD interactions, KHSRP binding can promote decay of ARE-containing mRNAs (22,23). Here, we show that KHSRP is one of several RBPs whose levels increase in PNS axons after injury and during regeneration. This increase in axonal KHSRP occurs rapidly after PNS nerve injury through translation of its mRNA in axons. A conventional knockout of the murine *KHSRP* gene increases axonal levels of the KHSRP target mRNAs, *Gap43*, *Snap25* and *Fubp1*. KHSRP contains 4 hnRNP K homology (KH) domains of about 70 amino acid residues each and these can bind to single strand DNA or RNA with varying degrees of specificity (22). Interestingly, the increase in *Gap43* mRNA requires an intact fourth KH domain in KHSRP that has been linked to promoting mRNA decay by interacting with components of the cytoplasmic exosome (22). In contrast, KHSRP's modulation of *Fubp1* mRNA levels does not require this domain of KHSRP. Selectively deleting KHSRP alleles from only neurons points to a neuron intrinsic mechanism driving

the accelerated axon regeneration. Together, our data indicate that neuronal KHSRP slows axon growth and emphasize that localized synthesis of KHSRP in axons provides a means to modulate axonal mRNA levels, which slows nerve regeneration.

MATERIALS AND METHODS

Animal use and survival surgery

The Institutional Animal Care and Use Committee of the University of South Carolina approved all animal procedures. Adult male Sprague Dawley rats (175–250 g) or both male and female *Khsrp* knockout (*Khsrp*^{-/-}) (24), wild type (*Khsrp*^{+/+}) or *Khsrp*^{fl/fl} mice on C57/Bl6 background were used for all experiments. Wild type animals were typically littermates, and heterozygous animals (*Khsrp*^{+/-}) were used in several experiments as indicated. Mice for conditional knockout of *Khsrp* were generated by Biocytogen (Wakefield, MA) using CRISPR/EGE™-based gene editing to insert loxP sites between exons 1 and 2 and exons 6 and 7 that would result in a frameshift upon Cre-driven recombination but were not predicted to affect splicing of the *Khsrp* RNA transcript prior to any recombination. Insertion of loxP was confirmed by sequencing, and expression of full length *Khsrp* was confirmed by RT-PCR in founder mice on C57Bl/6 background. Founders were bred to homozygosity after crossing with wild type C57Bl/6 mice.

Isoflurane inhalation was used for anesthesia in all survival surgery experiments (see below). Animals were euthanized by CO₂ asphyxiation as indicated in results. For peripheral nerve injury, anesthetized male rats or mice were subjected to sciatic nerve crush at mid-thigh level as previously described (25). Briefly, the nerve at ~2.5 cm from its origin but proximal to its trifurcation was crushed with #2 fine jeweler's forceps, twice for 15 s. each; axotomy was monitored by the initial contraction of the hind limb upon applying pressure to the nerve, and then lack of hind paw extension during and upon recovery from anesthesia. For 'double crush' injury experiments, a unilateral peripheral sciatic nerve crush was performed at mid-thigh level on day 0, as a 'conditioning lesion', and a second crush injury was performed at 0.5 cm proximal to the first crush site following the same procedure. For consistency between animals, a single experimenter performed the crush injuries within each series of animals.

For the *Khsrp*^{fl/fl} mice, *in vivo* deletion of the *KHSRP* was accomplished by injecting 3 μl consisting of 1.32 × 10⁹ AAV2-CMV-Cre-GFP viral particles (Univ. North Carolina Vector Core, Chapel Hill, NC) diluted to 600 mM NaCl into the proximal sciatic nerve just distal to the sciatic notch and at least 1 cm from the crush sites (26).

Sciatic nerve ligations were performed in male rats as described previously as the larger size of these animals provided greater precision in ligation and subsequent crush injuries (27). Briefly, rat sciatic nerve was ligated approximately 1 cm proximal to planned mid-thigh nerve crush site. Immediately after applying 4-0 suture around the nerve and tying the suture to constrict the nerve, the sciatic nerve was crushed distal to the ligation site as above and then animals were euthanized 3–16 h later. Efficacy of the ligation was

assessed by immunofluorescence for anterogradely transported amyloid precursor protein (APP) and retrogradely transported signal transducer and activator of transcription 3 α (Stat3 α).

Cell Culture

Dissociated cultures of adult dorsal root ganglia (DRG) were prepared as described (28). For experiments with naïve DRG neurons, all lumbar, thoracic, and lower cervical DRGs were collected. To study effects of *in vivo* injury conditioning, L4-6 DRGs were used from ipsi-lateral (injury-conditioned) or contra-lateral (naïve) to the crush injury. DRGs were harvested in Hibernate-A medium (Brain-Bits, Springfield, IL) and then dissociated with collagenase as described. After centrifugation and three washes in DMEM/F12 (Life Technologies, Grand Island, NY), dissociated ganglia were cultured in complete medium containing DMEM/F12, 1 \times N1 supplement (Sigma, St. Louis, MO), 10% fetal bovine serum (Hyclone, Logan, UT), and 10 μ M cytosine arabinoside (Sigma) on poly-L-lysine (Sigma) plus laminin (Millipore, Burlington, MA) coated substrates. For imaging, dissociated DRGs were cultured on coated glass coverslips. For analyses of axonal RNA levels or *in vitro* regeneration assay (see below), dissociated ganglia were cultured on polyethylene-tetrate (PET) membrane inserts (1 μ m pores; Falcon-Corning, Tewksbury, MA) (29). Axons and CB were isolated from DRGs cultured on PET membranes as described (30).

For transfection, dissociated ganglia were pelleted at 100 \times g for 5 min and resuspended in 100 μ l 'Nucleofector solution' (Rat Neuron Nucleofector kit; Lonza, Alpharetta, GA). 4–6 μ g of plasmid was electroporated using the AMAXA Nucleofector device (Neurons Rat DRG, G-013 program; Lonza) before plating and maintained for 48 h.

For *in vitro* Cre-driven recombination, dissociated DRGs from *Khsrp^{fl/fl}* mice were incubated with 4.4 \times 10¹² particles/ml of AAV2-CMV-Cre-GFP or AAV2-CMV-GFP (UNC Vector Core) for 24 h after initial plating. Cultures were analyzed 3 d later. Loss of KHSRP was confirmed at both the mRNA and protein levels.

Plasmid constructs

pAc-GFP-KHSRP and pAc-GFP-KHSRP Δ KH4 (KHSRP with deleted KH4 domain) constructs have been published (21). All fluorescent reporter constructs for analyses of RNA translation were based on eGFP with myristoylation element (GFP^{MYR}; originally provided by Dr Erin Schuman, Max-Planck Institute, Frankfurt) (31). cDNAs for the 5'UTR and 3'UTRs of *Khsrp* mRNA were custom synthesized by Integrated DNA Technologies (Coralville, IA) and GenScript Biotech (Piscataway, NJ), respectively. The 5'UTR was engineered with 5' NheI and 3' BamHI restriction sites and cloned into pGFP^{MYR}5'camk2a/3'actg (30), replacing the 5'UTR of CamK2a [GFP^{MYR}5'khsrp/3'actg]. The 3'UTR sequence was engineered with 5' NotI and 3' XhoI restriction sites and used to replace the *Actg* 3'UTR in pGFP^{MYR}5'khsrp/3'actg plasmid [pGFP^{MYR}5'/3'khsrp].

Mass spectrometry for axonal RBPs

Axoplasm from 2 cm segments of sciatic nerve immediately proximal to crush site was extruded into nuclear transport buffer (20 mM HEPES [pH 7.3], 110 mM potassium acetate, 5 mM magnesium acetate) supplemented with protease/phosphatase inhibitor cocktail (Roche) and RNasin Plus (Promega, Madison, WI). Contralateral (uninjured) sciatic nerve of comparable level and length was used for control. Three animals were used for each time point and both naïve and injured sciatic nerve axoplasm. Preparations were cleared by centrifugation at 20 000 \times g, 4°C for 30 min, supernatants were diluted in 0.5 ml of TRIzol LS reagent (Invitrogen) and protein was extracted per the manufacturer's protocol. Protein pellets were digested with trypsin as previously described (32).

Parallel reaction monitoring (PRM) was performed on *Q Exactive Plus Mass Spectrometer* (Thermo-Fisher) online with *nanoAcquity UPLC System* (Waters, Milford, MA). Digested peptides samples (0.5 μ g) were injected onto 200 cm monolithic silica-C18 column (GL Sciences, Tokyo, Japan) and separated using a 6 h reversed phase chromatography gradient as previously described (32). The mass spectrometer was operated in PRM mode with the following parameters: positive polarity, $R = 17\,500$ at 200 m/z , AGC target 1e6, maximum IT 190 ms, MSX count 1, isolation window 3.0 m/z , NCE 35%. PRM data were analyzed in Skyline v. 3.5 (33). Skyline PRM document has been uploaded to *PanoramaWeb Public* (34) and can be accessed at <https://panoramaweb.org/axon-rbps.url>.

RNA isolation and PCR analyses

RNA was isolated from dissociated DRG neurons or cell body/axon compartments collected from insert cultures using *RNeasy Microisolation kit* (Qiagen, Hilden, Germany). Sciatic nerve was cut in small pieces and digested with collagenase at 37°C for 30 min with intermittent trituration. RNA was isolated from the collagenase-treated nerve using Trizol LS reagent (Invitrogen, Carlsbad CA) according to manufacturer's instructions. RNA concentration was measured by fluorimetry with *Ribogreen* (Life Technologies) and 10–50 ng of RNA was reverse transcribed with *Sensifast cDNA synthesis kit* (Bioline, London, UK). DRG axonal purity was assessed by RT-PCR, performed with primers designed to detect cell body-restricted mRNAs (*cJun* and *Map2*) and glial cell-specific mRNAs (*Gfap*). Droplet digital (dd) PCR was performed according to manufacturer's procedure with *Evagreen* detection (Biorad, Hercules, CA). Mitochondrial 12S ribosomal RNA (*Mtrnr1*) and Hmgb1 mRNA levels were used for normalizing RNA yields across different isolates. Following primers were used RT-PCR and RT-ddPCR (all from IDT, listed as 5' to 3'): *Mtrnr1*, sense – GGCTACACCTTGACCTAACG and antisense – CCTTACCCCTTCTCGCTAATTC; *Actb*, sense – CTGTC-CCTGTATGCCTCTG and antisense – ATGTCACG-CACGATTTCC; *cJun*, sense – GCAAAGATGGAAAC-GACCTTCTAC and antisense – AAGCGTGTCTGGC-TATGC *Gfap*, sense – AGTTACCAGGAGGCACTTG and antisense – GGTGATGCGGTTTTCTTCG; *Hmgb1*, sense – CATGGCAAAGGAGATCC and

antisense –CTCTGAGCACTTCTTGGAG; *Gap43*, sense – CAGGAAAGATCCCAAGTCCA and antisense – GAGGAAAGTGGACTCCACACA; *Map2*, sense – CTGGACATCAGCCTACTCA and antisense – AATAGGTGCCCTGTGACCTG; *Snap25*, sense – CAAATTTAACCCTTCCAGCA and antisense –CAGAATCGCCAGATCGACAG; *Fubp1*, sense – GCACCAGCTACAACCCAA and antisense – GCCTTTGTATAATCAACCTGTCC ; and *Khsrp*, sense – CCAGTTGAGAACCAATCGAGTC and antisense – CACCGTGAATAACAACACTCTCT.

Immunofluorescent staining

Immunofluorescence was performed as previously described (35) with all steps at room temperatures unless specified otherwise. Coverslips were fixed with 4% paraformaldehyde (PFA) in phosphate-buffered saline (PBS) for 15 min at room temperature and washed 3 times in PBS. PBS washed neurons were permeabilized with 0.3% Triton X-100 in PBS for 15 min and then blocked in 5% BSA for 1 h. Neurons were incubated with primary antibodies overnight in humidified chambers at 4°C. Primary antibodies consisted of chicken anti-NFH, -NFM plus -NFL cocktail (1: 500; Aves Lab, Tigard, OR, NFH # AB_2313552, NFM # AB_2313554, and NFL # AB_2313553), RT97 mouse anti-NF (1:500; Devel. Studies Hybridoma Bank, Iowa City, IA), and rabbit anti-KHSRP (1:200; Novus Biologicals, Centennial, CO, #NBP1-18910). After washes in PBST, coverslips were incubated with combination of FITC-conjugated donkey anti-mouse, Cy5 conjugated donkey anti-chicken (both at 1:500; Jackson ImmunoRes., West Grove, PA) as secondary antibodies for 1 h. After 1 h, coverslips were washed 3 times in PBS, rinsed with distilled H₂O, and mounted with *Prolong Gold Antifade with DAPI* (Thermo-Fisher, Waltham, MA).

For regeneration studies on mouse sciatic nerve and quantifying axonal content of KHSRP *in vivo*, sciatic nerve segments were fixed for 4 h in 4% PFA and then cryoprotected overnight in 30% sucrose in PBS at 4°C. 10 µm cryostat sections for rat sciatic nerve and 20 µm cryostat sections for mouse sciatic nerve were processed for immunostaining as previously described (35). Primary antibodies consisted of RT97 mouse anti-NF (1:500), rabbit anti-KHSRP (Novus Biologicals, #NBP1-18910), and rabbit anti-Stathmin-2/SCG10 (1:500; Novus Biologicals, #NBP1-49461). Stathmin-2/SCG10 immunofluorescence was used to detect regenerating mouse sciatic nerve axons (36). Cy3-conjugated donkey anti-rabbit and FITC-conjugated donkey anti-mouse in combination were used as secondary antibodies for rat sciatic nerve (both at 1:500, Jackson ImmunoRes.). Cy3-conjugated donkey anti-rabbit antibodies were used on mice sciatic nerve (1:500, Jackson ImmunoRes).

Immunofluorescence for neuromuscular junctions (NMJs) was performed as previously published with minor modifications (37). Briefly, all steps were carried out at room temperature. Gastrocnemius muscle was cleared of any connective tissue, washed in PBS, fixed in 4% PFA, washed in PBS 3 times for 5 min each. Muscle was then dissected into smaller pieces and incubated with 1 µg/ml of

Alexa Fluor 488 conjugated α -Bungarotoxin for 4 h with rocking (Thermo-Fisher, #B13422). Tissues were washed with PBS 3 times for 5 min each, treated with methanol at –20°C for 5 min, and rinsed in PBS 3 times for 5 min each with rocking. Tissues were blocked for 1 h with 2% BSA, 0.4% Triton X-100 in PBS. Tissues were then incubated overnight with the following cocktail of primary antibodies to presynaptic components diluted in blocking solution with rocking: rabbit anti-NF 200 (1:200; Millipore-Sigma, #N4142), mouse anti-synaptophysin (1:300; Millipore-Sigma, #MAB5258), and rabbit anti-synapsin-I (1:200; Millipore-Sigma, #AB1543P). The following day, tissues were rinsed 3 times 5 min each in PBS while rocking. Samples were then incubated in Cy3-conjugated donkey anti-rabbit and Cy3-conjugated donkey anti-mouse antibodies (1:500; Jackson ImmunoRes) for 4 h. After rinsing in PBS, muscle fibers were spread into monolayers under a stereomicroscope and affixed to slides using *Prolong Gold Antifade*; coverslips were sealed with clear nail polish.

All samples were mounted with *Prolong Gold Antifade* and imaging was performed at room temperature. Samples were analyzed by either epifluorescent or confocal microscopy. Leica DMI6000 epifluorescent microscope with ORCA Flash ER CCD camera (Hamamatsu) was used for epifluorescent imaging. Confocal imaging for immunofluorescence was performed on a Leica SP8X microscope (DMI6000 M platform; Buffalo Grove, IL) fitted with a galvanometer Z stage and HyD detectors; HC PL Apo 63x/1.4 NA objective (oil immersion) was used with acquisition parameters matched for individual experiments using LAS-X software. Z-stack images were post-processed by Leica *Lightning Deconvolution* integrated into *LASX* software. Deconvolved image stacks were projected into single plane images. For visualizing NMJs, sequential scanning was used to separate the green and red channels and Z stack at 200 nm intervals.

Fluorescence in-situ hybridization

Single molecule Fluorescence *in situ* hybridization (smFISH) plus IF was used to detect *Khsrp* mRNA in DRG and sciatic nerve. We used custom designed Cy3-labelled Stellaris probes (LGC Biosearch Tech, Middlesex, UK) for mouse *Khsrp* mRNA (Genbank ID # NM_010613.3) with Cy3-labelled scramble probe for control. RT97 mouse anti-NF (1:200) and chicken anti-GFAP (1:200; Millipore-Sigma, #AB5541) were used as primary antibodies; FITC- and Cy5-conjugated donkey anti-mouse and anti-chicken (1:200 each) were used as secondary antibodies. Samples processed without addition of primary antibody served as control for antibody specificity. Samples were mounted with *Prolong Gold Anti-fade with DAPI*.

smFISH/IF on DRG cultures was performed as described previously (38). Briefly, coverslips were rinsed in PBS and then fixed in buffered 2% PFA for 15 min, with all steps carried out at room temperature unless specified otherwise. Coverslips were rinsed 2 times in PBS, then permeabilized in 0.3% Triton X-100 in PBS for 5 min. Samples were equilibrated for 5 min in hybridization buffer (50% dextran sulphate, 10 µg/ml *E. coli* tRNA, 10 mM ribonucleoside vanadyl complex, 80 µg BSA, and 10% formamide

in $2\times$ SSC), and then incubated with 12.5 μ M probe plus mouse anti-NF (1:200) for 12 h at 37°C. Coverslips were then washed in PBS + 0.3% Triton X-100 3 times, followed by incubation with FITC-conjugated donkey anti-mouse for 1 h. After rinse in PBS, post fixation in 2% PFA for 15 min, and a second PBS wash, coverslips were inverted and mounted on glass slides.

Detection of mRNA in mouse tissues was done as previously described (38), with all steps carried out at room temperature unless specified otherwise. Briefly, sciatic nerve segments were fixed for 4 h in 2% PFA, cryoprotected overnight in 30% buffered sucrose at 4°C, and then cryosectioned at 20 μ m thickness (sections were stored at -20°C until used). Sections were brought to room temperature, washed three times in PBS for 5 min each, and then treated with 20 mM glycine and fresh 0.25 M NaBH₄ in PBS (3 times, 10 min each for both) to quench autofluorescence. Sections were quickly rinsed in 0.1 M Triethylamine (TEA) and then incubated in 0.1 M TEA + 0.25% acetic anhydride for 10 min. Sections were dehydrated in 70, 95 and 100% ethanol (3 min each) and then delipidated in chloroform for 5 min followed by 100 and 95% ethanol (3 min each). After washing in $2\times$ SSC, sections were incubated overnight at 37°C in a humidified chamber with 12.5 μ M probe and RT97 anti-NF (1:100) in hybridization buffer. The following day, sections were washed in $2\times$ SSC + 10% formamide at 37°C for 30 min, followed by two incubations in $2\times$ SSC for 5 min each. Sections were then briefly rinsed in PBS + 1% Triton-X100, and then incubated with donkey secondary antibodies diluted in 10 X blocking buffer (1:100; Roche, Penzberg, Germany) + 0.3% Triton-X100 for 1 h. Sections were finally washed in PBS for 5 min, post-fixed in 2% PFA for 15 min, washed 3 times in PBS (5 min each), rinsed in DEPC-treated water, and mounted under glass coverslips.

smFISH and IF signals were imaged using Leica SP8X as above. 63 \times /NA 1.4 oil immersion objective and pulsed white light laser was used for imaging RNA in both culture and tissue samples. Scramble probe was used to set the image acquisition parameter that would not acquire any nonspecific signal from scramble probe. Taking XYZ image stacks at least two locations in each section scanned nerve sections.

Detection of nascently synthesized proteins

Rat sciatic nerve from three animals per condition was either left naïve or *in vitro* crushed and incubated in DMEM medium containing, 10% FBS + Cyclosporin A (20 μ M; Sigma) + 1% penicillin/streptomycin. Nerves were treated with 200 μ g/ml anisomycin or vehicle (0.1% DMSO) for 3 h at 37°C, followed by adding 100 μ g/ml *O*-propargyl-puromycin (OPP; Invitrogen) for 1 h at 37°C. Axoplasm was extruded in 1 ml of transport buffer (20 mM HEPES [pH 7.4], 100 mM sodium acetate, 5 mM magnesium acetate), after extrusion SDS was added to 1%. Protein concentration was quantified by BCA assay and 350 μ g of total protein was used for biotin conjugation by click chemistry (100 μ M biotin-PEG3-azide) according to manufacturer's instruction. The reaction mix was incubated for 2 h at room temperature on a rotator. Five volumes ice-cold acetone was added to precipitate the protein. Protein pellets were resus-

pended in PBS containing 1% SDS. Streptavidin pull-down was carried out overnight at 4°C in 1 ml volume containing 60 μ l of streptavidin magnetic beads (Thermo-Fisher), 1% NP40, 0.1% SDS and 1 \times Complete EDTA-free protease inhibitor cocktail (Roche) in PBS. 10% of protein used for pull-down was taken to generate input samples. Beads were washed three times for 10 min with 1% NP40, 0.1% SDS in PBS at room temperature. Proteins were eluted from streptavidin beads by boiling for 10 min in 2 \times Laemmli sample buffer and then adjusted to 1 \times with PBS for denaturing polyacrylamide gel electrophoresis (SDS/PAGE).

Immunoblotting

Adult mouse DRG cultures (3 days *in vitro*, ~80 000 neurons/well) were lysed in Laemmli sample buffer and denatured by boiling at 95°C \times 5 min. Rat axoplasm from naïve and crushed sciatic nerves was extruded in nuclear transport buffer (20 mM HEPES [pH 7.3], 110 mM potassium acetate, 5 mM magnesium acetate, supplemented with protease inhibitors) as previously described (39). Lysates were cleared of debris by centrifugation at 15 000 \times g for 15 min at 4°C and then normalized for protein content using Bradford assay (Bio-Rad). Normalized protein lysates were fractionated by 10% SDS/PAGE and transferred onto a PVDF membrane (GE Healthcare Life Sciences, Marlborough, MA). After blocking in 5% non-fat dried milk powder (Bio-Rad) diluted in Tris-buffered saline with 1% Tween 20 (TBST), membranes were probed overnight at 4°C with rabbit anti-KHSRP (1:1000; Novus, # NBP1-18910), rabbit anti-GFP (1:1000; Enquire BioReagents, Littleton, CO, # QAB10298), rabbit anti-GAPDH (1:2000; Cell Signaling Tech, Beverly, MA, # 2118), rabbit anti- α -tubulin (1:1000; Cell Signaling Tech, # 2125); mouse anti-eIF2 α (1:1000; Cell Signaling Tech, # 2103) and rabbit anti-eIF2 α ^{PS51} (1:1000; Cell Signaling Tech, # 9721) antibodies or streptavidin-HRP (1:10 000, Abcam, # Ab_7403) diluted in blocking buffer. Blots were washed in TBST and then incubated with horseradish peroxidase (HRP)-conjugated anti-rabbit IgG (1:2000; Cell Signaling Tech.) diluted in blocking buffer for 1 h at room temperature. Blots were washed in TBST and signals were detected by ECL Prime™ (GE Healthcare Life Sciences).

Fluorescence recovery after photobleaching

Fluorescence recovery after photobleaching (FRAP) experiments were conducted at 37°C, 5% CO₂ as previously described (40). Briefly, dissociated adult mouse DRG cultures transfected with the GFP^{MYR}5'/3'khsrp were equilibrated in culture medium as above except phenol red was excluded. A region of interest (ROI) in the most distal axon of dissociated DRG neurons was photobleached with 488 nm argon laser set at 100% power for 80 frames at 0.65 s each. Pre-bleach and post-bleach signals were captured using 70% power for 488 nm laser line every 30 s (2 for pre-bleach and 30 for post-bleach). Translation dependence for recovery was tested by pre-treating DRG cultures with 100 μ g/ml anisomycin (Sigma) or 150 μ g/ml cycloheximide (Sigma) 20 min prior to photobleaching. For testing Ca²⁺-dependent translation by FRAP, transfected DRG

cultures were pretreated with 1 μ M thapsigargin (Sigma), 3 μ M BAPTA-AM (Sigma), 90 μ M GSK260614 (Bio-Techne Corp/Tocris, Minneapolis, MN) or 50 μ M Sephin1 (Apexbio, Houston, TX). Leica SP8X confocal microscope was used for imaging at 37°C, 5% CO₂ with 63X/NA 1.4 oil immersion objective. Pinhole was set to 3 Airy units for pre-bleach, bleach, and post-bleach sequences to ensure full thickness excitation of the axon. ROIs were 40 \times 40 μ m and at least 250 μ m from the soma.

Image analyses

ImageJ was used to quantify protein and RNA levels in sciatic nerve tissues from optical planes of XYZ scans. Axon only signal was extracted via *Colocalization plug-in* that extracted only protein or RNA signals that overlapped with axonal marker (NF) in each plane. Extracted 'axon only signal' was projected as a separate channel (38). Signal intensities were then calculated from each XY plane of these axon only channels. NF immunoreactivity area was used to normalize signal intensities across the individual XY planes. The relative signal intensity was then averaged for all tiles in each biological replicate.

To assess regeneration *in vivo*, tile scans were post-processed by *Straighten plug-in* for *ImageJ* (<http://imagej.nih.gov/ij/>). SCG10 fluorescence intensity was measured along the length of the nerve using *ImageJ*. Regeneration index was calculated by measuring the average SCG10 intensity in bins across at least 3 mm distal to the crush site. The crush site was defined by the position along the nerve length with maximal SCG10 intensity (secondarily confirmed by DAPI signals and DIC images). For analyses of axon growth *in vitro*, dissociated DRGs were immunostained with NF antibodies as described above. Images from 36 or 48 h cultures were used for neurite length and branching parameters (neurites/cell body and branch density) using WIS-Neuromath (41). For NMJs, confocal Z stacks from muscle were projected as single XY images using *ImageJ*. Nerve terminal and endplate (AChR) areas were calculated by *ImageJ* and fractional occupancy of NMJ was calculated by dividing nerve terminal area to endplate AChR area as described (37).

For FRAP image analyses, raw images sequences were analyzed for recovery in the bleached ROI using Leica confocal software package. Recovery was determined relative to pre-bleach and post-bleach signals, which were set at 100 and 0% to allow for comparisons between experiments and between neurons.

Statistical analyses

GraphPad Prism software package (La Jolla, CA) was used for statistical analyses. One-way ANOVA with Tukey post-hoc was used to compare between data points and Student's t-test was used to compare two independent groups for most experiments. For FRAP studies, two-way ANOVA with Tukey post-hoc was used, where control values were compared to cycloheximide- and anisomycin-treated cultures for each time point. All experiments were performed in at least triplicate. $P \leq 0.05$ was considered as statistically significant.

RESULTS

Peripheral nerve injury changes the axonal RNA binding protein population

Protein synthesis in PNS axons has been shown to facilitate nerve regeneration after injury (5). Transport and translation of the mRNA templates needed for this intra-axonal protein synthesis are driven by proteins interacting with those mRNAs (4). We recently showed that many RBPs that were thought to have exclusively nuclear roles localize into PNS axons by an RNA affinity mass spectrometry (RAMS) approach (42). The levels of some of these RBPs increased after axotomy, which pointed to functions in growing axons. As this RAMS assay focused on interactions with axonal mRNA localization motifs, we wanted to gain a more systematic and unbiased view of the axonal RBP population. With the limited amount of protein obtained from PNS axoplasm coupled with the likely low abundance of RBPs relative to cytoskeletal components in the axons, we turned to a targeted mass spectrometry approach using PRM to profile axonal RBPs in naïve, injured and regenerating sciatic nerve axoplasm. By mining the *UniProt* database (<https://www.uniprot.org>) for proteins with 'RNA binding' in functional or domain descriptions and validated expression in the nervous system (cross-referencing each for RNA expression in neurons using the *GeneCards* database (<https://www.genecards.org>), we arrived at 357 proteins to test. Of these, 196 were represented in a reference MS library that had been generated from rat nervous system tissues (including sciatic nerve, dorsal root ganglion (DRG), and spinal cord) and were included in a PRM-MS method that targeted 511 unique precursor peptides (26). Using this PRM method, we quantified the abundance of 84 RBPs represented by 184 precursor peptides in adult rat sciatic nerve axoplasm. Several of the RBPs showed increased or decreased axoplasm levels over 3–28 days post injury compared to the contralateral uninjured sciatic nerve (Figures 1A, B; Supplemental Figure S1). Initial validation by immunoblotting from naïve and 7 days post-crush injury sciatic nerve axoplasm showed that FXR1, hnRNP A3, hnRNP AB, hnRNP H1 and KHSRP increased in the 7 days samples (Figure 1C). We have previously reported axonal levels of hnRNP H1 similarly increase following nerve crush injury (42).

It is appealing to speculate that the increased levels of these RBPs in injured and regenerating peripheral nerve axoplasm would serve to promote regeneration. For example, hnRNP H1 was identified by the RAMS approach as binding to *Hmgb1* mRNA's axonal localization motif (42), and local translation of *Hmgb1* mRNA promotes axons growth (43). The increase in KHSRP after injury (Figure 1D-E) was surprising as this RBP has been shown to promote mRNA decay (22,23), including targeting *Gap43* mRNA, whose encoded protein has long been linked to axon growth promotion, for degradation (21). KHSRP also has other cytoplasmic functions including RNA transport and microRNA (miRNA) processing that could support axon regeneration (44,45). Thus, we examined KHSRP more closely to determine which function the increased axoplasmic KHSRP might serve in the injured sciatic nerve. Intriguingly, the axoplasm KHSRP levels were significantly

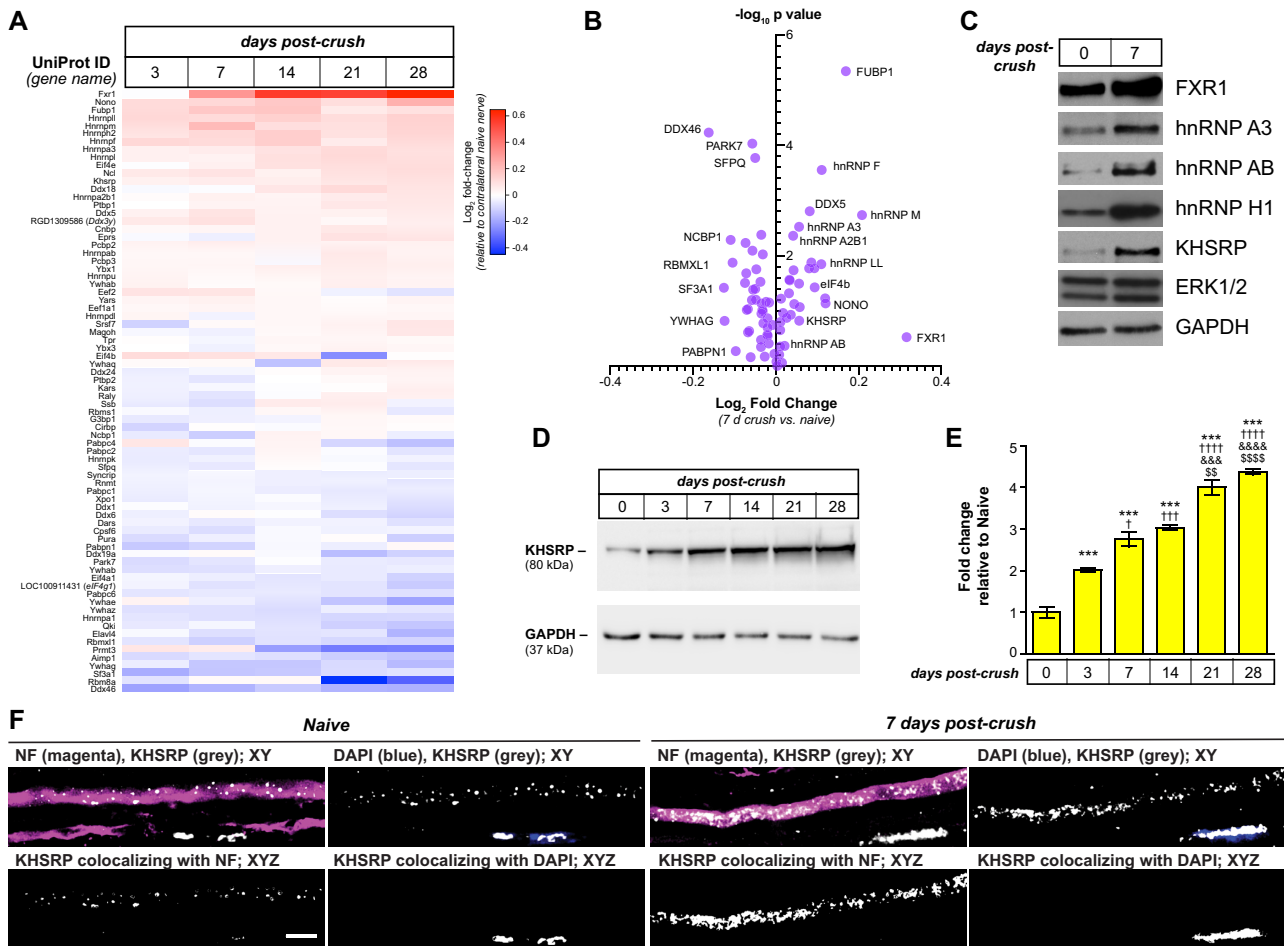


Figure 1. Peripheral nerve injury changes the axonal RNA binding protein populations. (A, B) Sciatic nerve axoplasm harvested proximal to the injury site from 3–28 days post-crush lesion was trypsin-digested and processed for liquid chromatography and mass spectrometry using parallel reaction monitoring (PRM) to detect proteins with known RNA binding activity. Levels of proteins from spectral counts relative to uninjured (naïve) axoplasm shown are Log_2 fold-change as indicated in (A) ($N = 3$ for each time point). (B) shows volcano plot for PRM results for 7 days crush versus naïve samples graphed as log_2 fold-change versus negative $\text{log}_{10}P$ value for differences. Also see Supplemental Figure S1 for graphical representation of full time course. (C) Representative immunoblot for naïve and 7 days injured sciatic nerve axoplasm confirms the increase in FXR1, hnRNP A3, hnRNP AB, hnRNP H1 and KHSRP. ERK 1/2 and GAPDH show relatively equivalent loading of the lysates. (D, E) Representative immunoblots for kinetics of KHSRP elevation in sciatic nerve axoplasm over 0–28 days post-crush lesion are shown in (D). Quantification of KHSRP immunoreactivity across multiple animals is shown in (E) as mean fold-change relative to naïve \pm standard error of the mean (SEM); $N = 3$ mice for each time point; *** $P \leq 0.001$ for versus 0 day, † $P \leq 0.05$, ††† $P \leq 0.005$, †††† $P \leq 0.001$ and ††††† $P \leq 0.0005$ versus 3 days, &&& $P \leq 0.001$ and &&&& $P \leq 0.0005$ versus 7 days, and \$\$ $P \leq 0.01$ and \$\$\$\$ $P \leq 0.0005$ versus 14 days by one-way ANOVA with Tukey's post-hoc analysis). (F) Representative confocal images for KHSRP protein in naïve and 7 days post-crush sciatic nerve. Upper panels of each pair show XY images of merged neurofilament (NF) + KHSRP and DAPI + KHSRP; lower panels show KHSRP signals overlapping with NF or DAPI in individual Z planes projected as an XYZ image [scale bar = 5 μm].

higher at 21–28 days post injury than earlier time points (Figure 1D, E). With the mid-thigh sciatic nerve crush used, rats here begin to regain lower hind limb function over 21–28 days post injury indicative of some target reinnervation. Thus, KHSRP levels appeared to be increased in axons after injury and continued to elevate across the duration of axon regeneration at least until target reinnervation begins.

While the axoplasm preparation used here is highly enriched for axonal proteins, the preparation does contain some glial constituents (46). Since KHSRP is ubiquitously expressed (47,48), we used confocal microscopy to determine if the elevations in KHSRP levels seen by PRM and immunoblotting derived from axonal KHSRP. Consistent

with the ubiquitous KHSRP expression, KHSRP protein immunofluorescence was seen in both axons and the adjacent non-neuronal cells; however, extraction of the KHSRP immunofluorescent signals overlapping with neurofilament signals across individual optical planes, showed markedly increased intra-axonal KHSRP signals at 7 days post injury compared to uninjured nerve (Figure 1F). Interestingly, DAPI co-staining showed that the signals in non-neuronal cells within the nerve were predominantly nuclear, likely representing Schwann cell nuclei in the nerve (Figure 1F). Thus, axotomy increases axonal KHSRP levels in both axons and glia in the PNS, but glial KHSRP is predominantly intra-nuclear compared to the cytoplasmic signals of the axons.

Loss of KHSRP enhances axon growth in the peripheral nervous system through an axon-intrinsic mechanism

Since axonal mRNA translation supports both axon development and regeneration (4) and KHSRP has multiple functions (22,24,45,48,49), we asked if axonal KHSRP might play a role in axon growth in adult neurons. To address this, we took advantage of a constitutive KHSRP knockout mouse line, where both *KHSRP* alleles are deleted (47). Cultures of dissociated lumbar (L) 4–6 DRG neurons from *KHSRP* knockout (*Khsrp*^{-/-}) mice showed significantly increased axon lengths compared to those from wild type (*Khsrp*^{+/+}) mice (Figure 2A, B), but there were no differences in axon branching between the genotypes (Supplemental Figure S2A, B). This raised the possibility that KHSRP attenuates axon growth in adult DRG neurons as we had previously reported for embryonic CNS neurons (21).

Cultures of dissociated L4–6 DRGs from mice that have been primed or conditioned by an *in vivo* sciatic nerve crush injury exhibit increased growth that is transcription-independent, but translation-dependent (25,50). The *in vivo* conditioning injury activates transcription of growth-associated genes, whose mRNA products are then translationally regulated after the second injury that is brought by the DRG dissociation at the time of culture (25). With KHSRP's roles in post-transcriptional regulation (51), we tested whether the *in vitro* axon growth from *in vivo* 'injury-conditioned' L4–6 DRGs is further increased in the absence of KHSRP. In contrast to the naïve DRG cultures, there was no significant difference in axon lengths comparing the injury-conditioned DRGs from *Khsrp*^{-/-} versus *Khsrp*^{+/+} mice (Figure 2A, B). To be certain that KHSRP was indeed absent from these neurons, we assessed KHSRP levels in soma and axon preparations from L4–6 DRG neurons cultured on a porous membrane for separation of axons (52). KHSRP immunoreactive bands were clearly present in the *Khsrp*^{+/+} soma and axon isolates, but were not detected in the *Khsrp*^{-/-} samples even with extended exposures (Figure 2C). These data suggest that removing KHSRP can increase axon growth, but *in vivo* injury conditioning appears to overcome KHSRP's growth attenuating effects when those neurons are dissociated and cultured.

The comparable axon growth from injury-conditioned DRG neurons from the *Khsrp*^{-/-} and *Khsrp*^{+/+} mice could reflect a ceiling effect for KHSRP deletion in the DRG neurons, with the growth promoting effects of KHSRP loss mitigated by the *in vivo* conditioning lesion. Our initial data from Figure 1 pointed to an increase of KHSRP within the axons *in vivo* after peripheral nerve injury, and those axons are notably stripped away from the soma by the dissociation for culturing the neurons. Thus, an alternative hypothesis is that axon-intrinsic functions of KHSRP underlie its growth-attenuating functions. That is, when the axons are sheared from the soma during *in vitro* culture preparation the intra-axonal increase in KHSRP seen in Figure 1 could have been lost or minimized. To test this alternate hypothesis, we compared *in vivo* axon regeneration in the *Khsrp*^{-/-} and *Khsrp*^{+/+} mice. Although, there was no significant difference in axonal profiles extending beyond the crush site comparing *Khsrp*^{-/-} and *Khsrp*^{+/+} mice at 7 days after crush

injury, the *Khsrp*^{-/-} mice showed higher regeneration indices than the *Khsrp*^{+/+} mice at 10 days post-injury and the *Khsrp*^{-/-} mice showed significantly higher regeneration indices 14 days post-injury (Figure 3A). This suggests that the growth-attenuating effects seen from the injury-induced increase in axonal KHSRP after axotomy can accumulate over time after injury, which is consistent with the increasing axoplasm levels of this protein shown in Figure 1.

Considering this delayed effect of axonal KHSRP on nerve regeneration, we asked if the growth attenuation from KHSRP might be seen sooner if axotomy occurred in the setting of already increased axonal KHSRP. To address this possibility, we used an *in vivo* injury-conditioning paradigm where an initial sciatic nerve crush injury is performed and then 7 days later the nerve is crushed a second time proximal to the initial injury site (53). For comparison, the contralateral sciatic nerve underwent single crush at the same time and same approximate level as the second lesion. Three days later, nerve regeneration contralateral to the conditioning lesion (i.e. single crush injured nerves) showed no significant difference between *Khsrp*^{-/-} and *Khsrp*^{+/+} mice (Figure 3B, C). In sharp contrast, the injury-conditioned *Khsrp*^{-/-} nerves (i.e. double crush injured nerves) showed a dramatic increase in axon regeneration compared to the injury-conditioned *Khsrp*^{+/+} mice (Figure 3B, C; Supplemental Figure S3A). To determine if this increased axon growth affected target reinnervation, we analyzed neuromuscular junctions (NMJ) in the gastrocnemius muscle 14 days after crush injury (Figure 3D). NMJ occupancy, based on percentage of presynaptic compared postsynaptic area, showed no differences between *Khsrp*^{+/+} and *Khsrp*^{-/-} mice at 14 days in the single crush injured nerves (Figure 3E). However, the double crush injured nerves showed significantly greater NMJ occupancy in *Khsrp*^{-/-} versus *Khsrp*^{+/+} at 14 days after the second crush lesion (Figure 3D, E).

The difference between *in vivo* nerve regeneration and *in vitro* axon growth after injury-conditioning is intriguing since the injury-conditioning is thought to drive a change in gene expression programs, with injury-conditioned neurons already having a growth-associated gene expression program active at the time of second injury (50,54). Since axons are stripped from soma in the cultures used in Figure 2, axon-intrinsic roles of KHSRP for dampening the full effects of these changes in the injury-conditioned neurons could explain the difference between *in vivo* and *in vitro* axon growth. To test this possibility, we developed an *in vitro* axotomy model where DRGs were initially allowed to extend axons and then the axon shafts were severed. By severing the axon shafts rather than stripping axons from the soma, this allowed us to assess regeneration initiating from the axon shaft rather than initiation of new axon growth from the soma. For this, DRG neurons from *Khsrp*^{-/-} and *Khsrp*^{+/+} mice were cultured on porous membrane filters for 36 h, and then the lower surface of the membrane was scraped to remove axons by severing the axon shafts as they exited the pores of the membrane (Supplemental Figure S3B). After an additional 72 h in culture, neurofilament immunostained axons were traced along the membrane's lower surface and total axon lengths were quantified (Supplemental Figure S3C). Consistent with increased regener-

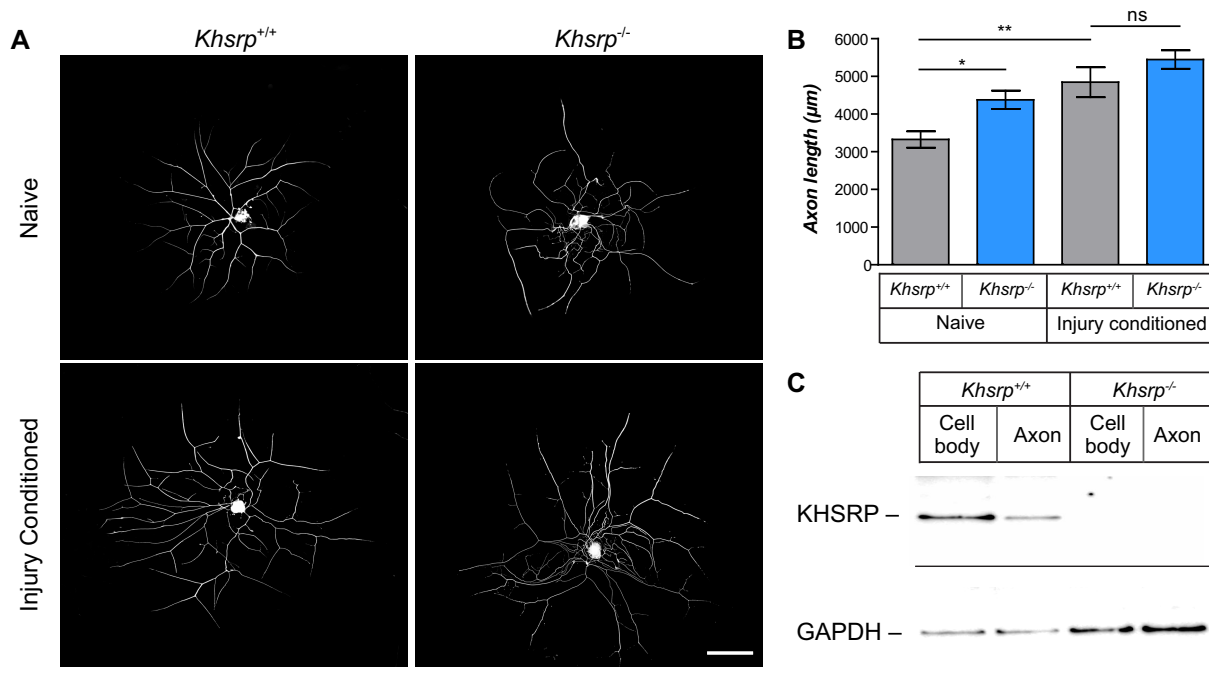


Figure 2. KHSRP depletion enhances axonal growth from naïve but not injury-conditioned DRG neurons. (A) Representative images of 24 h DRG neuron cultures from naïve and 7 days injury-conditioned *Khsrp*^{+/+} and *Khsrp*^{-/-} mice immunostained for NF are shown [scale bar = 100 μm]. (B) Quantification of total axon length per neuron from cultures as in (A) shows significantly greater axon growth in *Khsrp*^{-/-} compared to *Khsrp*^{+/+} DRGs. However, there is no significant difference in axon length between the genotypes when cultures were prepared 7 days after nerve crush injury (i.e. ‘injury-conditioned’ neurons). Data are expressed as mean ± SEM; refer to Supplemental Figure S2 for axon branching analyses ($N \geq 75$ neurons analyzed per condition in three independent culture preparations; * $P \leq 0.05$ and ** $P \leq 0.01$ by one-way ANOVA with Tukey’s post-hoc analysis). (C) Representative immunoblot shows signal for KHSRP in cell body and axonal preparations from DRG cultures from indicated mice. There is no KHSRP band in the *Khsrp*^{-/-} cell body or axonal preparations, despite GAPDH showing more protein in those samples than in the *Khsrp*^{+/+} samples.

ation from injury-conditioned nerves *in vivo*, the *Khsrp*^{-/-} DRGs showed more axon regeneration than the *Khsrp*^{+/+} DRGs after their axon shafts were severed *in vitro* (Supplemental Figure S3D). Taken together, these data suggest that the increase in axonal KHSRP slows extension of the regenerating axon.

The axotomy-induced increase in axonal KHSRP occurs via axon-intrinsic mRNA translation

We next asked what mechanisms might underlie the increase in axonal KHSRP upon injury. mRNA translation in axons provides a means to rapidly change protein content in the axons (4). For example, injury-induced translation of axonal *Calr* mRNA was recently shown to support early regrowth of severed axons (55) and translation of axonal *Kpnbl* mRNA generates a retrograde signal that triggers transcriptional changes in the neuronal soma (54). Thus, we wondered whether the change in axonal KHSRP levels might be driven by localized translation of its mRNA. We had previously detected *Khsrp* mRNA in RNA-seq analyses of sciatic nerve axoplasm (42); however, these axoplasm preparations obviously can contain non-neuronal contents (46), particularly for mRNAs isolated from injured nerve as Lee *et al.* used (42). To overcome this limitation, we used smFISH/IF to ask whether sensory axons contain *Khsrp* mRNA. Dissociated DRG cultures from *Khsrp*^{+/+} mice showed prominent *Khsrp* mRNA signals in soma and

axons, with granular appearing axonal *Khsrp* mRNA signals (Figure 4A–C). There was no *Khsrp* mRNA signal in the axons of the *Khsrp*^{-/-} DRG cultures (Figure 4C). smFISH signals for the soma of the DRGs from *Khsrp*^{-/-} mice did show a faint but consistent signal with the *Khsrp* mRNA probes that was significantly greater than with scrambled probes (Figure 4A, B). The *Khsrp*^{-/-} mice were generated by deleting exons 1–13 (of 18 exons) of the murine *KHSRP* gene (24), and the *Stellaris* smFISH *Khsrp* mRNA probes used here hybridize to sequences across the full exons comprising the mature *Khsrp* mRNA. Considering the absence of KHSRP protein in these cultures by immunoblotting (Figure 2C), we suspect that the faint *Khsrp* mRNA signal in soma of the *Khsrp*^{-/-} DRGs reflects RNA transcription of exons 14–18 remaining in the *Khsrp*^{-/-} mice or portions thereof, as confirmed by RNA-sequencing of *Khsrp*^{-/-} adult mouse brain and embryonic cortical neuron cultures (data not shown). The absence of *Khsrp* mRNA signals in the axons of *Khsrp*^{-/-} DRGs here and lack of any detectable KHSRP by immunoblotting in the *Khsrp*^{-/-} DRG lysates shown in Figure 2C suggest that any transcript from remaining exons 14–18 of *KHSRP* in the *Khsrp*^{-/-} mice does not localize into axons or generate detectable protein.

smFISH/IF performed on sciatic nerve sections also showed prominent *Khsrp* mRNA signal in the axons of the *Khsrp*^{+/+} mice that was completely absent in *Khsrp*^{-/-} mice (Figure 4D, E). Axonal *Khsrp* mRNA signals showed no significant differences in *Khsrp*^{+/+} mice comparing naïve

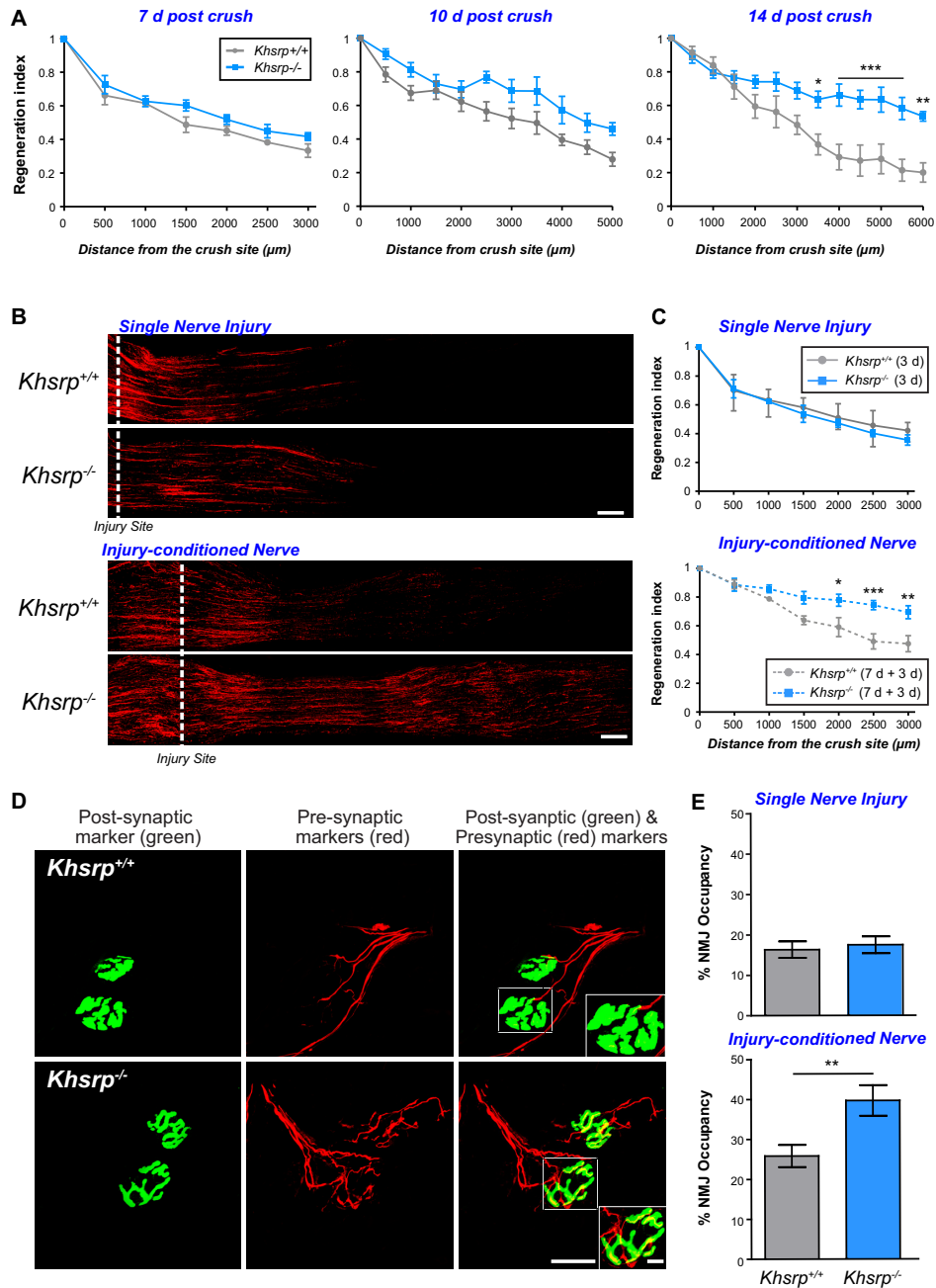


Figure 3. KHSRP deletion increases *in vivo* axon regeneration after a conditioning sciatic nerve injury. (A) Regeneration indices calculated as fraction of SCG10 axonal profiles at injury site (0 µm) are shown as mean ± SEM as indicated. There was no significant differences between *Khsrp*^{-/-} and *Khsrp*^{+/+} mice at 7 and 10 days post-injury, but significant differences are seen between the genotypes at 14 days post-injury ($n = 6$ mice per genotype and timepoint, * $P \leq 0.05$, ** $P \leq 0.01$ and *** $P \leq 0.001$ by two-way ANOVA with Bonferroni post-hoc analysis). (B) Representative SCG10 immunostained images of sciatic nerves after single (3 days; upper panels) or injury-conditioned (7 + 3 days; lower panels) sciatic nerve crush injuries for *Khsrp*^{+/+} and *Khsrp*^{-/-} are shown. Proximal is on the left and distal on the right; the dashed line indicates the injury site, with the second injury for the double crush injured animals placed at ~0.5 cm proximal to the initial injury site [scale bar = 500 µm]. (C) Regeneration indices calculated as fraction of SCG10 axonal profiles relative to the injury site (0 µm) are shown as mean ± SEM as indicated. There was no significant difference in the regeneration after the single injury, but the injury-conditioned *Khsrp*^{-/-} mice show significantly higher regeneration indices. Refer to Supplemental Figure S3B for regeneration index comparisons of naïve versus injury-conditioned nerves within genotypes ($N = 5$ mice per genotype and condition; * $P \leq 0.05$, ** $P \leq 0.01$ and *** $P \leq 0.001$ by two-way ANOVA with Bonferroni post-hoc analysis). (D) Confocal XYZ images of gastrocnemius muscles of the injury-conditioned *Khsrp*^{+/+} and *Khsrp*^{-/-} mice at 14 days after second nerve crush are shown. NMJs are detected by post-synaptic (α-bungarotoxin; green) and pre-synaptic markers (cocktail of anti-NF, -synapsin I and -synaptophysin; red) signals showing higher matching of pre- and post-synaptic markers in *Khsrp*^{-/-} than *Khsrp*^{+/+} mice. Inset panels on lower right of both rows show higher magnification of the NMJs outlined by dashed boxes [scale bars = 20 µm for main panels and 5 µm for insets]. (E) Quantification of NMJ occupancy (% presynaptic area/postsynaptic area) shows significantly greater occupancy in the injury-conditioned *Khsrp*^{-/-} than in *Khsrp*^{+/+} mice but no difference between genotypes was seen with the single crush lesion (injury-conditioned = 7 + 14 days; single nerve crush = 14 days). Data are expressed as mean ± SEM ($N \geq 15$ NMJs quantified in three animals per condition per genotype; ** $P \leq 0.01$ by Student's *t* test).

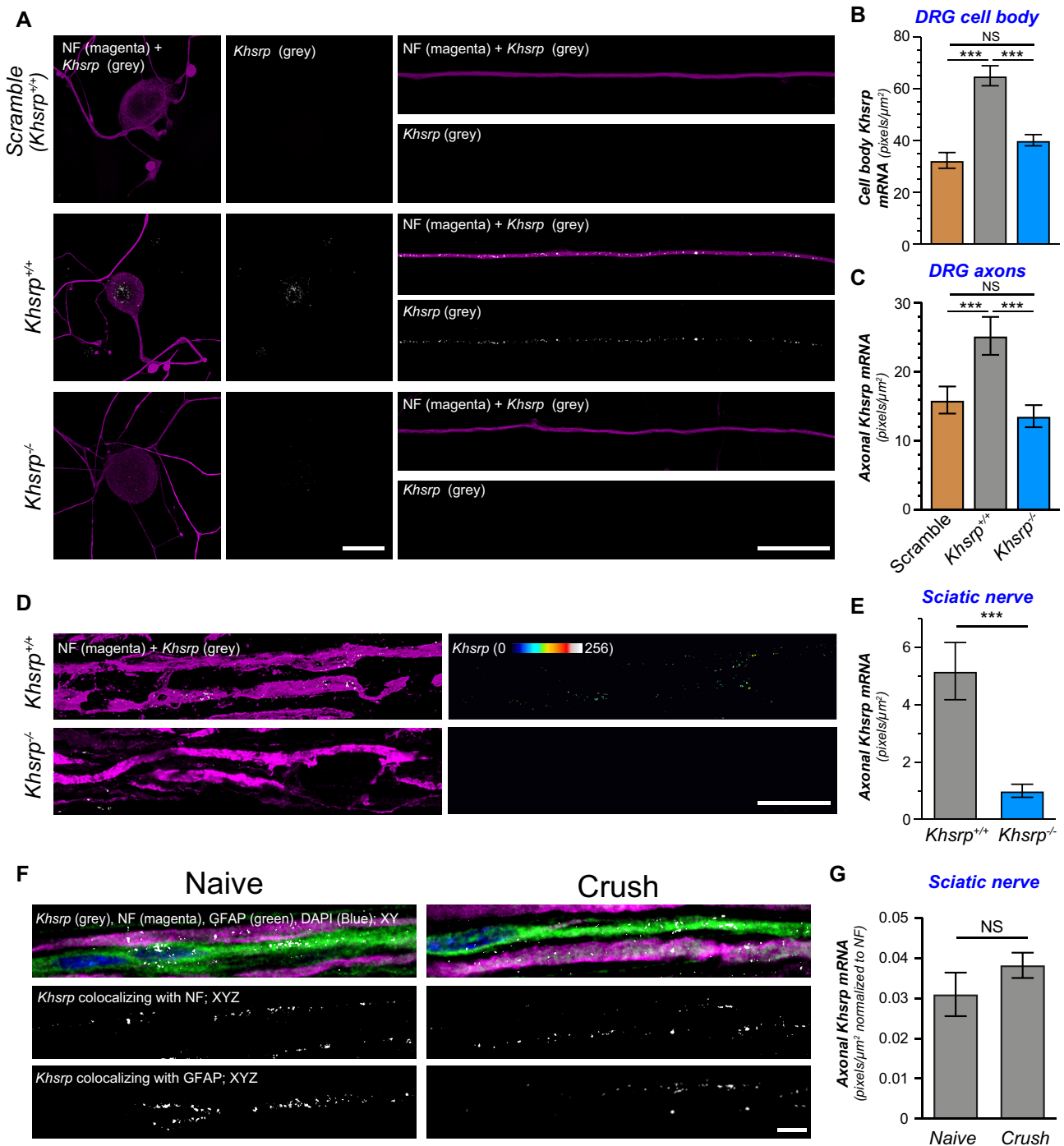


Figure 4. *Khsrp* mRNA is transported into PNS axons. (A) Representative confocal images for smFISH/IF for *Khsrp* mRNA (grey) and NF (magenta) in dissociated DRG cultures are shown as indicated. There is a clear signal for *Khsrp* mRNA in the cell body (left and middle columns) and distal axons (right column) of DRGs from *Khsrp*^{+/+} mice. Axons of *Khsrp*^{-/-} DRGs show *Khsrp* mRNA signals comparable to the scrambled probe; however, there was faint *Khsrp* signal in the soma of the *Khsrp*^{-/-} cultures that was consistently above the scrambled probe signal [scale bar = 10 μm]. (B, C) Quantification smFISH signals for *Khsrp* mRNA in soma (B) and axons (C) is shown as mean \pm SEM for scramble (*Khsrp*^{+/+}) and *Khsrp* mRNA (*Khsrp*^{+/+} and *Khsrp*^{-/-}) probes; in each case, scramble probe was hybridized to *Khsrp*^{+/+} DRG cultures as in panel A ($N \geq 16$ neurons over three separate cultures; * $P \leq 0.05$, ** $P \leq 0.01$ and *** $P \leq 0.001$ by one way ANOVA with Tukey's post-hoc analysis). (D) Representative confocal images for smFISH/IF for *Khsrp* mRNA (grey) and NF (magenta) in uninjured sciatic nerve are shown as indicated. Left column shows XYZ projections from eight optical planes taken at 0.2 μm Z step intervals; right column shows 'axon only' *Khsrp* mRNA signals generated by extracting FISH signals overlapping with NF in individual Z sections and projecting those as an 'Axonal *Khsrp* mRNA' XYZ image [scale bar = 5 μm]. (E) Quantification of axonal *Khsrp* mRNA signals from (D) are shown as mean \pm SEM ($N = 6$ animals per genotype; ** $P \leq 0.01$ by Student's *t*-test). (F) Representative matched exposure smFISH/IF images for *Khsrp* mRNA, NF, GFAP and DAPI in naive versus 7 days post-crush injured sciatic nerves as indicated. Upper row shows merged signals as single XY planes; lower two rows show XYZ projections for *Khsrp* mRNA colocalizing with NF (middle row) and GFAP (lower row). Representative matched exposure images for scramble probe are shown in Supplemental Figure S3E [scale bar = 5 μm]. (G) Quantification of axonal smFISH signals for *Khsrp* mRNA in naive and 7 days regenerating sciatic nerve axons ($N = 6$ animals; no significant differences detected by Student's *t*-test).

and 7 days injured sciatic nerve (Figure 4F, G; Supplemental Figure S3E), suggesting that the elevation of KHSRP protein seen in Figure 1 is not from increased transport or survival of *Khsrp* mRNA following sciatic nerve crush injury. Notably, *Khsrp* mRNA was also detected in Schwann cells (Figure 4F), which is consistent with the strong nuclear KHSRP signals seen in the Schwann cells in Figure 1F.

The RNA analyses above suggest that KHSRP can be synthesized locally in axons. To determine if the PNS nerve injury-induced increase in KHSRP is intrinsic to the nerve or results from KHSRP transported from proximal nerve and soma, we ligated the sciatic nerve to restrict anterograde transport and performed a crush injury distal to the ligation (Figure 5A). Adult rats were used for these analyses since the larger sciatic nerve could be ligated more consistently than in the mouse. Immunofluorescence for amyloid precursor protein (APP) and signal transducer and activator of transcription 3 α (Stat3 α) confirmed that the ligation attenuated both anterograde and retrograde transport, as APP accumulated proximal to and Stat3 α accumulated distal to the ligation site (Supplemental Figure S4). As expected, KHSRP signals were detected both in axons and adjacent Schwann cells (Figure 5B, C). Colocalization with DAPI indicated that the non-axonal KHSRP is predominantly nuclear as seen in Figure 1F (data not shown). Axonal KHSRP Immunofluorescence showed increased signals at both the proximal and distal ligation sites as well as the crush site relative to axons in naïve nerve; however, the increased axonal KHSRP signals at the crush site were significantly greater than the proximal and distal ligation sites and continued to elevate over time after injury (Figure 5D). These results are consistent with local synthesis of KHSRP in the injured nerve, as opposed to anterograde transport of KHSRP from the soma or more proximal axons where the signal would have been greater at the proximal ligation site than crush and distal ligation sites. Of note, ligation causes a degree of nerve injury, and the increase in KHSRP levels both proximal to and distal to the ligation are consistent with such an injury response.

To more directly test if KHSRP is translated in sciatic nerve axons, we exploited an *ex vivo* nerve injury model where we used puromycin incorporation to detect newly synthesized peptides in axoplasm extruded from the nerve (56). For this, we excised segments of rat sciatic nerve and placed these into culture medium with cyclosporin A included to delay Wallerian degeneration (57) and OPP for puromycinylation of nascently synthesized polypeptides (58,59). There was a significant increase in puromycinylation of KHSRP in the axoplasm isolates at 4 h after *ex vivo* nerve crush that was attenuated by the protein synthesis inhibitor anisomycin (Figure 5E, F). Notably, an increase in overall KHSRP levels was also seen in the crushed nerve axoplasm and this was similarly attenuated by pretreatment with anisomycin (Figure 5E), confirming that axonal injury increases KHSRP in sciatic nerve axoplasm through mRNA translation.

To further test for translation of *Khsrp* mRNA in axons, we fused 5' and 3' untranslated regions (UTR) of rodent *Khsrp* mRNA to the coding sequence of a diffusion-limited GFP reporter cDNA (GFP^{MYR}5'/3'khsrp; Figure 5G) as a surrogate for axonal localization and translation

of *Khsrp* mRNA. Co-translational myristoylation of this GFP reporter allows for membrane attachment to limit the protein's diffusion (31,40), and the UTRs of axonal mRNAs typically contain motifs for axonal mRNA targeting and translational regulation (4). By fluorescence recovery after photobleaching (FRAP), DRG neurons expressing GFP^{MYR}5'/3'khsrp showed rapid fluorescent recovery that was significantly attenuated by pretreatment with anisomycin or a second protein synthesis inhibitor cycloheximide (Figure 5G; Supplemental Figure S5A). Importantly, the bleached regions of interests (ROI) were separated from the soma by >250 μ m, indicating that this fluorescence recovery occurs faster than can be accounted for by anterograde transport of reporter synthesized in the soma.

The above data support that the axotomy-induced increase in axonal KHSRP is derived from localized translation of *Khsrp* mRNA in axons, but these did not address the mechanism underlying this translational increase. Axon injury is known to increase axoplasmic Ca²⁺ and activate a localized intrinsic stress or unfolded protein response (60). Although increased cytoplasmic Ca²⁺ is well known to block generalized protein synthesis through an inhibitory phosphorylation of the translation factor eIF2 α (Figure 5H), some injury-response mRNAs show a paradoxical increase in their translation with Ca²⁺ despite eIF2 α phosphorylation (61,62). We used thapsigargin to simulate the injury-induced increase in axoplasmic Ca²⁺ (59). Recovery of axonal GFP^{MYR}5'/3'khsrp fluorescence was significantly increased by thapsigargin treatment; conversely, chelating Ca²⁺ with BAPTA-AM attenuated recovery of the axonal GFP^{MYR}5'/3'khsrp fluorescence (Figure 5I). Inhibition of protein kinase R-like ER kinase (PERK) prevented the increase in axonal GFP^{MYR}5'/3'khsrp fluorescence recovery seen with thapsigargin treatment (Figure 5H, I). Conversely, pretreatment with Sephin1 that prevents dephosphorylation of eIF2 α partially reversed the BAPTA-AM dependent decrease in axonal GFP^{MYR}5'/3'khsrp fluorescence recovery (Figure 5H-I). Immunoblotting confirmed that the increase in eIF2 α ^{PS51} in response to thapsigargin was prevented by the GSK260614 PERK inhibitor and decrease in basal eIF2 α ^{PS51} seen with BAPTA-AM was reversed by treatment with Sephin1 to inhibit eIF2 α phosphatase (Supplemental Figure S5B). Thus the axotomy-induced increase in axonal KHSRP is derived through activation of *Khsrp* mRNA via Ca²⁺ \rightarrow PERK \rightarrow eIF2 α ^{PS51}.

KHSRP's ARE-binding KH4 domain attenuates axon growth

KHSRP was previously shown to bind to the ARE in *Gap43* mRNA's 3'UTR and promote decay of the transcript (21). The *GAP43* gene is transcriptionally activated following PNS nerve injury (63), and axonal *Gap43* mRNA levels increase in regenerating PNS axons (20). Given that *Gap43* mRNA and protein expression are typically associated with axon growth (64), we tested the effect of *Khsrp* knockout on the sciatic nerve levels of this KHSRP target mRNA. RTddPCR from sciatic nerve samples of *Khsrp*^{+/+} mice showed an increase in *Gap43* mRNA at 7 days after nerve crush injury and this was significantly greater in the *Khsrp*^{-/-} versus *Khsrp*^{+/+} mice (Figure 6A). Given that

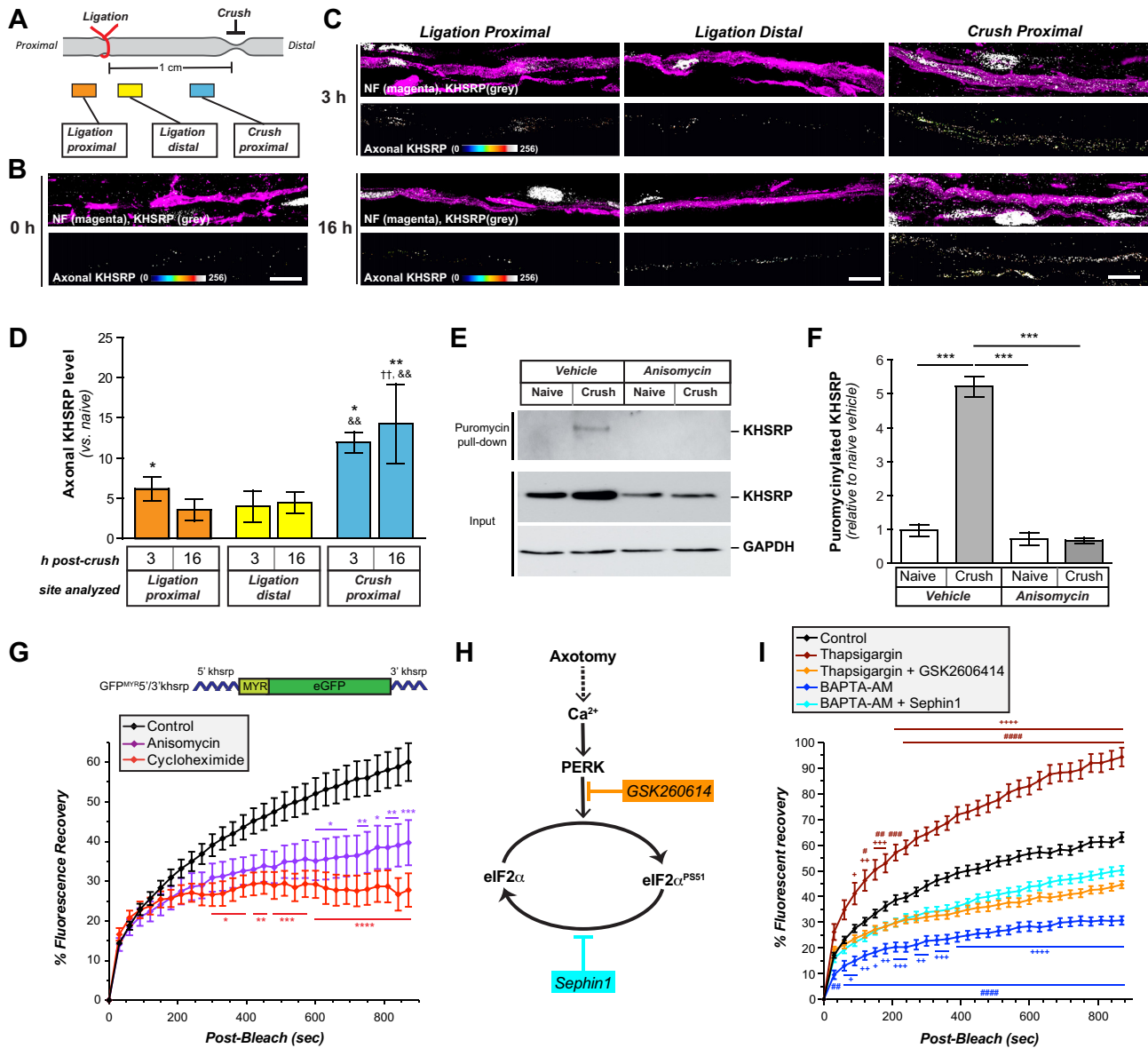


Figure 5. Khsrp mRNA is rapidly translated in axons after injury. (A) Schematic of nerve ligation model used for panels B and C. Proximal nerve is on the left and distal on the right as indicated. The nerve was ligated and then immediately crushed ~1 cm distal to the ligation. (B, C) Confocal images for KHSRP protein in naïve (B) and post-crush injury (3 and 16 h; C). Upper rows of image pairs show XYZ projections of merged signals for KHSRP (grey) and NF (magenta), while lower rows show ‘axonal KHSRP’ signals as from individual Z planes that were projected as an XYZ image. As in Figure 1E, the strong signals that are outside of the axons are in Schwann cell nuclei based on DAPI co-labeling (data not shown). Representative images for ligation efficiency and KHSRP signals proximal and distal to ligation are shown in Supplemental Figure S4 [scale bar = 5 μm]. (D) Quantification of the axonal KHSRP immunoreactivity from ligation proximal and distal and crush sites are shown as mean ± SEM (N = 3 animals per time point; * P ≤ 0.05 and ** P ≤ 0.01 for indicated time points versus naïve nerve, †† P ≤ 0.01 for indicated time points versus ligation proximal, and && P ≤ 0.01 for indicated time points versus ligation distal by Student’s *t*-test; ligation proximal versus distal have no significance). (E) Representative immunoblots for *ex vivo* puromycinylated naïve versus crushed sciatic nerve segments are shown as indicated. Protein synthesis inhibition with anisomycin completely blocks the puromycinylation of KHSRP in axoplasm samples extruded from the nerve segments, and GAPDH shows relatively equivalent protein loading across the lanes. Note that total KHSRP levels also increases with crush injury and this was attenuated by anisomycin. (F) Quantification of puromycinylated KHSRP signals from (D) is shown as mean ± SEM. Crush injury significantly increases axonal KHSRP synthesis and this blocked by anisomycin (N = 3; *** P ≤ 0.001 by one way ANOVA with Tukey’s post-hoc analysis). (G) FRAP analyses for distal axons of neurons transfected with GFP^{MYR}5'/3'khsrp (schematic above graph) is shown as normalized average % recovery ± SEM. Pre-treatment with anisomycin or cycloheximide significantly attenuates the GFP recovery, indicating protein synthesis dependent recovery for GFP^{MYR} fluorescence in the axons (N ≥ 10 neurons over three culture preparations; * P ≤ 0.05, ** P ≤ 0.01 and *** P ≤ 0.005 by two-way ANOVA with Bonferroni post-hoc analyses for indicated time points versus control). Representative images sequences for FRAP are shown in Supplemental Figure S5. (H) Schematic for proposed signaling pathway for axotomy induced increase in axoplasmic Ca²⁺ leading to eIF2α phosphorylation is shown. GSK260614 was used as a specific PERK inhibitor and Sephin1 as an inhibitor of eIF2α dephosphorylation. (I) FRAP analyses for distal axons of neurons transfected with GFP^{MYR}5'/3'khsrp reporter is as normalized average % recovery ± SEM. Treatment with thapsigargin increased and BAPTA-AM decreased recovery over control conditions. The thapsigargin-induced increase was blocked by GSK260614, while the BAPTA-AM-induced decrease was partially blocked by Sephin1 (N ≥ 17 neurons over three culture preparations; # P ≤ 0.05, ## P ≤ 0.01, ### P ≤ 0.005 and #### P ≤ 0.001 for thapsigargin or BAPTA-AM versus corresponding control time points and +P ≤ 0.05, ++P ≤ 0.01, +++P ≤ 0.005 and ++++P ≤ 0.001 for thapsigargin or BAPTA-AM versus corresponding thapsigargin + GSK260614 or BAPTA-AM + Sephin1 time points by two-way ANOVA with Bonferroni post-hoc analyses for indicated time points versus control; for data points appearing to error bars, the SEM is too small to show).

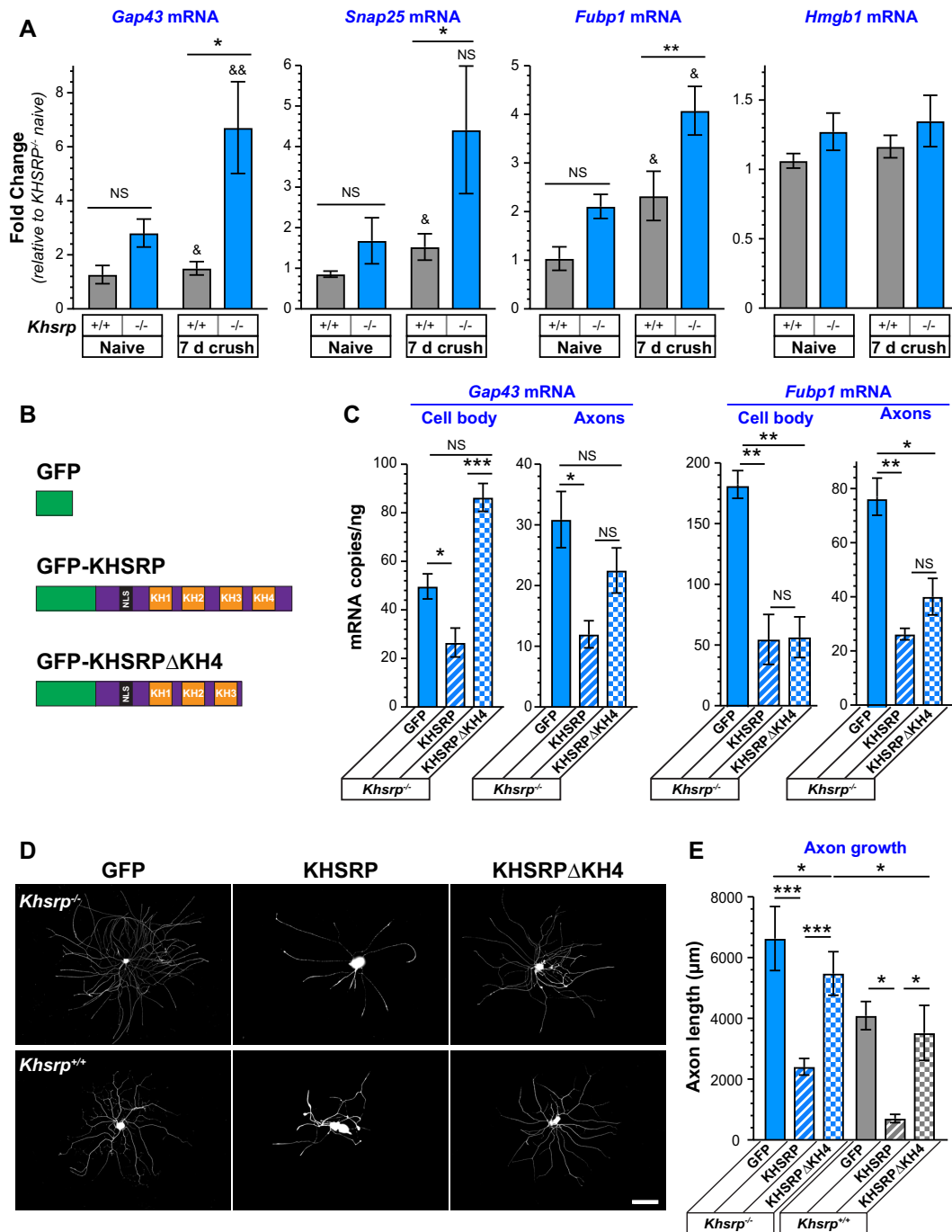


Figure 6. Increased levels of KHSRP target mRNAs in *Khsrp*^{-/-} neurons. (A) Sciatic nerve levels of *Gap43*, *Snap25* and *Fubp1* mRNAs are increased 7 days after sciatic nerve crush in *Khsrp*^{-/-} compared to *Khsrp*^{+/+} mice. In contrast, sciatic nerve *Hmgb1* mRNA levels show no change comparing the *Khsrp*^{-/-} versus *Khsrp*^{+/+} mice. Values shown as mean \pm SEM ($N = 5$ mice per genotype; * $P \leq 0.05$, ** $P \leq 0.01$ and NS for indicated pairs across genotype within condition and & $P \leq 0.05$ and && $P \leq 0.01$ for crush versus naive within genotype by Student's *t*-test). (B) Schematic of expression constructs used for rescue experiments shown in panels C–F. (C) Analyses of soma and axon *Gap43* and *Fubp1* mRNA levels in *Khsrp*^{-/-} DRG neurons transfected with GFP, GFP-KHSRP and GFP-KHSRP Δ KH4 is shown as mean mRNA copies/ng of total RNA \pm SEM after normalization to mitochondrial 12S rRNA. Supplemental Figure S6A shows expression levels for GFP, GFP-KHSRP and GFP-KHSRP Δ KH4 and Supplemental Figure S6B shows RNA levels for transfected *Khsrp*^{+/+} DRG cultures ($N = 3$ per condition; * $P \leq 0.05$, ** $P \leq 0.01$, *** $P \leq 0.001$ and NS = not significant for indicated pairs by Student's *t* test). (D, E) Representative images of DRG neurons transfected with GFP, GFP-KHSRP or GFP-KHSRP Δ KH4 are shown for *Khsrp*^{+/+} and *Khsrp*^{-/-} (schematics for constructs above image columns) are shown in (D) as indicated. Quantification of total axon length/neuron for transfected DRG neurons is shown in (E) as mean \pm SEM. Expression of GFP-KHSRP decreases axon outgrowth in both *Khsrp*^{+/+} and *Khsrp*^{-/-} DRGs, but GFP-KHSRP Δ KH4 had no significant effect on *Khsrp*^{+/+} and only a modest decrease in axon length in *Khsrp*^{-/-} neurons ($N \geq 30$ neurons over three different culture preparations; * $P \leq 0.05$, ** $P \leq 0.01$ and *** $P \leq 0.001$ by one-way ANOVA with Tukey's post-hoc analysis for indicated comparisons) [scale bar = 100 μ m].

transcription of the *GAP43* gene is increased during axon regeneration, there were only modest, non-significant increases in nerve *Gap43* mRNA of the uninjured *Khsrp*^{-/-} than *Khsrp*^{+/+} mice (Figure 6A). Since KHSRP can potentially bind to many different axonal mRNAs, we asked if axonal *Snap25* and *Fubp1* mRNAs might be affected by *Khsrp* knockout. Both mRNAs are elevated in cortex and hippocampus of *Khsrp*^{-/-} mice and the neurites of cortical neurons cultured from embryonic day 18 *Khsrp*^{-/-} versus *Khsrp*^{+/+} mice (65). The Hengst lab reported translation of *Snap25* mRNA in central nervous system (CNS) axons promotes synaptogenesis (66), but to our knowledge, potential roles for FUBP1 in neurons have not been tested. *Snap25* and *Fubp1* mRNAs were detected in sciatic nerve, and their levels were significantly increased in the injured sciatic nerves of *Khsrp*^{-/-} compared to *Khsrp*^{+/+} mice (Figure 6A). In contrast, *Hmgb1* mRNA (also called Amphoterin), an mRNA that we have previously shown localizes to axons and whose local translation also supports axon growth (43), did not show any significant difference between *Khsrp*^{-/-} and *Khsrp*^{+/+} nerves (Figure 6A). Notably, *Hmgb1* mRNA did not show binding to KHSRP in RIP-Seq analyses (65). Thus, increased levels of some mRNAs encoding proteins linked to axon growth accompany the increased axon regeneration seen in injury-conditioned *Khsrp*^{-/-} mice, but this does not extend to all mRNAs encoding growth-associated proteins indicating a level of specificity to KHSRP's role in modulating target neuronal mRNA levels.

Previous work has shown that the KH3 and KH4 domains of KHSRP are needed for ARE binding and to promote decay of ARE-containing mRNAs via the cytoplasmic exosome complex (22), and deletion of KH4 attenuated the effects of KHSRP on neurite growth in embryonic CNS neuron cultures (21). To determine if loss of KHSRP's function in promoting mRNA decay is responsible for the elevations in KHSRP target mRNAs in the *Khsrp*^{-/-} mice, we transfected *Khsrp*^{-/-} DRG cultures with full length GFP, GFP-KHSRP, GFP-KHSRPΔKH4 (Figure 6B) and evaluated levels of *Gap43* and *Fubp1* mRNAs in soma and axon preparations. *Gap43* and *Fubp1* mRNAs were chosen for these analyses since they showed relatively consistent changes in the sciatic nerve with KHSRP deletion. Immunoblotting confirmed the presence of bands at the predicted molecular weights for GFP, GFP-KHSRP and GFP-KHSRPΔKH4 proteins (data not shown) and comparable expression levels in the transfected DRG neurons based on GFP fluorescence (Supplemental Figure S6A). *Gap43* mRNA levels in the *Khsrp*^{-/-} DRG axons and soma significantly declined with GFP-KHSRP expression, but no significant change was seen in the axons with GFP-KHSRPΔKH4 expression while there was a significant increase in soma *Gap43* mRNA levels with GFP-KHSRPΔKH4 expression (Figure 6C). Similarly, both axonal and soma *Fubp1* mRNA levels were decreased in *Khsrp*^{-/-} DRG neurons expressing in GFP-KHSRP. In contrast to *Gap43* mRNA, GFP-KHSRPΔKH4 expression significantly decreased both axonal and soma *Fubp1* mRNA levels in the *Khsrp*^{-/-} DRG cultures (Figure 6C). *Gap43* and *Fubp1* mRNA levels in cell bodies and axons of transfected *Khsrp*^{+/+} DRGs followed similar trends with

Gap43 mRNA decreased by transfection GFP-KHSRP but not GFP-KHSRPΔKH4 and *Fubp1* mRNA decreased by both GFP-KHSRP and GFP-KHSRPΔKH4 (Supplemental Figure S6B). These data indicate that KHSRP promotes *Gap43* mRNA decay in both axons and soma, but other functions of KHSRP account for its depletion of *Fubp1* mRNA in the DRG neurons.

Since KHSRP's KH4 domain destabilized the growth associated *Gap43* mRNA, we asked if the KH4 domain is needed for KHSRP's axon growth attenuating effects. While GFP-KHSRP expression reversed the axon growth promoting effects of KHSRP deletion in the *Khsrp*^{-/-} DRGs, GFP-KHSRPΔKH4 expression only modestly decreased axon growth in the *Khsrp*^{-/-} DRGs (Figure 6D). Consistent with effects of these constructs on KHSRP target mRNA levels in the *Khsrp*^{+/+} DRGs, GFP-KHSRP expression decreased axon growth in the *Khsrp*^{+/+} DRGs but expression GFP-KHSRPΔKH4 had no effect on axon growth in these wild type DRG neurons (Figure 6E). Taken together these data indicate that the RNA degradation activity of KHSRP contributes to slowing of axon growth by KHSRP and decreased axonal levels of *Gap43* mRNA but not *Fubp1* mRNA.

Effects of KHSRP knockout on axon regeneration are neuron-intrinsic

The studies above suggest a neuron-intrinsic role for KHSRP in slowing PNS axon regeneration. However, we relied on a constitutive KHSRP knockout mouse (24), where *Khsrp* expression is lost in every tissue and cell type. Since KHSRP is ubiquitously expressed and previous studies have shown altered interleukin and cytokine expression in *Khsrp*^{-/-} mice in non-neuronal cells (24,47), we could not completely exclude that the accelerated regeneration in *Khsrp*^{-/-} mice derived through KHSRP's roles in non-neuronal cells. To test for neuronal specific functions *in vivo*, we generated a conditional *Khsrp* knockout mouse with LoxP sites surrounding exons 2–6 of the mouse *KHSRP* gene (*Khsrp*^{fl/fl}). DRGs cultured from the *Khsrp*^{fl/fl} mice were transduced with AAV2-CMV-Cre virus *in vitro* for KHSRP deletion. This resulted in approximately 50% reduction in *Khsrp* mRNA in the DRG cultures by RTddPCR (Figure 7A). In our hands, the AAV2 preparations show preferential transduction of the neurons in these murine DRG cultures that contain sensory neurons, Schwann cells, and some fibroblasts. Consistent with this, immunofluorescence of the DRG cultures showed near complete loss of KHSRP in the neurons but continued expression of KHSRP in the non-neuronal cells (Figure 7B, C). In keeping with findings in the *Khsrp*^{-/-} DRG cultures shown in Figure 2, the AAV2-CMV-Cre-GFP transduced *Khsrp*^{fl/fl} DRG cultures showed significantly increased axon lengths compared to AAV2-CMV-GFP transduced DRGs but no change in axon branching (Figure 7D, E). Thus, selectively depleting KHSRP from adult sensory neurons increases axon growth, indicating neuron-intrinsic effects for KHSRP in slowing axon growth rates *in vitro*.

We used an *in vivo* AAV2-CMV-Cre-GFP transduction to determine if deletion of neuronal *Khsrp* alleles from adult mice affects nerve regeneration. For this, the AAV2 prepara-

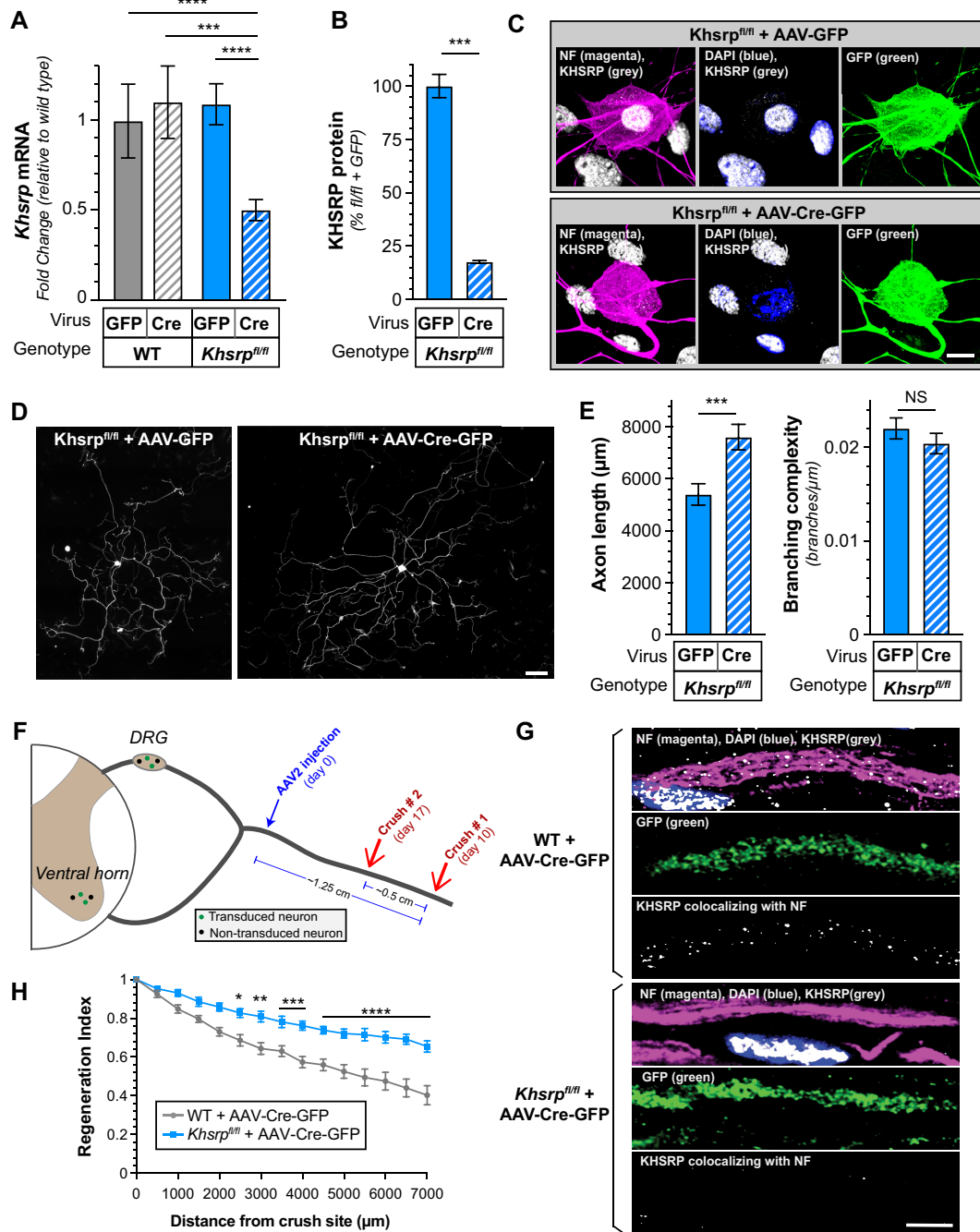


Figure 7. Axon growth promotion from loss of neuronal KHSRP. (A–C) DRGs cultured from *Khsrp^{f/f}* mice in (A) show reduced *Khsrp* mRNA by RTddPCR after transduction with AAV2-CMV-GFP-Cre compared to AAV2-CMV-GFP. AAV2-CMV-GFP-Cre transduction of wild type DRGs had no effect on *Khsrp* mRNA levels. By immunofluorescence where only neuronal KHSRP levels were assessed in (B), the AAV2-CMV-GFP-Cre transduced *Khsrp^{f/f}* DRGs showed greater than 80% reduction in KHSRP signals. Representative immunofluorescent images in (C) show relative absence of KHSRP signals in neuronal cell body and axons of DRGs, but prominent signals in adjacent Schwann cells of the AAV2-CMV-GFP-Cre transduced *Khsrp^{f/f}* dissociated DRG culture ($N = 6$ culture preparations for each condition for (A) and $N = 29$ neurons in three separate transfections for (B); $*** P \leq 0.005$ and $**** P \leq 0.001$ for indicated pairs by Student's t -test) [scale bar = 10 μm]. (D, E) Representative immunofluorescent images for NF in AAV2-CMV-GFP versus AAV2-CMV-GFP-Cre transduced *Khsrp^{f/f}* mice is shown in (D). Quantification of axon length and axon branching (E) as mean \pm SEM indicate show significantly increased axon growth with Cre-driven deletion of *KHSRP* in the *Khsrp^{f/f}* mouse DRGs ($N \geq 75$ neurons over three separate culture preparations/transductions for each condition; $*** P \leq 0.005$ and NS = not significant by Student's t -test) [scale bar = 100 μm]. (F) Schematic with time line for viral transduction of *Khsrp^{f/f}* mice followed by double sciatic nerve crush lesion as used in Figure 3. Note that the AAV2-CMV-GFP-Cre injection site is separated from both nerve crush sites by ~ 0.75 and ~ 1.25 cm. The animals were transduced on day 0, underwent distal nerve crush on day 10 (crush # 1), and underwent proximal nerve crush on day 17 (crush # 2). Regeneration was evaluated 3 days later. (G, H) Representative confocal images for KHSRP and NF with DAPI staining of sciatic nerve from AAV2-CMV-GFP-Cre transduced wild type and *Khsrp^{f/f}* mice after nerve crush injury # 2 are shown in (G) as indicated. (H) shows regeneration indices for AAV2-CMV-GFP-Cre transduced wild type and *Khsrp^{f/f}* mice 3 days after nerve crush injury # 2. Supplemental Figure S7 shows SCG10 and GFP immunofluorescence to compare regeneration between the AAV2-CMV-GFP-Cre transduced wild type and *Khsrp^{f/f}* mice ($N = 6$ mice per genotype; $* P \leq 0.05$, $** P \leq 0.01$, $*** P \leq 0.005$ and $**** P \leq 0.001$ by two-way ANOVA with Bonferroni post-hoc analysis) [scale bar = 10 μm].

tions were injected into the proximal sciatic nerve in a high salt solution that facilitates retrograde transport of the viral particles (26). We used the same conditioned-crush lesion as shown in Figure 3, with initial crush injury performed 10 days after AAV transduction and placing the first crush site ~1.25 cm from the transduction site plus the second crush site at ~0.75 cm distal from the transduction site (Figure 7F). Immunofluorescence showed prominent KHSRP signals in the NF⁺/GFP⁻ axons and no detectable KHSRP signals in the SCG10⁺/GFP⁺ axons; notably, Schwann cells in the nerve showed prominent KHSRP signals in their nuclei indicating the AAV2-CMV-Cre-GFP did not transduce these non-neuronal cells distant from the injection site (Figure 7G). Consistent with findings in the *Khsrp*^{-/-} mice shown in Figure 3, axons of the Cre-GFP expressing neurons of the *Khsrp*^{fl/fl} mice showed significantly higher regeneration indices compared to wild type mice transduced with the same AAV2-CMV-Cre-GFP preparations (Figure 7H; Supplemental Figure S7). Taken together, these data indicate that the local increase in axonal KHSRP attenuates PNS axon regeneration selectively through neuron-intrinsic mechanisms.

DISCUSSION

RBPs play critical roles in determining neuronal protein levels through post-transcriptional control mechanisms. Since one mRNA can generate multiple copies of a protein, RNA stability needs to be tightly regulated. HuD and KHSRP both bind to ARE-containing mRNAs, with HuD increasing mRNA stability and KHSRP promoting mRNA decay (51). Here, we show that PNS axons contain many different RBPs whose levels change after injury, including KHSRP. Axonal KHSRP protein levels rapidly increase in peripheral nerves following axotomy through intra-axonal translation of *Khsrp* mRNA. Since *Khsrp*^{-/-} mice show increased axonal levels of *Gap43*, *Snap25* and *Fubp1* mRNAs compared to *Khsrp*^{+/+} mice, the increase in KHSRP levels following injury decreases the levels of KHSRP target mRNAs in injured sciatic nerves. Interestingly, we find that the decline in *Gap43* mRNA levels with KHSRP expression requires an intact KH4 domain that is essential for the RNA degradation role of KHSRP (22), while *Fubp1* mRNA levels appear to be regulated by other functions of KHSRP. Deletion of the *Khsrp* gene increases axon growth. Axon regeneration was further accelerated *in vivo* in the absence of neuronal KHSRP comparing *Khsrp* deleted to wild type mice under injury-conditioned settings, where KHSRP is elevated in axons. Neuronal specific knockout confirmed that KHSRP's growth effects are neuron intrinsic. The growth attenuating functions of KHSRP also require an intact KH4 domain, pointing KHSRP's RNA decay promoting functions for regulating axon growth. Together, our data point to injury-induced increase in axonal KHSRP as an axon-intrinsic mechanism that serves to slow axon regeneration by affecting mRNA stability.

The targeted proteomics approach used here uncovered 84 RBPs in sciatic nerve axoplasm, with many beyond KHSRP showing increased or decreased levels after nerve injury and during regeneration. Since the axoplasm isolates used here rely on extrusion in detergent-free conditions (46),

we likely missed many RBPs that are associated with cytoskeleton or the axoplasmic membrane. HuD/ELAVL4 has previously been reported to fractionate with the cytoskeleton in rat hippocampus (67). So, this may explain the decrease in HuD/ELAVL4 levels detected in the sciatic nerve axoplasm after injury shown in Figure 1 compared to our previous immunofluorescent analyses showing that axonal HuD/ELAVL4 levels increase during regeneration (20). Despite limitations in methodology and amount of starting materials for the axoplasm, we have substantially increased the number of known axonal RBPs with the PRM approach used here. Other axoplasm RBPs identified here as increasing or decreasing with sciatic nerve injury and during regeneration could contribute to axon growth. For example, fragile X-related protein 1 (FXR1) showed a remarkable increase after sciatic nerve crush in Figure 1A-C, and this protein has been shown to associate with RNA granules containing fragile x mental retardation protein (FMRP) in CNS axons (68,69). Though not identified here, FMRP is well characterized as a translational modulator in dendrites and it regulates translation of microtubule-associated protein 1b (MAP1b) in axonal growth cones (70). Additionally, a number of hnRNPs linked to RNA splicing were previously shown to localize into axons and bind to axonal mRNA localization motifs, including hnRNP H1 and F that bind to *Hmgbl* mRNA localization motif and whose depletion from adult DRGs decreases axon outgrowth (42). Our PRM data also show a decline in Splicing Factor Proline and Glutamine Rich (SFPQ) that has been linked to survival of axons through its role in transport of *Bclw* mRNA into axons (71,72). It will be of high interest to determine how the different RBPs impact axon growth after injury as well as the mechanisms underlying their change in axonal levels. Notably analyses of axoplasm hnRNP H1 and F immunoprecipitates pointed to existence of mRNA regulons encoding growth-associated proteins binding to axonal hnRNP H1 and F (42). So, it is likely that the axonal RBPs reported here can post-transcriptionally regulate many axonal mRNAs. It is appealing to hypothesize that those increased RBPs in injured and regenerating axons support axon growth, so it is intriguing that KHSRP does the opposite. It has been reported that intra-axonal translation decreases as axons reach their target tissues and form synapses (73). Further, ribosome profiling for axonal mRNAs has shown changes in which mRNAs are translated in retinal ganglion cell axons as they reach the optic tectum and form synapses (1). Thus, one can speculate that the elevated KHSRP levels contribute to changes in axonal mRNA populations associated with different stages of regeneration, with the nascently synthesized axonal KHSRP locally interacting with target mRNAs and possibly displacing other RBPs from those transcripts.

Much interest has been devoted to understanding how mRNAs are transported into axons and dendrites. RBPs that drive subcellular mRNA localization sometimes also regulate translation of those mRNA cargos (4). To our knowledge, KHSRP is the first example for localized translational regulation of an RBP mRNA whose protein product goes on to modify axon growth potential. This emphasizes that localized translation can be used to modify the fate of other subcellular mRNAs. The initial increase

in axonal KHSRP protein through translational upregulation of its axonal mRNA shows similar kinetics to translational induction of injury-associated mRNAs following axotomy. mRNAs encoding Importin β 1, Vimentin, RanBP1, Stat3, mTor and Calreticulin (CALR) are translated in axons within the first six hours following PNS nerve crush injury (39,40,56,74,75). Some of those locally synthesized injury-associated proteins help to signal retrogradely to the soma for changing gene expression, including transport of transcription factors, to support the injured neuron's survival and axon regeneration (41). A retrograde calcium wave has also been shown to support axon regeneration by altering neuronal gene expression after PNS injury through epigenetic mechanisms (76). Together, these changes in gene expression are thought to contribute to the enhanced regeneration seen in injury-conditioned neurons. Consistent with this, the accelerated axon growth seen upon culture of *in vivo* injury-conditioned DRG neurons requires mRNA translation but not new gene expression (25,50). Notably, translation of *Calr* mRNA in injured axons is needed for initiating axon growth so the axonal CALR protein serves a localized function in injured axons (55). Our data show that axonal *Khsrp* is an injury-response mRNA whose intraxonal translation is activated through $\text{Ca}^{2+} \rightarrow \text{PERK} \rightarrow \text{eIF2}\alpha^{\text{PSS1}}$ pathway similar to *Calr* mRNA (55,77). In contrast to axonal *Calr* mRNA, the axonal *Khsrp* protein product attenuates rather than facilitates axon regeneration. This nerve regeneration slowing effect of KHSRP undoubtedly requires other factors in the nerve including components of the cytoplasmic exosome for RNA degradation and possibly other mediators interacting with KHSRP.

The difference between *in vivo* axon regeneration for the *Khsrp* deleted compared to wild type mice was most apparent in the injury-conditioned setting where axonal KHSRP levels are already increased in the wild type mice at the time of the second injury. Though axon growth from the cultured DRGs is overall increased by neuronal *Khsrp* deletion, *in vivo* injury-conditioning diminished the differences between the *Khsrp* deleted and wild type DRGs when cultured. This could represent a ceiling or threshold effect, where the transcriptional changes occurring after injury-conditioning are able to overcome growth-attenuating effects of KHSRP. Consistent with this notion, exogenously manipulating neuronal levels of several growth-associated proteins that are part of the transcriptional response to injury-conditioning has been shown to induce corresponding changes in axon growth potential, with the injury-conditioning effect blocked by genetic knockouts and depletions or increased axon growth from naive neurons upon overexpression of growth-associated proteins (78). For example, overexpression of GAP43 can increase axon growth in naive neurons (79). Notably, increasing axonally targeted *Gap43* mRNA levels in naive DRG neurons shifts them to an elongating growth phenotype typical of injury-conditioned DRG neurons (80). Though axonal levels of *Gap43*, *Snai2* and *Fubp1* mRNAs all increased in wild type mice after nerve injury, this clearly was not sufficient to accelerate axon growth considering that loss of KHSRP exaggerated these increases and accelerates axon growth above what is seen in injury-conditioned mice.

Rather than a transcriptional effect, our data support the hypothesis that axonal KHSRP levels accumulate after axotomy through post-transcriptional mechanisms. This constitutes a novel axon-intrinsic mechanism to slow axon regeneration over time. Consistent with this idea of accumulating effects of KHSRP, there was no significant difference between single injury *Khsrp*^{-/-} and wild type mice at 7 days but the single injury *Khsrp*^{-/-} mice showed significantly greater regeneration at 14 d than the wild type mice (Figure 3A). Additionally, *in vitro* axon regeneration was much greater in *Khsrp*^{-/-} and wild type DRG cultures when axon shafts were severed (Supplementary Figure S3B, C) as opposed to initiation of axon growth that is seen after dissociation of the ganglia. Naive DRG neurons are known to transition to the elongating axon growth phenotype typical of the injury-conditioned neurons if provided a 12–24 h *in vitro* period for new gene expression after culturing (50). The assessment for regeneration of sheared axons in cultured DRGs used here in Supplementary Figure S3B-C was designed with this *in vitro* priming period in mind. Others have used a similar *in vitro* priming and then replated the DRGs after the priming from dissociation and culture (81). In contrast to our approach, replated neurons reinitiate axon growth from the soma. So, the replated neurons are starting over without the effect of axons with elevated KHSRP, similar to the culture of injury-conditioned neurons used in Figure 2. Taken together, our data support a model where accumulation of KHSRP in the axonal compartment slows axon regeneration through post-transcriptional mechanisms. The increase in axon regeneration from injury-conditioned KHSRP deleted mice raises the possibility of targeting localized post-transcriptional mechanisms to increase axon regeneration rates beyond the accelerated rates normally seen in injury-conditioned neurons.

KHSRP has been shown to play roles in RNA splicing, trafficking and degradation as well as miRNA biogenesis (22,44,45,48). Our analyses of KHSRP target mRNA levels in sciatic nerves of the *Khsrp*^{-/-} mice indicate that loss of KHSRP increases axonal levels of those mRNAs. miRNAs are clearly linked to stability of mRNAs, and their precursors (pre-miRNAs) have been shown to localize into PNS axons, with PNS nerve injury triggering localized processing of some axonal pre-miRNAs into mature miRNAs (82). So, the increase in axonal KHSRP could affect axonal mRNA levels by promoting miRNA maturation. In our hands, levels of previously reported KHSRP target miRNAs (45) showed no changes when comparing DRG cultures from *Khsrp*^{-/-} vs. *Khsrp*^{+/+} mice (data not shown). However, our data argue for a direct effect of KHSRP on mRNA stability rather than an indirect effect through miRNAs or other mediators underlying the axon growth attenuation by KHSRP. KHSRP's KH3 and KH4 are implicated in promoting mRNA decay by binding to ARE-containing mRNAs and recruiting components of the cytoplasmic exosome complex used for RNA degradation (22). We find that KHSRP's function in slowing axon regeneration and decreasing axonal *Gap43* mRNA in adult DRG neurons require an intact KH4 domain that has been shown to bind to ARE-containing mRNAs. As we see with *Fubp1*

mRNA levels being depleted by KHSRP Δ KH4 expression, KHSRP can clearly affect neuronal mRNA levels through other mechanisms. A recent combination of RNA profiling studies of *Khsrp*^{-/-} vs. *Khsrp*^{+/+} mice brains with mRNAs co-immunoprecipitating with KHSRP from mouse brain uncovered 527 KHSRP bound mRNAs whose levels are elevated in KHSRP knockout brain, including *Gap43*, *Fubp1* and *Snap25* mRNAs (65). Thus, KHSRP likely binds to many other axonal mRNAs beyond those tested here to regulate their stability, and the proteins encoded by those mRNAs could also impact axon growth capacity. Recent work in neonatal DRG neurons showed that KHSRP binds to the long non-coding RNA *ALAE* in axons and this interaction prevents KHSRP's binding to *Gap43* mRNA. In contrast to our observations, translation of axonal *Gap43* mRNA was altered by the KHSRP-*ALAE* RNA competition (83) rather than increased *Gap43* mRNA levels that we see with deletion of KHSRP. We cannot completely exclude secondary effects from KHSRP deletion used here as contributing to axon growth promotion. Consistent with this possibility, the work from Olguin *et al.* reported ~1400 mRNAs showing altered levels in brains of *Khsrp*^{-/-} but no binding to KHSRP in RIP assays (65). Some of these mRNAs may represent indirect effects from the protein products altered by KHSRP mRNA decay promotion. Also, it should be noted that KHSRP deletion does not affect levels of all regeneration-associated mRNAs, since it did not affect *Hmgbl* mRNA levels and intra-axonal translation of *Hmgbl* enhances axon growth (43). RNA profiling studies will be needed to determine the RNA regulons that are directly regulated by KHSRP's decay-promoting domains vs. those that are regulated by other domains of KHSRP or indirectly modified by changing levels of proteins encoded by KHSRP mRNA targets. Nonetheless, our data point to the KHSRP's role in promoting mRNA decay as a key determinant of axon regeneration *in vivo*.

DATA AVAILABILITY

Proteomics data for axonal RBPs has been uploaded to PanoramaWeb Public and can be accessed at <https://panoramaweb.org/axon-rbps.url>.

SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online.

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