Local Translation of Extranuclear Lamin B Promotes Axon Maintenance

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INTRODUCTION

Wiring the central nervous system (CNS) requires the precise guidance of axons to distant synaptic targets and their subsequent maintenance. En route, the growth cone (GC) at the tip of a growing axon and remote from the cell body encounters numerous extracellular signals to which it must promptly respond to guide the axon to its synaptic target. Developing axons possess a considerable degree of functional autonomy, and axons severed from their cell bodies can respond to guidance cues in culture and navigate accurately in vivo. This autonomous signal processing in the axon often involves rapid regulation of protein levels in the GC, which is achieved by local protein synthesis and degradation. For example, local translation of β-actin is required for chemotropic responses of GCs to guidance cues in cultured neurons, and local translation of ephA2 mRNA is required for midline crossing of chick spinal commissural axons in vivo.

An unexplored question is whether axons also draw on local translation for their survival. Distal axons require a constant supply of energy to maintain their integrity, and axonal degeneration is often associated with mitochondrial disorders. One of the first mitochondrial diseases to be understood at a molecular level was Leber’s Hereditary Neuropathology (LHON), which results from a mutation in mitochondrial DNA and is characterized by degeneration of the optic nerve. Mitochondrial diseases share common pathologies, including axonal degeneration, with diseases caused by mutations in lamin genes, collectively known as laminopathies. Interestingly, lamin mRNAs have recently been found in various axons.

Lamins, encoded by three genes, lamins A, B1, and B2, are type V intermediate filament proteins, which comprise the nuclear lamina inside the inner nuclear membrane. Unlike other types of intermediate filaments such as neurofilaments and nestin, lamin filamentous lamins contain a nuclear localization signal (NLS) and in most case a CaaX motif at their C termini. Posttranslational modification of the CaaX motif is responsible for a nonspecific membrane targeting of lamins, which are then imported to the nucleus by a mechanism involving the NLS.

SUMMARY

Local protein synthesis plays a key role in regulating stimulus-induced responses in dendrites and axons. Recent genome-wide studies have revealed that thousands of different transcripts reside in these distal neuronal compartments, but identifying those with functionally significant roles presents a challenge. We performed an unbiased screen to look for stimulus-induced, protein synthesis-dependent changes in the proteome of Xenopus retinal ganglion cell (RGC) axons. The intermediate filament protein lamin B2 (LB2), normally associated with the nuclear membrane, was identified as an unexpected major target. Axonal ribosome immunoprecipitation confirmed translation of lb2 mRNA in vivo. Inhibition of lb2 mRNA translation in axons in vivo does not affect guidance but causes axonal degeneration. Axonal LB2 associates with mitochondria, and LB2-deficient axons exhibit mitochondrial dysfunction and defects in axonal transport. Our results thus suggest that axonally synthesized lamin B plays a crucial role in axon maintenance by promoting mitochondrial function.
of these cells (Toesca, 1992), hinting at an extranuclear role for lamins.

Here we show by using a proteomics approach that LB2 is locally synthesized in cultured Xenopus retinal ganglion cell (RGC) axons in response to guidance cue stimulation. Accordingly, lb2 mRNA localizes to and is translated in retinal axons in vivo. Furthermore, LB2 proteins localize to mitochondria in axons, and axonal synthesis of LB2 proteins and their extranuclear actions are required to maintain mitochondrial function and axonal integrity, implicating a role for local translation in preventing axon degeneration. The results show that the functional autonomy of developing axons requires a sustained local supply of LB2 proteins to mitochondria. This mechanism may provide a therapeutic insight into laminopathies by revealing an unexpected link between lamins and axonal mitochondria.

RESULTS

Subcellular Profiling using DIGE-NCAT: Identification of Cue-Induced Proteins

Several genome-wide screens using axons/GCs severed from neurons grown in culture have identified thousands of axonally localized mRNAs (Andreassi et al., 2010; Gumy et al., 2011; Taylor et al., 2009; Zivraj et al., 2010)—a puzzlingly large number to provide functionally relevant insights. We sought to identify which of these axonally localized mRNAs are actually translated in response to guidance cues. First, we used 2D difference gel electrophoresis (2D-DIGE), a method that permits quantitative comparative analysis between two proteomes, to ask whether global proteomic differences could be detected after cue stimulation in cultured embryonic Xenopus retinal neurons. (Lilley et al., 2002). This method alone, however, was not sufficiently sensitive to detect minute changes in axonal proteomes (Figure S1 available online). Therefore, we developed a DIGE-NCAT (noncanonical amino acid tagging) strategy that combines 2D-DIGE with bioorthogonal noncanonical amino acid tagging (BONCAT) and used it to investigate cue-induced changes in the subcellular proteome of RGC axons. BONCAT uses amino acid analogs (e.g., L-azidohomoalanine: AHA) that are incorporated into de novo polypeptides. These analogs can be linked to a fluorescent tag (e.g., tetramethylrhodamine: TAMRA) by “click chemistry” (Dieterich et al., 2007) and detected after 2D-DIGE. In DIGE-NCAT, axonally synthesized proteins are first labeled with AHA and TAMRA, and the axonal lysates are mixed with CyDye-labeled (used as the internal standard) and unlabeled lysates (used to enrich proteins per spot). The combined lysate is then run on one gel. Using CyDye-labeled spots as reference points, AHA-labeled axonal proteins from different conditions can be compared (Figure S2). Mass spectroscopy (MS) analysis of minute quantities of axonal, AHA-labeled protein is also possible because of retinal lysate enrichment.

We first confirmed that AHA selectively labels newly synthesized polypeptides in retinal cultures in a protein synthesis inhibitor-sensitive manner (Figure 1A). Coomassie staining of the same gel showed that the amounts of total proteins in these groups were indistinguishable (Figure 1B). We used the ratio of AHA to Coomassie intensity as an index of de novo protein synthesis (Figure 1C) and compared axonal protein synthesis induced by different guidance cues (Figure 1D). Engrailed-1 (En-1) elicited the greatest increase (~80%) (Figure 1D), consistent with previous reports showing that extracellular Engrailed causes rapid translation-dependent guidance responses in RGC axons (Brunet et al., 2005; Wizenmann et al., 2009). Therefore, we chose En-1 as a stimulus to investigate the changes in the axonal proteome in cultured RGC axons.

To examine specific changes taking place exclusively in the axonal proteome upon En-1 stimulation, we prepared axon-only cultures by culturing whole embryonic eyes, allowing the RGC axons to grow out, and then removing the eyes leaving only the distal axons attached to the substrate (Figure 2A). Typically, 1,000 embryonic eyes were cultured for each condition. PCR amplification of genomic DNA containing the β-actin gene...
in each fraction showed that the axonal fraction was free of cell body contamination (Figure 2A). In contrast, \(\beta\)-actin mRNA, which is known to localize to RGC axons in Xenopus (Leung et al., 2006), was detected by RT-PCR from the same fraction. These primers flank one intron, and therefore the RT-PCR product is shorter than the genomic DNA PCR product. En-1 stimulation of RGC axons and visualization of newly synthesized proteins with DIGE-NCAT revealed clear differences between control and En-1 conditions (Figure 2B). The overall rate of protein synthesis was significantly increased (Figure S2G), and quantitative analysis revealed that out of 909 putative protein spots detected in the two conditions, 259 (28.5%) spots increased and 335 (36.8%) decreased in intensity by 1.5-fold or more with En-1 stimulation (Figures 2C and 2D). Representative spots from the increased, decreased, or unchanged groups were analyzed by MALDI-TOF MS (Figure S2H and Table S1). The two
spots that showed the most robust increase were identified as LB2 (gene name—\textit{lmnb2}; MASCOT scores—254 and 351; Figures 2B, 2C, and S2H; Table S1).

LB2 Is Locally Synthesized in RGC Axons In Vitro and In Vivo

Identification of LB2 synthesis in axons was unexpected because lamins are major constituents of the nuclear envelope (Hutchison, 2002), and thus our results raised the intriguing possibility of an extranuclear role for locally synthesized LB2 in neurons. If LB2 were axonally synthesized, its transcripts and proteins would be expected to reside in the axons. Using RT-PCR and fluorescent in situ hybridization (FISH), we determined that \textit{lb2} transcripts localize to retinal axons and GCs in vitro (Figures 3A–3C). In situ hybridization (ISH) of retinal sections showed strong staining in the optic nerve head (ONH), a nucleus-free, axon-only structure where the RGC axons collect to exit the eye (Figure 3G), indicating that \textit{lb2} mRNA localizes
to axons in vivo (Figure 3D). Abundant signal was also observed in the inner and outer plexiform layers (IPL and OPL, respectively), suggestive of localization in the dendrites of RGCs and in the axons of other retinal neurons such as bipolar cells. RGCs express little or no mRNAs encoding other lamins (A and B1) (Figures 3E and 3F). In sharp contrast to lb2, pax6 mRNA, which encodes a nuclear protein also highly expressed by RGCs, does not localize to the ONH (Figure 3G), suggesting that specific axonal transport rather than diffusion accounts for the axonal localization of lb2 mRNA. Axonal lb2 mRNA is also detected outside the eye, including the neuropil of the optic tectum where RGC axons terminate (Figures S3A–S3D). To examine the detected outside the eye, including the neuropil of the optic tectum where RGC axons terminate (Figures S3A–S3D). To examine the expression of LB2 protein, we used a specific antibody against LB2, which detected a single band of the predicted molecular weight on a western blot (Figure S3K) and a specific nuclear signal in immunocytochemistry (Figure S3H). Immunostaining revealed LB2 expression in axons and GCs in vitro (Figure 3H), which showed a dose-dependent reduction when lb2 mRNA translation was inhibited with a translation-blocking antisense morpholino (LB2MO), verifying specificity (Figures 3H, 3I, 3J, and S3J). Experiments with two additional antibodies, which specifically recognize LB2, confirmed this finding (Figures S3K–S3M). Axonal LB2 protein was also detected in vivo in the ONH, optic nerve, and tectal neuropil (Figures 3J, 3K, and S3E–S3G). Of note, fibroblast-like cells in culture exhibited specific LB2 immunostaining in the cytoplasm in addition to a strong nuclear signal (Figure S3H), suggesting that the extranuclear localization of LB2 is not limited to RGCs.

We next used a different method, quantitative immunofluorescence (QIF), to examine En-1-induced LB2 synthesis in axons. Indeed, 1 hr stimulation with En-1 resulted in a 34% translation-dependent increase of LB2 proteins in GCs severed from their cell bodies (Figures 4A and 4B). This increase was not a secondary result of differences in GC morphology, as the GC area was not affected (Figure 4C). In contrast, En-1 did not stimulate the axonal translation of β-actin mRNA, which localizes to these GCs (Leung et al., 2006) (Figures S4A–S4C). Next, we investigated whether lb2 mRNA is axonally translated in vivo. We employed translating ribosome affinity purification (TRAP), which enables isolation of actively translating mRNAs (i.e., mRNAs bound to ribosomes) by expressing a GFP-tagged ribosomal protein, L10a, in cells of interest (Heiman et al., 2008). To examine LB2 synthesis exclusively in retinal axons, we transplanted GFP-L10a-expressing eye primordia to unlabeled hosts (Figure 4D). The grafted eye sent out GFP-L10a-expressing RGC axons to the contralateral optic tectum of the host embryo, and the host brains containing the GFP-L10a-expressing RGC axons were collected (without eyes) and affinity purified against GFP. Because RGCs are the only neurons sending a projection out of the eye and into the brain, this enabled us to examine mRNAs associated with GFP-L10a (i.e., ribosomes) exclusively from developing RGC axons in vivo (axon-TRAP). RT-PCR from axon-TRAP confirmed the presence of β-actin mRNA—a molecule known to be locally synthesized in RGC axons (Leung et al., 2006)—and lb2 mRNA in the axonal translome (Figures 4E and 4F). In contrast, pax6 and bnn mRNAs, which encode nuclear proteins and are expressed in the eyes at similar levels to lb2 mRNA, were not present (Figure S4D). Lamin A and B1 mRNAs were not isolated by axon-TRAP (Figure S4E), as expected from their negligible levels of expression in RGCs (Figures 3E and 3F). These results suggest that lb2 mRNA is indeed locally translated in RGC axons in vivo.

Local Synthesis of LB2 Is Critical for Axon Survival In Vivo

We next assessed retinal axons, in which lb2 translation is blocked by MO, by lipophilic dye (DiI) labeling (Figures 5A and 5B). Although RGC axons in LB2 morphants appeared to be relatively normal when they reached their target in the optic tectum at stage 39/40, a severe axonal loss was observed 2 days later at stage 45 (Figure 5B). Approximately 70% of the brains had fewer than 15 Dil-labeled axons at stage 45, whereas less than 5% of control morphant brains did (Figure 5D). Degenerating axons, distinguished by their beaded and discontinuous morphology (Figure 5C), were evident along the pathway in LB2 morphant brains at stage 45, suggesting that LB2 may be required for axon maintenance. Morphants appeared normal with no overt differences in embryo morphology and Dil filling of the eye (data not shown). Furthermore, HRP filling of LB2 morphant eyes also showed a similar loss of retinal axons at stage 45 (data not shown), indicating that the reduction in the number of retinal axons is not due to differences in Dil uptake. Next, in order to rule out the possibility of nonspecific off-target effects of the MO (Heasman, 2002), we performed rescue experiments by coinjecting a MO-resistant LB2-GFP expression vector (Prüfert et al., 2004). LB2-GFP almost completely rescued the axon degeneration phenotype (Figures 5B and 5D), demonstrating that the phenotype is specific to the loss of LB2 function. We next asked whether this phenotype was an axonal event, rather than a consequence of cell body death, which is predominantly mediated by the caspase-3 pathway (Nikolaev et al., 2009). Examination of cell body death using active caspase-3 as a marker showed no difference between CoMO- and LB2MO-injected embryos at stages 41–45 (Figures 5E and 5F).

Blastomere injection delivers the MO into the prospective brain as well as the eye (Figure S5A). To confirm the RGC-autonomous effect of LB2MO, we used targeted electroporation to deliver the MO specifically into the eye, without affecting the brain (Figures 6A and 5B). As with blastomere injection, the RGC axon projection was normal at stage 40 but became significantly less dense by stage 45 (Figure 6B), as quantified by the mean Dil signal intensity of the distal 150 μm portion of the projection (Figures 6C and 6D). A more central question, however, is whether LB2 synthesis in the axon is specifically required for axon survival. To inhibit lb2 mRNA translation specifically in axons, we took advantage of the fact that distal RGC axons are anatomically compartmentalized away from the soma and navigate close to the lateral surface of the brain, making these axons easily accessible to electroporation (Chien et al., 1993; Falk et al., 2007; Holt, 1984). This allows delivery of LB2MO directly into distal axons (Figures 6E, 6F, SSC, and 5A), similar in principle to inhibition of axonal proteins using compartmentalized chambers in vitro (Hengst et al., 2009). We found that axonal delivery of LB2MO led to a significant axon loss (Figure 6G).
Diffusion of the MO retrogradely into the soma was not observed at stage 45 in RGC bodies in the contralateral eye (absence of FITC-labeled MO; data not shown), and the nuclear LB2 intensity in the RGC layer was not reduced, ruling out the possibility of a nuclear LB2 knockdown by axonal LB2MO delivery (Figure 6H).

To further examine whether axonally synthesized LB2 functions...
locally or in the nucleus after retrograde transport to the nucleus, we asked whether an LB2 lacking the NLS (LB2ΔNLS) could rescue the axon degeneration phenotype. This mutant form of LB2 does not enter the nucleus and instead localizes to cytoplasmic structures (Figure 6I). We introduced RNAs encoding LB2ΔNLS into RGCs by eye electroporation in LB2-depleted embryos that had been injected with a reporter plasmid to label RGCs with RFP (ath5::RFP) along with the LB2MO (Figures 6J and S5D). Strikingly, LB2ΔNLS almost completely rescued the LB2 loss-of-function phenotype of RGCs (Figures 6K and 6L).

Figure 5. Inhibiting LB2 Translation Results in Degeneration of RGC Pathway In Vivo
(A) MO injection and RGC axon labeling using Dil (green: MO; Tel: telencephalon; Ch: optic chiasm; OT: optic tract; Tec: optic tectum; and red: Dil).
(B) Dil-labeled RGC axons in MO-injected embryos with or without LB2-GFP RNA.
(C) Characteristic beaded morphology of dying axons in LB2 morphants. The boxed area is shown in the right panel.
(D) Quantitative analysis of the LB2MO-induced reduction in RGC axons and its rescue by LB2-GFP (n = no. of brains; 3 replicates; ****p < 0.0001; Fisher’s exact).
(E and F) The frequency of active caspase-3-positive cells in retinal sections (mean ± SEM; n = sections analyzed; Mann-Whitney).
Scale bars: 65 μm in (B) and (C), 100 μm in (C) right panel, and 25 μm in (E). See also Figure S5.
suggesting that extranuclear LB2 is sufficient for axon survival. Taken together, these results suggest that axonally synthesized LB2 is required and sufficient for RGC axon survival.

**LB2 Knockdown Leads to Mitochondrial Dysfunction**

How does LB2 regulate axon survival? We concentrated on mitochondria because the mitochondrial proteome contains LB2 (Fountoulakis et al., 2002; Rezaul et al., 2005; Taylor et al., 2002), and mitochondrial dysfunction is strongly associated with axon degeneration in pathological conditions including multiple sclerosis (Su et al., 2009) and amyotrophic lateral sclerosis (Shi et al., 2010). Furthermore, maintenance of mitochondrial membrane potential and function is critical in caspase-3-independent neurite degeneration (Ikegami and Koike, 2003).
a phenotype we observe in LB2 morphants (Figures 5B and 5E). Coimmunocytochemistry using antibodies against LB2 and CoxIV, a cytochrome c oxidase subunit localized to the inner mitochondrial membrane, revealed a significant overlap of LB2 and CoxIV in axons and GCs (Figure 7A; Pearson’s coefficient = 0.800 ± 0.01, n = 7), which was confirmed by the proximity ligation assay (PLA) that produces a positive signal only when two different antibodies are within 40 nm (Figure S7C)—a distance consistent with a potentially direct molecular interaction (Fredriksson et al., 2002). LB2 also colocalizes with VDAC2, a mitochondrial outer membrane protein (Figures 7B and S7B), as well as with mitochondria-targeted GFP (Mito-GFP) (Figure 7C). OMX super-resolution microscopy analysis showed that LB2 localizes inside the mitochondria in a punctate pattern (Figure 7C). In line with these results, LB2ΔNLS that cannot be imported into the nucleus localizes to cytoplasmic structures that include mitochondria (Figures S6B–S6H). These results suggest that LB2 synthesized in distal axons where nuclear import machinery is not abundant may be imported into mitochondria.

We next examined how LB2 depletion affects mitochondria. First, we measured the mitochondrial membrane potential (ΔΨm) using two indicators and found that it is significantly reduced in LB2-depleted axons (Figures 7D and 7E). We also analyzed mitochondrial morphology and found that LB2 morphant axons contained strikingly elongated mitochondria although the average number was not different from the control (Figures 7F–7H). Dynamic image analysis showed that the motile axonal mitochondria (approximately 20% of total) exhibited a slight but statistically nonsignificant decrease in motility in both directions in LB2-depleted axons (1.3 versus 1.1 anterograde movements/50 μm axon in 5 min, p = 0.14; 0.8 versus 0.4 retrograde movements/50 μm axon in 5 min, p = 0.13; n = 14 for CoMO and n = 17 for LB2MO; Mann-Whitney). A reduced mitochondrial potential and elongated morphology are associated with dysfunctional mitochondria (Ikegami and Koike, 2003; Yoon et al., 2006). Mitochondrial dysfunction impairs axonal transport of organelles such as lysosomes, which is a typical feature of degenerating axons (Kaasik et al., 2007), and causes optic neuropathies in humans (Carelli et al., 2004). Therefore, we analyzed the dynamics of lysosome movements in cultured retinal axons and found that LB2-depleted axons showed a significantly reduced number of directional movements (i.e., ≥3 μm movements occurring without a pause or a change of direction) compared to the control axons (Figure 7I). The reduction was observed in both anterograde and retrograde directions.

**DISCUSSION**

Here we have developed a proteomics strategy, DIGE-NCAT, to visualize subtle changes in the axonal proteome. The key advantage of proteomics-based approaches in detecting cue-induced
mRNA translation and protein synthesis is the accumulation of signal throughout the cue stimulation period. Cue-induced local mRNA translation is a dynamic process regulated in a narrow time window. For example, the BDNF-induced increase in local mRNA translation activity in dendrites of cultured hippocampal neurons shows complex kinetics in which different parts of the same dendrite reach a peak at different times (Aakalu et al., 2001). Therefore, many mRNAs whose translation is induced by a cue may evade ribosome immunoprecipitation protocol (i.e., TRAP) performed at a fixed time point after the cue stimulation. The main advantage of DIGE-NCAT over other proteomics approaches is its sensitivity. We were able to detect subtle changes in axonal protein levels, both up and down, suggesting that this technique may prove to be a versatile tool to study dynamic changes of the axonal proteome including translational repression and/or protein degradation. Metabolic labeling represents another sensitive technique, but the use of nonradioactive material and an internal standard for more reliable comparison between multiple conditions give DIGE-NCAT more advantages.

This proteomics approach revealed, unexpectedly, that LB2 is axonally synthesized in response to extrinsic cue stimulation. LB2, a known nuclear protein, has previously been shown to be involved in regulating nuclear translocation and neuronal migration. In Drosophila, LB2 regulates the apical migration of nuclei in differentiating photoreceptors (R cells) by connecting the microtubule-organizing center (MTOC) to the nucleus to regulate nuclear translocation, a critical process for neuronal nuclear migration (Patterson et al., 2004). A recent study showed that lb2 knockout mice exhibit specific defects in neuronal migration in the cerebral cortex and cerebellum (Coffinier et al., 2010), suggesting that LB2’s role in regulating nuclear translocation may be evolutionarily conserved. However, our finding indicates that axonally synthesized LB2 functions in mitochondria to support axon survival. The local synthesis of LB2 in a distal process remote from the nucleus may enable the protein to escape nuclear localization and take on cytoplasmic functions. Indeed, we show that deleting the 6 amino acid NLS at the C terminus enhances the mitochondrial targeting of LB2 (Figures S6B–S6G). Intriguingly, MitoProt, Software that predicts mitochondrial proteins with high accuracy (Claros and Vincens, 1996), shows that Xenopus LB2 has an extremely high probability of mitochondrial localization (p = 0.97). Other nuclear-encoded mitochondrial proteins such as CoxIV, Cox17, CoQ7, and Hsp70 are also locally synthesized in the presynaptic nerve terminal (Aschrafi et al., 2010; Gioio et al., 2001), and we have identified such proteins in this proteomics study (Table S1), suggesting that axonal mitochondria require a local supply of new proteins. It will be interesting in the future to investigate whether lb2 mRNA contains an axonal localization element as recently shown for coxIV mRNA (Aschrafi et al., 2010). Exactly how LB2 modulates mitochondrial function still remains to be elucidated.

Mitochondrial function is essential for numerous processes required to maintain axonal health and stability (Knott et al., 2010), suggesting that LB2’s role in regulating nuclear translocation and neuronal migration in the cerebral cortex and cerebellum (Coffinier et al., 2010), embryonic mouse DRG neuronal axons (Gumy et al., 2000; Nguyen et al., 2004; Sasaki et al., 2000; Zhang et al., 2009). Because defective retrograde transport is strongly associated with axon degeneration (Jing and Malicki, 2009), one plausible cytoplasmic role for LB2 would be direct regulation of axonal transport through its interaction with Nudel. We did not, however, detect a strong colocalization between LB2 and Nudel in axons and GCs (Figure S7A: Pearson’s coefficient 0.417 ± 0.03, n = 5), and we observed a generally decreased cargo transport activity instead of a specific defect in retrograde transport (Figure 7I). Nonetheless, the possibility that extranuclear LB2 interacts with Nudel to regulate axonal cargo transport should not be excluded.

A general reduction in axonal transport prompted us to investigate mitochondria, the main source of ATP in axons, and our data indicate that axonally synthesized LB2 functions in mitochondria to support axon survival. The local synthesis of LB2 in a distal process remote from the nucleus may enable the protein to escape nuclear localization and take on cytoplasmic functions. Indeed, we show that deleting the 6 amino acid NLS at the C terminus enhances the mitochondrial targeting of LB2 (Figures S6B–S6G). Intriguingly, MitoProt, Software that predicts mitochondrial proteins with high accuracy (Claros and Vincens, 1996), shows that Xenopus LB2 has an extremely high probability of mitochondrial localization (p = 0.97). Other nuclear-encoded mitochondrial proteins such as CoxIV, Cox17, CoQ7, and Hsp70 are also locally synthesized in the presynaptic nerve terminal (Aschrafi et al., 2010; Gioio et al., 2001), and we have identified such proteins in this proteomics study (Table S1), suggesting that axonal mitochondria require a local supply of new proteins. It will be interesting in the future to investigate whether lb2 mRNA contains an axonal localization element as recently shown for coxIV mRNA (Aschrafi et al., 2010). Exactly how LB2 modulates mitochondrial function still remains to be elucidated.
and accumulating evidence points to mitochondrial dysfunction as one of the main causes in some neurodegenerative diseases, including familial Parkinson’s disease (Lin and Beal, 2006) and LHON (Wallace et al., 1988). Our data showing that LB2 depletion causes a reduction in mitochondrial membrane potential (Figures 7D and 7E) suggest an intriguing model whereby extrinsic cues secreted from the optic tectum, such as En-1, stimulate in distal RGC axons LB2 synthesis, which then acts locally to regulate mitochondrial function in order to meet high metabolic demands of these axons. Intriguingly, a recent study showed that En-1 also stimulates translation of mRNAs encoding mitochondrial complex I subunits and protects against degeneration of midbrain dopaminergic neurons in an animal model of Parkinson’s disease (Alvarez-Fischer et al., 2011), suggesting that the translational control of mRNAs encoding nuclear-encoded mitochondrial proteins in axons is a conserved mechanism to regulate axonal survival.

In summary, the results presented here suggest that LB2 is an axonally synthesized protein with an extranuclear role in mitochondria that is necessary to maintain the integrity of distal axons. Our findings may provide insight into the mechanisms underlying the cell type-specific nature of some laminopathies.

**EXPERIMENTAL PROCEDURES**

**Embryos and Blastomere Injection**

*Xenopus laevis* embryos were fertilized and raised as described (Drinjakovic et al., 2010). EGFP-L10a, EGFP-LB2, and Mito-XFP constructs were kind gifts from N. Heintz (Rockefeller University), G. Krohne (University of Würzburg), and M. Coleman (Babraham Institute), respectively. Ath5:RFP was previously described (Zolesi et al., 2008). LB2ΔNLS was made by inverse PCR from EGFP-LB2. Capped RNAs were made using mMessageMachine SP6 kit (Ambion). Translation blocking MOs were designed and supplied by GeneTools (Philomath, OR, USA). Embryos were injected at the 8-cell stage with MO, DNA, or RNA in dorsal animal blastomeres (presumptive CNS) as described previously (Drinjakovic et al., 2010).

**Electroporation**

Eye electroporation was performed as described (Falk et al., 2007). For pathway electroporation, the lateral side of the brain at stage 40 was exposed, and MO was electroporated close to the lateral side of the brain.

**RGC Pathway Visualization and Analysis**

Dil labeling of RGC axons was performed as described (Drinjakovic et al., 2010). Dil intensities were measured using OpenLab (Improvision). Injections of ath5:RFP (200 pg per blastomere) resulted in a mosaic expression of RFP in RGCs, which labeled fewer retinal axons compared to the Dil method. For this reason, the brains with fewer than 15 and 3 axons were counted when quantitating the axon degeneration phenotype by Dil and ath5:RFP labeling, respectively.

**TRAP**

Eye transplantation (Holt, 1984) and immunoprecipitation (Heiman et al., 2008) were performed as described. One nanogram of GFP-L10a or GFP RNA (negative control) was delivered by blastomere injections. Ten pooled brain hemispheres were used per group, and the presence of candidate mRNAs was examined by RT-PCR.

**Retinal/Axon Cultures and Stimulation**

For all the retinal culture experiments, eye primordia from stage 35/36 embryos were dissected and cultured at 20 °C for 40 hr as described (Drinjakovic et al., 2010). This stage corresponds to a stage between 42 and 43 in vivo, which is approximately 1–2 days prior to stage 45 when the LB2MO-induced axon degeneration phenotype appears. External cues used are provided in the Extended Experimental Procedures. Axon cultures were prepared as described (Campbell and Holt, 2001) by separating distal axons from the eye and removing eyes/loose cells.

**Mitochondrial Membrane Potential (ΔΨm) Measurement**

Live retinal cultures were incubated with 20 nM tetramethylrhodamine, methyl ester (TMRM), and MitoTracker red CM-H2XRos (MitoTracker) for 20 min and washed with culture medium three and six times, respectively. ΔΨm was expressed by fluorescence ratio of mitochondria (Fm) to cytoplasm (Fc) calculated using Openlab. Fc was measured no more than 3 μm away from the mitochondrion of interest.

**PCR and RT-PCR**

RNA-only and DNA/RNA copurification were performed using RNasea micro kit (QIAGEN) and DNaseasy blood and tissue kit (QIAGEN), respectively. For quantitative RT-PCR, cDNA was prepared from total RNA extracted from stage 41 embryonic eyes and amplified on a Rotor-Gene 3000 (Corbett) using QIAGEN kits.

**ISH**

ISH on retinal sections and FISH of cultured axons were performed using digoxigenin (DIG)-labeled riboprobes generated from IMAGE clones as previously described (Zivraj et al., 2010).

**Immunofluorescence**

Immunostaining, imaging and analysis were performed as described (Piper et al., 2006). Pearson’s coefficients were calculated using Velocity (Perkin Elmer). For experiments shown in Figure 6I, HEK293T cells were transfected with LB2ΔNLS plasmid using Lipofectamine 2000 (Invitrogen), labeled by MitoTracker, fixed, and imaged using a laser-scanning confocal microscope (Leica) and a 60 x 1.4NA oil immersion objective (Leica). Optical sections at 0.3–1.0 μm separation were taken and analyzed using Velocity. Immunostaining of sections was performed as described (Drinjakovic et al., 2010). For LB2 staining of embryonic sections, unfixed embryos were anesthetized, embedded in OCT, rapidly frozen in liquid nitrogen, sectioned by a cryostat, and then fixed in methanol. CellProfiler (Broad Institute) was used for quantification of nuclear LBZ. For OMX imaging, retinal cultures obtained from Mito-GFP (200 pg per blastomere) injected embryos were fixed, immunostained for LB2, and imaged using DeltaVision OMX 3D-SIM System V3 controlled by softWoRx 5.0.0 (Applied Precision) and a 100 x 1.4NA oil objective.

**Live Imaging**

Cultured axons were imaged under a Perkin Elmer Spinning Disk UltraVIEW ERS, Olympus IX81 Inverted microscope, and a 60 x 1.2NA water immersion objective. Mitochondria were visualized by blastomere injection of Mito-RFP plasmid (200 pg per blastomere), and lysosomes were labeled with Lysotracker Red (Invitrogen). Images were taken for 5 min at a 1 to 3 s interval and analyzed using Velocity.

**DIGE-NCAT**

AHA (Invitrogen) (50 nM and 500 nM for retinal eye cultures and DIGE-NCAT, respectively) containing samples were labeled using Click-IT TAMRA kit (Invitrogen), gel-separated, fixed, and visualized using a TyphoonTM 9410 imager (GE Healthcare). The total lane intensity in 1D gel was measured using ImageJ (NIH). For DIGE-NCAT experiments, 1,000 eyes were used per condition for axon-only culture. Some of the eyes were lysed and labeled with Cy3 or Cy5 (Minimal labeling kit; GE Healthcare), and the rest of the eyes were lysed without labeling. Axon-only cultures were stimulated with AHA/cue. RNA and protein were copurified using PARIS (Ambion), and RNA concentrations were used to equilibrate samples. The protein lysate was then labeled with TAMRA, mixed with 50 μg of CyDye-labeled + 150 μg of unlabeled eye lysates, 2D-DIGE separated, visualized, and analyzed by biological variation analysis. Cy5 was used as a standard for multiplex comparison. MS analyses were performed at the Cambridge Centre for Proteomics (University of Cambridge) and Alphalyse A/S (Denmark).
SUPPLEMENTAL INFORMATION

Supplemental Information includes Extended Experimental Procedures, seven figures, and one table and can be found with this article online at doi:10.1016/j.cell.2011.11.064.

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