



β II-spectrin promotes mouse brain connectivity through stabilizing axonal plasma membranes and enabling axonal organelle transport

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β II-spectrin is the generally expressed member of the β -spectrin family of elongated polypeptides that form micrometer-scale networks associated with plasma membranes. We addressed in vivo functions of β II-spectrin in neurons by knockout of β II-spectrin in mouse neural progenitors. β II-spectrin deficiency caused severe defects in long-range axonal connectivity and axonal degeneration. β II-spectrin-null neurons exhibited reduced axon growth, loss of actin-spectrin-based periodic membrane skeleton, and impaired bidirectional axonal transport of synaptic cargo. We found that β II-spectrin associates with KIF3A, KIF5B, KIF1A, and dynactin, implicating spectrin in the coupling of motors and synaptic cargo. β II-spectrin required phosphoinositide lipid binding to promote axonal transport and restore axon growth. Knockout of ankyrin-B (AnkB), a β II-spectrin partner, primarily impaired retrograde organelle transport, while double knockout of β II-spectrin and AnkB nearly eliminated transport. Thus, β II-spectrin promotes both axon growth and axon stability through establishing the actin-spectrin-based membrane-associated periodic skeleton as well as enabling axonal transport of synaptic cargo.

neuronal cytoskeleton | axonal transport | axonal growth | spectrin | STORM

Spectrins are elongated proteins that form extended plasma membrane-associated networks cross-linked by short actin filaments and cooperate with ankyrins to coordinate functionally related proteins within plasma membrane domains (1, 2). A spectrin-actin network was first discovered in mammalian erythrocytes, which contain only a plasma membrane, but spectrin assemblies are also associated with specialized plasma membranes in epithelial cells, neurons, cardiomyocytes, and skeletal muscle (1, 2). Plasma membrane-associated spectrin-actin networks in erythrocytes form a 2D polygonal structure, as first observed by electron microscopy (3). Superresolution light microscopy has revealed similar structures in neuronal cell bodies and parts of the dendrites, as well as a quasi-1D periodic structure in axons and dendrites (4, 5). These structures are based on the principle of 190-nm spectrin tetramers cross-linked by multivalent short actin filaments that are capped by actin-capping proteins such as adducin (1, 2, 6) (see Fig. 3D). The flexible nature of spectrin results in either polygonal networks, where spectrin tetramers connect individual short actin filaments at the nodes of the polygons (3, 7), or ring-like structures of aligned actin filaments that wrap the circumference of axons and parts of the dendrites evenly spaced by spectrin tetramers (4, 5). Both configurations confer mechanical stability on a cellular scale: Deficiency of spectrin results in fragility of erythrocytes and fragmentation of axons (8, 9). Noteworthily, other actin networks coexist with actin rings and polygons in neurons (10).

Spectrins and ankyrins also have been implicated in microtubule-dependent transport of intracellular organelles (11–15). The dynein/dynactin motor complex drives neuronal microtubule-based transport in the retrograde direction, while a diverse group of kinesin motors move cargo anterogradely (16). Unlike dynein,

kinesins belong to a superfamily of proteins (KIFs) that include over 40 genes, although not all members are involved in neuronal transport (16). Both dynein and kinesin are involved in neuronal disease pathogenesis (16). Initial evidence of spectrins' involvement in transport emerged from the observation that a pair of high-molecular-weight polypeptides, termed fodrin, comigrated with multiple classes of axonal organelles following metabolic pulse labeling of axonal cargo in the optic nerve (11). Subsequent identification of these polypeptides as α - and β -spectrin led to a focus on their plasma membrane roles (17). Nevertheless, other reports have supported the functional connections of both spectrin and ankyrin with organelle trafficking. For example, α II-spectrin associates with the kinesin motor KIF3 and with axonal organelles in rats, and it moves with similar velocities as KIF3 in pulse-chase experiments (12). Moreover, the dynactin complex binds spectrin through Arp1 (18) and ankyrin-B (AnkB) through p62/dynactin-4 (19). Expression of *Drosophila* β -spectrin bearing human β III-spectrin mutations causing spinocerebellar ataxia type 5 (SCA5) results in impaired neuronal organelle transport in flies (13). AnkB associates with endolysosomal organelles through PI3P lipids and is required for their axonal transport (14) and for polarized recycling of α 5/ β 1 integrin dimers in migrating fibroblasts (15).

These divergent functions of spectrins and ankyrins raise the question of whether specialized family members and isoforms are dedicated to either plasma membrane structure or organelle transport, or if the same polypeptides perform both roles. In the

Significance

Establishment and maintenance of long axonal projections are essential for brain development and function of the nervous system. Plasma membrane-associated structural proteins promote axon stability and efficient organelle transport, which is thought to be critical in supporting axonal elongation, while both activities are essential for long-term survival of neurons. This study demonstrates that β II-spectrin in axons enables organelle bidirectional transport in addition to its previously established role in forming a periodic membrane-associated actin-spectrin skeleton. β II-spectrin thus promotes both axon stability and axon growth through biochemically distinct functions.

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case of ankyrins, AnkB (encoded by the *ANK2* gene) associates with intracellular organelles and this activity depends upon a single exon that diverged from ankyrin-G (AnkG, encoded by the *ANK3* gene) early in vertebrate evolution (20). Four vertebrate β -spectrins share ankyrin-binding activity, and three of them exhibit tissue-specific expression. β I-spectrin is primarily found in erythrocytes, striated muscle, and a subset of neurons. β III-spectrin is expressed at highest levels in the cerebellum, where it is enriched in dendrites of Purkinje neurons, but it is also expressed in other areas of the brain (21). β IV-spectrin similarly exhibits cell-type-specific expression in neurons and pancreatic β -cells (1, 2, 22). Consistent with these restricted expression patterns, mice and humans lacking these β -spectrins are viable, although with serious phenotypes, including severe anemia for β I-spectrin (23), cerebellar ataxia for β III-spectrin (24), and abnormal neural development for β IV-spectrin (25). β II-spectrin, in contrast to other members of the β -spectrin family, is expressed in most cell types (although not in mature erythrocytes), and whole-body β II-spectrin knockout (KO) mice die by embryonic day (E)12 (26).

Here, we find that KO of β II-spectrin targeted to neuronal progenitors results in major loss of long-range axonal connectivity as well as axonal degeneration. β II-spectrin-deficient neurons have reduced axonal growth in vitro and exhibit loss of periodic actin rings. However, loss of β II-spectrin also markedly impairs bidirectional transport of synaptic organelles in axons. We determined that β II-spectrin associates with the dynein/dynactin motor complex, and several kinesins. Our results suggest that assembly of the periodic actin-spectrin-based membrane skeleton is not required to enable the intracellular transport machinery. Finally, we establish that the deficits in transport caused by loss of β II-spectrin are independent of its interaction with AnkB. Together these findings demonstrate that β II-spectrin multitasks in vivo as a critical structural component of axonal plasma membranes as well as an enabler of axonal organelle transport.

Results

Loss of Central Nervous System β II-Spectrin Causes Loss of Brain Connectivity. To determine functional roles of β II-spectrin in neurons in vivo, we crossed mice homozygous for the β II-spectrin floxed allele (β II $Sp^{flx/flx}$) developed by Rasband and coworkers (27) with Nestin-Cre mice, which express Cre-recombinase in neural progenitor cells. We confirmed over 90% reduction of

brain β II-spectrin levels in β II $Sp^{flx/flx}$ /Nestin-Cre (β II-SpKO) mice by immunofluorescence staining (Fig. 1A) and by immunoblot analysis of total brain homogenates and culture neurons (SI Appendix, Fig. S1A). The residual β II-spectrin signal in β II-SpKO mice is likely contributed by brain endothelial cells and a small population of neurons that escaped Cre recombination. Depletion of neural β II-spectrin slightly reduced levels of α II-spectrin, its obligatory partner, and 440-kDa and 220-kDa isoforms of AnkB (SI Appendix, Fig. S1B and C). In contrast, we detected no changes in the expression of β III-spectrin, AnkG, or β IV-spectrin Σ 1 but significant elevations in β IV-spectrin Σ 6 isoform and ankyrin-R (AnkR) at postnatal day (PND)25 (SI Appendix, Fig. S1B and C). Together, these data suggest that other cytoskeletal proteins do not compensate for the embryonic loss of β II-spectrin, except for β IV-spectrin Σ 6, which is predominantly expressed in the postnatal brain and localized to axon initial segments (AIS) (28), and AnkR, which is expressed in a subset of forebrain and cerebellar neurons (29, 30).

Conditional β II-SpKO mice were born at expected ratios but were smaller than β II $Sp^{flx/flx}$ (β II-Sp-WT) littermate controls (Fig. 1B), with motor coordination deficits and multiple seizures (SI Appendix, Fig. S2B). β II-SpKO mice experienced early postnatal lethality between PND15 and PND45 (SI Appendix, Fig. S2A), as previously reported (27). Histological characterization of PND0 and PND6 β II-SpKO brains revealed either complete absence or significant reduction of long axonal bundles connecting cerebral hemispheres, including the corpus callosum (CC) (Fig. 1A and SI Appendix, Fig. S2C). We next implemented an MRI- and diffusion tensor imaging (DTI)-based protocol with compressed sensing, which delineates white matter structures in the brain and allows 3D reconstruction of axonal tracts (31). Fractional anisotropy (FA) maps derived from DTI images from β II-SpKO PND25 brains show loss or complete absence of long axonal projections, including those in the CC (Fig. 1C and D, yellow inset). Analysis of 3D-reconstructed fibers revealed that β II-SpKO brains have a major reduction in the average length of axons (Fig. 1G) and in FA values, which is indicative of loss of axonal integrity (Fig. 1H). β II-spectrin deficiency thus leads to widespread deficits in axonal length and long axon tracts.

Loss of β II-Spectrin Results in Larger-Caliber Axons and Axonal Degeneration. To determine if loss of β II-spectrin compromises axon survival, we examined the ultrastructural organization of

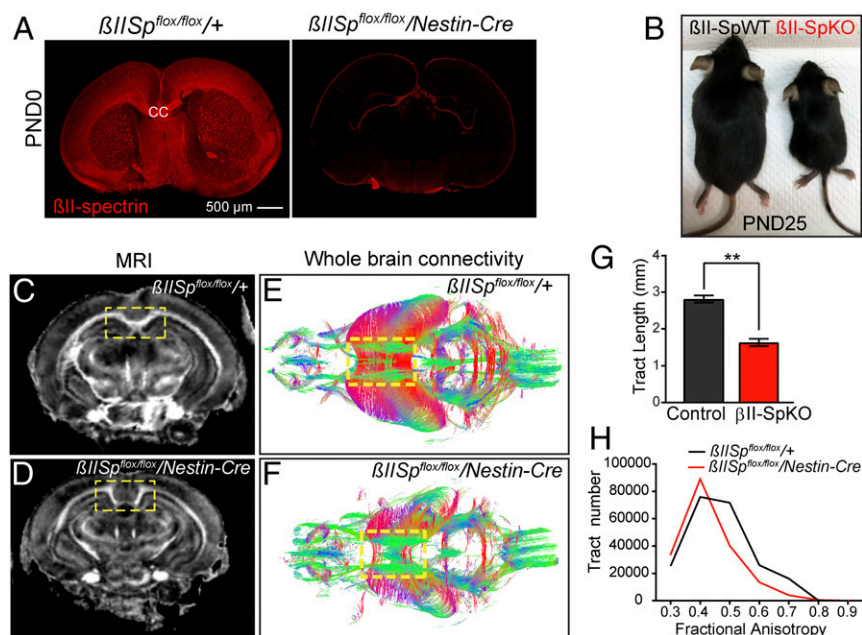


Fig. 1. Brain β II-spectrin is required for axonal integrity and long-range axonal connectivity. (A) β II $Sp^{flx/flx}/+$ and β II $Sp^{flx/flx}$ /Nestin-Cre PND0 brains stained for β II-spectrin. (Scale bar, 500 μ m.) (B) PND25 β II $Sp^{flx/flx}/+$ (β II-SpWT) and β II $Sp^{flx/flx}$ /Nestin-Cre (β II-SpKO) mice. (C and D) FA maps of β II $Sp^{flx/flx}/+$ (F) and β II $Sp^{flx/flx}$ /Nestin-Cre (G) PND21 brains derived from DTI. (E and F) Three-dimensional images of white matter tracts from β II $Sp^{flx/flx}/+$ (E) and β II $Sp^{flx/flx}$ /Nestin-Cre (F) mice colored by FA values. The yellow box indicates the position of the CC, which is absent in β II $Sp^{flx/flx}$ /Nestin-Cre mice. (G) Analysis of white matter tract length in β II $Sp^{flx/flx}/+$ ($n = 6$) and β II $Sp^{flx/flx}$ /Nestin-Cre ($n = 7$) brains. Data represent mean \pm SEM. Student's t test, $***P < 0.01$. (H) FA distribution of white matter tracts in β II $Sp^{flx/flx}/+$ ($n = 6$) and β II $Sp^{flx/flx}$ /Nestin-Cre ($n = 7$) brains.

long-range axons, using transmission electron microscopy (TEM) of cross-section of the CC. PND25 β II-SpKO brains showed a significant increase in the diameter of myelinated axons relative to controls (Fig. 2 *A–E* and *SI Appendix*, Fig. S3 *B* and *C*). Furthermore, brains from β II-SpKO mice showed signs of axonal degeneration, including myelin sheath decompaction and vacuolization (Fig. 2*D*, red asterisks) and dense material in the cytosol, likely reflecting breakdown of cytoskeletal elements (Fig. 2*D*, blue asterisk), and the appearance of neuroinflammatory microglia (Fig. 2*B* and *SI Appendix*, Fig. S3*A*, red arrowhead). The g-ratio (ratio of diameters of inner axon and outer fiber) also increased, indicating thinning of the myelin sheet (Fig. 2*F*). These changes in myelin organization were predominantly found in large-diameter axons (Fig. 2*F*, red circle). Loss of myelin was not associated with a decrease in myelin basic protein expression (*SI Appendix*, Fig. S1 *B* and *C*). These observations suggest that β II-spectrin is required for maintenance of myelinated axons, which form and subsequently undergo degeneration. Interestingly, loss of α II-spectrin in myelinated peripheral sensory neurons in mice also resulted in preferential loss of larger-diameter axons as the mice aged, which was attributed to a protective role of α II-spectrin in larger-diameter axons (32). Neurons lacking α -adducin, which is also periodically incorporated into the periodic axonal structure (4), form an axonal actin periodic structure but develop axons with slightly enlarged actin rings and calibers (33).

β II-Spectrin Promotes Axonal Growth of Cultured Neurons. To investigate the contribution of β II-spectrin to axon growth, we evaluated cortical neurons cultured from wild-type (WT) and β II-SpKO E15.5 embryos. Loss of β II-spectrin did not affect axonal specification, including formation of the AIS. We detected normal clustering of AnkG at the AIS (*SI Appendix*, Fig. S4 *A*, asterisk and *SI Appendix*, Fig. *C*). The length of the AIS in both day in vitro (DIV)11 β II-SpKO cultured neurons (*SI Appendix*, Fig. S4 *A* and *B*) and in cortical slices of PND6 β II-SpKO brains (*SI Appendix*, Fig. S4 *C* and *D*) was also normal. Similarly, loss of β II-spectrin does not affect the normal polarized distribution of axonal versus dendritic proteins in neurons, as indicated by the normal localization of transferrin receptor (TfR-mCherry) and the trans-Golgi network protein 38 (TGN38-YFP) in dendrites of DIV11 β II-SpKO neurons (*SI Appendix*, Fig. S5 *A* and *B*), where they preferentially localized normally. Consistent with the reduction in axonal length observed in β II-SpKO brains, loss of β II-spectrin caused significant reduction in the length of the axons in cortical (Fig. 3 *A–C*) and hippocampal (*SI Appendix*, Fig. S6) cultured neurons. Rescue with mScarlet-tagged β II-spectrin resulted in restoration of axon length to WT values

(Fig. 3 *B* and *C*). β II-spectrin, thus, is not required for initial stages of axonal formation and neuronal polarization but rather promotes the growth and maintenance of axonal processes once they are established.

The Periodic Actin Ring Structure in Axons Is Disrupted in β II-Spectrin-Deficient Neurons. Loss of β II-spectrin in cultured neurons using acute short hairpin RNA knockdown disrupts the periodic actin arrangement (4, 5). We conducted 3D stochastic optical reconstruction microscopy (STORM) imaging (34, 35) of hippocampal neuronal cultures derived from E15.5 mice lacking neuronal β II-spectrin through embryonic development. Loss of β II-spectrin largely disrupted the 190-nm-based periodic distribution of actin and adducin in axons of DIV14 neurons (Fig. 4 *A–E*), with a higher level of residual periodic structures observed in neurons at DIV20 (Fig. 4 *F–J*), suggesting the possibility of a gradual, partial compensation. In general, the level of residual periodic structures in β II-spectrin-depleted neurons was higher in axon regions proximal to the soma. Loss of β II-spectrin also substantially disrupted the periodic structures in dendrites (*SI Appendix*, Fig. S7 *A–E*). These results confirm that β II-spectrin is important for the formation of the periodic membrane skeleton structure. The periodic membrane skeletal network likely confers stability to axons, and loss of this structure likely contributes to the axonal degeneration noted in β II-spectrin-deficient brains (Fig. 2).

Loss of β II-Spectrin Does Not Directly Affect Microtubule Dynamics. Recent studies using actin-destabilizing drugs suggest that cortical actin plays microtubule-maintaining roles in axons, and that the abundance of axonal actin rings directly correlates with axonal microtubule organization (36). To determine whether loss of β II-spectrin and the resulting disruption of the actin-spectrin periodic network impairs microtubule dynamics, we followed the localization of the microtubule plus end-associated protein EB3-tdTomato in live neurons. Movies from transfected DIV5 cortical neurons and kymograph analysis show a similar average number of EB3-tdT comets (Fig. S8*A*, asterisks) moving toward microtubule growing ends in control and β II-SpKO cells (*SI Appendix*, Fig. S8*B*). Similarly, we found that β II-SpKO axonal microtubules have normal velocity and run length of EB3-tdT comets (*SI Appendix*, Fig. S8 *C* and *D*). TEM cross-sections of non-degenerating CC axons from control and β II-SpKO mouse brains where we were able to detect microtubule structures did not reveal clear differences in microtubule arrangement or numbers between phenotypes, although reliable quantitative analysis was not possible because the microtubule bundles were not always unequivocally distinguishable in our samples (*SI Appendix*, Fig. S8*E*). These data

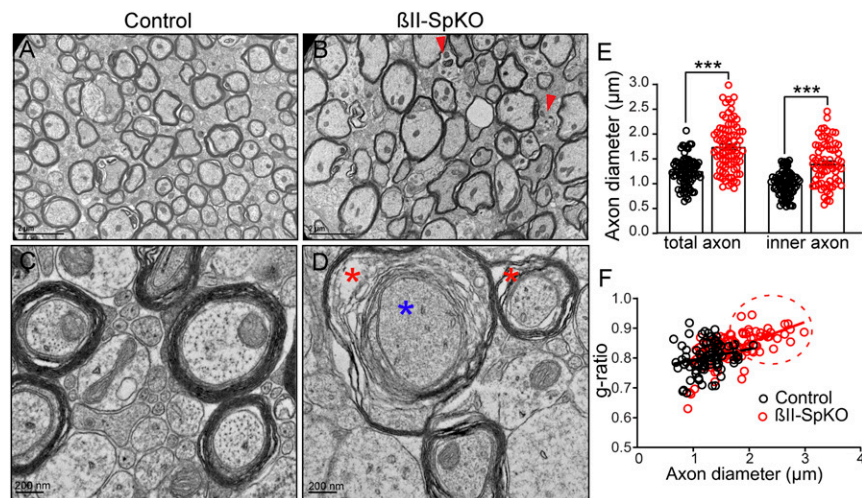


Fig. 2. β II-spectrin maintains the caliber and integrity of myelinated CC axons. (*A* and *B*) TEM images of cross-sections through the CC from β II-Sp^{fllox/fllox}+/+ (*A*, control) and β II-Sp^{fllox/fllox}/Nestin-Cre (*B*, β II-SpKO) PND25 mice ($n = 3$ mice). Red arrowheads indicate inflammatory microglia. (Scale bars, 2 μ m.) (*C* and *D*) Higher-magnification images of CC axons from control (*C*) and β II-SpKO (*D*) mice. Asterisks indicate degenerating axons with decompaction of the myelin sheath and vacuolization (red asterisks) and debris (blue asterisk). (Scale bars, 200 nm.) (*E*) Comparison of myelinated axon diameter from control ($n = 85$) and β II-SpKO ($n = 84$) CC axons. Data represent mean \pm SEM. Student's t test, *** $P < 0.001$. (*F*) g-ratio versus axon diameter from CC of PND25 control and β II-SpKO mice. Red dotted circle area represents a population of β II-SpKO large-diameter axons with decreased myelin content that is absent in control brains.

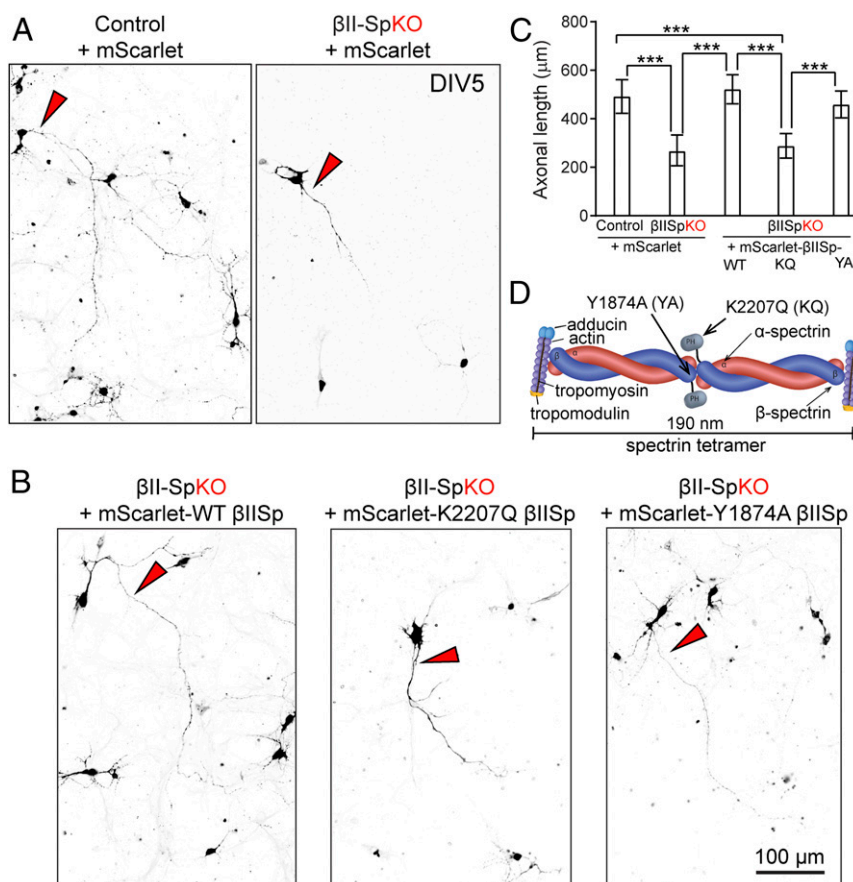


Fig. 3. β II-spectrin is required for axonal elongation of cultured neurons. (A) DIV5 control or β II-SpKO cortical neurons transfected at DIV0 with pmScarlet. (B) DIV5 β II-SpKO cortical neurons transfected at DIV0 with pmScarlet-tagged WT β II-Spectrin or indicated mutants. Red arrowheads in A and B indicate axons. (Scale bar, 100 μ m.) (C) Axonal length in control ($n = 12$), β II-SpKO ($n = 8$), and β II-SpKO neurons rescued with WT- β II-Sp ($n = 7$), K2207Q- β II-Sp ($n = 7$), and Y1874A- β II-Sp ($n = 7$) constructs. Data represent mean \pm SEM. One-way ANOVA with Dunnett's multiple comparisons test, $***P < 0.001$. (D) Schematic of spectrin tetramer organization in axons comprising dimers of α -spectrin and β -spectrin that cross-link short actin filaments capped by adducin and tropomodulin and determine their periodic distribution every 200 nm in axons. The Y1874A and K2207Q mutations respectively abrogate binding to ankyrin or PIP lipids.

suggest that loss of β II-spectrin does not directly alter microtubule dynamics and may not affect microtubule stability. However, other downstream progressive effects caused by loss of β II-spectrin, such as axonal degeneration (Fig. 2B and *SI Appendix*, Fig. S3, red arrowhead, and Fig. 2D, red asterisks), likely disrupt microtubule tracks (Fig. 2D, blue asterisk).

β II-Spectrin Promotes Bidirectional Multiorganelle Transport in Axons. Neurons rely on the transport of organelles to grow and maintain dendrites and axons (37, 38). Spectrins have been suggested to function as cargo adaptors in axonal transport, as discussed above. Therefore, we hypothesized that the reduced axonal length phenotypes observed in β II-SpKO brains and cultured neurons are associated with impaired axonal cargo motility. We tested this hypothesis by tracking the dynamics of yellow fluorescent protein (YFP)-tagged synaptophysin (a synaptic vesicle marker) in hippocampal neuron cultures from control and β II-SpKO brains using time-lapse video microscopy. Loss of β II-spectrin in neurons impaired bidirectional synaptic vesicle transport. We detected significant reduction in the percentage of motile cargo and overall flux in β II-SpKO axons (Fig. 5A–C). Similarly, we observed differences in both velocity and displacement of motile synaptic vesicles (Fig. 5D and E), with the most severe effects observed on anterograde motion. Loss of β II-spectrin also impaired the bidirectional flux of LAMP1-green fluorescent protein-positive endosome/lysosome vesicles (*SI Appendix*, Fig. S9A, D, and E) and caused significant deficits in

their run length and retrograde velocity (*SI Appendix*, Fig. S9B and C). The differences in the motility of synaptic versus endolysosomal cargo in β II-SpKO axons likely reflect known intrinsic bias in transport direction for each of these cargos and potential selective effects of loss of β II-spectrin on particular anterograde motors required for their motion. Nonetheless, deficits in transport of both synaptic cargo and lysosomes were rescued by expression of mScarlet-tagged WT β II-spectrin (Fig. 5A–E and *SI Appendix*, Fig. S9).

To evaluate whether β II-spectrin functions as a motor effector, we first evaluated its association with both anterograde and retrograde motor proteins via immunoprecipitation from whole-brain protein homogenates. We found that, similar to α II-spectrin (12), β II-spectrin associates with KIF3A (Fig. 6D). In addition, β II-spectrin also coimmunoprecipitates with KIF1A and KIF5B as well as the p150^{Glued} subunit of the dynein complex (Fig. 6D). Second, we confirmed that β II-spectrin associates with intracellular membranes and found strong enrichment of β II-spectrin in cellular fractions containing synaptic vesicles isolated through subcellular fractionation of crude total brain homogenates (Fig. 6A, membrane). Strikingly, loss of β II-spectrin in β II-SpKO brains reduced the association of KIF1A, KIF5B, KIF3A, and the p150^{Glued} subunit of the dynein complex with intracellular membranes (Fig. 6A and B), while the total expression of these motors in brain postnuclear supernatant fractions was unchanged (Fig. 6C). Taken together, these data indicate that β II-spectrin associates with both motor protein and

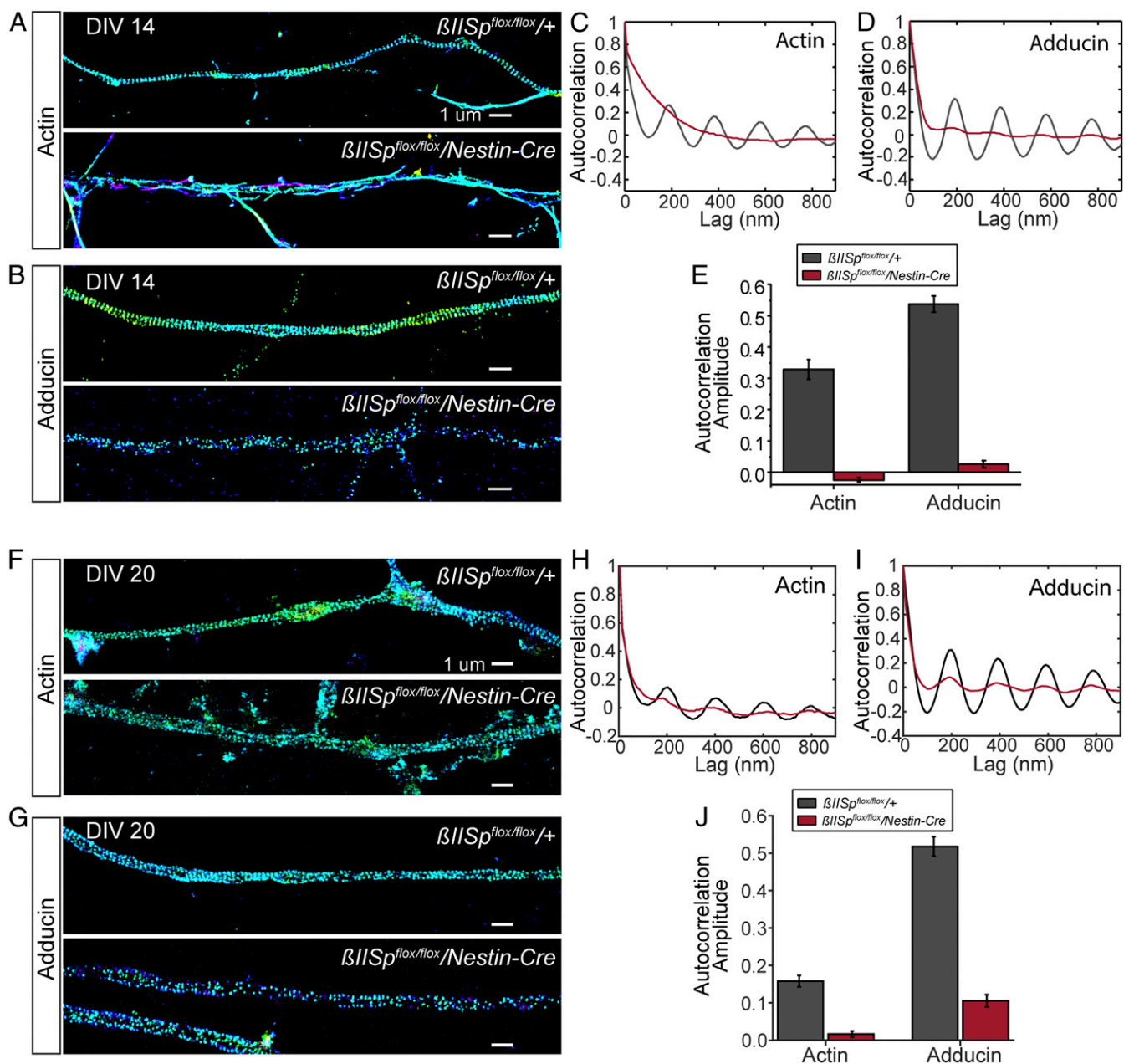


Fig. 4. β II-spectrin is important for the assembly of the actin–spectrin-based periodic membrane skeleton structure in axons. (A and B) Three-dimensional STORM images of actin (A) and adducin (B) in axons of DIV14 control and β II-SpKO neurons. (Scale bars, 1 μ m.) (C and D) Average autocorrelation functions of actin (C) and adducin (D) calculated from multiple, randomly selected axonal regions. Actin: $n = 96$ for β II $Sp^{fllox/fllox}/+$ samples and ($n = 128$) for β II $Sp^{fllox/fllox}/Nestin-Cre$ samples. Adducin: $n = 95$ for β II $Sp^{fllox/fllox}/+$ samples and ($n = 148$) for β II $Sp^{fllox/fllox}/Nestin-Cre$ samples. (E) Average autocorrelation amplitudes corresponding to actin (C) and adducin (D) periodicity functions in axons. (F and G) Three-dimensional STORM images of actin (F) and adducin (G) in axons of DIV20 control and β II-SpKO neurons. (Scale bars, 1 μ m.) (H and I) Average autocorrelation functions of actin (H) and adducin (I) calculated from multiple, randomly selected axonal regions. Actin: $n = 167$ for β II $Sp^{fllox/fllox}/+$ samples and $n = 199$ for β II $Sp^{fllox/fllox}/Nestin-Cre$ samples. Adducin: $n = 130$ for β II $Sp^{fllox/fllox}/+$ samples and $n = 130$ for β II $Sp^{fllox/fllox}/Nestin-Cre$ samples. (J) Average autocorrelation amplitudes corresponding to actin (H) and adducin (I) periodicity functions in axons. Data represent mean \pm SEM. The lower autocorrelation amplitude observed for actin in DIV20 versus DIV14 control neurons is likely because a stronger fixative (glutaraldehyde), which better preserve actin filaments, was used for DIV14 neurons, whereas DIV20 neurons were fixed with paraformaldehyde.

cargoes and may play a prominent role in coupling motor proteins to membranes.

β II-Spectrin Promotes Normal Organelle Axonal Transport in the Absence of Periodic Actin Rings. Loss of β II-spectrin largely disrupts the organization of periodic actin rings in axons imaged at DIV14 (Fig. 4) and results in impaired axonal transport in DIV7 neurons (Fig. 5 and *SI Appendix, Fig. S9*). Loss of

β II-spectrin also causes significant reduction in the length of the axons in cultured neurons, which is apparent as early as DIV5 (Fig. 3 B and C). Axons lacking β II-spectrin are prone to deteriorate, as evidenced by signs of degeneration observed in electron microscopy (EM) sections of callosal axons from PND25 β II-SpKO brains (Fig. 2 B and D and *SI Appendix, Fig. S3*). The onset of axonal degeneration in β II-spectrin KO neurons likely happens much earlier, and as noted above, and it is

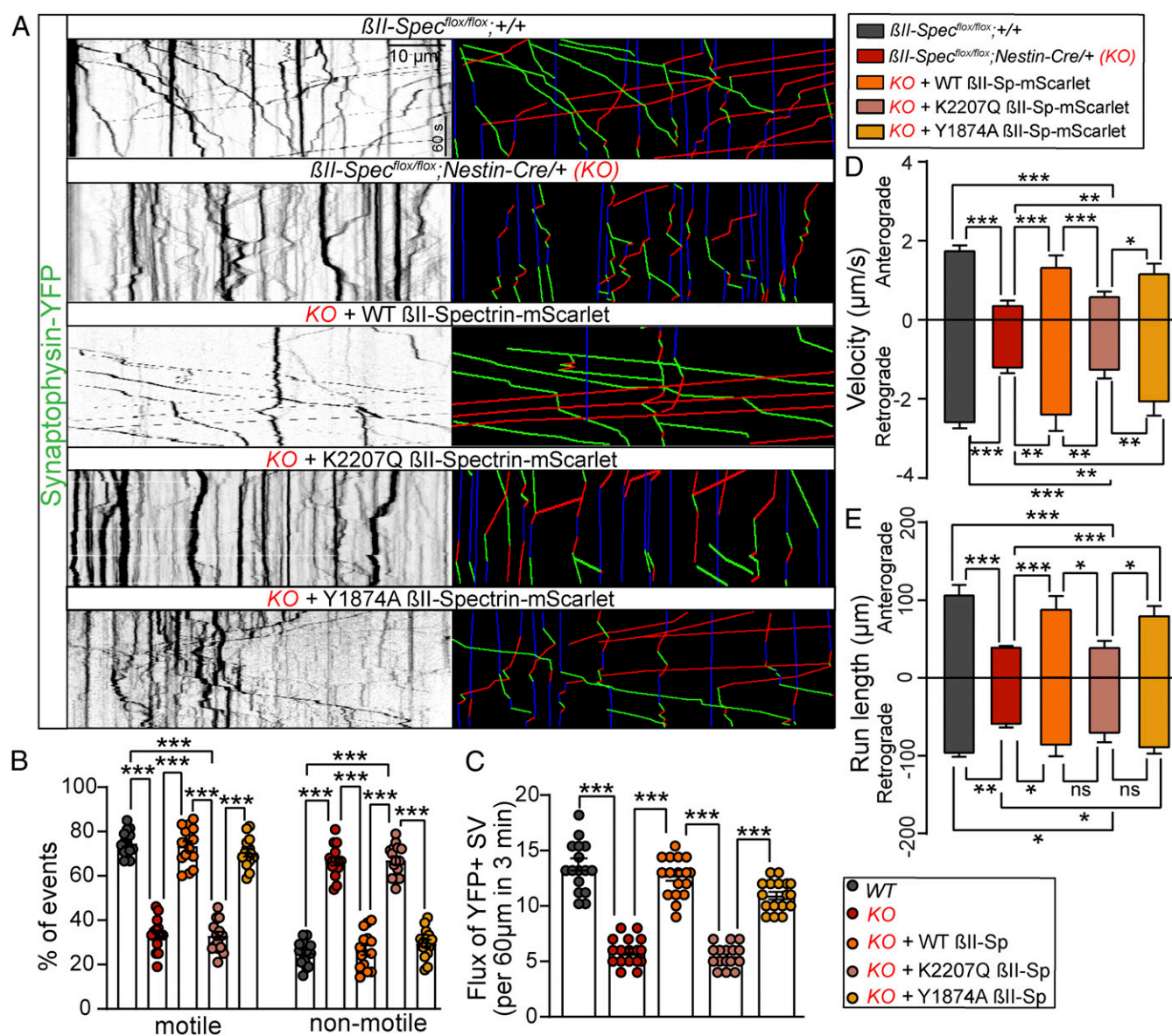


Fig. 5. β II-spectrin promotes bidirectional axonal transport. (A) Kymographs showing motility of YFP-tagged synaptophysin-positive synaptic vesicles in axons of DIV7 neurons from the indicated genotypes. Trajectories are shown in green for anterograde, red for retrograde, and blue for static vesicles. (Scale bar, 10 μm and 60 s.) (B–E) Relative number of motile and nonmotile synaptic vesicles (B), quantification of flux (C), anterograde and retrograde velocity (D), and run length (E) of synaptic vesicles for each genotype and rescue in $\beta II-Sp^{flox/flox}; +/+$ ($n = 15$), $\beta II-Sp^{flox/flox}; Nestin-Cre$ (KO) ($n = 17$), KO + WT $\beta II-Sp$ -mScarlet ($n = 12$), KO + K2207Q $\beta II-Sp$ -mScarlet ($n = 11$), and KO + Y1874A $\beta II-Sp$ -mScarlet ($n = 10$) axons of DIV7 neurons. Data represent mean \pm SEM. One-way ANOVA with Dunnett's multiple comparisons test, *** $P < 0.001$, ** $P < 0.01$, * $P < 0.05$. ns, not significant.

expected to cause disruptions in the transport machinery. Thus, an important question is whether the transport defects observed in β II-spectrin KO maturing neurons may be indirectly caused by the deficits in axonal outgrowth. To address this question, we tracked the dynamics of YFP-tagged synaptophysin in DIV2 control and β II-SpKO cortical neuronal cultures, when differences in axonal length are not very pronounced (Fig. 7A, B, and E). Like in DIV7 β II-spectrin KO neurons, bidirectional synaptic vesicle transport is already impaired in DIV2 axons lacking β II-spectrin, as reflected by the significant reduction in number of motile vesicles and overall flux (Fig. 7D, G, and H). These results indicate that the deficits in axonal transport caused by loss β II-spectrin precede overt deficits in axonal outgrowth and degeneration.

As shown above, disruption of the axonal actin–spectrin periodic structure caused by loss of β II-spectrin does not directly

affect microtubule dynamics. However, it is not clear whether the presence of the axonal actin–spectrin periodic structure is a prerequisite for the enablement of normal axonal transport. To begin to answer this question, we took advantage of the proximal-to-distal axon, progressive formation of the actin–spectrin periodic structure (39) to correlate organelle transport with actin rings along the axon. As previously reported (39), we detected an actin–spectrin periodic structure in the proximal (Fig. 7A, A1) but not in the distal (Fig. 7A, A2) axonal region of DIV2 control neurons, which is consistent with the rate of propagation of the periodic structure along the axons (39). As expected, actin–spectrin periodicity was lacking along the length of DIV2 β II-spectrin KO axons (Fig. 7B, B1 and B2). We imaged synaptic vesicle dynamics in both the proximal and the distal axon regions of control neurons and noticed no difference in their motilities or flux between those regions (Fig. 7C, G, and

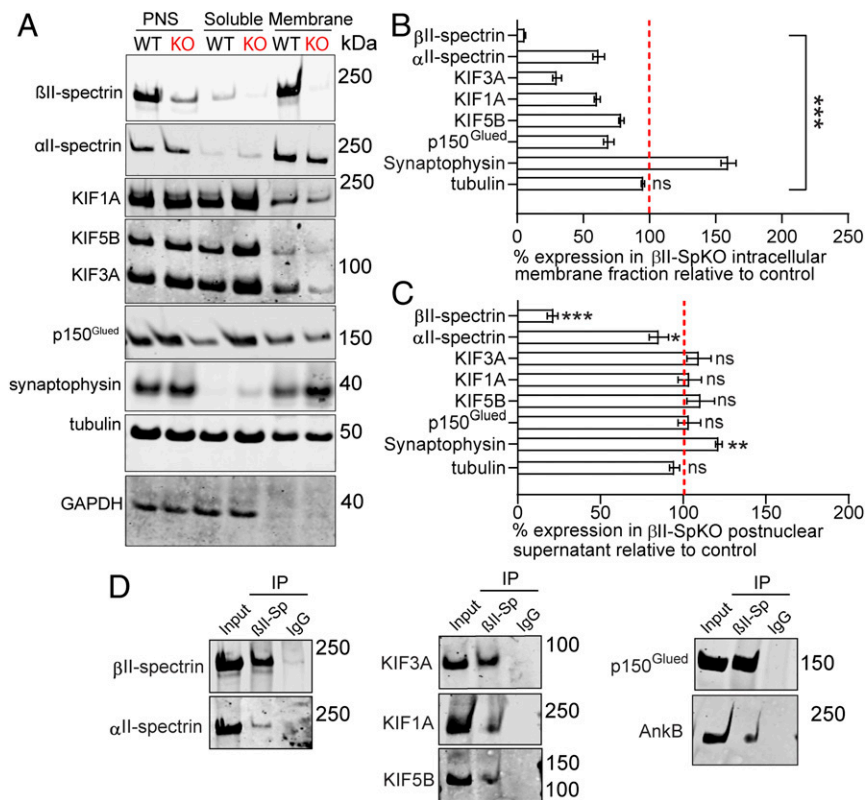


Fig. 6. β II-spectrin forms protein complexes with motor proteins and is required for their association with intracellular membranes. (A–C) Immunoblots (A) and corresponding quantification of protein levels of β II-spectrin, α II-spectrin, dynactin (p150^{Glued}), KIF1A, KIF3A, KIF5B, and synaptophysin in intracellular membrane (B) and postnuclear supernatant fractions (C) evaluated after subcellular fractionation of brain homogenates. Synaptophysin is a fractionation control. Tubulin and GAPDH are loading controls. Data represent mean \pm SEM. One-way ANOVA with Dunnett's multiple comparisons test, *** P < 0.001, ** P < 0.01, * P < 0.05. ns, not significant. (D) Western blot of the input and elution fraction from brain homogenates immunoprecipitated with a rabbit β II-spectrin antibody or isotype IgG control and immunoblotted with antibodies specific for β II-spectrin, α II-spectrin, AnkB, p150^{Glued}, KIF1A, KIF3A, and KIF5B.

H). As observed in older neurons, synaptic vesicle dynamics in DIV2 β II-spectrin KO were deficient relative to control neurons, but like in control neurons, organelle dynamics were similar between the proximal and the distal axon regions (Fig. 7 D, G, and H). These observations suggest that assembly of the periodic actin-spectrin skeleton is not required to enable the intracellular transport machinery. Thus, β II-spectrin independently assembles the neuronal actin periodic structure and promotes axonal organelle transport.

β II-Spectrin and AnkB Independently Enable Axonal Organelle Transport.

The axonal transport and growth deficits of β II-SpKO neurons and the loss of long-range axonal connectivity in β II-SpKO mouse brains resemble phenotypes observed upon loss of AnkB (14, 40); 220-kDa AnkB promotes axonal organelle transport via dual interactions with PI3P phosphoinositide lipids on membranes and the p62/dynactin4 subunit of the dynein/dynactin motor complex (14). Therefore, we asked whether β II-spectrin requires AnkB binding to promote axonal transport and growth. To our surprise, we found that mScarlet-tagged Y1874A β II-spectrin, which is unable to bind AnkB (Fig. 3D) (41), rescued both synaptic vesicle transport (Fig. 5 A–E) and axonal length (Fig. 3 B and C) deficits in neurons lacking β II-spectrin. The strong effects of loss of β II-spectrin on anterograde cargo motion, which are not observed in AnkB-deficient neurons, together with results from our rescue experiments, suggest that β II-spectrin promotes axonal transport independently of AnkB.

Previous studies using reconstituted motility assays using squid axon's cytoplasmic material and liposomes coated with acidic phospholipids showed that a dominant-negative construct containing the pleckstrin homology (PH) domain of β II-spectrin affected organelle and liposome retrograde motility, suggesting that β -spectrin's association with membrane lipids is required for dynein/dynactin-driven axonal transport (42). The β II-spectrin PH domain interacts with PI(4,5)P, PI(3,4)P2, and PI(3,4,5)P3 lipids through a highly conserved binding pocket (43). Mutation of a basic PH domain residue (K2207Q) (Fig. 3D) eliminated binding of

Drosophila β -spectrin as well as mammalian β II-spectrin to these phosphoinositides (41, 44). Thus, we next evaluated synaptophysin–YFP dynamics in β II-SpKO neurons transfected with mScarlet-tagged K2207Q β II-spectrin and found that this mutant failed to restore both axonal transport (Fig. 5 A–E) and axon length (Fig. 3 B and C). These data indicate that β II-spectrin binding to PIP lipids is required for its functions in axonal transport and growth.

Finally, we employed a genetic approach to determine whether β II-spectrin and AnkB promote axonal transport through independent pathways. If this were the case, dual loss of these proteins would cause a more severe impairment of transport dynamics than their individual deficits. To test this prediction, we developed mice bearing homozygous copies of both β II-spectrin and AnkB floxed alleles (*AnkB*^{lox/flox}; *β II-Spec*^{lox/flox}) and crossed them to Nestin-Cre to eliminate both proteins in all neuronal precursors. While individual KO of either AnkB or β II-spectrin in the brains causes postnatal lethality, their combined loss results in early embryonic death, with no *AnkB*^{lox/flox}; *β II-Spec*^{lox/flox}; Nestin-Cre/+ animals recovered. However, mice with total loss of one of these two genes and 50% reduction in the other (*AnkB*^{lox/flox}; *β II-Spec*^{lox/+}; Nestin-Cre/+ or *AnkB*^{lox/+}; *β II-Spec*^{lox/flox}; Nestin-Cre/+) showed earlier lethality and more severe morphological deficits and motor impairments than either each of the single gene KO or the double-heterozygous mice (Fig. 8 A and B). To overcome the embryonic lethality associated with double KO of neuronal AnkB and β II-spectrin, we cultured hippocampal neurons from embryonic *AnkB*^{lox/flox}; *β II-Spec*^{lox/flox} animals and cotransfected them at plating (DIV0) with synaptophysin–YFP to assess synaptic vesicles dynamics, and with either control pBFP or pBFP-Cre plasmids. Kymograph analysis from time-lapse movies of synaptophysin–YFP in axons reveal a dramatic reduction in motile synaptic cargo (red and green lines) in Cre-BFP-expressing *AnkB*^{lox/flox}; *β II-Spec*^{lox/flox} axons (Fig. 8 C, F, and G). Simultaneous loss of AnkB and β II-spectrin in neurons almost entirely stalled synaptic vesicle transport. The few synaptophysin–YFP-positive vesicles that retained motility exhibited further reductions in both anterograde and retrograde

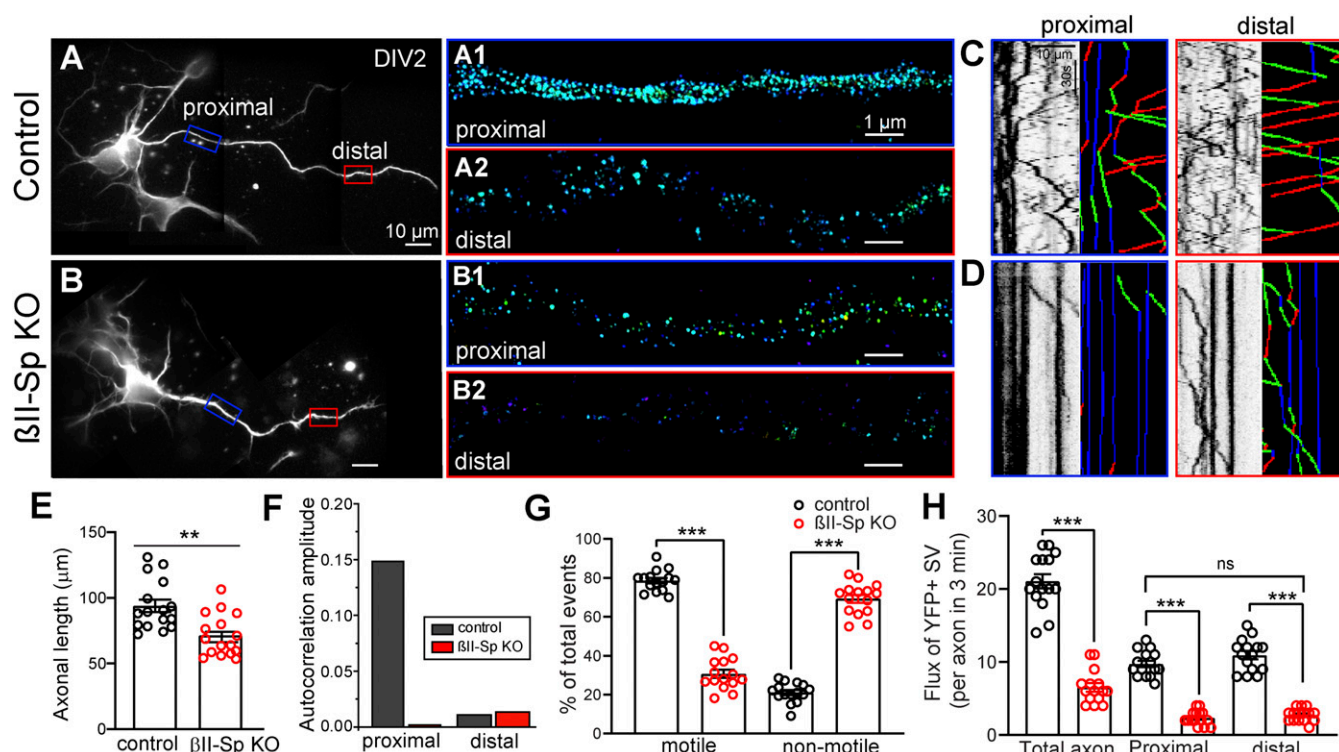


Fig. 7. Axonal transport deficits in β II-spectrin KO neurons appear early in axonal development. Normal axonal transport precedes the assembly of the actin-spectrin periodic skeleton. (A and B) DIV2 control (A) and β II-SpKO (B) cortical neurons labeled with β -tubulin. Rectangles demark proximal (blue) and distal (red) regions of the axon. (Scale bars, 10 μ m.) Three-dimensional STORM images of β II-spectrin in the proximal (control, A1 and β II-SpKO, B1) and the distal (control, A2 and β II-SpKO, B2) axonal regions of DIV2 cortical neurons. (Scale bars, 1 μ m.) (C and D) Kymographs showing the mobility of YFP-tagged synaptophysin-positive synaptic vesicles in the proximal (blue outlines) and distal (red outlines) axon regions in control (C) and β II-SpKO (D) cortical neurons. Trajectories are shown in green for anterograde, red for retrograde, and blue for static vesicles. (Scale bar, 10 μ m and 30 s.) (E) Axonal length in control ($n = 16$) and β II-SpKO ($n = 16$) DIV2 cortical neurons. (F) Average autocorrelation amplitude of β II-spectrin calculated from multiple, randomly selected cortical neurons. Axon proximal ($n = 23$) and distal ($n = 40$) regions for β II-Sp^{fllox/fllox} samples. Axon proximal ($n = 39$) and distal ($n = 37$) regions for β II-Sp^{fllox/fllox}/Nestin-Cre samples. (G) Relative number of motile synaptic vesicles in DIV2 β II-Sp^{fllox/fllox} (+) ($n = 21$) and β II-Sp^{fllox/fllox}/Nestin-Cre (KO) ($n = 23$) axons. (H) Quantification of flux of synaptic vesicles along the total length, the proximal, and the distal regions of the axons of in DIV2 β II-Sp^{fllox/fllox} (+) ($n = 21$), β II-Sp^{fllox/fllox}/Nestin-Cre (KO) ($n = 23$) cortical neurons. Data represent mean \pm SEM. Student's *t* test, *** $P < 0.001$, ** $P < 0.01$. ns, not significant.

velocity and run length parameters (Fig. 8 D and E). That dual loss of these transport facilitators causes a more severe impairment of transport dynamics than their individual deficits supports the conclusion that AnkB and β II-spectrin promote axonal transport of synaptic cargo through independent pathways.

Discussion

In this study we present direct *in vivo* evidence that β II-spectrin multitasks in mouse brain, with important roles in both establishing the membrane-associated periodic skeleton as well as enabling axonal transport of synaptic cargo. These biochemical activities, although involving distinct molecular partners and cellular locations, have a concerted effect in promoting both axon growth and stability. Our finding that β II-spectrin promotes axonal transport was foreshadowed by the identification of β -spectrin polypeptides comigrating with axonal cargo in the optic nerve nearly 40 y ago (11), and more recently by observations that α -II spectrin associates with and comigrates with kinesin (12), that spectrin is required to reconstitute dynein-dependent motility of liposomes (42), and that introduction of human SCA5 mutations into *Drosophila* β -spectrin impairs axonal organelle transport (13). Interestingly, AnkB, which is a β II-spectrin partner *in vitro* and *in vivo* (18, 45, 46), also promotes axonal organelle transport and is required for formation of long axon tracts in the developing brain (14). However, we find that β II-spectrin and AnkB promote axonal transport through distinct mechanisms. β II-spectrin lacking ankyrin-binding activity restores transport, while mice lacking both β II-spectrin and AnkB

exhibit more severely impaired axonal transport than those missing either protein alone. Instead of utilizing AnkB to associate with organelle membranes, β II-spectrin likely connects through interaction with PIP lipids, since mutation of its PH domain eliminating binding to PI(3,4)P2, PI(4,5)P2, and PI(3,4,5)P3 also prevents rescue of axonal transport and axon length. Strikingly, neurons lacking both β II-spectrin and AnkB exhibit a nearly complete loss of both anterograde and retrograde axonal transport of synaptic cargo, and brain-specific, double β II-spectrin and AnkB KO mice die embryonically.

The finding that axonal transport of synaptic cargo is nearly eliminated by dual loss of β II-spectrin and AnkB was unexpected, given the diversity of kinesins, kinesin binding-proteins, and partners for dynein/dynactin (47, 48). We, therefore, experimentally evaluated alternatives to roles of these proteins as transport facilitators whereby they enable the coupling of organelles to motors. For example, β II-spectrin and the periodic actin rings have been proposed to be required for microtubule stability (36). However, deficiency of β II-spectrin (SI Appendix, Fig. S8 A–D) or AnkB (14) had no effect on microtubule dynamics or polarity by direct imaging of EB3-tdT-labeled microtubule fast-growing ends. Additionally, as we showed, organelle transport proceeds normally through regions in the axon before the actin-spectrin periodic structure forms. Thus, we propose that β II-spectrin and AnkB are versatile motor cofactors that promote the long-range axonal transport of multiple organelles. The orchestration of the tethering and precise delivery of cargo along the axon are not fully understood. The prevailing view consists of a model where multiple activators, adaptors, and

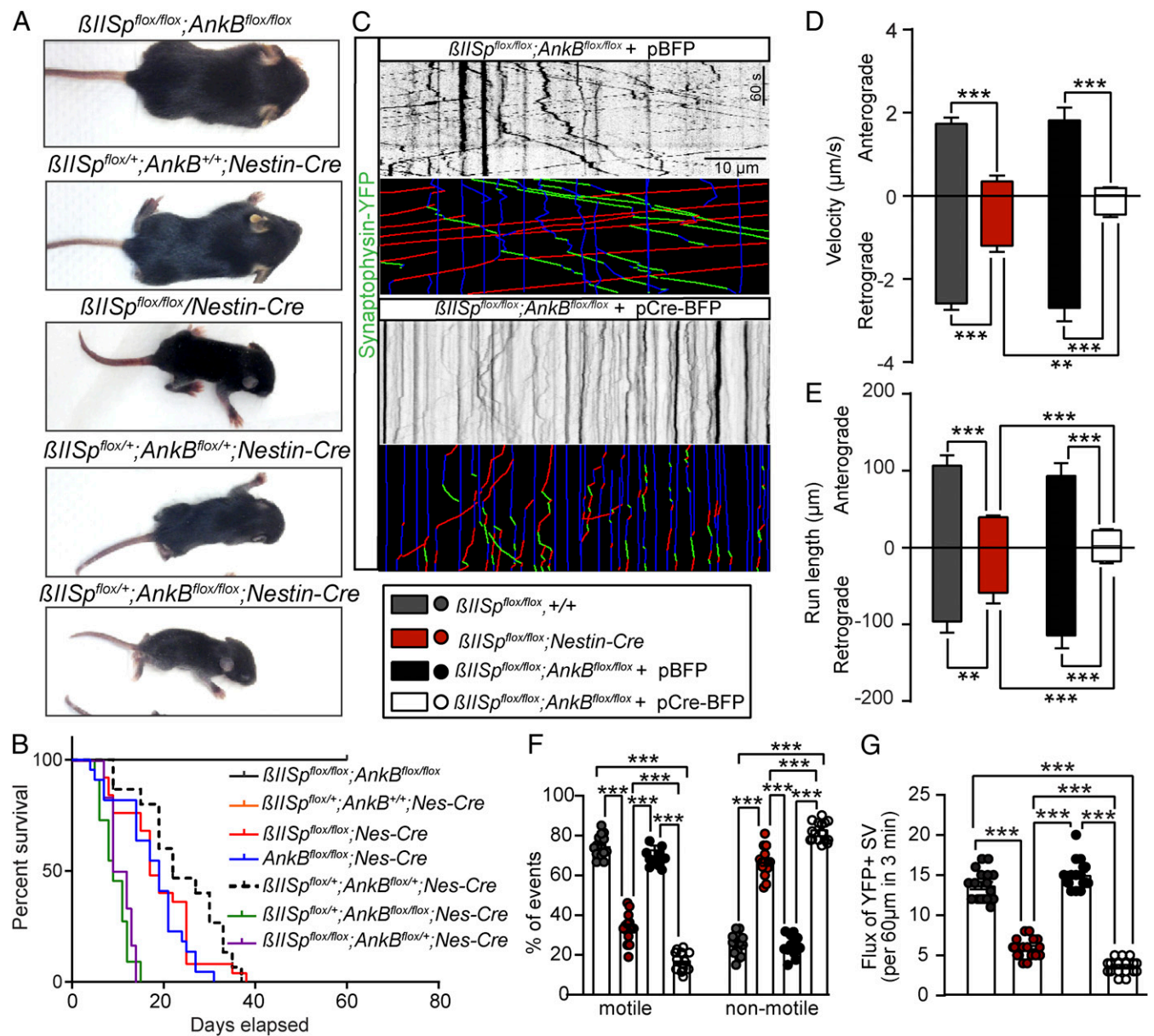


Fig. 8. Independent β II-spectrin and AnkB pathways promote axonal transport. (A) PND8 mice of the indicated genotypes. (B) Mice survival curves. A Mantel–Cox test was used to compare survival. (C) Kymographs showing the mobility of YFP-tagged synaptophysin-positive synaptic vesicles in axons from β II $Sp^{flx/flx}; AnkB^{flx/flx}$ neurons transfected with either pBFP or pCre-BFP plasmids. Trajectories are shown in green for anterograde, red for retrograde, and blue for static vesicles. (Scale bar, 10 μ m and 60 s.) (D–G) Anterograde and retrograde velocity (D), run length (E), relative number of motile and nonmotile synaptic vesicles (F), and quantification of flux (G) of synaptic vesicles in β II $Sp^{flx/flx}$ (anterograde $n = 76$, retrograde $n = 94$), β II $Sp^{flx/flx}/Nestin-Cre$ (anterograde $n = 55$, retrograde $n = 68$) axons, and in axons of β II $Sp^{flx/flx}; AnkB^{flx/flx}$ neurons transfected with BFP (anterograde $n = 75$, retrograde $n = 91$) or Cre-BFP (anterograde $n = 127$, retrograde $n = 119$) of DIV7 neurons. Data represent mean \pm SEM. One-way ANOVA with Dunnett’s multiple comparisons test, *** $P < 0.001$, ** $P < 0.01$.

cofactors facilitate the processivity and selective coupling of motors to specific membrane cargos (47, 49, 50). Our current data support β II-spectrin and AnkB as principal motor cofactors for at least synaptic cargo in axons. Going forward it will be critical to elucidate the full scope of organelles that depend on β II-spectrin for axonal transport and determine at a molecular level how β II-spectrin interacts with motor complexes.

Recent cryo-EM structures of dynein propose that its p150^{Glued} subunit is sterically less hindered than the Arp1 subunit (47). While Arp1 has been shown to interact with spectrin in vitro and under conditions of overexpression (18), the cryo-EM data support our results suggesting that β II-spectrin may also bind p150^{Glued}. Similarly, it is not clear whether β II-spectrin interacts with KIF3A, KIF5B, KIF1A, and perhaps other kinesins, directly or through

intermediary protein(s). Finally, it will be of interest to determine the roles of β -spectrins and AnkB in cellular domains in addition to axons, which are highly specialized for long-range transport. β -spectrin homologs are not evident in genomes of fungi or plants. Kinesins and dynein/dynactin thus were present nearly a billion years before spectrins. Therefore, interactions of spectrins with motor proteins evolved in the context of multicellularity, and likely will turn out to include features that are cell type- and/or cell domain-specific.

Materials and Methods

Detailed materials and methods, including statistical analysis, are described in *SI Appendix, Materials and Methods*. Experiments were performed in accordance with the guidelines for animal care of the institutional animal care

and use committee at Duke University and the University of North Carolina at Chapel Hill. All mice were housed at 22 °C ± 2 °C on a 12-h-light/12-h-dark cycle and fed regular chow and water for ad libitum consumption. For full details about the generation of the mice used in these studies and the EM, STORM, DTI, and intracellular transport experiments see *SI Appendix, Materials and Methods*.

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