Agrin regulates CLASP2-mediated capture of microtubules at the neuromuscular junction synaptic membrane

Nadine Schmidt,1 Sreya Basu,1 Stefan Sladecek,1 Sabrina Gatti,1 Jeffrey van Haren,2 Susan Treves,3,4 Jan Pielage,5 Niels Galjart,2 and Hans Rudolf Brenner1

1Department of Biomedicine, Institute of Physiology, University of Basel, CH-4056 Basel, Switzerland
2Department of Cell Biology, Erasmus Medical Center, 3015 GE Rotterdam, Netherlands
3Department of Anesthesia and 4Department of Biomedicine, Basel University Hospital, CH-4031 Basel, Switzerland
5Friedrich Miescher Institute for Biomedical Research, CH-4058 Basel, Switzerland

Agrin is the major factor mediating the neuronal regulation of postsynaptic structures at the vertebrate neuromuscular junction, but the details of how it orchestrates this unique three-dimensional structure remain unknown. Here, we show that agrin induces the formation of the dense network of microtubules in the subsynaptic cytoplasm and that this, in turn, regulates acetylcholine receptor insertion into the postsynaptic membrane. Agrin acted in part by locally activating phosphatidylinositol 3-kinase and inactivating GSK3β, which led to the local capturing of dynamic microtubules at agrin-induced acetylcholine receptor (AChR) clusters, mediated to a large extent by the microtubule plus-end tracking proteins CLASP2 and CLIP-170. Indeed, in the absence of CLASP2, microtubule plus ends at the subsynaptic muscle membrane, the density of synaptic AChRs, the size of AChR clusters, and the numbers of subsynaptic muscle nuclei with their selective gene expression programs were all reduced. Thus, the cascade linking agrin to CLASP2-mediated microtubule capturing at the synaptic membrane is essential for the maintenance of a normal neuromuscular phenotype.

Introduction

The function of the dynamic cytoskeleton in synapse formation and maintenance is poorly understood. At the neuromuscular junction (NMJ) the array of synaptic proteins such as the acetylcholine receptors (AChRs) is determined by the specific set of genes induced by the nerve selectively in the muscle nuclei underlying the synapse. However, although the nerve-induced synaptic gene expression program can explain the set of proteins expressed in the synaptic region, it does not account for their focal insertion into the postsynaptic muscle membrane where the density of, e.g., AChRs declines sharply from ~10,000/µm² to <5/µm² within a few micrometers of muscle fiber length. One possibility is a focal transport involving microtubules (MTs) oriented toward and captured at the subsynaptic muscle membrane. Indeed, the NMJ is associated with dense, subsynaptic networks of cortical actin filaments and microtubules (Jasmin et al., 1990; Ralston et al., 1999). Actin filaments are thought to be involved in anchoring AChRs along with other proteins that play a part in NMJ formation and maintenance (Dai et al., 2000; Borges and Ferns 2001), and they may be involved in MT capturing. However, the generation of the subsynaptic actin networks has not been investigated. Likewise, it is not known how MTs are organized, and whether and how they are involved in the formation of the postsynaptic membrane.

The behavior of MTs and their interactions with other intracellular components are largely regulated by MT plus-end tracking proteins (+TIPs), which specifically associate with the growing, or “plus” ends of the MTs (Galjart 2010). In motile fibroblasts, a paradigm for studying MT behavior in cultured cells (Akhmanova et al., 2001), a subset of MTs is oriented toward the leading edge and becomes selectively stabilized.

Abbreviations used in this paper: AChR; acetylcholine receptor; GSK3, glycogen synthase kinase-3; MT, microtubule; NMJ, neuromuscular junction; PI3-K, phosphatidylinositol 3-kinase; +TIP, plus-end tracking protein.
Subsynaptic MT induction can be mediated by agrin in vivo. Specifically, agrin secreted from muscle fibers upon intracellular injection of expression plasmid into nonsynaptic regions of innervated muscle fibers (Jones et al., 1997) induces ectopic AChR clusters, which have a MT network associated with them similar to that observed in vivo (Fig. 1 c). This suggests that the nerve normally induces the subsynaptic MTs, at least in part, by the secretion of agrin.

To examine the orientation of subsynaptic MTs at the NMJ, muscle fibers of knock-in mice expressing GFP-tagged CLIP-170 (Akhamanova et al., 2005) at the plus ends of the MTs were fixed in a way that preserves dynamic MTs (see Materials and methods). Labeling with an antibody against GFP showed CLIP-170 enriched at synaptic AChR clusters (Fig. 1 d). 3D reconstruction of NMJs from confocal image stacks showed that the plus ends of the MTs were preferentially colocalized with the synaptic AChR cluster. In contrast, in many instances they were completely absent from the synaptic gutter containing the nerve terminal (Fig. 1 d). These findings indicate that the plus ends of MTs probing the interior of the cell cortex become preferentially associated with synaptic AChR clusters.

In cultured cells, MT stabilization is promoted by several +TIPs, including CLASP2, which are directly or indirectly linked to the actin network (Galjart 2010). Cortical actin is highly enriched at the NMJ (Dai et al., 2000; Bruneau et al., 2008), and is thought to be involved in the anchoring of the AChRs in the subsynaptic muscle membrane. Immunolabeling of CLASP2 suggested that it was organized in a pattern similar to that of CLIP-170, with an enrichment at the crests of the synaptic folds (i.e., between troughs marked by AChR-rich lines [Fig. 2 a], as well as along the edge of the AChR cluster [Fig. 2 c]). Interestingly, the AChR cluster edge is where new AChRs are inserted into the synaptic membrane of developing NMJs (Kummer et al., 2004). Imaging NMJs after 3 d of denervation, i.e., when nerve terminals were degenerated, produced similar results (not depicted). Thus, CLIP-170 and CLASP2 immunoreactivities were postsynaptic.

Double stainings of CLIP-170 and CLASP2 at the synapse revealed significant colocalization of these proteins at the NMJ (Fig. 2 b), indicating that subsynaptic MT ends contain both +TIPs. However, MT capture and stabilization can still occur in the absence of either of these proteins as indicated by the synaptic MT networks at Clasp2−/− (Fig. 2 d) and at Clip1−/−;Clasp2−/−/NMJs (not depicted) visualized by Tyr-tubulin and Glu-tubulin stainings. Although no impairment of stable synaptic MTs could be resolved by tubulin stainings in either of these mutants due to variable staining conditions, MT capture as revealed by CLIP-170 staining was reduced by the deletion of CLASP2 (see Fig. 4). Taken together, these findings suggest that agrin/MuSK triggers downstream signals that capture MTs at synaptic AChR clusters by a process that includes CLIP-170 and CLASP2, and that this, in turn, promotes the insertion of AChRs into the synaptic membrane.

NMJs are impaired in Clasp2−/− mice

To test this hypothesis, we investigated NMJs in mice in which CLASP2 had been genetically deleted. The phenotype of NMJs in adult Clasp2−/− mice was abnormal in several ways.
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subsynaptic nuclei which, through their muscle activity–resistant expression of AChR subunit and other synaptic genes, contribute to the maintenance of the NMJ.

If both CLASP2 and CLIP-170 are involved in subsynaptic MT capturing, as is also suggested by their colocalization at the NMJ (Fig. 2 b), one would expect a similar NMJ phenotype in mice lacking CLIP-170 as in the CLASP2-deficient mice. We therefore measured synaptic AChR density in CLIP-170−/− mice, and found a reduction to 80% of control (Fig. 3 c), similar to what we observed in the Clasp2−/− mice. These results suggest that the CLASP2–CLIP-170 interaction that was documented in vitro serves to regulate synaptic function in vivo.

CLASP2 promotes attachment of MT plus-ends to synaptic membranes of mature NMJs

To test for a role of CLASP2 in MT stabilization at the NMJ, we compared the localization of MT plus-ends as visualized by...
Figure 2. Synaptic localization of CLASP2, Tyr-tubulin, and Glu-tubulin in wt and Clasp2−/− NMJs. (a) Staining of CLIP-170 and CLASP2 at NMJ of wild-type muscle. Like CLIP-170 (left, green), CLASP2-decorated MT plus-ends (right, green) are enriched at synaptic AChR clusters primarily at their edges and at the crests of synaptic folds (arrowheads; troughs of folds marked by the AChRs). Bar, 5 µm. Note that images in panel a are from the level of the synaptic folds, i.e., deeper in the synaptic gutter; in contrast, maximum intensity projections from optical slices taken at the level of the edge of the synaptic AChR cluster and thus of the synaptic gutter (such as in Fig. 1 d, top images) reveal pronounced CLIP-170 enrichment compared with adjacent perisynaptic membrane. This explains the apparent difference in CLIP-170 intensity along the edge of the clusters in the two images (Fig. 2 a vs. 1 d). (b) Double staining of CLIP-170 and CLASP2 at NMJ of wild-type muscle. Accumulation of both +TIPs at edge of synaptic AChR cluster (blue) and overlap of the two +TIPs (marked by asterisks) is consistent with cooperation between CLIP-170 and CLASP2 at MT plus-ends. Note that some of the CLIP-170 and CLASP2 staining does not overlap, which may reflect differences in antibody accessibility and staining, or, alternatively, independent functions of these proteins in the NMJ. For localization of CLIP-170 in Clasp2−/− NMJ, see Fig. 4. Bar, 2.5 µm. (c) Top: CLASP2 is enriched at edge of synaptic AChR cluster at wild-type NMJ. Bottom: absence of CLASP2 staining of NMJ from Clasp2−/− muscle shows specificity of antibody used. Bars, 5 µm. (d) Both Tyr-tubulin (top) and Glu-tubulin (bottom) are enriched at NMJs of Clasp2−/− muscle. Bars, 10 µm.
CLIP-170 labeling in wild-type and in Clasp2−/− mice. As described above, CLIP-170 was prominent at the edges of the synaptic AChR clusters at wild-type NMJs but was also present throughout the postsynaptic membrane in the primary synaptic fold. In Clasp2−/− mice the synaptic localization of CLIP-170 was reduced. This was most readily seen from the decrease of CLIP-170 immunoreactivity along the edges of the synaptic AChR clusters (Fig. 4 a).

Because the resolution of confocal microscopy is limited, in particular in the z-direction, some synapses were imaged by using structured illumination microscopy, which increases resolution both in x-y and in z-directions by a factor of 2 compared with conventional confocal microscopy (Gustafsson et al., 2008). In individual optical slices taken through synaptic AChR clusters oriented en face, AChR labeling appeared as contour lines following the synaptic membrane at that respective z-level. In wild-type animals, puncta of CLIP-170 immunoreactivity precisely followed these lines. This indicates that in wild-type mice the plus ends of the MTs were present exactly, within the limit of z-resolution, at the level of the synaptic membrane. In contrast, at Clasp2−/− synapses CLIP-170 puncta were largely absent from the AChR contour lines (Fig. 4 b). Thus, in the absence of CLASP2, the incidence of MT plus-ends at the level of the AChRs was markedly reduced. This observation implicates CLASP2 in the capturing of CLIP-170–labeled MT ends (and thus of MTs) at the agrin-induced AChR cluster.

Agrin phosphorylates GSK3β through PI3-K

Next we examined how agrin regulates CLASP2-mediated MT capturing. At the leading edge of motile fibroblasts CLASP2-mediated MT capturing is regulated by phosphatidylinositol 3-kinase (PI3-K) and glycogen synthase kinase-3 β (GSK3β; Akhmanova et al., 2001). In many instances, PI3-K activates AKT through its recruitment to inositol lipids and subsequent phosphorylation by PDK-1; AKT in turn phosphorylates GSK3β, which is active by default but is rendered inactive by phosphorylation on Serine 9 (Ser9). The inactivation of GSK3β triggered by PI3-K prevents phosphorylation of CLASP2 and, as a consequence, enhances its affinity to MT plus-ends (Akhmanova et al., 2005; Kumar et al., 2009; Watanabe et al., 2009). Blockade of number of nuclei underlying the AChR cluster (marked by arrowheads). Bar, 10 µm. Graphs show mean values of respective parameters ± SEM, n = 47 wild-type and 52 Clasp2−/− NMJs from 3 muscles of each genotype analyzed [*; P < 0.05; **; P < 0.01, two-sided t test]. (b) AChR density and AChR insertion rates are reduced in Clasp2−/− muscle. To determine AChR densities sternomastoid muscles were saturated with α-BTX–Alexa 594. Bar, 25 µm. Means ± SEM, n = 63 and 72 NMJs from 3 wild-type and 3 Clasp2−/− muscles analyzed [*; P < 0.05; **; P < 0.01, two-sided t test]. To estimate AChR insertion rates sternomastoid muscles were denervated to increase AChR turnover; 7 d later AChRs in superficial endplates were saturated with α-BTX–Alexa 488. After another 7 d (to allow AChR turnover), endplates were saturated with α-BTX–Alexa 594, and the average intensity of Alexa 488 fluorescence in Clasp2−/− NMJs was normalized to that in wild-type muscles. Bar, 25 µm. Means ± SEM, n = 28 wild-type and 25 Clasp2−/− endplates analyzed [*; P < 0.05; **; P < 0.01, two-sided t test]. (c) AChR density is reduced in Clip1−/−, Clip2−/− muscle. Analysis similar as in b. Given are means ± SEM, n = 30 wild-type and 31 Clip1−/−,Clip2−/− [*; P < 0.05; **; P < 0.01, two-sided t test]. Bar, 25 µm.
with agrin, which more closely resembles physiological agrin presentation (Fig. 5 a, right; for full blots see Fig. S2). Thus agrin, by activating PI3-K and AKT, phosphorylates (i.e., inactivates) GSK3β in cultured myotubes.

We next examined whether activated PI3-K and p-GSK3β are present at the NMJ in vivo. PI3-K activity at NMJs was assessed by measuring relative synaptic GFP fluorescence levels in fibers electroporated with expression constructs encoding either a wild-type or a mutant fragment of Bruton’s tyrosine kinase (BTK). Both constructs comprised a GFP tag coupled to the respective wild-type and mutant BTK pleckstrin homology domains, of which the wild type, but not the mutant, binds to PtdIns(3,4,5)P3 formed by active PI3-K (Várnai et al., 1999). Fig. 5 b shows that the synaptic GFP fluorescence signal was greatly reduced when fibers expressed the mutant BTK-GFP construct, indicating that PI3-K is activated at the NMJ.

PI3-K is known to inhibit agrin-induced AChR clustering in cultured myotubes (Nizhynska et al., 2007). We therefore tested the hypothesis that agrin, by activating PI3-K, induces phosphorylation of GSK3β at Ser9 (p-GSK3β).

To this end, C2C12 and primary wild-type myotubes were exposed to 5 nM of either agrin isoforms that are selectively expressed by motor neurons (neural agrin) or those expressed by nonneural tissues such as muscle. Of these, only neural agrin can induce synaptic membranes in muscle (Meier et al., 1998). Western blotting of lysates showed a pronounced but transient increase in both p-AKT and p-GSK3β within a few minutes in response selectively to neural, but not to nonneural agrin. Both AKT and GSK3β phosphorylation could be blocked by pretreating cells with PI3-K inhibitors (LY294002 or ZSTK474) or with AKT inhibitor (A6730; Fig. 5 a, left). Similar results were seen when myotubes were grown on a substrate impregnated with agrin, which more closely resembles physiological agrin presentation (Fig. 5 a, right; for full blots see Fig. S2). Thus agrin, by activating PI3-K and AKT, phosphorylates (i.e., inactivates) GSK3β in cultured myotubes.

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Figure 5. Agrin phosphorylates PI3-K, AKT, and GSK3β in muscle cells. (a) Neural, but not nonneural, agrin phosphorylates GSK3β via phosphorylation of PI3-K and AKT in cultured myotubes. Western blots of lysates from C2C12 myotubes to which soluble neural (5 nM) or nonneural (5 nM) agrin was added. Blots probed with antibodies as described in Materials and methods. Panels showing p-GSK3β and p-AKT are derived from same blot exposed for different times. Note that (1) AKT and GSK3β are not phosphorylated by nonneural agrin (top left panel, right lane); (2) their phosphorylation by neural agrin is inhibited by the PI3-K blocker LY294002 (50 µM) and the AKT inhibitor A6730 (0.5 µM, bottom left panel), respectively. Neural agrin also phosphorylates AKT and GSK3β when presented attached to culture substrate (top right panel, see Materials and methods). Full blots are shown in Fig. S1. Bar, 20 µm. (b) PI3-K is activated at the postsynaptic membrane of the NMJ. Soleus muscles of wild-type mice were electroporated with expression
Consistent with our hypothesis for the agrin-induced signaling cascade, we further observed by immunofluorescence that p-GSK3β was colocalized with the synaptic membrane of normal (Fig. 5 c; for specificity of the antibody used, see Fig. S2) and denervated NMJs (not depicted), excluding that the p-GSK3β immunoreactivity was presynaptic. p-GSK3β was not distributed evenly but appeared focally localized at the crests of the synaptic folds between the regions carrying the AChRs (Fig. 5 d), i.e., similar to CLIP-170 and CLASP2 (Fig. 2 a). When combined, these results indicate that agrin, by activating PI3-K, phosphorylates and thus inactivates GSK3β at the NMJ.

MTs are captured at agrin-induced AChR clusters by a process involving PI3-K and CLASP2
If agrin induces the trapping of plus ends of dynamic MTs at differentiating synaptic spots, it should be possible to visualize this in real time using GFP-tagged +TIPs. To mimic the stable presentation of agrin to the muscle at mature NMJs, we deposited neural agrin in small patches (20–60 µm in diameter) onto a laminin substrate. Myotubes that contact these agrin deposits form large AChR clusters that include stabilized AChRs and have associated with them a cluster of myonuclei that express the synapse-specific form of AChRs (Fig. 6 a; Jones et al., 1996). We therefore compared the dynamics of MTs in primary GFP-CLIP-170-expressing myotubes, inside and outside agrin-induced AChR clusters, using TIRF microscopy.

Analysis of individual frames of time-lapse movies (see Video 1) of wild-type myotubes suggested a much higher density of GFP-CLIP-170–labeled MT plus-ends inside the AChR clusters than outside (Fig. 6 b, center). However, maximum intensity projections (MIPs) of all 160 frames of the time-lapse imaging experiment showed that also outside the cluster many MT ends do reach the membrane (Fig. 6 b, right). The fact that inside the cluster the difference in MT plus-end density between single frames and MIPs is less pronounced than outside suggests that MT ends reaching the cell membrane inside the cluster are more likely to be captured than they are outside the cluster.

Analysis of changes in GFP-CLIP-170 labeling and distribution in distance versus time plots (so-called kymographs) revealed two types of behavior (Fig. 6 c). One type (here called type I) consisted of largely continuous movement of GFP-CLIP-170 “comets” during the time they remained within the penetration range of the evanescent wave. These comets had an average speed of 0.15 ± 0.01 µm/s (SEM, n = 73 tracks) and represent the continuous advance of the plus ends of growing MTs. In the other type of behavior (here called type II), the GFP-CLIP-170 fluorescence appeared stable over a period of 160 s or longer, with fluorescence flashing up at discrete spots along the track for tens of seconds before fading. Type II behavior suggests reduced freedom of GFP-CLIP-170 movement, and the capture of GFP-CLIP-170–labeled MT ends. Conspicuously, GFP-CLIP-170 dynamics was different inside and outside agrin-induced AChR clusters. Inside clusters, ~75% of GFP-CLIP-170 tracks observed were of type II, and the remaining 25% of type I. In contrast, this ratio was reversed outside receptor clusters, where only 25% were of type II (Fig. 6 f).

As suggested above for adult NMJs, MT stabilization at agrin-induced AChR clusters in cultured myotubes was mediated by PI3-K activation and CLASP2. Specifically, when MT dynamics were examined in primary GFP-CLIP-170 myotubes treated with the PI3-K inhibitor ZSTK474 (Fig. 6 d), or in myotubes derived from Clasp2−/−;GFP-Clip-170′/′ (i.e., knockout/knock-in) mice (Fig. 6 e; see Video 2), type I and II comets were equally common in both cluster and noncluster regions (Fig. 6 f), suggesting nonspecific MT plus-end capturing both outside and inside the clusters. These experiments demonstrate that agrin enables the capture of CLIP-170–coated MT ends through a process requiring CLASP2, and that this process depends on PI3-K activation.

CLIP-170 is increased at plus ends of MTs in the proximity of agrin-induced AChR clusters
The GFP fluorescence of MT plus-ends imaged inside agrin-induced AChR clusters in control wild-type myotubes appeared brighter by eye than those in the same myotubes outside the AChR cluster. This difference could be due to MTs coming closer to the internal myotube membrane inside clusters than outside and/or to an increased load of CLIP-170 molecules on MT tips induced by agrin. To examine these possibilities, myotubes were labeled for CLIP-170 and EB3, a member of the EB1-family of “core” +TIPs that is expressed in myotubes (Fig. 7). Myotubes were then imaged by confocal microscopy, and the intensity of CLIP-170 fluorescence on MT plus-ends, identified by their EB3 immunoreactivity, was assayed inside and outside AChR clusters by line scans. At MT plus-ends inside AChR clusters in control myotubes the intensity of CLIP labeling was significantly higher than outside clusters, whereas EB3 was not changed (Fig. 7 a). This suggests that pathways activated by agrin increase the affinity of CLIP-170 for MT plus-ends.

Next, we examined whether the increase in CLIP-170 load was mediated through PI3K and CLASP2 by repeating the
Figure 6. Capturing of dynamic MTs at agrin-induced AChR clusters in cultured myotubes is abolished by pharmacological inhibition of PI3-K and by genetic elimination of Clasp2. (a) Scheme illustrating focal impregnation of culture substrate with neural agrin. Transfected COS1 cells secrete and locally deposit neural agrin on a laminin substrate, and the cells are then lysed. Stable AChR clusters form where GFP-Clip-170ki/ki myotubes contact agrin deposits. (b) MT plus-ends decorated with GFP-CLIP-170 as observed by TIRF microscopy at 1-s intervals for 160 s. Shown are AChR cluster, surface reflective interference contrast (SRIC) image of same myotube, first frame and maximum intensity projection of all 160 images of the stack. Note higher density and brighter GFP signal at AChR cluster (marked in red). Bars, 10 µm. (c) Representative examples of kymograms of type I and type II comet behavior. See text for discussion of comet behavior and criteria for comet classification. (d) Same as in b, except that myotubes had been incubated for 90 min in ZSTK474 (5 µM) before GFP analysis. Note that “cluster”-specific comet behavior is abolished. Bars, 10 µm. (e) Same as in b, except that myotubes derived from Clasp2−/−;GFP-Clip-170ki/ki mice were used. Note that “cluster”-specific comet behavior is abolished. Bars, 10 µm. (f) Quantification of comet behavior inside and outside agrin-induced AChR clusters. Comets are grouped from kymograms into type I (mobile) and type II (stable) comets as described in the text, and the percentages of stable comets, both inside and outside agrin-induced AChR clusters, is given (means ± SEM, n = 4 myotubes for each condition; number of comets analyzed: wild type: 374; wild type, ZSTK474-treated: 342; Clasp2−/−: 323; *, P < 0.05; **, P < 0.01, two-sided t test).
above experiment in wild-type myotubes treated with ZSTK474 (Fig. 7 b) and in Clasp2\(^{−/−}\) mutant myotubes (not depicted). In both cases the enrichment of CLIP-170 was reduced to extrasynaptic levels (Fig. 7 c). Thus, in addition to CLASP2 deficiency, blockade of PI3-K inhibited CLIP-170 enrichment. These data strongly suggest that agrin leads to CLASP2-mediated MT capture and regulates the load of CLIP-170 at MT plus-ends, whereas EB3 remains unchanged; they provide further evidence for the in vivo relevance of a CLIP–CLASP interaction.

MTs are required for insertion of AChRs into agrin-induced AChR clusters in cultured myotubes in a manner dependent on PI3-K and CLASP2

Having found that CLASP2 deletion reduces the area and density of synaptic AChR clusters at NMJs in vivo, we examined the role of MT capturing in the maintenance of agrin-induced AChR clusters in cultured myotubes. In a first set of experiments, we asked whether MT integrity was essential for the maintenance of agrin-induced AChR clustering. Myotubes were treated with 10 µM nocodazole for 3 h, sufficient to depolymerize most MTs. At the end of the drug treatment, the AChR clusters were labeled with α-BTX, and their area measured. The clusters of AChR in nocodazole-treated myotubes were markedly (60%) smaller than those in untreated myotubes (Fig. 8 a). The density of AChRs was, however, unchanged. These data indicate that the number of AChRs in the clusters was decreased by nocodazole. This decrease might occur through reduced AChR insertion into, or accelerated AChR removal from, the clusters. To distinguish between these possibilities the experiment was repeated, but with AChR clusters stained at the beginning of the drug incubation period. With this protocol, AChR cluster size was similar in control and in nocodazole-treated myotubes (Fig. 8 b). Thus, once AChRs had been inserted into the cluster, their removal was not affected by nocodazole. These experiments thus demonstrate that nocodazole reduces AChR cluster size by impairment of AChR insertion, and further, that in these experiments, measuring AChR cluster size can be used to differentiate between AChR insertion and removal.

Finally, we tested the effects of inhibitors of PI3-K, AKT, and GSK3\(^{β}\), on AChR insertion. Using the criteria just described, inhibition of PI3-K and AKT decreased and inhibition of GSK3\(^{β}\) increased AChR insertion into AChR clusters (Fig. 8 a). Likewise, genetic deletion of Clasp2 (using Clasp2\(^{−/−}\) myotubes) reduced AChR insertion (Fig. 8 c). In these Clasp2\(^{−/−}\) myotubes, in contrast to wild-type myotubes, inhibition of PI3-K

Figure 7. The CLIP-170 load relative to EB3 at MT plus-ends is greater inside than outside AChR clusters and is reduced by PI3-K inhibition and by deletion of Clasp2. (a) Comets in AChR clusters in GFP-Clip-170\(^{ki/ki}\) myotubes revealed by immunolabeling of EB3 and of GFP, to visualize CLIP-170. Regions outlined by boxes within and outside AChR cluster are shown enlarged at bottom. Agrin-induced AChR clusters in blue, GFP-CLIP-170 in green, EB3 in red. Note increased load of CLIP-170 at comets in AChR cluster (arrowheads) compared with comets outside cluster lacking increased CLIP-170 (asterisks). Bars: (top) 10 µm; (bottom) 5 µm. (b) Same as panel a but after blockade of PI3-K with ZSTK474. Note reduction in CLIP-170 load in PI3-K–blocked compared with nonblocked myotubes. Bars: (top) 10 µm; (bottom) 5 µm. Asterisks same as in panel a. (c) Quantification of CLIP-170 staining on EB3-stained comets inside relative to that of comets outside AChR clusters. The elevated CLIP-170 load at agrin-induced AChR clusters observed in wild-type myotubes is abolished by PI3-K inhibition and by genetic deletion of Clasp2. EB3 staining is similar inside and outside AChR clusters. Percentages are means ± SEM from 84 to 245 comets analyzed in wild-type, ZSTK474-blocked, and Clasp2\(^{−/−}\) myotubes (4 cells each). *, P < 0.05; **, P < 0.01, two-sided t test.
Figure 8. Impairment of MT function by nocodazole, by inhibition of PI3-K and AKT, and by genetic deletion of CLASP2 impairs AChR insertion into agrin-induced AChR clusters in cultured myotubes. (a) Areas of agrin-induced AChR clusters in primary myotubes appear smaller when AChRs are stained after 3 h of treatment with the following drugs: nocodazole (10 µM); inhibitors of AKT (A6730, 0.5 µM); PI3-K (ZSTK474, 5 µM; only the effect of ZSTK474 is illustrated). However, the density of AChRs in clusters remains unchanged. Conversely, inhibiting GSK3 activity (CHIR99021, 0.1 µM) increases AChR cluster size. Bar, 30 µm. Means ± SEM. Total numbers of AChR clusters analyzed in two independent experiments: Control (wild type): 78; nocodazole: 67; A6730: 68; ZSTK474: 38; CHIR99021: 54 (*, P < 0.05; **, P < 0.01, two-sided t test). (b) In contrast, AChR clusters appear unchanged after 3 h of drug treatment when AChRs are stained at the time of drug addition; again, AChR densities in clusters remain unchanged. Bar, 60 µm. Means ± SEM. Total numbers of AChR clusters analyzed in two independent experiments: Control (wild type): 30; nocodazole: 46; ZSTK474: 27; CHIR99021: 21 (*, P < 0.05; **, P < 0.01, two-sided t test). (c) AChR clusters are smaller in Clasp2−/− than in wild-type muscle, and unlike in wild-type muscle (see panel a), cluster size is not affected by ZSTK474 (5 µM) or CHIR99021 (0.1 µM). AChRs were stained after drug treatment. Bar: 30 µm. Graphs give means ± SEM, 20–78 clusters analyzed from 2 cultures for each condition (*, P < 0.05; **, P < 0.01, two-sided t test). AChR clusters marked by arrowheads.
Here, we have used the NMJ of a GFP-CLIP-170 knock-in mouse mutant as a model for studying the role of MTs at the synapse. This model has two major advantages. First, synaptic differentiation and maintenance of the NMJ, including the regulation of a stable subsynaptic MT network, are controlled by one major organizer, agrin. In our in vitro paradigm, agrin was presented in a locally stable fashion, similar to that in the muscle fiber’s synaptic basal lamina in vivo. Second, the use of myotubes derived from GFP-CLIP-170 knock-in mice ensures normal expression levels of this GFP-tagged +TIP, which functions in vivo like nontagged CLIP-170 (Akhmanova et al., 2005). This allows us to observe GFP-CLIP-170–decorated dynamic MTs at agrin-induced AChR clusters under completely physiological conditions.

Discussion

The role of dynamic MTs in synaptic function is only beginning to be understood. It has only recently been reported that dynamic MTs enter dendritic spines transiently in a way that appears to be dependent on synaptic activity and correlated with LTP (Gu et al., 2008; Hu et al., 2008; Jaworski et al., 2009). However, neither the mechanisms that regulate MT entry into spines nor their role in modulating synaptic plasticity have been established.

(ZSTK474) did not reduce AChR insertion any further (Fig. 8 c), nor did inhibition of GSK3β with CHIR 99021 increase receptor insertion, indicating that PI3-K, p-GSK3β, and CLASP2 are part of the same signaling cascade. Taken together, these experiments are consistent with our observations at mature NMJs (Fig. 3 b) and strongly support our model (see Fig. 9) that agrin, acting through PI3-K and p-GSK3β, promotes MT capturing at the synapse (Fig. 4 b and Fig. 6), and that this enhances AChR insertion into agrin-induced AChR clusters.

Figure 9. Regulation and function of CLIP-170/CLASP2-dependent microtubule stabilization at the postsynaptic membrane of the neuromuscular junction. (a) Working model summarizing the role of neural agrin to capture MTs at the postsynaptic membrane of the NMJ. Agrin-induced MT capturing at AChR clusters is regulated through: [1] local GSK3β inactivation promoting the recruitment of unphosphorylated CLASP2 to MT plus-ends and lattice; [2] recruitment of CLIP-170 through CLIP/CLASP and CLIP/MT interactions; [3] local, PI3-K–dependent recruitment and immobilization of LL5β in the synaptic membrane. Molecules and their interactions inferred from work published in other systems rather than from present data in muscle cells are marked by broken symbols. (b) At developing NMJs, constitutively expressed synaptic AChRs are delivered to synaptic membrane by lateral diffusion in the plane of the cell membrane (arrows). At adult NMJs, MTs stabilized by mechanisms outlined in panel a ensure focal delivery of synaptic AChRs to the crests of synaptic folds. Cartoon in panel b is modified from Ruegg, 2005.

Here, we have used the NMJ of a GFP-CLIP-170 knock-in mouse mutant as a model for studying the role of MTs at the synapse. This model has two major advantages. First, subsynaptic differentiation and maintenance of the NMJ, including the regulation of a stable subsynaptic MT network, are controlled by one major organizer, agrin. In our in vitro paradigm, agrin was presented in a locally stable fashion, similar to that in the muscle fiber’s synaptic basal lamina in vivo. Second, the use of myotubes derived from GFP-CLIP-170 knock-in mice ensures normal expression levels of this GFP-tagged +TIP, which functions in vivo like nontagged CLIP-170 (Akhmanova et al., 2005). This allows us to observe GFP-CLIP-170–decorated dynamic MTs at agrin-induced AChR clusters under completely physiological conditions.

The classical function of agrin is to cluster AChRs (and other synaptic proteins) in the subsynaptic muscle membrane. This involves the induction of a scaffold for anchoring these proteins at high density to the subsynaptic actin cytoskeleton. The present experiments, both at NMJs in vivo and in cultured...
myotubes, now reveal an additional aspect of agrin function in the clustering process, i.e., in the insertion of the AChRs into the synaptic receptor cluster. AChR insertion into the synaptic muscle membrane requires MTs captured at the synaptic membrane via a pathway downstream of agrin regulating, through PI3-K and GSK3β, the CLASP2-mediated MT capture at the subsynaptic membrane. Thus, synaptic MT capturing at the NMJ might be regulated similarly as at the leading edge of migrating cells where, by inactivation of GSK3β, CLASP2 is dephosphorylated, which increases its association with MTs (Kumar et al., 2009; Watanabe et al., 2009).

MTs can be linked through nonphosphorylated CLASP2 to cortical sites in two nonmutually exclusive ways: (1) through the PIP3 sensor LL5β (Lansbergen et al., 2006; Hotta et al., 2010), and (2) through direct binding of CLASP2 to actin (Tsvetkov et al., 2007). In addition, as suggested by the increase of the CLIP load on MTs at AChR clusters, increased (de-phosphorylated) CLASP2 at the clusters might increase binding of CLIP-170 via direct CLASP2–CLIP interactions. Whether increased CLIP also participates in MT capturing at the NMJ, e.g., through the Rac effector IQGAP1 to actin (Fukata et al., 2002; Watanabe et al., 2004) or through potentiating further CLASP2 binding, is not known, but would be consistent with the reduced AChR density at Clip115−/--;Clip170−/− NMJs (Fig. 3 c).

Of the potential CLASP2 interactors, LL5β is expressed in the endplate membrane and is localized at the crests between the AChR-rich regions at the mouth of the folds (Kishi et al., 2005), i.e., where p-GSK3β, CLIP-170, and CLASP2 are enriched (Figs. 1, 2, and 4). Further, its depletion inhibits AChR clustering (Kishi et al., 2005). LL5β is thus a strong candidate for mediating the GSK3-β/CLASP2–dependent MT capturing observed here. Our working model for CLASP2-mediated aspects of synaptic MT capturing is illustrated in Fig. 9.

MT capturing at the NMJ might also be mediated via the +TIPs APC (Zumbrenn et al., 2001) and ACF7 (Wu et al., 2011). Of these, APC is enriched at the NMJ, and overexpression of a dominant-negative APC truncation mutant inhibits agrin-dependent AChR clustering (Kishi et al., 2005). LL5β is thus a strong candidate for mediating the GSK3-β/CLASP2–dependent MT capturing observed here. Our working model for CLASP2-mediated aspects of synaptic MT capturing is illustrated in Fig. 9.

The directed transport of synaptic components to the end-plate membrane as proposed here for CLASP2-mediated synaptic MT capturing may be more important at mature NMJs than during early stages of their development. Early in NMJ development, muscle fibers express high levels of synaptic proteins, including AChRs, along their entire surface as part of their developmental program. This allows the recruitment of proteins to the developing synapse by lateral diffusion in the plane of the cell membrane (Anderson and Cohen 1977; Flanagan-Steet et al., 2005). In contrast, at later stages of NMJ maturation, when nerve-induced electrical muscle activity has down-regulated extrasynaptic AChR expression, the supply of synaptic molecules from nonsynaptic membrane is no longer available. This necessitates the transport of AChRs and other synaptic proteins from the TGN below the synaptic membrane where synapse-specific gene expression is maintained by the agrin-induced transcription from the fundamental muscle nuclei underlying the synapse (Brenner et al., 1990). The mechanisms described here can explain the highly focal nature of this transport to the synaptic muscle membrane through mechanisms similar to those controlling polarity and directed migration in motile cells.

**Materials and methods**

**Animals**

Generation of the Clasp2 single knockout strain and of the Clip1;Clip2 double knockout mouse strain will be described elsewhere. In brief, the genes encoding CLIP-170 (Clip1 gene) and CLIP-115 (Clip2 gene) were targeted by homologous recombination in embryonic stem (ES) cells. In the case of the Clip1 gene we inserted a GFP-loxP-Neo-loxP cassette (where Neo indicates the neomycin resistance gene) into the exon containing the ATG translation initiation codon. This yielded the Clip1 knockout allele. Removal of the neomycin resistance gene in ES cells by Cre recombinase yielded the GFP-Clip170 knock-in [ki] allele (Akhmanova et al., 2005). In the case of the Clip2 gene we inserted a loxP-neo-loxP cassette at the 5′ end of the gene and a loxP-Puro-loxP-LacZ cassette (where Puro indicates the puromycin resistance gene, and LacZ the β-galactosidase gene) at the 3′ end of the gene. Cre-mediated recombination in ES cells yielded the Clip2 knockout allele (Hoogenraad et al., 2002). Single Clip1 and Clip2 knock-out and GFP-Clip170+ mice were obtained by germline transmission of the modified alleles in chimeric mice, which were in turn obtained by injecting ES cells carrying the modified alleles into recipient blastocysts. The Clip1 and Clip2 single knockout mice were crossed to generate the double knockout line.

For electroporation and in vivo stainings, animals were anesthetized with ketamine (87 mg per kg body weight) and xylazine (13 mg per kg body weight). Postoperative analgesia was by 4 injections of buprenorphine at 12h intervals. Mice were sacrificed with CO2. Animal handling was approved by the Cantonal Veterinary Office of Basel-Stadt.

Depending on experimental suitability, soleus (for electroporations), epitrochleo-anconeus (ETA; for optimal preservation of dynamic microtubules) or sternomastoideus (for estimates of AChR density from whole BTX-Alexa 488–stained NMJs) muscles were used.

**Chemicals and antibodies**

ZSTK474 (LC Laboratories) was used at 1 or at 5 µM on cultured myotubes analyzed biochemically or by imaging, respectively. All other reagents were applied in the following concentrations: nocodazole (10 µM; Sigma-Aldrich), LY294002 (50 µM; Sigma-Aldrich), A6730 (500 nM; Sigma-Aldrich), CHIR99021 (100 nM; Axon Medchem) α-protein phosphatase (2,000 U; New England Biolabs, Inc.), and rapamycin (100 nM; LC Laboratories). AChRs were labeled with α-BTX-Alexa 488, 594, or 647 (Invitrogen). GFP-CLIP-170 (by staining for GFP), EB1, and CLASP2 were visualized with antibodies from Invitrogen, BD, and Absea, respectively. Polyclonal antibodies against AKT, p-AKT (S473), GSK3β, pGSK3β (S9; no. 9336), and Vinculin (S161, S173) (BD) were used. Antibodies against α-BTX and α-AChR (3 μg) were obtained by immunostaining of muscle sections and 2.5 µg of antibodies was used for immunoblot analyses. ZSTK474 and LY294002 were custom made at Absea, using a bacterially expressed and purified GST-EB3 fusion protein as antigen, as described previously (Stepanova et al., 2003). Antibodies against AKT, p-AKT (S473), GSK3β, pGSK3β (S9; no. 9336), and Vinculin (S161, S173) (BD) were used.
and actin were purchased from Cell Signaling Technology. All other primary antibodies were obtained from Sigma-Aldrich (anti-tyrosinated α-tubulin, anti-neurofilament), Millipore (anti-detyrosinated α-tubulin), Santa Cruz Biotechnology, Inc. (anti-myosin), and Dako (anti-S100, Z0311). Secondary anti-
body were goat anti-rabbit, goat anti-mouse, or goat anti-rat antibodies conjugated to Alexa 488 (Invitrogen) as well as donkey anti-chicken Cy2 (Jackson ImmunoResearch Laboratories, Inc.). HRP-conjugated secondary antibodies were from Santa Cruz Biotechnology, Inc. Singer’s solution was obtained from Braun. Phosphatase inhibitors PCK 1 and 2 as well as Collage-
nase type 1A were purchased from Sigma-Aldrich, protease inhibitors and 
Fugene HD from Roche, and PDGF-BB from Peprotech. Rapamycin was a gift from M.N. Hall (Biozentrum, University of Basel, Basel, Switzerland).

Estimation of synaptic PI3-K activity
40 µg of pHBK-TGF or pHBK-R2BCGFP (Varnó et al., 1999) in 10 µl 0.9% NaCl was injected with a Hamilton syringe into the soleus muscle of anesthetized C57BL/6 mice (8–10 wk old). After suturing the skin, 8 pulses (20 ms, 1 Hz, 200 V/cm) were applied to the leg using an ECM 830 elec-
trotraction system. 10–14 d later, electroporated soleus muscles were dis-
sected, AChRs stained with 1 µg/ml α-BTX-Alexa 594 for 1 h at RT, fixed in
d 4% paraformaldehyde (PFA) for 2 h, placed in 30% sucrose/PBS over-
night, and frozen. Muscles were then embedded and sectioned at 12 µm thickness in a cryostat (CM 1950; Leica). Sections were mounted in Citifluor 
and a Dynal-MPC-L magnetic particle concentrator (Blanco-Bose et al.,
2011). Before proceeding, the sections were blocked with 5% donkey serum in PBS, washed 3x in PBS, and then incubated in the primary antibody for 2 h at RT. The sections were washed 3x in PBS and then incubated with HRP-conjugated secondary antibody (1:200 dilution) for 1 h at RT. After washing, the sections were incubated in DAB solution for 2–5 min to visualize the AChR staining. After Collagenase type 1A treatment (0.5%, 37°C, 15 min), myotube culture dishes and recovered ETA muscles were rapidly transferred to −80°C methanol for 2 h, then transferred to a 1:1 mixture of methanol and 4% PFA at −20°C for 1 h, followed by 30-min periods at 4°C and at RT. Final fixation was in 4% PFA (RT, 10 min). Teased bundles of 4–10 muscle fibers were then incubated at 4°C for 15 min in 100 mM glycine and perm-
ibilized in 20% normal goat serum (NGS)/PBS/2% Triton X-100 at RT for 2 h. In the case of cultured myotubes, glycine treatment was fol-
lowed by permeabilization in 20% NGS/PBS/0.5% Triton X-100 (RT, 1 h). Primary and secondary antibodies were diluted in 5% NGS/PBS/0.1% Triton X-100 and were applied at 4°C overnight (primaries) or at RT for 45 min (secondaries).

Quantification of GFP-CIPL-170 load on EB3 comets
The mean fluorescence of GFP-CIPL-170 and EB3, visualized with the re-
spective fluorescent antibodies, was determined in ImageJ (Plugin “Measure RGB”) in and outside of the AChR cluster area in nonnaturated, nonproc-
essed confocal images. The ratio of GFP-CIPL-170 and EB3 fluorescence 
served as a measure for the load of GFP-CIPL-170 on EB3 comets. For EB3 fluorescence quantitation, only the intensities of EB3 comets were taken into account.

Estimation of number of subsynaptic myonuclei
Soleus muscles from wild-type or clasp2−/− mice of similar weight were stained with 1 µg/ml of α-BTX-Alexa 594 in buffered 1L5 medium (RT, 1 h), fixed with 4% PFA for 2 h, and individual fibers were teased and permea-
bilized in 20% NGS and 2% Triton X-100. Staining with S100 was performed overnight at 4°C. After incubation with secondary antibody, fibers were mounted in Prolong Gold containing DAPI. Enface synapses were imaged in 1 µm stacks using an ACS APO 63x/1.3 NA objective at the SPE confocal microscope (DMI 4000B; Leica). For synapse area measure-
ments, synapses were outlined in the maximum intensity projection for each 
image manually and the area of the outlined region was calculated using 
ImageJ. Schwann cell nuclei marked by S100 staining were discounted from the total nuclear count as visualized by DAPI. For measurement of muscle fiber areas, muscles were treated and sectioned as described in Estimation of synaptic PI3-K activity. To delineate muscle fiber plasma mem-
branes, cryosections were postfixed for 5 min in 2% PFA, washed, stained with 
Wheat Germ Agglutinin Oregon Green 488 overnight, and mounted. The analysis software from Olympus was used to outline fiber recters and calculate 
cross-sectional area.

Structured illumination microscopy (SIM)
Fixed samples were imaged using the ELYRA S.1 structured illumination microscopy (Carl Zeiss). Images were acquired using a 63x/1.40 oil Plan Apochromat objective and an EMCCD camera (iXon 885; Andor Technol-
ogy). Stacks of images of NMJs en face were taken through the entire depth of the synapse at 0.125 µm. Image processing was performed using the Carl Zeiss Zen software to achieve a maximal resolution of 110 nm in
Dynamics of GFP-CLIP-170 comets were measured in myotubes derived from GFP-CLIP-170<sup>+/+</sup> mice. Agrin-induced AChR clusters were stained with α-BTX–Alexa 594 (1 µg/ml, 30 min), and after washing with prewarmed DME, cells were incubated at 37°C for 1 h before being mounted on a thermostat perfusion chamber. Cells were imaged at 37°C in Krebs Ringer's solution (140 mM NaCl, 5 mM KCl, 1 mM Mg<sup>2+</sup>, 2 mM Ca<sup>2+</sup>, 20 mM Hepes, 1 mM NaH<sub>2</sub>PO<sub>4</sub>, and 5.5 mM glucose) at pH 7.4. Online fluorescence images were acquired using an inverted TIRF microscope (TE2000; Nikon) equipped with an oil immersion CFI Plan Apochromat 100× TIRF objective (1.49 NA) and an electron multiplier CCD camera (91100-13, Hamamatsu Photonics; Treves et al., 2010). The focal plane corresponding to the coversglass/cell membrane contact was adjusted with a surface reflective interference contrast (SRIC) cube and maintained throughout the experiment with the help of a perfect focus system. GFP-CLIP-170 dynamics were imaged for 160 s with a rate of 1 frame per second (MetaMorph; Molecular Devices).

Quantitative analysis of GFP-CLIP-170 comet dynamics

Dynamics of GFP-CLIP-170 comets were analyzed inside and outside of agrin-induced AChR cluster with ImageJ (National Institutes of Health). The maxprojection of all acquired 160 frames of the live imaging experiment was used to mark all the comet traces within the imaged area. Kymographs (time–space plots) of all the comets visualized in the TIRF plane were generated with the Kymograph plugin for ImageJ from J. Rietdorf (Friedrich Miescher Institute, Basel, Switzerland) and A. Seitz (EMBL Heidelberg, Heidelberg, Germany). To study the effect of PI3-K signaling as well as CLASP2 on GFP-CLIP-170 comet dynamics, comets from myotubes either treated with ZSTK474 (37°C, 60–80 min) or deficient in CLASP2 were used for the analysis.

Quantitative analysis of AChR cluster size

An effect of the drugs indicated or of CLASP2 deficiency on AChR cluster size was determined by incubating the cells for 3 h at 37°C in the absence or presence of pharmacological inhibitors. Cells were then fixed (4% PFA) and stained with α-BTX–Alexa 594 (1 µg/ml, 30 min) to label agrin-induced AChR clusters. To elucidate AChR removal rates in culture, agrin-induced AChR clusters in primary cells were stained with α-BTX–Alexa 594 (1 µg/ml, 30 min) before the cells were incubated for 3 h at 37°C to allow removal of fluorescent receptors in the absence or presence of pharmacological inhibitors. Cells were then fixed (4% PFA) and imaged (THC Plan Apo 20x/0.70 objective; inverted microscope [DMI 6000B; Leica]; 1394 ORCA-ERA camera [Hamamatsu Photonics]; Veloci6.0.1 [PerkinElmer]). Cluster sizes after 3 h were determined by measuring the area of the remaining AChR fluorescence in ImageJ (National Institutes of Health).

Quantification of AChR density and insertion in vivo

To determine the density of AChRs in CLASP2-deficient and age-matched control littersmates, the sternomastoid muscles of anesthetized mice were exposed and bathed for 90 min in Ringer’s solution containing 5 µg/ml α-BTX–Alexa 594 (α dose that has previously been demonstrated to be sufficient to saturate all receptors; Bruneau et al., 2005). Mice were sacrificed, and muscles were re-anaesthetized, sternomastoid muscles were exposed, and newly inserted AChRs were stained with α-BTX–Alexa 594 (5 µg/ml, 90 min). Mice were then sacrificed. Fixation, mounting, and image acquisition, and quantification was done as described above.

Immunoblotting

Myotubes were lysed in 3x SDS-lysis buffer (150 mM Tris, pH 6.8, 300 mM DTT, 6% SDS, 0.2% bromphenol blue, and 30% glycerol) supplemented with phosphatase inhibitors (1:100), protease inhibitors (1/10 tablet per ml lysis buffer), and 100 mM DTT. Lysates were denatured at 95°C for 5 min and loaded on a 10% SDS-PAGE. Gels were transferred onto PVDF membranes and developed with ECL after incubation with primary and secondary antibodies.

Statistical analyses

Data are given as mean ± SEM. The Shapiro–Wilks test was used to test if datasets belong to a normally distributed population. If so, quantitative comparisons of numerical datasets were tested for statistical significance by means of the parametric, nonpaired, two-sided Student’s t test. Otherwise, the nonparametric, two-sided U-test (Mann & Whitney) was applied. One asterisk marks P values below 5% (P < 0.05), whereas two asterisks correspond to P values below 1% (P < 0.01).

Online supplemental material

Fig. S1 shows the full blots from which Fig. 5 A was extracted. They show the effect of neural vs. nonneural agrin on phosphorylation of GSK3β and AKT and the effects of AKT, mTOR, and PI3-K inhibitors. Fig. S2 shows the tests used to demonstrate the specificity of the anti-phospho-GSK3β antibody (no. 9336; Cell Signaling Technology), using myc-tagged GSK3β wt and 9A mutant transfected into COS cells. Video 1 shows capturing of GFP-CLIP-170 comets at agrin-induced AChR clusters in primary of GFP-CLIP-170<sup>+/+</sup> myotubes, using TIRF microscopy. Video 2 shows inhibition of capturing of GFP-CLIP-170 comets at agrin-induced AChR clusters in Clasp2<sup>−/−</sup>/GFP-CLIP-170<sup>+/+</sup> primary myotubes, using TIRF microscopy.

Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.201111130/DC1.

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References


of neuregulin receptors.


Figure S1. Full scans of blots shown in Fig. 5a. Sections irrelevant for the images shown in the figure are crossed out.
Figure S2. Antibody no. 9336 (Cell Signal Technology) specifically detects GSK3β-(Ser9)P. (a) Parallel Cos1 cell cultures were left untransfected or they were transfected with expression constructs coding for myc-GSK3β-Ser9 (wt) or for myc-GSK3β-Ser9Ala (S9A). Cultures were either stained with DAPI and anti-myc or anti-GSK3β-(Ser9)P (a), or lysates were subject to WB (b) with same antibodies and those indicated in (c). In untransfected control cultures both immunofluorescence and WB show minor myc-GSK3β-(Ser9)P immunoreactivity (green) localized in the Golgi adjacent to the nuclei (blue). Transfected cells visualized by myc immunoreactivity (red) show strong myc-GSK3β-(Ser9)P immunoreactivity (green), but only when transfected with constructs for myc-GSK3β-Ser9 (wt, due to autophosphorylation), but not with myc-GSK3β-Ser9Ala (S9A). Bars, 20 µm. (c) Full scans of blots shown in b. Sections irrelevant for the images shown in the figure are crossed out.
Video 1. **GFP-CLIP-170 comets are captured at agrin-induced AChR clusters.** Dynamics of GFP-CLIP-170 decorated MT plus ends in a myotube derived from a GFP-Clip-170<sup>ki/ki</sup> mouse. Note the immobilization and increased CLIP-170 loads within compared with outside of the agrin-induced AChR cluster (marked in Fig. 6 b). Frames are acquired over a period of 160 s (1 frame per second) in the TIRF plane [inverted TIRF microscope [TE2000; Nikon] equipped with an oil immersion HCX PL APO 100x TIRF objective [NA = 1.49]].

Video 2. **Deletion of CLASP2 impairs capturing of GFP-CLIP-170 comets at agrin-induced AChR clusters.** Dynamics of GFP-CLIP-170 decorated MT plus ends in a myotube derived from a GFP-Clip-170<sup>ki/ki</sup> mouse deficient in CLASP2. Note that deletion of CLASP2 abolishes the difference in MT plus-end dynamics within and outside the agrin-induced AChR cluster [marked in Fig. 6 e]. Image acquisition as in Video 1.