

Par3 regulates polarized convergence between APP and BACE1 in hippocampal neurons



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ABSTRACT

The convergence between amyloid precursor protein (APP) and its β -secretase β -site APP cleaving enzyme 1 (BACE1) is a prerequisite for the generation of β -amyloid peptide, a key pathogenic agent for Alzheimer's disease. Yet the underlying molecular mechanisms regulating their convergence remain unclear. Here, we show that the polarity protein partitioning-defective 3 (Par3) regulates the polarized convergence between APP and BACE1 in hippocampal neurons. Par3 forms a complex with BACE1 through its first PDZ domain, which is important for regulating BACE1 endosome-to-TGN trafficking. In the absence of Par3, there is an increase in the convergence between internalized APP and BACE1. In hippocampal neurons, loss of Par3 leads to increased APP and BACE1 convergence in axons but not in dendrites. This polarized convergence mainly occurs in retrograde or stalled axonal late endocytic organelles and is likely due to compartment-specific regulation of APP trafficking by Par3. Together, our data show a novel function for Par3 in regulating polarized convergence between APP and BACE1 in hippocampal neurons.

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1. Introduction

A key hallmark of Alzheimer's disease (AD) is the accumulation of β -amyloid ($A\beta$) plaques. $A\beta$ is generated by sequential cleavage of the amyloid precursor protein (APP) by β - and γ -secretases (Haass et al., 2012; Tang, 2005). The β -site APP cleaving enzyme 1 (BACE1) is the main β -secretase responsible for $A\beta$ generation, with the cleavage of APP by BACE1 being the rate-limiting step (Cole and Vassar, 2007; Thinakaran and Koo, 2008). Thus, physical convergence of APP and BACE1 is a prerequisite for $A\beta$ generation during AD progression (Das et al., 2013, 2016; Sun and Roy, 2018). However, mechanisms regulating the subcellular convergence between APP and BACE1 are unclear.

After biogenesis at the Golgi, both APP and BACE1 are delivered to the plasma membrane through secretory vesicles. At the plasma membrane, APP and BACE1 are internalized through different endocytic pathways. Although APP internalization is through a

classic clathrin-dependent pathway (Kyriazis et al., 2008), BACE1 endocytosis is clathrin-independent (Sannerud et al., 2011). In wild type hippocampal neurons under physiological conditions, APP and BACE1 are largely trafficked through different organelles, which helps preclude large amounts of $A\beta$ generation under normal conditions (Das et al., 2013). Yet, the pathological changes that occur in the AD brain that induce the convergence between APP and BACE1 remain unclear.

We recently showed that in the human AD brain, there is a significant loss of the polarity protein Par3 (Sun et al., 2016). In addition, we found that Par3 regulates the subcellular trafficking of both APP and BACE1 (Sun et al., 2016; Sun and Zhang, 2017). Par3 functions together with atypical PKC (aPKC) (Sun and Zhang, 2017), which forms a trimolecular complex with Par3 and another polarity protein Par6 (Joberty et al., 2000). To further understand the underlying molecular mechanisms by which Par3 regulates BACE1 trafficking and determine the effects of Par3 on APP/BACE1 convergence, we examined the domains of Par3 responsible for regulating BACE1 trafficking. Here, we show that Par3 forms a complex with BACE1 through its first PDZ domain. In the absence of Par3, we found a significant increase in the convergence of BACE1 with internalized APP. Interestingly, in cultured hippocampal neurons, loss of Par3 results in polarized convergence of BACE1 with internalized APP in axons but not in dendrites. Together, our studies show novel mechanisms for regulating the convergence between APP and BACE1.

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2. Materials and methods

2.1. Plasmids and reagents

pcDNA3-APP (human wild type APP695) was a generous gift from Dr Sangram S. Sisodia (University of Chicago). pcDNA3-GFP-APP was generated by inserting EGFP near the N-terminus of APP after the signal peptide at the KpnI site (Collin and Martens, 2006). mRFP-tagged human BACE1, GFP-tagged Rab7, myc-tagged human full length Par3b and Par3c, and shRNA constructs against luciferase or Par3 have been described previously (Sun et al., 2016; Sun and Zhang, 2017; Zhang and Macara, 2006). Myc-tagged Par3b and Par3c fragments were obtained from Drs. Syed Mukhtar Ahmed and Ian Macara at Vanderbilt University and have been described (Chen and Macara, 2005; Joberty et al., 2000; Zhang and Macara, 2006). Neuro2a cells (N2a) stably expressing wild type human APP695 was a generous gift from Dr Gopal Thinakaran (University of Chicago).

2.2. Hippocampal neuronal culture and transfection

Procedures involving animals have been approved by the Institutional Animal Care and Use Committee (IACUC) at the Rutgers Robert Wood Johnson Medical School (16–022). Primary hippocampal neurons were established from embryonic day 18 Sprague-Dawley rats as described previously (Bernard and Zhang, 2015; Sun et al., 2013; Wu et al., 2017). At DIV6–7, neurons were transfected using calcium phosphate precipitation as described previously (Sun et al., 2013).

2.3. Coimmunoprecipitation and Western blotting

Coimmunoprecipitation was performed as described previously (Sun and Zhang, 2017). Briefly, N2a cells were transfected with the indicated constructs, lysed, and incubated with myc 9E10 antibody followed by IgG sepharose beads preblocked with 5% BSA. Beads were washed 3 times with lysis buffer. Bound proteins were eluted by Laemmli sample buffer and analyzed by Western blotting. Primary antibodies used in Western blotting include mouse anti-FLAG antibody (1:2000, M2, Sigma-Aldrich, Cat. No. F1804), rabbit anti-FLAG (1:2000, Cell Signaling Technology, Cat. No. 2368), and mouse anti-Myc (9E10). Secondary antibodies include horseradish peroxidase-conjugated goat anti-mouse and anti-rabbit antibodies (1:10,000, Jackson ImmunoResearch).

2.4. Immunocytochemistry and live confocal imaging

Immunocytochemistry was performed as described previously (Sun and Zhang, 2017). Briefly, N2a cells were transfected with the indicated constructs and fixed in 4% paraformaldehyde with 4% sucrose in PBS. After permeabilization in 0.2% Triton X-100 and blocking with 5% goat serum, cells were incubated with primary antibodies followed by Alexa fluorophore-conjugated secondary antibodies. Primary antibodies used include mouse anti-APP (1:100, 6E10, Signet, Cat. No. SIG-39320), FLAG (mouse monoclonal, 1:500, Sigma-Aldrich, Cat. No. F1804; rabbit polyclonal, 1:100, Cell Signaling Technology, Cat. No. 2368), TGN38 (mouse monoclonal, 1:800, Thermo Scientific, Cat. No. MA3-063), and presenilin 1 (1:200, Millipore, Cat. No. MAB5232). Secondary antibodies used include Alexa Fluor 405, 488, or 594-conjugated goat anti-mouse or anti-rabbit secondary antibodies (1:500, Invitrogen).

For immunostaining of internalized APP, N2a cells stably expressing WT APP695 were transfected with BACE1-FLAG along with other indicated constructs. Seventy-two hours later, live cells were stained with 6E10 for 1 hour at 4 °C, and then incubated at 37 °C for 60 minutes. Hippocampal neurons were transfected with WT

APP695, BACE1-GFP, along with other indicated constructs. Seventy-two hours later live neurons were stained with 6E10 for 30 minutes at 4 °C, and then incubated at 37 °C for 60 minutes. The 60 minutes time point was chosen because our previous studies indicate that loss of Par3 increases APP trafficking to late endosomes at 60 minutes after internalization (Sun et al., 2016). The 6E10 antibody reacts with amino acid 1–16 of the A β region within the extracellular domain of APP and thus can be used for live labeling of surface APP to follow its internalization (Tang et al., 2015; Xiao et al., 2012). Cells were fixed in 4% paraformaldehyde with 4% sucrose in PBS for 15 minutes at room temperature, and then blocked by HRP-conjugated secondary antibody for 1 hour at room temperature. After washing by PBS, cells were permeabilized with 0.2% Triton X-100 in PBS for 5 minutes at room temperature and then immunostained with the indicated primary or secondary antibodies.

Confocal images were obtained using an Olympus FV1000MPE microscope with a 60 × water immersion objective (1.0 numerical aperture) with sequential-acquisition setting. For live cell imaging, neurons were transferred into a 37 °C heated chamber. Cells were visualized with 488 nm excitation for eGFP and 559 nm for mRFP. Time-lapse images were collected under 5% of laser intensity with a 100 μ m pinhole size. A total of 200–500 frames were captured through continuous scanning with a frame rate of around 0.8 frames per second. The stacks of images were imported into ImageJ. For tracing anterograde or retrograde movement of different vesicles in live neurons, single and separated axons were selected, and kymographs were generated in ImageJ with the Image Stabilizer and Multi Kymograph plug-ins. The height of kymographs represents time (200–500 frames), whereas the width represent the length (μ m) of the axon imaged. Vesicles movement were counted and quantified by unpaired Student's *t*-tests. A membranous organelle was classified as immobile if it was stationary for the entire recording period; a motile one was counted only if the displacement was at least 10 μ m. Colocalization of vesicles was quantified by ImageJ with the Colocalization Pipeline plug-in.

2.5. Statistics

All experiments were repeated at least 3 times. The number of biological variables examined and the significance levels are presented in each figure legend. Power analysis was used to determine sample size at 90% power and $\alpha = 0.05$. Two-tailed, unpaired Student's *t*-tests or one way ANOVA were used to calculate the *p* values, using GraphPad Prism. Error bars represent the S.E.M.

3. Results

3.1. Par3 regulates BACE1 trafficking through an interaction mediated by its first PDZ domain

Our recent studies show that Par3 regulates BACE1 trafficking through interacting with BACE1 and recruiting aPKC to promote phosphorylation of Ser498 at the C-terminus of BACE1 (Sun and Zhang, 2017). To further understand the interaction between Par3 and BACE1, we performed coimmunoprecipitation of BACE1 with different fragments of Par3. We found that N-terminal Par3 fragments containing the first PDZ domain were sufficient for mediating the interaction with BACE1, whereas fragments lacking the first PDZ domain did not interact with BACE1 (Fig. 1A and B). This suggests that the interaction of BACE1 with Par3 is mediated through the first PDZ domain of Par3.

Intriguingly, the BACE1 C-terminus does not contain any typical PDZ-binding motifs. Thus, to further confirm that the Par3-BACE1 interaction is through the first PDZ domain, we performed

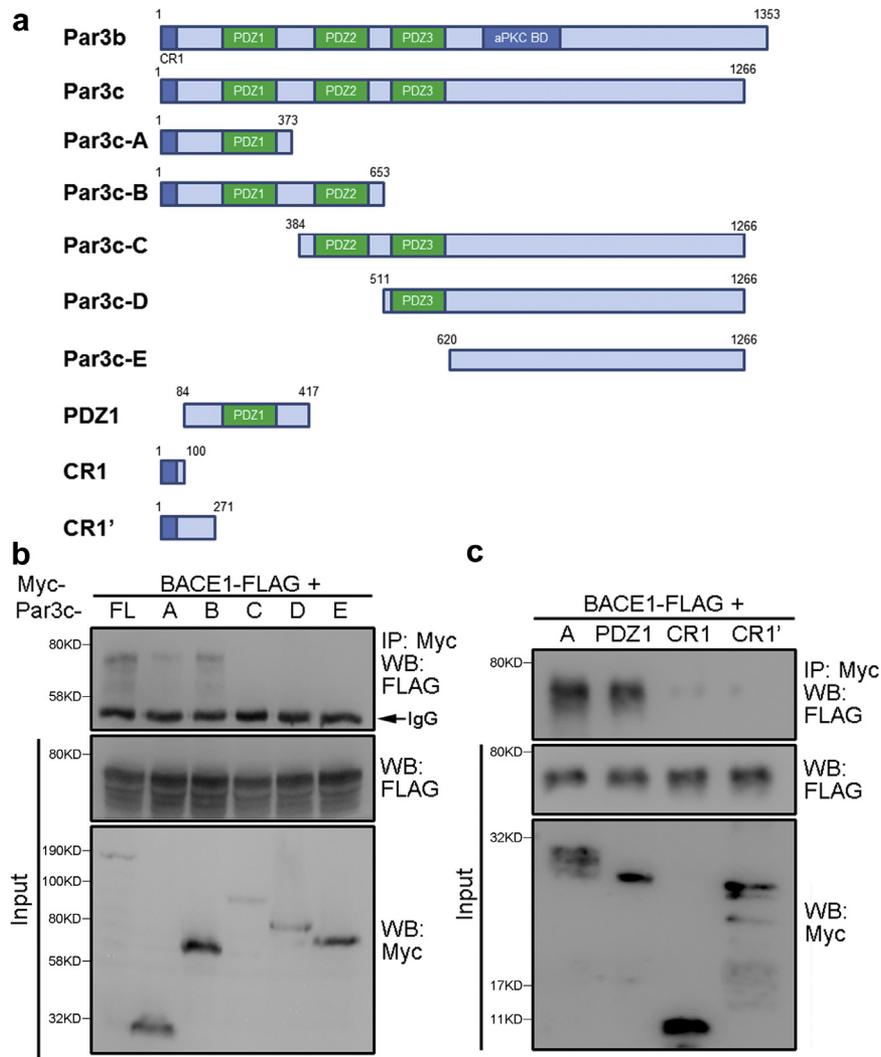


Fig. 1. The PDZ1 domain of Par3 mediates the interaction with BACE1. (A) Schematic diagram for the Par3 constructs. (B, C) N2a cells were transfected with the indicated constructs, lysed, and immunoprecipitated with the myc 9E10 antibody. Immunoprecipitated complexes were analyzed by Western blotting.

coimmunoprecipitation of BACE1 with smaller N-terminal Par3 fragments. As seen in Fig. 1C, Par3-PDZ1, which contains the intact PDZ1 domain, but missing the N-terminal CR1 domain, interacts with BACE1, whereas 2 other N-terminal fragments that contain the CR1 domain, but lack the PDZ1 domain, do not interact with BACE1. These results further confirm that the interaction between Par3 and BACE1 is through the first PDZ domain.

Next, we explored the domains of Par3 that mediate its effects on BACE1 trafficking. We recently showed that Par3 and aPKC promote BACE1 endosome-to-TGN trafficking in primary hippocampal neurons (Sun and Zhang, 2017). In the absence of Par3 or aPKC activity, BACE1 accumulates in the late endosome/lysosome pathway. This effect was observed in hippocampal neurons as well as N2a cells (Sun and Zhang, 2017). To see which domains of Par3 mediate this effect, we depleted endogenous Par3 in N2a cells and re-expressed different Par3 fragments. Both Par3b and Par3c isoforms can efficiently restore the TGN localization of BACE1 (Fig. 2). This suggests that direct interaction between Par3 and aPKC is not necessary, as Par3c lacks the aPKC-binding domain (Gao et al., 2002). This is further confirmed by the ability of the N-terminal Par3c-B fragment to rescue the TGN localization of BACE1. By contrast, a C-terminal fragment of Par3, which lacks the N-terminal

PDZ1 domain, cannot rescue BACE1 localization (Fig. 2). Together, these data suggest that Par3 interaction with BACE1 through the PDZ1 domain is necessary for its effect on BACE1 trafficking. Because direct interaction between Par3 and aPKC is not necessary for the effects on BACE1 trafficking (Fig. 2), it is likely that Par3 recruits aPKC through Par6, which forms direct interactions with both Par3 and aPKC (Joberty et al., 2000).

3.2. Depletion of Par3 increases colocalization of internalized APP with BACE1

The convergence of APP and BACE1 is a necessary step for A β generation. Because APP cleavage by BACE1 is believed to occur after it is internalized from the plasma membrane (Haass et al., 2012; Sun and Roy, 2018), we examined the colocalization of BACE1 with internalized APP by live antibody feeding in N2a cells stably expressing wild type human APP (Thinakaran et al., 1996). We found that the internalized APP show significantly increased convergence with BACE1 on Par3 depletion (Fig. 3A and C). In addition, we observed a significant increase in the colocalization of BACE1 and the γ -secretase subunit presenilin 1 (PS1) in Par3-depleted cells (Fig. 3B and D). Together, these results suggest that

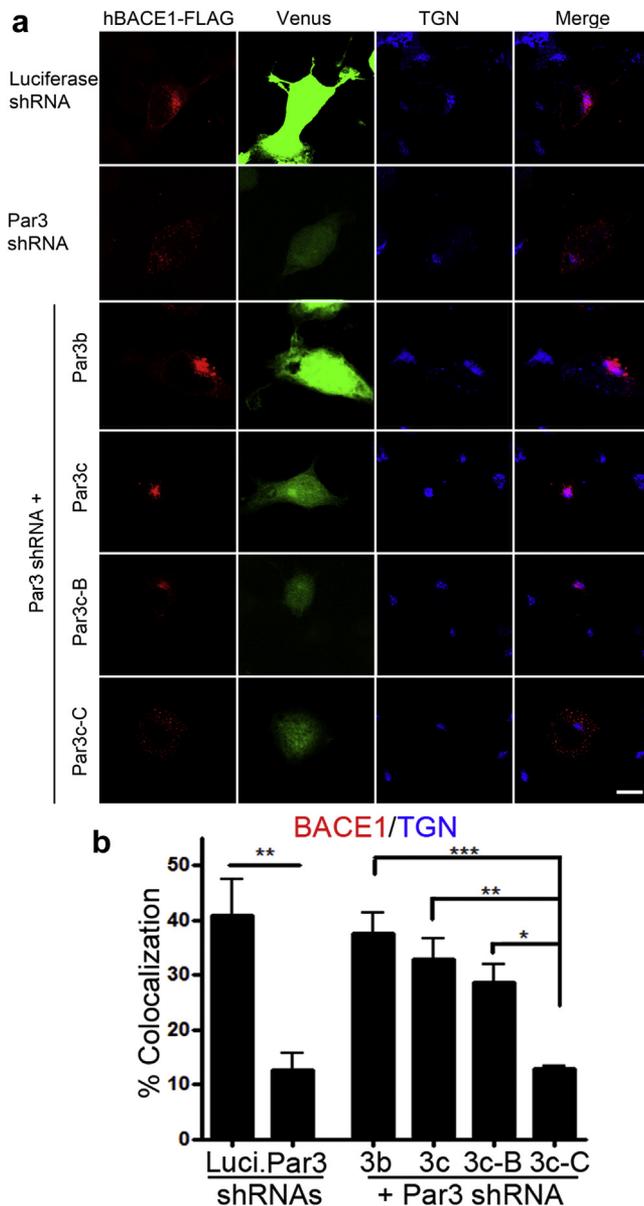


Fig. 2. The PDZ1 domain of Par3 is necessary for its effect on BACE1 trafficking. (A) N2a cells were transfected with indicated constructs together with hBACE1-FLAG. Cells were immunostained for FLAG (red) and TGN38 (blue). Venus (green) indicates positive cells. (B) Quantification for percentages of BACE1 colocalized with TGN. Data were expressed as Mean \pm SEM with Student's *t*-test, **p* < 0.05; ***p* < 0.01; ****p* < 0.001. N (cells) = 15 (Luciferase shRNA), 15 (Par3 shRNA), 15 (Par3b), 15 (Par3c), 19 (Par3c-B), 23 (Par3c-C), from 5 independent experiments. Scale bar: 10 μ m. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)

Par3 depletion increases the convergence of internalized APP with BACE1 and PS1, which will lead to increased A β generation.

3.3. Depletion of Par3 results in polarized convergence of APP and BACE1 in axons

Previous studies have shown that both APP and BACE1 trafficking are differentially regulated in dendrites and axons of neurons (Buggia-Prevot et al., 2013, 2014; Ubelmann et al., 2017). In addition, A β generation is known to be polarized with axonal pre-synaptic terminals being the preferred site (Buggia-Prevot et al.,

2013; Buxbaum et al., 1998; DeBoer et al., 2014; Lazarov et al., 2002; Lyckman et al., 1998; Niederst et al., 2015; Yu et al., 2018). Thus, we examined the cotrafficking of APP and BACE1 in dendrites versus axons of DIV11 hippocampal neurons. In axons under control conditions, we observed limited cotrafficking of APP and BACE1. However, in axons of Par3-depleted neurons, we observed a significant increase in the cotrafficking of BACE1 with APP in retrograde and stalled vesicles (Fig. 4A–D). Remarkably, we observed limited convergence between APP and BACE1 in dendrites with or without Par3 (Fig. 4E–H), suggesting that the effect of Par3 on APP and BACE1 convergence is restricted to axons.

Interestingly, the effects of Par3 on BACE1 trafficking are similar for both dendrites and axons, with loss of Par3 inhibiting BACE1 retrograde trafficking and increasing the percentage of stalled vesicles (Fig. 4C and G). By contrast, the effects of Par3 on APP trafficking are different in dendrites and axons. Although Par3 does not significantly affect APP retrograde trafficking in axons, loss of Par3 significantly inhibits APP retrograde trafficking in dendrites (Fig. 4B and F). Thus, the polarized regulation of APP/BACE1 convergence in axons versus dendrites is likely because of the differential effects of Par3 on axonal and dendritic APP retrograde trafficking. Taken together, these results suggest that Par3 depletion enhances polarized A β generation by increasing the convergence of APP and BACE1 within neuronal axons.

Finally, because the increase in APP and BACE1 convergence was observed in retrograde and immobile vesicles, we would like to determine the identity of these vesicles. Thus, we performed live imaging analysis in DIV 11 hippocampal neurons expressing BACE1-RFP and endosomal markers. In the absence of Par3, there is a significant increase in the percentage of BACE1 containing vesicles that are positive for Rab7, a marker for late endosomes (Yap et al., 2018). In particular, there is a significant increase in stalled vesicles that are dual-positive for BACE1 and Rab7 (Fig. 5). No significant changes were observed in the colocalization of BACE1 and Rab11, a recycling endosome marker (Supplementary Information, Figure S1) (Buggia-Prevot et al., 2014).

To confirm that BACE1 is converging with internalized APP in the axons, we performed live antibody feeding in hippocampal neurons. Similar to the results in N2a cells, we found that loss of Par3 significantly increased the percentage of internalized APP that converges with BACE1 in the axons (Fig. 6). Taken together, these data suggest that in the absence of Par3, there is an increase in BACE1 trafficking in late endocytic organelles where it converges with internalized APP.

4. Discussion

The convergence of APP and BACE1 is a rate-limiting step for the generation of A β in AD pathogenesis. Yet, the mechanisms regulating APP/BACE1 convergence in the subcellular trafficking pathways remain unclear. Here, we reveal mechanistic insight into the role of Par3 in regulating BACE1 trafficking and APP/BACE1 convergence. We show that Par3 regulates BACE1 trafficking by forming a complex with BACE1 through its first PDZ domain. In addition, we show that loss of Par3 leads to the convergence of internalized APP with BACE1 and PS1. Live imaging experiments in primary hippocampal neurons reveal that loss of Par3 leads to APP/BACE1 convergence during endosomal trafficking in axons but not in dendrites of hippocampal neurons.

Interestingly, the C-terminus of BACE1 does not contain any known PDZ-binding motifs, which raises the question regarding the nature of Par3-BACE1 interaction. It is possible that the interaction between Par3 and BACE1 is not direct, but rather through another binding partner of Par3, such as Par6, which binds PDZ1 of Par3 (Joberty et al., 2000). Alternatively, BACE1 may form a complex with

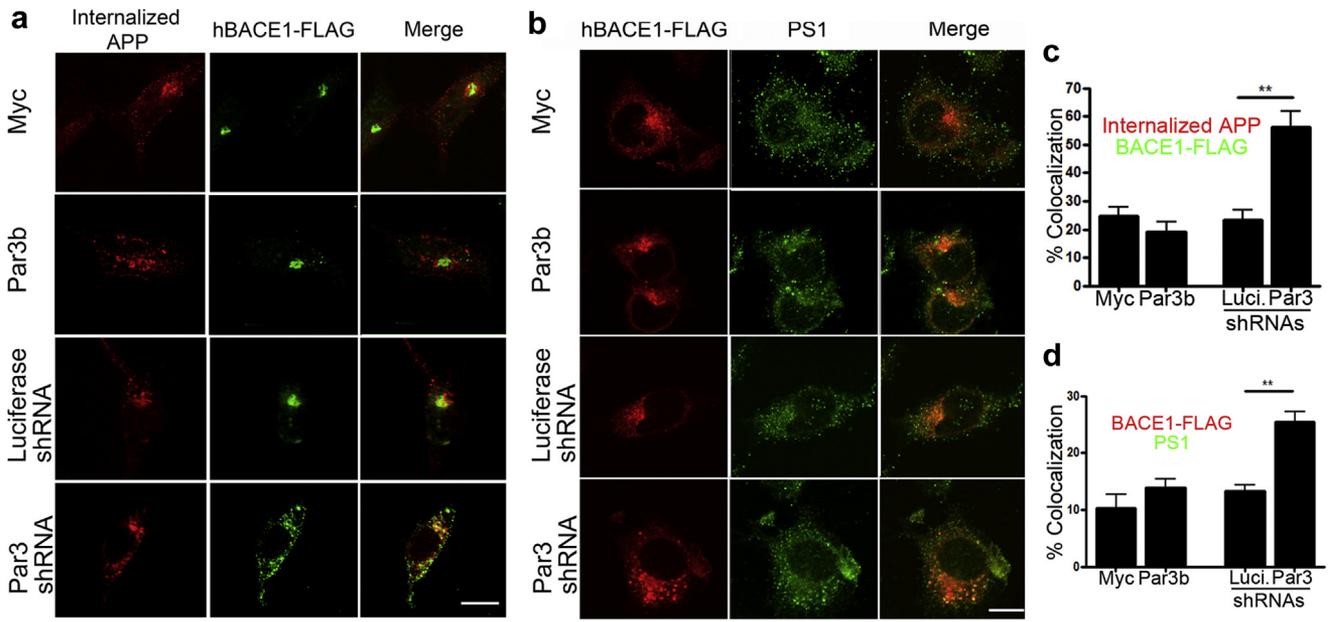


Fig. 3. Depletion of Par3 increases colocalization of internalized APP with BACE1. (A) APPwt N2a cells were transfected with indicated constructs together with hBACE1-FLAG. Seventy-two hours after transfection, internalized APP (60 minutes) was immunostained (red) together with FLAG (green) to visualize colocalization of internalized APP and BACE1. (B) N2a cells were transfected with indicated constructs together with hBACE1-FLAG. Seventy-two hours after transfection, cells were immunostained for FLAG (red) and Presenilin-1 (PS1, green). Scale bar: 10 μ m. (C) Quantification of colocalization of internalized APP with BACE1-FLAG. N (cells) = 19 (myc), 17 (Par3b), 19 (Luciferase shRNA), 20 (Par3 shRNA) from 5 independent experiments. (D) Quantification of the colocalization between BACE1-FLAG and PS1. N (cells) = 12 (myc), 12 (Par3b), 12 (Luciferase shRNA), 11 (Par3 shRNA) from 3 independent experiments. Data were expressed as Mean \pm SEM with Student's *t*-test: ** *p* < 0.01. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)

Par3 through another membrane protein that interacts with PDZ1 of Par3, such as Nectin or junctional adhesion molecules (Ebnet et al., 2003; Itoh et al., 2001; Takekuni et al., 2003).

Our previous studies show that the Par3/aPKC complex regulates retrograde endosome-to-TGN trafficking of BACE1 through the phosphofurin acidic cluster sorting protein 1 (PACS1). The

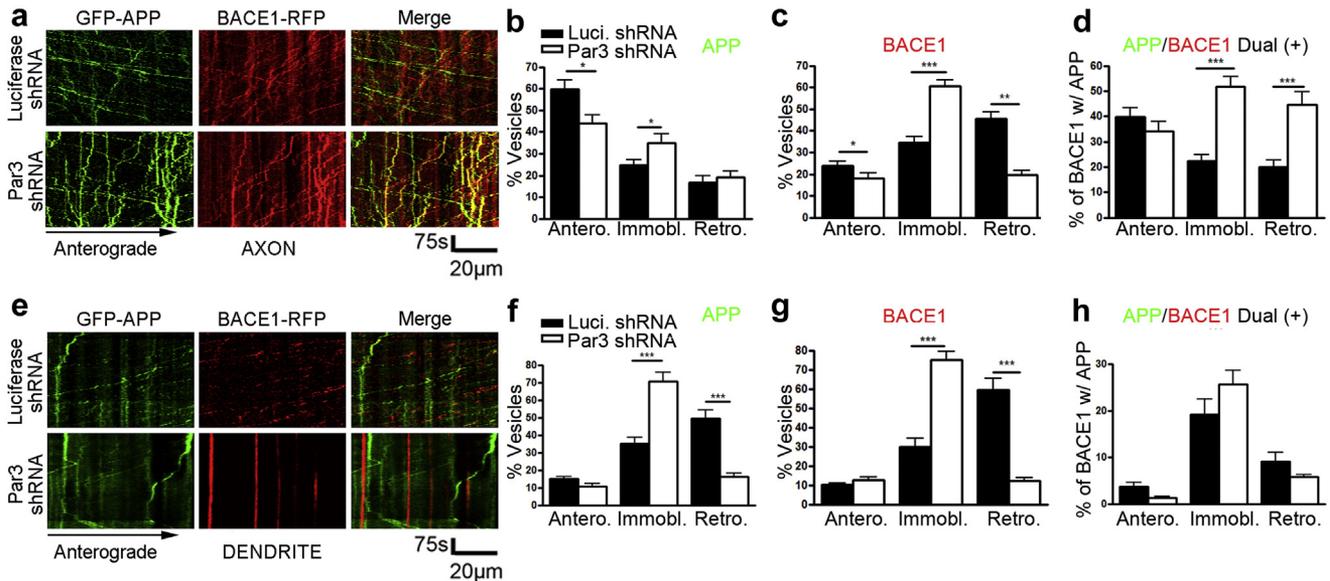


Fig. 4. Loss of Par3 increases APP and BACE1 convergence in axons. (A) Hippocampal neurons were transfected with indicated constructs together with GFP-APP and hBACE1-RFP. Live images were captured on DIV 11 and kymographs were generated to trace APP and hBACE1 trafficking in axons. (B) Quantification of the mobility of APP containing vesicles in axons (anterograde, immobile, and retrograde). (C) Quantifications of the mobility of BACE1 containing vesicles in axons. (D) Quantifications of the percentage of BACE1 vesicles that are co-trafficking with APP vesicles in axons. N (cells) = 17 (Luciferase shRNA), 14 (Par3 shRNA) from 6 independent experiments, with 20–50 vesicles analyzed per movie. (E) Hippocampal neurons were transfected with indicated constructs together with GFP-APP and hBACE1-RFP. Live images were captured on DIV11 and kymographs were generated to trace APP and hBACE1 trafficking in dendrites. (F) Quantification of the mobility of APP containing vesicles in dendrites (anterograde, immobile, and retrograde). (G) Quantifications of the mobility of BACE1 containing vesicles in dendrites. (H) Quantifications of the percentage of BACE1 vesicles that are co-trafficking with APP vesicles in dendrites. N (cells) = 14 (Luciferase shRNA), 16 (Par3 shRNA) from 4 independent experiments, with 10–30 vesicles analyzed per movie. Data were expressed as Mean \pm SEM with Student's *t*-test: * *p* < 0.05; ** *p* < 0.01; *** *p* < 0.001. Scale bar: 75s/20 μ m.

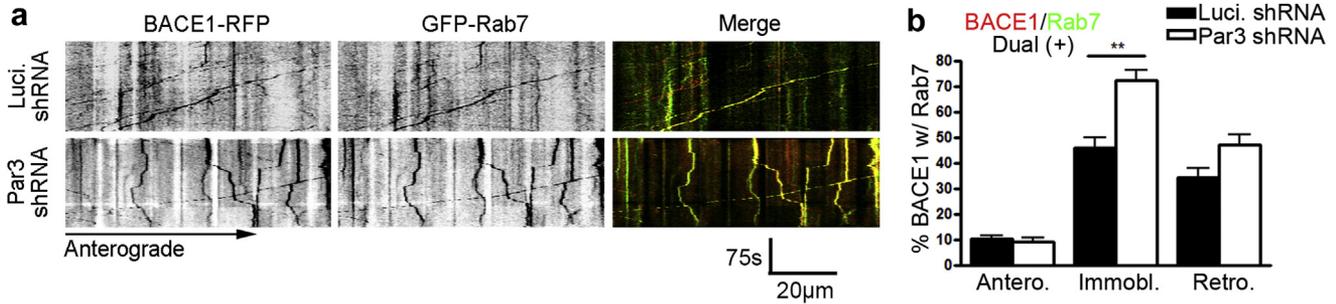


Fig. 5. Loss of Par3 increases retention of BACE1 in late endocytic organelles in axons. (A) Hippocampal neurons were transfected with indicated constructs together with GFP-Rab7 and hBACE1-RFP. Live images were captured on DIV11 and kymographs were generated to trace Rab7 and hBACE1 trafficking in axons. Scale bar: 75s/20 μ m. (B) Quantification of the percentages of BACE1 vesicles colocalized with Rab7 that are trafficking anterograde, immobile or retrograde, respectively. N (cells) = 19 (Luciferase shRNA), 23 (Par3 shRNA) from 8 independent experiments, with 20–50 vesicles analyzed per movie. Data were expressed as Mean \pm SEM with Student's *t*-test: ** $p < 0.01$.

Par3/aPKC complex promotes BACE1 phosphorylation at Ser498, which facilitates its interaction with PACS1 and enhances retrograde endosome-to-TGN trafficking (Sun and Zhang, 2017). Together with our current data, these studies point to the following working model for Par3 in regulating BACE1 trafficking: Par3 forms a complex with BACE1 through its first PDZ domain. aPKC is recruited to this complex likely through its interaction with Par6, as direct interaction between Par3 and aPKC is not required (Fig. 2). The Par3/aPKC complex promotes BACE1 phosphorylation at Ser498. Phosphorylated BACE1 binds to PACS1, which facilitates retrograde trafficking of BACE1. In the absence of Par3, there is a reduction in BACE1 phosphorylation, leading to an inhibition of retrograde trafficking.

Our data show that loss of Par3 leads to APP and BACE1 convergence in axons of primary hippocampal neurons. In particular, we found that there is a significant increase in APP/BACE1 dual positive vesicles that are either retrogradely trafficking or immobile. We further found that most of the retrograde or immobile BACE1 vesicles are Rab7 positive showing that they are predominantly late endocytic organelles. Our data are consistent with recent reports suggesting that axonal late endocytic retention of BACE1 leads to increased A β generation in the Alzheimer's neurons (Ye and Cai, 2014; Ye et al., 2017). Together these data point to the late endosomes as an important subcellular site for A β generation, along with other reported sites such as early endosomes and recycling endosomes (Das et al., 2013, 2016; Grbovic et al., 2003; Kinoshita et al., 2003; Sannerud et al., 2011; Ubelmann et al., 2017). It is likely that A β is generated at multiple subcellular locations within the neuron, and that diverse molecular pathways may differentially regulate A β generation at these locations. Par3 appears to function at the late endosome level as loss of Par3 does not affect APP or BACE1 localization to early endosomes or recycling endosomes

(Sun et al., 2016; Sun and Zhang, 2017). The axonal late endocytic retention of BACE1 is also consistent with our previous data showing increased intracellular A β in neurons depleted of Par3 (Sun et al., 2016). It will be interesting to determine whether the observed axonal convergence of APP and BACE1 is occurring at presynaptic terminals.

In addition, we found that in the absence of Par3, there is a polarized convergence between APP and BACE1 in axons but not dendrites. These data add to the growing literature that APP and BACE1 trafficking are differentially regulated in dendrites and axons of neurons. For example, the endocytic regulators BIN1 and CD2AP differentially regulate APP and BACE1 trafficking in dendrites and axons of hippocampal neurons (Guimas Almeida et al., 2018; Ubelmann et al., 2017). Internalized BACE1 undergoes unidirectional retrograde trafficking in dendrites, which is regulated by EHD family proteins, whereas in axons, BACE1 undergoes bidirectional trafficking (Buggia-Prevot et al., 2013, 2014). These data are consistent with the observation that A β preferentially accumulates in the presynaptic terminals (Yu et al., 2018) and underscores the importance of understanding APP and BACE1 trafficking and convergence within the context of neurons because of their highly compartmentalized nature (Winckler et al., 2018). Because the polarized effects of Par3 is likely due to differential regulation of APP retrograde trafficking in axons versus dendrites (Fig. 4), it will be interesting to determine whether Par3 functions together with some of the known regulators of polarized APP trafficking, such as CD2AP (Ubelmann et al., 2017).

In summary, our data show a novel function for Par3 in regulating polarized APP and BACE1 convergence in neuronal axons. We also provide further mechanistic insight into the regulation of BACE1 trafficking by Par3. Our data are consistent with a growing body of literature highlighting the compartmentalized nature of APP and BACE1 trafficking and convergence and provide further

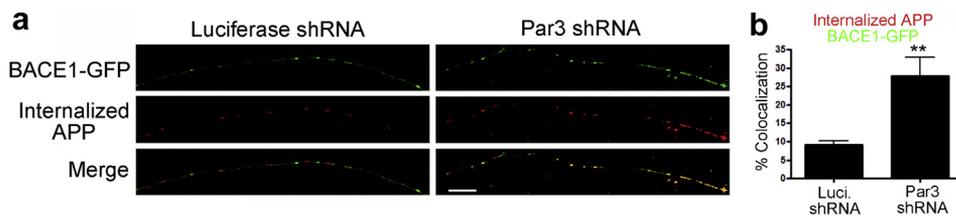


Fig. 6. Loss of Par3 increases the convergence of internalized APP with BACE1 in axons. (A) Hippocampal neurons were transfected with WT APP695, BACE1-GFP and the indicated shRNA constructs. Seventy-two hours after transfection, internalized APP (60 minutes) was immunostained (red) to visualize colocalization of internalized APP and BACE1 in axons. Scale bar: 10 μ m. (B) Quantification of colocalization of internalized APP with BACE1 in axons. N (cells) = 8 (Luciferase shRNA), 13 (Par3 shRNA), from 3 independent experiments. Data were expressed as Mean \pm SEM with Student's *t*-test: ** $p < 0.01$. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)

mechanistic evidence pointing to a key role for the Par polarity complex in AD pathogenesis.

Disclosure

The authors declare they have no competing financial interests.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.neurobiolaging.2019.01.023>.

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