Title:

Synaptic Location Is a Determinant of the Detrimental Effects of α -Synuclein Pathology to Glutamatergic Transmission in the Basolateral Amygdala

Chetan Nagaraja*, Liqiang Chen*, Samuel Daniels*, Lindsay Meyerdirk, Jennifer A Steiner, Martha L. Escobar Galvis, Michael Henderson, Patrik Brundin, Hong-Yuan Chu

Department of Neurodegenerative Science, Parkinson's Disease Center, Van Andel Institute

* These authors contributed equally to this work

Address

333 Bostwick Ave NE, Grand Rapids, MI, 49503, USA

Correspondence

Hongyuan.chu@vai.org

Author contributions.

C.N., L.C., and S.D., investigation, formal analysis; L.M., M.H., J.A.S, and P.B, resources; H.Y.C., conceptualization; H.Y. C: writing– original draft; M.L.E.G, M.H., P.B. and H.Y.C., writing– review and editing; H.Y.C., supervision and funding acquisition. All authors reviewed and approved the manuscript.

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Conflicts of interest

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Abstract

α -Synuclein (α Syn) is a soluble protein enriched at presynaptic boutons and its abnormal aggregation is a hallmark of Parkinson's disease (PD). The amygdala shows selective vulnerability to the formation of insoluble α Syn aggregates, and its circuit dysfunction may contribute to prevalent psychiatric deficits in PD. Yet, how α Syn aggregates affect the amygdala function remains elusive. In the present study, we examined the presence of α Syn in glutamatergic axon terminals and how its aggregation affects glutamatergic transmission in the basolateral amygdala (BLA). Using confocal microscopy, we showed that α Syn is primarily present in the axon terminals expressing vesicular glutamate transporter 1 (vGluT1), but not in those expressing vGluT2. Using electrophysiology and optogenetics, we found that glutamatergic neurotransmission from vGluT1⁺ axon terminals are functionally more vulnerable than vGluT2⁺ axon terminals to the toxic effects of pathological α Syn aggregation triggered by α Syn preformed fibrils (PFFs). Using the PFFs models and mouse genetics, we showed that loss of α Syn promotes short-term depression of cortico-BLA synapses responding to repetitive stimuli, leading to an input-selective filtering of cortical inputs to the BLA. Together, we demonstrate that α Syn expresses specifically in a subset of synapses in mouse amygdala, which causes an input-specific decrease of cortical inputs to the BLA as α Syn aggregation forms. These results might be relevant to the reduced cortical control of amygdala function that has been associated with psychiatric deficits in PD.

Introduction

α -Synuclein (α Syn) is a soluble protein abundant at presynaptic axon terminals, where it regulates the dynamics of synaptic vesicles through interaction with synaptic proteins and presynaptic membranes (Burré et al., 2010; Runwal and Edwards, 2021; Vargas et al., 2017). α Syn is also prone to aggregation and aggregated α Syn is the major protein component of Lewy pathology seen in synucleinopathies like Parkinson's disease (PD) (Mezey et al., 1998; Spillantini et al., 1997). Moreover, increasing evidence supports the notion that pathologic α Syn propagates between synaptically interconnected brain regions and underlies PD progression (Angot et al., 2010; Luk et al., 2012; Uemura et al., 2020).

The amygdala is a key limbic structure for emotion regulation (Janak and Tye, 2015). Compelling clinical evidence indicates that cortical control of the amygdala activity is impaired in PD, leading to an inappropriate encoding of emotion valence or deficits in linking emotion to proper behavior (Bowers et al., 2006; Hu et al., 2015; Yoshimura et al., 2005). Moreover, the amygdala consistently shows selective vulnerability to Lewy pathology (Harding et al., 2002; Nelson et al., 2017; Sorrentino et al., 2019), thus an impaired amygdala network function has been proposed to underlie the disrupted emotion processing in PD patients (Harding et al., 2002). Still, the normal function of α Syn and how its aggregation can impair circuit function in the amygdala remain poorly understood. Here, we show that α Syn is primarily present in vesicular glutamate transporter 1-expressing (vGluT1⁺) cortical axon terminals, being barely detectable in vGluT2-expressing (vGluT2⁺) thalamic axon terminals in the basolateral amygdala (BLA). Using an α Syn preformed fibrils (PFFs) model of synucleinopathy, we

also show that development of α Syn pathology disrupts vGluT1⁺ cortico-BLA glutamatergic transmission, without affecting the vGluT2⁺ thalamo-BLA neurotransmission. Combining optical imaging with physiological approaches, we demonstrate that a partial (secondary to α Syn aggregation) or complete depletion of soluble α Syn (*Snca* KO mice) from the axon boutons promotes short-term depression at cortico-BLA synapses in response to prolonged stimulation, leading to an impaired gain control of cortical inputs to the BLA. Therefore, we conclude that changes in α Syn function decrease the cortical inputs to the BLA in an input-specific manner. Our data support clinical observations of an impaired cortical control of the amygdala activity that could contribute to psychiatric deficits of PD patients.

Results and Discussion

α Syn localizes preferentially in vGluT1⁺ axon terminals in mouse brain

Compelling evidence from PD patients suggests that the amygdala exhibits an impaired responsiveness to sensory stimuli, arising mainly from the cerebral cortex and the thalamus (Bowers et al., 2006; Hu et al., 2015). Since the cortical and thalamic glutamatergic axon terminals express vGluT1 and vGluT2, respectively (Freneau et al., 2001; Kaneko and Fujiyama, 2002; Vigneault et al., 2015), we examined the presence of α Syn in vGluT1⁺ and vGluT2⁺ axon buttons in WT mouse brains. Figure 1 shows that α Syn immunoreactivity colocalizes with vGluT1⁺ puncta but is absent where there are vGluT2⁺ puncta, in the BLA, the cerebral cortex, and the striatum. It suggests that α Syn preferentially presents at vGluT1⁺ axon terminals arising from cortical regions. This observation is consistent with previous *in situ* hybridization studies showing higher α Syn mRNA expression in the cerebral cortex and the hippocampus than in the thalamus (Abeliovich et al., 2000; Ziolkowska et al., 2005).

Figure 1. α Syn is selectively localized at vGluT1⁺ axon terminals in mouse brain. **A)** Representative confocal images showing colocalization of α Syn with vGluT1 (left), but not with vGluT2 (right) in the BLA (top), cerebral cortex (middle), and the striatum (bottom). Simultaneously collected confocal images from each brain region were split into vGluT1/ α Syn and vGluT2/ α Syn channels for the purpose of illustration. **B)** Zoomed z series of images of the boxed area from the BLA in (A) showing colocalization and correlated changes in the immunoreactive intensities of vGluT1 and α Syn. Such colocalization and correlation is absent between vGluT2 and α Syn within the same region. Images were taken and shown from a 0.75 μ m z-depth with an inter-section interval of 0.25 μ m. **C)** Bar graphs showing Pearson correlation coefficient between vGluT1 and α Syn, as well as between vGluT2 and α Syn, in the BLA (α Syn/vGluT1 = 0.82 ± 0.007 , α Syn/vGluT2 = 0.17 ± 0.004 ; n = 18 slices/4 mice; p < 0.05, MWU), cerebral cortex (α Syn/vGluT1 = 0.82 ± 0.006 , α Syn/vGluT2 = 0.12 ± 0.009 ; n = 8 slices/3 mice; p < 0.05, MWU), and the striatum (α Syn/vGluT1 = 0.83 ± 0.006 , α Syn/vGluT2 = 0.17 ± 0.02 ; n = 8 slices/3 mice; p < 0.05, MWU).

Glutamate release from vGluT1⁺ axon terminals is selectively disrupted by α Syn pathology

Considering that α Syn is selectively present in vGluT1⁺ axon terminals, and that the endogenous α Syn is a determinant of the propensity to form α Syn aggregation, one can hypothesize that these terminals are more susceptible to the detrimental effects of α Syn pathology compared to vGluT2⁺ terminals (Erskine et al., 2018; Vasili et al., 2022). To test this hypothesis, we triggered α Syn pathology in the brain using the intrastriatal PFFs seeding model (Luk et al., 2012). One-month post-injection, we detected robust α Syn pathology in vGluT1⁺ cerebral cortical regions (e.g., the temporal association cortex (TeA) and the perirhinal cortex), but we barely observed any cytoplasmic aggregates in vGluT2⁺ thalamic regions (data not shown), which is consistent with earlier reports (Burtscher et al., 2019; Henderson et al., 2019; Luk et al., 2012; Stoyka et al., 2019). These findings are also expected considering the low level, or even

absence of α Syn mRNA expression in vGluT2⁺ thalamic regions (Abeliovich et al., 2000; Ziolkowska et al., 2005).

The cerebral cortex and thalamus provide major excitation to the BLA for emotion regulation, thus we further examined whether vGluT1⁺ cortico-BLA and vGluT2⁺ thalamo-BLA synaptic transmission are differentially affected by α Syn pathology. We activated cortical and thalamic inputs of the BLA selectively by stimulating the external and internal capsules, respectively. One-month post-injection, the amplitude of electrically evoked cortico-BLA EPSCs was greatly decreased in the slices from PFFs-injected WT mice relative to those from controls (Figure 2A, B). In contrast, there was no difference in the amplitude of thalamo-BLA EPSCs between the PFFs-injected mice and controls (Figure 2C, D).

To avoid potential technical issues in the use of electrical stimulation, we employed optogenetic approach to confirm the above results (Figure 2E-J). One-month post-injection, the amplitude of optogenetically-evoked cortico-BLA EPSCs in slices from PFFs-injected mice was decreased relative to those from PBS-injected controls (Figure 2K, L). By contrast, we detected no difference in the amplitude of optogenetically-evoked thalamo-BLA EPSCs between groups (Figure 2M, N). Altogether, we demonstrated that the presence of α Syn makes vGluT1⁺ axon buttons more vulnerable to α Syn pathology after injection of PFFs and that this pathology is associated with detrimental effects on synaptic transmission.

Figure 2. α Syn aggregates preferentially disrupt vGluT1⁺ cortico-BLA transmission. **A-B)** Representative traces of EPSCs evoked by electrical stimulation of the external capsule (A) and summarized results (B) showing a reduced cortico-BLA transmission in slices from PFFs- versus PBS-injected mice. $n = 17$ neurons/4 mice for each group. **C-D)** Representative traces of EPSCs evoked by electrical stimulation of the internal capsule (C) and summarized results (D) showing unaltered thalamo-BLA transmission in slices from PFFs- versus PBS-injected wildtype mice. $n = 17$ cells/5 mice for controls, and 13 cells/4 mice. **E-G)** Representative images showing viral infection site in the TeA and nearby regions (E), and the axon terminal field in the BLA under low (F) and high (G) magnification. **H-J)** Representative images showing viral infection site in the midline thalamus (H), and the axon terminal field in the BLA under low (I) and high (J) magnification. **K-L)** Representative traces of optogenetically-evoked EPSCs (K) and summarized results (L) showing a reduced amplitude of cortico-BLA EPSCs in slices from PFFs- versus PBS-injected mice. $n = 35-37$ neurons/4 mice. **M-N)** Representative traces of optogenetically-evoked EPSCs (M) and summarized results (N) showing unaltered thalamo-BLA transmission in slices from PFFs- versus PBS-injected mice. $n = 28$ neurons/4 mice. *, $p < 0.05$, MWU followed by Bonferroni-Dunn correction for multiple comparisons.

Consistent with the development of α Syn pathology following PFFs injection requiring the presence of endogenous α Syn (Luk et al., 2012; Volpicelli-Daley et al., 2011), we did not detect difference in the cortico-BLA transmission between PFFs- versus PBS-injected α Syn KO mice (data not shown). Because of the lack of detrimental effect in α Syn KO mice, we conclude that the impaired cortico-BLA transmission in the PFFs model is likely to be caused by the gained toxic properties of α Syn as it aggregates (Cookson and Brug, 2008; Volpicelli-Daley et al., 2011).

These results further highlight the dependence of regional and cellular vulnerability on the endogenous α Syn, i.e., those neurons or axon terminals that express high levels of α Syn are prone to be functionally impacted by α Syn pathology relative to those which lack of or express low levels of α Syn (Surmeier et al., 2017; Thakur et al., 2019; Vasili et al., 2022). Of particular interest is that vGluT2⁺/TH⁺ midbrain dopamine neurons and their axon terminals in the striatum are more resilient to neurodegeneration in postmortem PD brains and animal models studies (Buck et al., 2021; Thomas et al., 2021). While several other mechanisms have been proposed (Buck et al., 2021), the absence of α Syn at vGluT2⁺ neurons/terminals could be critical for such resilience.

Pathological aggregation decreases α Syn levels at axon terminals and impairs short-term synaptic plasticity

Formation of cytoplasmic aggregates is believed to move soluble α Syn away from the presynaptic buttons, affecting its role in regulating synaptic vesicle pools (Benskey et al., 2016; Volpicelli-Daley et al., 2011). Consistent with the earlier

predictions, the intensity of α Syn immunoreactivity within vGluT1⁺ puncta decreased dramatically as α Syn pathology develops in our PFFs model, leading to an increased proportion of vGluT1⁺ buttons that lack detectable level of α Syn (Figure 3). These data suggest that α Syn pathology reduces the amount of α Syn present at the cortical axon terminals in the BLA, and in that way could affect its physiological function.

Figure 3. Decreased α Syn levels at axon terminals as it forms aggregates. A1-B3) Representative confocal images showing immunoreactivity of α Syn that is localized within vGluT1 immunoreactive puncta in BLA slices from a control (A1-A3) and PFFs-injected mouse (B1-B3). **C)** Summarized graph showing a reduced α Syn immunoreactivity per vGluT1 immunoreactive puncta in PFFs-injected mice relative to controls (control = 1049 ± 50 ; PFFs = 617 ± 18 , $n = 100$ puncta/group, $p < 0.0001$, MWU). **D)** Summarized graph showing a reduced percentage of vGluT1 immunoreactive puncta associated with detectable α Syn immunoreactivity in PFFs-injected mice relative to controls (controls = $85 \pm 5.2\%$; PFF = $26 \pm 1.7\%$, $n = 6$ slices/group, $p = 0.002$, MWU).

α Syn modulates the dynamics of synapse vesicle pools (Sulzer and Edwards, 2019), which plays a key role in regulating synaptic plasticity and the computational function of neural circuits (Abbott and Regehr, 2004; Alabi and Tsien, 2012). Thus, we stimulated cortico-BLA synapses repetitively to assess the impact of the observed reduction of α Syn levels on short-term synaptic plasticity. We detected a progressive

depression of cortico-BLA EPSCs in control mice (Figure 4A1, A2), reflecting mainly a progressive depletion of presynaptic vesicles (Alabi and Tsien, 2012; Cabin et al., 2002). Interestingly, the cortico-BLA EPSCs in the slices from PFFs-injected mice showed a greater depression relative to those from controls (Figure 4A1, A2). These results are in line with the key role of α Syn in sustaining and mobilizing synaptic vesicle pools (Cabin et al., 2002; Sulzer and Edwards, 2019; Vargas et al., 2017), and suggest the significant impact of the loss of α Syn function on synaptic plasticity and circuit computation.

Figure 4. Loss of α Syn impairs short-term plasticity of the cortico-BLA inputs.

A1) Representative cortico-BLA EPSCs traces from control and PFF-injected mice in response to repetitive stimulation (300 stimuli at 12.5 Hz). **A2)** Summarized graph showing the temporal profiles of cortico-EPSCs depression in slices from control and PFFs-injected mice. The cortico-BLA EPSCs from PFFs-injected mice exhibited greater reduction in the amplitude towards the end of repetitive stimulation. $n = 13-15$ neurons/ 3 mice. **B1)** Representative cortico-BLA EPSCs traces from WT control and α Syn KO mice in response to repetitive stimulation (300 stimuli at 12.5 Hz). **B2)** Summarized graph showing the temporal profiles of cortico-EPSCs depression in slices from WT control and α Syn KO mice ($n = 8-9$ neurons/3 mice. *, $p < 0.05$, MWU followed by Bonferroni-Dunn correction for multiple comparisons).

Because both gain of toxic properties and loss of normal α Syn function were induced in the PFFs models, we employed the α Syn KO mice to further assess the impact of a loss of α Syn on the short-term synaptic plasticity of glutamatergic synapses. Consistently, the amplitude of cortico-BLA EPSCs from α Syn KO mice also exhibited greater depression in response to repetitive stimulation relative to littermate WT controls (Figure 4B1-B2), indicating an impaired mobilization of synaptic vesicle pools. It is worth noting that the earlier onset and greater magnitude of cortico-BLA EPSC depression in α Syn KO mice versus that in PFFs-injected mice (Figure 4A-B). Thus, it is plausible that the different temporary profiles of short-term synaptic dynamics are linked to difference in the amount of α Syn present at presynaptic terminals between the PFFs model and KO mice. Last, we did not detect difference in the temporal profiles of thalamo-BLA EPSCs in slices from α Syn KO mice and those from WT controls (Figure S1), which indicates a neglectable impact of α Syn depletion on synaptic vesicle dynamics and is consistent with the lack of α Syn presence at vGluT2⁺ axon terminals (Figure 1). Together, these results suggest that pathological aggregation decreases the levels of soluble α Syn at the axon terminals, leading to a greater synaptic vesicle depletion and impaired short-term plasticity.

Figure S1. α Syn KO does not affect short term plasticity profiles of thalamo-BLA inputs. **A1)** Representative thalamo-BLA EPSCs traces from WT control and α Syn KO mice in response to repetitive stimulation (300 stimuli at 12.5 Hz). **A2)** Summarized graph showing the temporal profiles of thalamo-BLA EPSCs depression in slices from WT control and α Syn KO mice (n = 9 neurons/3 mice. No statistical significance was detected between groups, MWU followed by Bonferroni-Dunn correction for multiple comparisons).

Short-term synaptic plasticity is an important “gain control” mechanism for neurons to properly balance their responsiveness to distinct afferents in an input-specific manner (Abbott et al., 1997). Cortical and thalamic afferents of the BLA exhibit different short-term synaptic plasticity profiles *in vivo*, which could reflect their different contributions to the formation of emotion related memory and behavior (Sigurðsson et al., 2010). Physiologically, synapse-specific presence of α Syn could underlie such difference in the short-term plasticity profiles of the two inputs. One can postulate that once aggregates form, a decreased synaptic strength associated with α Syn toxicity (Figure 2) and an enhanced synaptic depression due to the loss of α Syn normal function at presynaptic buttons (Figure 4) make the BLA neurons less likely to respond to sustained and repetitive sensory inputs from cortical regions. These circuit changes might explain studies showing a decreased functional connectivity of cortico-amygdala, but not the thalamo-amygdala network in PD patients, which further leads to a failed

amygdala responsiveness to the aversive sensory inputs (Hu et al., 2015; Yoshimura et al., 2005).

Materials and Methods

Animals

Wild type (WT) C57Bl/6 mice of both sexes (3 – 4 month-old) were obtained from the Van Andel Institute vivarium internal colony and used in the study. α Syn knockout (*Snca*^{-/-}) mice were originally purchased from Jackson laboratories (Bar Harbor, ME) and maintained in C57Bl/6 background. Experimental *Snca*^{-/-} mice and littermate WT controls were generated from heterozygous *Snca*^{+/-} breeder pairs and were genotyped by Transnetyx (Cordova, TN, USA). Mice were housed up to four animals per cage under a 12/12 h light/dark cycle with access to food and water *ad libitum* in accordance with NIH guidelines for care and use of animals. All animal studies were reviewed and approved by the Institutional Animal Care and Use Committee at Van Andel Institute.

Preparation and validation of α Syn preformed fibrils

Purification of recombinant mouse α Syn and generation of α Syn preformed fibrils (PFFs) was conducted as described elsewhere (Luk et al., 2009; Volpicelli-Daley et al., 2014). The pRK172 plasmid containing the gene of interest was transformed into BL21 (DE3) RIL-competent *E. coli* (230245, Agilent Technologies). A single colony from this transformation was expanded in Terrific Broth (12 g/L of Bacto-tryptone, 24 g/L of yeast extract 4% (v/v) glycerol, 17 mM KH₂PO₄ and 72 mM K₂HPO₄) with ampicillin. Bacterial pellets from the growth were sonicated and the sample was boiled to precipitate undesired proteins. The supernatant was dialyzed with 10 mM Tris, pH 7.6, 50 mM NaCl, 1 mM EDTA, 1 mM phenylmethylsulfonyl fluoride (PMSF) overnight. Protein was filtered with a 0.22 μ m filter and concentrated using Vivaspin 15R 10K centrifugal filters

(VS15RH01, Sartorius). Protein was then loaded onto a Superdex 200 column and 2 mL fractions were collected. Fractions were run on SDS-PAGE and stained with InstaBlue protein stain (50-190-5499, Fisher Scientific) to select fractions that were highly enriched in α Syn. These fractions were combined and dialyzed in 10 mM Tris, pH 7.6, 50 mM NaCl, 1 mM EDTA, 1 mM PMSF overnight. Dialyzed fractions were applied to a MonoQ column (HiTrap Q HP 17115401, Cytiva) and run using a NaCl linear gradient (0.025-1 M). Collected fractions were run on SDS-PAGE and stained with InstaBlue protein stain. Fractions that were highly enriched in α Syn were collected and dialyzed into DPBS (Gibco). Protein was filtered through a 0.22 μ m filter and concentrated to 7 mg/mL (α Syn) with Vivaspin 15R 10K centrifugal filters. Monomer was aliquoted and frozen at -80°C. For preparation of α Syn PFFs, α Syn monomer was shaken at 1,000 rpm for 7 days. Conversion to PFFs was validated by sedimentation at 100,000 $\times g$ for 60 minutes and by thioflavin S staining.

Stereotaxic surgery

Mice were placed in a stereotaxic frame (Kopf) under 2% isoflurane anesthesia and were supported by a thermostatic heating pad. Recombinant α Syn fibrils were diluted in phosphate-buffered saline (PBS) to a concentration of 5 mg/ml. Prior to stereotaxic injection, PFFs were pulse sonicated at medium intensity with 30 seconds on and 30 seconds off for 10 minutes using a sonicator (Biorupter Pico). To induce the formation of α Syn aggregation in mouse brain (Luk et al., 2012), sonicated PFFs (2.0 μ l) were injected bilaterally into the dorsal striatum (anteroposterior, +0.2 mm from bregma; mediolateral, $\pm$ 2.0 mm from the midline; dorsoventral, -2.6 mm from the brain surface)

using a 10- μ l syringe (Hamilton) driven by a motorized microinjector (Stoelting) at a rate of 0.2 μ l/min. Given the neuronal heterogeneity of the BLA, retrobeads (Lumafluor Inc) were diluted 10x into α Syn fibrils or PBS to label the projection neurons in the BLA for physiological studies. The final injection volume of PFFs with beads mixture was adjusted to keep a constant amount of α Syn injected across animals. Control mice received 2.0 μ l PBS or PBS with retrobeads injections in the same location. For optogenetics studies, AAV vectors encoding ChR2(H134R)-eYFP (titer = 1×10^{12} vg/ml, Addgene 26973) were stereotactically injected into and centered at the temporal association cortex (TeA) (anterioposterior -3.3 mm from the bregma, mediodorsal ± 4.1 mm, dorsoventral -1.5 mm from the brain surface) and the midline thalamus (anterioposterior -1.5 mm from the bregma, mediodorsal 0 mm, dorsoventral -3.3 mm from the brain surface) (Ahmed et al., 2021; Amir et al., 2019). Animals were housed in their home cages before being euthanized for immunohistochemical or physiological studies at one-month post-injection.

Immunohistochemistry. Animals received overdosage of avertin intraperitoneally (i.p.) and were subsequently subjected to transcardial perfusion with PBS and 4% paraformaldehyde (PFA, pH 7.4). Brains were removed and post-fixed in 4% PFA overnight, before being re-sectioned at 70 μ m using a vibratome (VT1000s, Leica Biosystems, Buffalo Grove, IL). Brain sections were then rinsed using PBS and treated with 0.5% Triton X-100 and 2% normal donkey serum (MilliporeSigma) in PBS for 60 min at room temperature, followed by incubation with primary antibodies overnight at room temperature or for 48 hours at 4 °C. The concentrations of primary antibodies

were rabbit anti-pS129 α Syn (1:10,000, cat#: EP1536Y, Abcam), mouse anti- α Syn (1:1,000, cat#: 610787, BD Biosciences), mouse anti-NeuN (1:2000, cat#: MAB377, MilliporeSigma), rabbit anti-vGluT1 (1:1,000, cat#: ABN1647, MilliporeSigma) and guinea pig anti-vGluT2 (1:1000, cat#: 135404, Synaptic Systems). After being thoroughly rinsed with PBS for 3 times, the sections were incubated with the secondary antibodies (1:500, AlexaFluor 594 or 647 donkey anti-mouse IgG, AlexaFluor 488 or 594 donkey anti-rabbit IgG, or AlexaFluor 488 donkey anti-guinea pig IgG, Jackson ImmunoResearch) for 90 min at room temperature. Brain sections were rinsed 3 times with PBS and mounted on glass slides using Vectorshield antifade mounting medium (H-1000, Vector Laboratories) and cover slipped.

Confocal imaging and analysis

Confocal images were acquired using a Nikon A1R confocal microscopy. pS129 α Syn aggregates in different brain regions were imaged under a 40x objective lens. For α Syn and vGluT1/vGluT2 colocalization analysis, three synaptic markers were immunostained simultaneously and z-stack images were acquired using an oil immersion 100x objective (NA = 1.45; x/y, 1024 X 1024 pixels; z step = 150 nm). Images were acquired using identical settings between treatment groups, including laser power, pinhole size, gain, and pixel size. Intensity-based colocalization analysis was performed using Imaris software (version 9.3, Oxford, UK). Background subtraction on z-stack images was conducted using ImageJ (NIH) prior to importing files into Imaris. Once imported, two arbitrary regions of interest were created using the surface function (drawing mode: circle; radius = 15 μ m; number of vertices = 30) and the three channels

(vGluT2, vGluT1, α Syn) were masked based on the surface reconstruction to isolate fluorescence within the ROIs. The Imaris “Coloc” function was used to measure the colocalization between vGluT2/ α Syn and vGluT1/ α Syn for each ROI. Briefly, either vGluT1 or vGluT2 was selected as channel A and α Syn was selected as channel B. The automatic threshold feature within “Coloc” was used to calculate the threshold values for each channel. Colocalization channels for vGluT1/ α Syn and vGluT2/ α Syn for each ROI were created by using the "Build Coloc Channel" function. Colocalization parameters were obtained for quantification from the colocalization channels. For quantification of the α Syn intensity within vGluT1⁺ axon terminals, background subtraction on z-stack images was conducted using ImageJ (NIH). After background subtraction, vGluT1 immunoreactive puncta were manually identified as a set of regions of interest (ROI, n = 20 puncta per image) from each BLA section and the mean gray values of α Syn immunoreactivity within the same ROI were then measured using ImageJ (NIH).

Slice preparation for physiology. For physiological studies, mice were deeply anesthetized with avertin (300 mg/kg, i.p.) and then were perfused transcardially with ice-cold, sucrose-based artificial cerebrospinal fluid (aCSF) containing (in mM) 230 sucrose, 26 NaHCO₃, 10 glucose, 10 MgSO₄, 2.5 KCl, 1.25 NaH₂PO₄, and 0.5 CaCl₂, 1 sodium pyruvate, and 0.005 L-glutathione. Next, coronal brain slices (300 μ m) containing BLA were prepared in the same sucrose-based aCSF solution using a vibratome (VT1200S; Leica Microsystems, Buffalo Grove, IL). Brain slices were kept in normal aCSF (in mM, 126 NaCl, 26 NaHCO₃, 10 glucose, 2.5 KCl, 2 CaCl₂, 2 MgSO₄,

1.25 NaH₂PO₄, 1 sodium pyruvate, and 0.005 L-glutathione) equilibrated with 95% O₂ and 5% CO₂ for 30 min at 35°C and then held at room temperature until use.

Ex vivo electrophysiology recording and optogenetics. Brain slices were transferred into a recording chamber perfused at a rate of 3 – 4 ml/min with synthetic interstitial fluid (in mM, 126 NaCl, 26 NaHCO₃, 10 glucose, 3 KCl, 1.6 CaCl₂, 1.5 MgSO₄, and 1.25 NaH₂PO₄) equilibrated with 95% O₂ and 5% CO₂ at 33–34°C via a feedback-controlled in-line heater (TC-324C, Warner Instruments) (Chen et al., 2021). SR-95531 (GABAzine, 10 μM) was routinely added extracellularly to block GABA_A receptor-mediated inhibitory synaptic transmission. Neurons were visualized and recorded under gradient contrast SliceScope 1000 (Scientifica, Uckfield, UK) with infrared illumination using an IR-2000 CCD camera (DAGE-MTI, USA) and motorized micromanipulators (Scientifica, Uckfield, UK). Individual BLA projection neurons labeled with retrobeads were identified using a 60x water immersion objective lens (Olympus, Japan) and targeted for whole-cell patch-clamp recording. Data were collected using a MultiClamp 700B amplifier and a Digidata 1550B digitizer at a sampling rate of 50 kHz under the control of pClamp11 (Molecular Devices, San Jose, USA). Borosilicate glass pipettes (O.D. = 1.5 mm, I.D. = 0.86 mm, item #BF150-86-10, Sutter Instruments) for patch clamp recordings (4 – 6 MΩ) were pulled using a micropipette puller (P1000, Sutter Instruments, Novato, CA).

To assess glutamatergic transmission in the BLA, glass pipettes were filled with a cesium methanesulfonate based internal solution of (in mM): 120 CH₃O₃SCs, 2.8 NaCl, 10 HEPES, 0.4 Na₄-EGTA, 5 QX314-HBr, 5 phosphocreatine, 0.1 spermine, 4 ATP-Mg,

and 0.4 GTP-Na (pH 7.3, 290 mOsm). BLA neurons were voltage clamped at -70 mV to assess the excitatory postsynaptic currents (EPSCs) in response to presynaptic electrical or optogenetic stimulations. Concentric bipolar electrodes (FHC, Bowdoin, ME) or glass pipettes ($\sim 2 \text{ M}\Omega$) filled with the extracellular solution were used as stimulating electrodes and placed on the external and internal capsules to evoke glutamate release in the BLA from cortical and thalamic axon terminals, respectively (Shin et al., 2010). A range of electrical pulses (intensities = 30 – 120 μA , duration = 100 μs) were delivered through a stimulator (Digitimer, UK) to evoke glutamate release at either cortical or thalamic inputs. To study the process of synaptic vesicle pool mobilization during repetitive glutamatergic transmission, we delivered 300 electrical pulses (duration = 100 μs) at 12.5 Hz and measured the amplitudes of EPSCs to quantify changes in the presynaptic glutamate release (Cabin et al., 2002). Only one neuron was recorded from each slice in the prolonged repetitive stimulation studies.

Optogenetic stimulation pulses (1 ms duration) were delivered through a 60x objective lens (Olympus, Japan) using a 470 nm LED light source (CoolLED, UK). To isolate monosynaptic cortico-BLA and thalamo-BLA EPSCs, optogenetically evoked EPSCs were recorded in the presence of TTX (1 μM) and 4-AP (100 μM). Series resistance (R_s , $< 20 \text{ M}\Omega$) was regularly monitored throughout the recording to ensure R_s changes were less than 15%. Liquid junction potential ($\sim 9 \text{ mV}$) was not corrected.

Data analysis and statistics.

Electrophysiology data were analyzed offline in Clampfit 11.1 (Molecular Devices). The peak amplitude of monosynaptic EPSCs in response to electric stimulation was

quantified from an average of three to five sweeps. Digital confocal images were analyzed using ImageJ (NIH) or Imaris (Oxford,UK). Statistics were performed using Prism9 (GraphPad Software). To minimize the assumption of the data normality, non-parametric, distribution-independent Mann–Whitney U (MWU) test was used for non-paired data comparisons between groups, followed by Bonferroni-Dunn correction for multiple comparisons. All tests were two-tailed, and p value < 0.05 was considered statistically significant. Summary results are reported as mean plus standard error of mean.

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