Dendritic Localization and Exocytosis of NAAG in the Rat Hippocampus

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Abstract

While a lot is known about classical, anterograde neurotransmission, less is known about the mechanisms and molecules involved in retrograde neurotransmission. Our hypothesis is that N-acetylaspartylglutamate (NAAG), the most abundant dipeptide in the brain, may act as a retrograde transmitter in the brain. NAAG was predominantly localized in dendritic compartments of glutamatergic synapses in the intact hippocampus, where it was present in close proximity to synaptic-like vesicles. In acute hippocampal slices, NAAG was depleted from postsynaptic dendritic elements during neuronal stimulation induced by depolarizing concentrations of potassium or by exposure to glutamate receptor (GluR) agonists. The depletion was completely blocked by botulinum toxin B and strictly dependent on extracellular calcium, indicating exocytotic release. In contrast, there were low levels of NAAG and no effect by depolarization or GluR agonists in presynaptic glutamatergic terminals or GABAergic pre- and postsynaptic elements. Together these data suggest a possible role for NAAG as a retrograde signaling molecule at glutamatergic synapses via exocytotic release.

Key words: dendrite, immunogold, neurotransmission, retrograde, synapse

Introduction

Retrograde neurotransmission is the release of a signaling molecule from the postsynaptic neuron that acts on receptors at the presynaptic nerve terminal, leading to alterations in synaptic transmission (Regehr et al. 2009). Thus, retrograde transmission provides postsynaptic neurons with the ability to control their own inputs. This mechanism has gained increasing focus recently, as a way of fine-tuning signaling between neurons. While a lot is known about how presynaptic terminals communicate to their postsynaptic counterparts, less is known about the mechanisms involved in retrograde neurotransmission.

The retrograde transmitters can be classified based on their mechanisms of release: 1) lipid derivatives or gasses, 2) neuropeptides, and 3) classic neurotransmitters. The most well-established retrograde transmitters, the endocannabinoids and nitric oxide, belong to the first group. Being lipophilic, they are not subjected to a vesicular release mechanism, and their release is regulated at the level of synthesis. Neuropeptides are released from dense-core vesicles as well as through
nonvesicular mechanisms from postsynaptic elements, dendrites, and somata (Cheramy et al. 1981). Some classical neurotransmitters, in particular glutamate and gamma-aminobutyric acid (GABA), have been suggested to also act as retrograde transmitters (Zilberter 2000; Jenstad et al. 2009).

In anterograde neurotransmission, a rapid rise in intracellular calcium causes synaptic vesicles filled with neurotransmitters to fuse with the cell membrane and release their contents into the synaptic cleft; a release mechanism known as exocytosis. A number of proteins are involved in exocytosis, in particular the soluble N-ethylmaleimide-sensitive attachment protein receptor (SNARE) family. Although vesicles can dock at the active zone without the direct involvement of these proteins, the priming of vesicles is mediated by SNAREs. Some SNARE proteins have been identified postsynaptically (Kennedy et al. 2010; Lau et al. 2010; Suh et al. 2010; Jurado et al. 2013; Hussain and Davanger 2015), opening for the possibility that retrograde transmitters can be released by exocytosis.

N-Acetylaspartylglutamate (NAAG) is the most abundant dipeptide in the brain, where it is present at high concentrations (μM–mM) (Miyamoto et al. 1966; Koller et al. 1984; Guarda et al. 1988). Several lines of evidence suggest that NAAG acts as a neurotransmitter (Bergeron and Coyle 2012; Watanabe et al. 2018). Biochemical studies have shown that NAAG is released Ca2+-dependently from brain slices (Zollinger et al. 1994) and synaptosomes (Pittaluga et al. 1988). However, as synaptosomes contain both pre- and postsynaptic elements (Suresh and Dunaevsky 2015; Volgyi et al. 2018), these studies do not distinguish between pre- and postsynaptic release. Further supporting a neurotransmitter function, NAAG binds to N-methyl-D-aspartate (NMDA) receptors and group II metabotropic glutamate receptors (mGluRs) (Valivullah et al. 1994; Schaffhauser et al. 1998; Shave et al. 2001). In particular, NAAG activates mGluR3 (Wroblewska et al. 1997; Neale 2011), which are localized on presynaptic nerve terminals (Sanabria et al. 2004) and inhibit the release of glutamate (Di Iorio et al. 1996; Zhong et al. 2006). This suggests that NAAG may function as a modulating neurotransmitter. The remaining question is from which synaptic compartment NAAG is released. Immunocytochemistry places NAAG in both principal neurons and interneurons in the brain (Anderson et al. 1986; Frondoza et al. 1990; Moffett et al. 1993; Tsai et al. 1993), but apart from one study in the retina (Williamson and Neale 1988), the subcellular localization of NAAG in the central nervous system has not been studied. Therefore, it is not known which neuronal compartment contains the highest concentration of NAAG, which compartment is responsible for the release of NAAG in the brain, or the release mechanism, or whether NAAG exhibits retrograde signaling. To address these questions, we used high resolution postembedding immunogold cytochemistry to reveal the in vivo localization of NAAG in the hippocampus. In hippocampal brain slices, we studied the redistribution of NAAG in response to neuronal stimulation, as well as its sensitivity for botulinum toxin B and dependency on Ca2+.

Materials & Methods
Preparation and Incubation of In Vitro Hippocampal Slices
Acute hippocampal slices were obtained and treated largely as previously described (Holten et al. 2008). Adult Wistar rats were anesthetized with halothane or isoflurane (Abbott), decapitated, and the brains were put in ice-cold normal Krebs’ solution (in mM): 125.5 NaCl (Sigma-Aldrich, S7653), 2.5 KCl (Sigma-Aldrich, P9541), 1.0 NaH2PO4 (Sigma-Aldrich, S8282), 24 NaHCO3 (Sigma-Aldrich, S5761), 2.5 CaCl2 (Sigma-Aldrich, C5670), 2.0 MgSO4 (Sigma-Aldrich, M7506), and 15 mg glucose (Sigma-Aldrich, D9434). Hippocampi were dissected out, sliced (300 μm), and incubated for 45 min at 30 °C (after preincubations at 30 °C for 60 min) in oxygenated (95% O2, 5% CO2) normal (2.5 mM K+), or depolarizing (55 mM K+) solutions. Some of the slices were incubated at 2.5 mM K+ in the presence of a cocktail of GluR agonists (500 μM (S)-α-amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid (AMPA) (Tocris Bioscience, 83643-88-3), 100 μM cyclothiazide (Sigma Aldrich, 2259-96-3), 100 μM N-methyl-D-aspartic acid (NMDA) (Tocris Bioscience, 6384-92-5) and 100 μM (S)-3,5-dihydroxyphenylglycine (DHPG) (Tocris Bioscience, 162870-29-3). To reveal the mechanism of NAAG release, other slices were incubated with GluR agonists (in 2.5 mM K+) in the absence of calcium, or in the presence of botulinum toxin B (BoNT/B, 133 nM) (Sigma Aldrich, 93384-44-2). In the latter experiments, the slices were preincubated for 120 min with or without BoNT/B (133 nM) to increase the penetration of the toxin into the tissue before the incubation.

The slices were kept on a nylon mesh in a glass beaker with a continuous flow of O2 in the medium. After incubation, the slices were fixed for 1 h at room temperature (~22 °C) in a mixture of 5% 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide (EDC) (Thermo Fisher, 22981) and 1 mM N-hydroxysuccinimide (NHS) (Sigma-Aldrich, 130672) in 10 mM (4-[(2-hydroxyethyl)-1-piperazinethanesulfonic acid (HEPES) (Sigma-Aldrich, H7006) buffer (pH 7.4) or 30 min at room temperature in a mixture of 5% EDC and 1 mM NHS in 10 mM HEPES buffer, followed by 1 h at room temperature in a mixture of 5% EDC, 1 mM NHS, 1% formaldehyde (FA) (Sigma-Aldrich, 158127), and 0.5% glutaraldehyde (GA) (AGAR scientific, R1010) or 2.5% GA and 1% FA in 0.1 M phosphate buffer (Sigma-Aldrich, S9638 and S7907) for 1 h at room temperature. The slices were kept overnight at 4 °C in the same fixative and then stored in the fixative diluted at 1:10.

In Vivo Perfusion Fixation
Adult Wistar rats were anesthetized by injection of pentobarbital (100 mg/kg, i.p.) and fixed by perfusion through the heart with 4% EDC plus 4% FA and 0.5% GA in 10 mM HEPES or 4% EDC, 1 mM NHS, and 1% dimethyl sulfoxide in 0.9% NaCl using a peristaltic pump.

Light Microscopic Immunocytochemistry
Slices used for staining of NAAG were allowed to sink in 30% sucrose (Sigma-Aldrich, S7903) before sectioning at 20 μm on a freezing microtome. Slices for glutamate staining were preincubated for 120 min with or without BoNT/B (133 nM) to increase the penetration of the toxin into the tissue before the incubation.
by the antibodies. The labeling was performed according to a three-layer biotinylated antibody-streptavidin-biotinylated peroxidase method (Hsu et al. 1981). High-resolution digital images of the NAAG staining were obtained using an automated Olympus BX52 microscope equipped with a 20× objective (Olympus Uplan Apo, NA 0.70), a motorized stage (LEP MAC5000, LUDL Electronic Products Ltd, Hawthorne, NY, USA), an Optronics MicroFire digital camera (Optronics Goleta, CA, USA), and NeuroLucida 7.0 Virtual Slice software (MBF Bioscience, Inc, Williston, VT, USA).

High-resolution digital images of the glutamate staining were obtained using an automated slide scanner system (Axioskop Z1, Carl Zeiss Microlmaging, Jena, Germany) with a 20×/0.8 plan apochromat objective and an Orca Flash 4.0 camera (Hamamatsu Photonics, Japan). Slides were scanned with extended depth of focus (EDF) for enhanced visualization of details from several focal planes.

Postembedding Electron Microscopic Immunocytochemistry

Slices from each experimental condition described above and specimens from perfusion-fixed animals were cryoprotected in glycerol (Sigma-Aldrich, G5516), quickly frozen in liquid propane, freeze-substituted with methanol, embedded in Lowicryl HM20 (polysciences), and further processed for immunogold labeling (Bergersen et al. 2008). Ultrathin sections (80–90 nm) were cut and mounted on nickel grids or formvar-coated one-hole grids. The slices were sectioned perpendicularly to the surface of the slice. Then the sections were processed in single labeling experiments with the NAAG antibodies (dilution 1:300, 1:500, or 1:600) or in double labeling experiments with NAAG (1:300) and GABA (1:100) or glutamine synthetase (GS) (BD transduction laboratories, #84458, 0.3 μg/mL) antibodies (Bergersen et al. 2008). In the double labeling experiments with GABA, the sections were first treated with the NAAG antibodies and then with the GABA antibodies to visualize NAAG in GABA terminals. Since both these primary antibodies were from rabbit, the sections were treated with FA vapor at 80°C for 1 h before the NAAG labeling and the GABA labeling to prevent interference between the sequential labelings (Wang and Larsson 1985; Ottersen et al. 1992). A secondary antibody coupled to gold particles with a diameter of 10 nm was used in the first step (NAAG), and a secondary antibody with 15 nm gold particles was used in the second step (GABA). In the double labeling with GS, the sections were treated with the NAAG and GS antibodies at the same time.

Sera and Specificity Controls

The NAAG antiserum was raised in rabbits against NAAG bound to thyroglobulin by EDC (Cangro et al. 1987) and purified by adsorption on a series of agarose columns bearing sepharose-EDC-thyroglobulin and -BSA. Both the 990 GABA and the 607 glutamate antisera were raised in rabbits as described (Storm-Mathisen et al. 1983) by immunizations with GA–FA–BSA conjugates of GABA and glutamate, respectively. The GABA antiserum (Gundersen et al. 2004) and the glutamate antiserum (Ottersen et al. 1990; Gundersen et al. 1991) have been characterized previously and proven to be specific. As a precaution against cross-reactivities, soluble GA–FA–BSA conjugates of GABA and glutamate, respectively. The GABA antiserum (Gundersen et al. 2004) and the glutamate antiserum (Ottersen et al. 1990; Gundersen et al. 1991) have been characterized previously and proven to be specific. As a precaution against cross-reactivities, soluble GA–FA complexes of GABA and glutamate (Sigma-Aldrich), glutamate (Sigma-Aldrich), and β-alanine (0.2 mM) (Sigma-Aldrich, A9920) were added to the diluted GABA antibodies, and soluble GA–FA complexes of aspartate, asparagine, and glutamine were added to the glutamate antibodies 3 h before applying it to the sections. The selectivity of the NAAG antibodies was tested before use, and selectivity controls were included in each individual experiment. As an intrinsic selectivity control for the light microscopic NAAG immunolabeling, conjugates of different acetylated molecules and carboxylates linked to brain macromolecules by EDC were spotted on cellulose nitrate–acetate filters, and these were processed together with the tissue sections. The NAAG antibodies selectively stained the NAAG spot (see Supplementary Fig. S1). Furthermore, the NAAG immunoreactivity of tissue sections and dot blots was blocked by adding 1.0 mM EDC-treated NAAG (Sigma-Aldrich A9590) to the NAAG antibodies (see Supplementary Fig. S1). As an in-experiment specificity test, some tissue sections were exposed to antibodies that had been pretreated with soluble NAAG EDC complexes for at least 3 h prior to immunolabeling. These preabsorption tests were performed along with both peroxidase (see Supplementary Fig. S1) and immunogold tissue labelings (Fig. 1 and Supplementary Fig. S2). The preabsorbed immunoperoxidase produced very low immunoreactivities in all experiments. As the immunogold method was performed on an ultrathin section after embedding in plastic-like resins, the possible problem of uneven antibody penetration between different subcellular structures is eliminated, as only molecules on the section surface are labeled.

Quantitative Immunogold Analysis

The ultrathin sections were imaged in a FEI Tecnai 12 electron microscope. Electron micrographs were taken along random trajectories across the thickness of the slice in the stratum radiatum of CA1 and in the dentate gyrus at ×26,500–43,000 magnification. The BoNT/B-treated slices were cut perpendicular to the section surface so that the observations could be made at defined distances from the slice surface. BoNT/B is a relatively large protein, which restricts its diffusion into the tissue. The electron micrographs were taken within 30 μm from the surface of the slice. Here, the texture of the tissue was loose, permitting penetration of BoNT/B. In the CA1 hippocampus NAAG immunogold particles were quantified in terminals making asymmetrical synapses, in their postsynaptic spines, and in dendritic shafts. In the dentate gyrus NAAG immunoreactivity was quantified in mossy fiber terminals, as well as in their dendritic thorns, and in GABA positive nerve terminals and their postsynaptic compartments (granule cell somata and dendritic shafts). For slice experiments background NAAG labeling was calculated in sections labeled with NAAG antibodies that were preneutralized with soluble NAAG–EDC complexes for all tissue profiles included in the quantitative analysis and subtracted from the averages of the immunogold labeling produced by the NAAG antibodies. Quantification from perfusion-fixed tissue labeled either with NAAG antibodies or with NAAG antibodies preneutralized with soluble NAAG–EDC complexes is shown side-by-side in Figure 1 and Supplementary Figure S2. The areas of the various cellular and subcellular elements were recorded by ImageJ (http://rsb.info.nih.gov/ij/) and the coordinates were submitted to a program written in Python (http://www.python.org) for computation of particle densities (number of gold particles per μm²; (Larsson et al. 2005). The source code of ImageJ plugin and the Python program is available at https://old.liu.se/medfak/forskning/larsson-max/software?1=en. The association between gold particles and postsynaptic vesicle-like structures was measured as the distance from the center of the vesicle-like structure to center of the gold particle using ImageJ, compared to the distance from random points (generated by the
Figure 1. NAAG is predominantly located in postsynaptic dendritic spines/thorns. Subcellular structures from different cell types were evaluated in perfusion-fixed rat hippocampus by electron microscopy to investigate in which structures NAAG is found above background levels. This was performed in three different labeling experiments, with or without double labeling with GS (astrocyte marker). Therefore, the data were standardized to the Schaffer collateral terminals in each rat using z-transformation (A). The bars represent standardized mean and standard deviation. NAAG labeling was compared with the labeling in the same structures with NAAG antibodies preabsorbed with NAAG complexes. The latter gives a measure of the background, unspecific, labeling produced by the antibodies and was low in all compartments. Most of the nonsynaptic structures, however, were at the same low level, indicating low or nonexistent NAAG levels. The data were used qualitatively to determine which areas to focus on in the more detailed analysis, and hence no statistical analysis was performed. The NAAG staining appeared above background only in Shaffer collateral spines (B vs. C) and mossy fiber thorns (B vs. G). Electron micrographs of perfusion-fixed tissue show that the density of gold particles signaling NAAG was much higher in dendritic spines (sc spine) forming excitatory synapses with Schaffer collateral terminals (sc term) (C) than in their presynaptic counterpart. The same was true for the dendritic thorns (mf thorn) compared to the mossy fiber terminals (mf term) forming excitatory synapses in dentate gyrus (E). Scale bars = 100 nm.
from hippocampal slices incubated at physiological (2.5 mM) K+ concentrations revealed that the most prominent NAAG labeling was in postsynaptic dendritic spines of CA1 excitatory synapses (Fig. 2A,D) and dendritic thorns of mossy fiber synapses (Fig. 2E,H), while their corresponding nerve terminals (Schaffer collateral terminals and mossy fiber terminals, respectively) (Fig. 2A,D,E,H) and CA1 stem dendrites (Fig. 2I) displayed much weaker labeling (P < 0.01, one-way ANOVA, Tukey's multiple comparisons test, GraphPad). To determine if NAAG could be released from spines/thorns, we exposed hippocampal slices to stimulation by high (55 mM) K+ or GluR agonists (100 μM DHPG, 100 μM NMDA, and 500 μM AMPA stabilized by 100 μM cyclothiazide). During K+ or GluR agonist-induced stimulation, immunogold labeling of NAAG was selectively depleted from the postsynaptic Schaffer collaterals (Fig. 2B,C vs. Fig. 2A, P < 0.01, one-way ANOVA, Tukey's post hoc test for multiple comparisons, GraphPad) and mossy fiber thorns (Fig. 2F,G vs. Fig. 2E, P < 0.01, one-way ANOVA, Tukey's multiple comparisons test, GraphPad). The stimulation did not affect the density of NAAG gold particle in the nerve terminals of these synapses (Fig. 2B,C,F,G, P > 0.05, one-way ANOVA, Tukey's multiple comparisons test, GraphPad), or in dendritic shafts (Fig. 2I–K, P > 0.05, one-way ANOVA, Tukey's multiple comparisons test, GraphPad).

NAAG is Located to Postsynaptic Vesicles

NAAG was localized at high densities in dendritic spines of the Schaffer collateral synapses (Fig. 3A,B) and thorns of the mossy fiber synapses (Fig. 3A,C). Interestingly, in the postsynaptic dendritic spines/thorns NAAG was associated with small, round, and clear structures (insets in Fig. 3B,C) resembling what we have previously observed for neurotransmitter labeling in synaptic vesicles in nerve terminals (Bergersen et al. 2008; Morland et al. 2013). The vesicle-association of NAAG gold particles was demonstrated statistically by comparing the proportion of NAAG gold particles that were located in close vicinity to a vesicle. To an expected distribution (random points inserted in the same image). We demonstrate that 90.6% and 89.7% of the NAAG particles are localized within 21 nm from the center of the vesicle in Schaffer collateral spines and mossy fiber thorns, respectively (Fig. 3D,E). This proportion was significantly higher than what was found for random points, where 50.1% and 49.8% of the particles were located within 21 nm from the center of the closest vesicle for the two postsynaptic compartments, respectively (Fig. 1C,F), (P = 9.1 × 10−43 for Schaffer collateral spines and P = 8.9 × 10−50 for mossy fiber thorns; Pearson's chisquare test (Fig. 3G,F).

Characterization of Postsynaptic NAAG Depletion

The accumulation of NAAG labeling over postsynaptic vesicles prompted the question whether NAAG underwent vesicular release from the postsynaptic spines/thorns. To elucidate the mechanism whereby NAAG escapes from dendritic spines/thorns, we stimulated slices with GluR agonists and inhibited exocytosis either by removing Ca2+ from the extracellular medium or by treatment with BoNT/B. BoNT/B blocks vesicular fusion by cleaving VAMP2 (Pellizzari et al. 1996), a SNARE protein reported to regulate postsynaptic exocytosis (Jurado et al. 2013). To determine the efficacy of BoNT/B in our set-up, we first evaluated the effect of BoNT/B on glutamate depletion during membrane stimulation. In hippocampal slices depolarized with

Results

Statistics

The results were statistically evaluated (GraphPad) by the Students t-test when only comparing two groups, or with one-way ANOVA, Tukey’s post hoc test for multiple comparisons, to investigate differences in mean gold particle densities between individual experimental groups. In Figure 1, three rats, from three different staining experiments and with two different fixation regimes, have been standardized to the levels in Schaffer collateral terminals in each rat, using z-transformation. Statistical tests between perfusion-fixed brains labeled with NAAG antibodies, the same brains labeled with NAAG antibodies that had been preabsorbed with NAAG complexes, were not performed, as the latter is represented by two animals. The raw values (not standardized) are shown in Supplementary Figure S2. The distribution of distances between NAAG immunogold particles or random points and the closest vesicle-like structure was first converted to categorical data using the maximal theoretical distance of 21 nm as a cut-off. The difference in vesicle-association (portion of particles within 21 nm from the center of the vesicle) between NAAG immunogold particles and random points was analyzed by the Pearson’s chi-squared test (Microsoft Excel).

Results

NAAG is Located to Postsynaptic Vesicles in the Brain

To unravel the subcellular compartment that contains the highest levels of NAAG, we performed peroxidase labeling. NAAG immunoreactivity was located in cell bodies of CA1 and CA3 pyramidal cells, granule cells, and cells in the dentate gyrus (see Supplementary Fig. S1A). It was evident that the NAAG staining appeared in parts of the dendritic tree of these cells, in addition to very small dots lining the dendritic trees (see Supplementary Fig. S1C,D), as expected for substances that have a synaptic localization. This synaptic-like labeling was completely absent in the hippocampal section labeled with the same antibodies preabsorbed with soluble NAAG–EDC complexes in excess (see Supplementary Fig. S1B,D,E).

Electron microscopy revealed high levels of NAAG in postsynaptic spine of the Schaffer collateral synapses in stratum radiatum, the postsynaptic thorns of the mossy fiber synapses in the dentate gyrus, and stem dendrites (Fig. 1). Other compartments, including presynaptic Schaffer collateral terminals and mossy fiber terminals, as well as perivascular astrocytes, myelin, and axons showed staining that was undistinguishable from the background (Fig. 1). Hence, we focused our further ultrastructural analysis on Schaffer collateral synapses in the hippocampal stratum radiatum and the mossy fiber synapses of the dentate gyrus.

The predominant localization of NAAG in postsynaptic spines/thorns opens up the possibility that NAAG is released as a retrograde transmitter from the postsynaptic compartment.

NAAG is Depleted From Dendritic Spines/Thorns During Neuronal Activation

As in the perfusion-fixed tissue, electron microscopic analysis from hippocampal slices incubated at physiological (2.5 mM) K+...
Figure 2. NAAG is depleted upon depolarization with high potassium or GluR agonists. Electron micrographs from slice experiments (A and E) confirmed what was found in the perfusion-fixed tissue: During basal conditions (2.5 mM K+), gold particles signaling NAAG were mainly present in dendritic spines (sc spine) and thorns (mf thorn) forming asymmetric synapses with Schaffer collateral terminals (sc term) or mossy fiber terminals (mf term). In response to membrane depolarization by K+ (55 mM; B, F and J) or by GluR agonists (C, G and K), NAAG immunogold particles were depleted from the postsynaptic sc spines (B; C compared to A) and mf thorns (F; G compared to E), without any change in the corresponding terminals or stem dendrites (J; K compared to I). Bar charts show the density of NAAG immunogold in Schaffer collateral presynaptic terminals (sc term) and postsynaptic spines (sc spine) (D) and stem dendrites (stem dend) (L) in the stratum radiatum, and in mossy fiber terminals (mf term) and thorns (mf thorn) of the dentate gyrus (H) in hippocampal slices incubated at 2.5 mM K+ (black bars), depolarizing conditions (55 mM K+; gray bars) or GluR agonists (white bars). The bars represent the average number of NAAG immunogold particles/μm2 ± SD. ** shows the difference from the same profile at basal conditions (P < 0.01, one-way ANOVA, Tukey’s multiple comparisons test, GraphPad). The total numbers of tissue profiles included in the quantifications were as follows (basal-K+–GluR): Schaffer collateral terminals and spines, 95-104-87; stem dendrites, 61-64-66, mossy fiber terminals, 62-69-61; and mossy fiber thorns, 174-156-180 (two Wistar rats, n = three slices in each condition).

K+ (55 mM) BoNT/B clearly inhibited the depletion of glutamate from nerve terminals (see Supplementary Fig. S3). Thus, having tested that the BoNT/B concentration used in our experiments was sufficient to block exocytosis, we proceeded to analyzing the effect of BoNT/B on the depletion of NAAG. In hippocampal slices stimulated with GluR agonists, BoNT/B or zero Ca2+ blocked depletion of NAAG from postsynaptic CA1 dendritic spines (Fig. 4B–E) and mossy fiber thorn synapses (Fig. 4F–I). Inhibiting exocytosis by BoNT/B did not affect the content of NAAG immunogold particles in the nerve terminals (Fig. 4D,H) or in dendritic shafts (data not shown) during membrane stimulation. Interestingly, in postsynaptic spines/thorns subjected to membrane depolarization combined with inhibited exocytosis (in particular BoNT/B treatment), the NAAG signal was increased above resting values (Fig. 4B vs. Fig. 4D; Fig. 4J,F vs. Fig. 4H,K), P < 0.01, one-way ANOVA, Tukey’s multiple comparisons test, GraphPad.

Postsynaptic Accumulation of NAAG and Depletion Upon Depolarization Did Not Occur in GABA Synapses

In GABAergic nerve terminals, K+–induced membrane depolarization led to depletion of GABA (P < 0.05, unpaired t-test, two tails, GraphPad) as predicted (Gundersen et al. 2004), but there was no depletion of NAAG (P > 0.05, unpaired t-test, two tails, GraphPad) (Fig. 5). Hence, postsynaptic release of NAAG appears to be restrained to the glutamatergic system.

High Ratio of NAAG Between Post- and Presynaptic Compartments was Consistent Between Experimental Conditions

Although the staining intensity may vary between different staining experiments, the staining pattern was stable. The high ratio between sc. spines and sc. terminals was high both from perfusion-fixed rat hippocampus 4.98 (95% confidence interval
Figure 3. NAAG is located in vesicular structures in both postsynaptic dendritic spines in stratum radiatum (A, left) and in postsynaptic mossy fiber thorns in dentate gyrus (A, right). In the dendritic spines and thorns, gold particles representing NAAG were closely related to vesicular structures, which in size and shape resemble presynaptic vesicles (insets in B and C). Scale bars in (B and C): 100 nm; scale bar in insets: 10 nm. The distribution of distances of gold particles and the closest vesicle-like structures was quantified. In both postsynaptic compartments, the sc spines and the mf thorns, a large proportion of the NAAG particles were localized within 21 nm from the center of a vesicle (D and F). For comparison, for the randomly inserted points (E and G), only about half of the points were found within the same range from the vesicle ($P = 9.1 \times 10^{-43}$ for sc spines; and $P = 8.9 \times 10^{-50}$ for mf thorns; Pearson chi square test). 21 nm represents the theoretical distance between the center of the gold particle and the epitope (the radius of the gold particle [5 nm] + the length of the primary-secondary complex).
Figure 4. NAAG is depleted by exocytosis from postsynaptic dendrites. The schematic drawing (A) shows the proposed mechanism of action and regulation of NAAG release [numbers in bold indicate mechanisms studied by us]. Presynaptic membrane fusion of transmitter-containing vesicles is mediated by members of the SNARE family: the v-SNARE, target membrane SNARE (t-SNARE), and synaptosomal-associated protein (SNAP), in addition to Ca\(^{2+}\) sensors. We investigated whether the same mechanism is responsible for the postsynaptic NAAG release. Glutamate released from the nerve terminal stimulates dendritic ionotropic glutamate receptors (GluR) (1), which triggers Ca\(^{2+}\)-dependent (2) and v-SNARE-dependent (3) release of NAAG. NAAG then acts on presynaptic mGluR3/postsynaptic NMDAR (mixed agonist/antagonist), the former inhibiting Glu release from the nerve terminal. The electron micrographs (B–I) show how inhibiting exocytosis during membrane depolarization affects the depletion of NAAG immunogold particles from postsynaptic dendritic spines/thorns. During basal conditions (2.5 mM K\(^{+}\)), gold particles representing NAAG were present postsynaptically in dendritic spines and thorns belonging to Schaffer collaterals (sc spine) and mossy fibers (mf thorn) and with very low levels in the presynaptic terminals (sc term and mf term). When postsynaptic depolarization was elicited by GluR agonists, NAAG was depleted from the postsynaptic spines (C) and thorns (G). When v-SNARE was degraded by BoNT/B, NAAG was trapped postsynaptically despite GluR agonist activation (D and H). Finally, in the absence of Ca\(^{2+}\) (zero Ca\(^{2+}\)), NAAG was also trapped postsynaptically despite the presence of GluR agonists (E and I). Scale bars, 100 nm. Quantitative results of NAAG immunogold particles in different tissue profiles are shown in (J) for the Schaffer collateral synapses of the stratum radiatum and (K) for the mossy fiber synapses of the dentate gyrus. Resting conditions: 2.5 mM K\(^{+}\) (basal, black bars), GluR agonists (GluR ag., white bars), GluR agonists in the presence of botulinum toxin B (GluR ag BoNT/B, light gray bars), and GluR agonists in the absence of Ca\(^{2+}\) (GluR ag zero Ca\(^{2+}\), dark gray bars). The bars represent the average number of NAAG immunogold particles/μm\(^2\) ± SD in Schaffer collateral spines (sc spines) and nerve terminals (sc term) belonging to Schaffer collateral synapses in the stratum radiatum (J), and thorns (mf thorns) and mossy fiber terminals (mf term) belonging to hilar mossy fiber synapses (K). ∗∗: P < 0.01, one-way ANOVA, Tukey’s multiple comparisons test, GraphPad. The numbers of tissue profiles per slice that was included in the quantifications were as follows: sc spines and sc terminals: 20–26 (basal), 23–29 (GluR agonists), 19–25 (GluR agonists zero Ca\(^{2+}\)), 19–25 (GluR agonists BoNT/B); mf thorns: 37–63 (basal), 41–71 (GluR agonists), 53–57 (GluR agonists zero Ca\(^{2+}\)), 47–73 (GluR agonists BoNT/B); mf terminals: 19–25 (basal), 20–29 (GluR agonists), 20–21 (GluR agonists zero Ca\(^{2+}\)), 19–22 (GluR agonists BoNT/B), (slice experiment from two rats (pooled), n = 4–5 slices in each condition).
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Figure 5. NAAG is neither released from GABA-positive nerve terminals nor their postsynaptic sides. In GABAergic synapses in the molecular layer in dentate gyrus of the hippocampal formation (A), NAAG was neither depleted from GABA-positive nerve terminals nor their postsynaptic sides (B and C) in response to depolarization. GABA synapses are colabeled for NAAG (small gold particles, arrowheads) and GABA (large gold particles). The figure shows representative electron micrographs of a slice incubated at 2.5 mM K⁺ (B) or at 55 mM K⁺ (C). Labeled nerve terminals (GABA term) contact granule cell bodies (postsynaptic side) (B and C). Scale bars, 100 nm. Bar charts show the density of NAAG (D) and GABA (E) labeling (average number of gold particles/μm² ± SD, n = 4 basal slices and six depolarized slices, slice experiment from one adult Wistar rat) in nerve terminals containing GABA gold particles (GABA terminal) and the opposing postsynaptic elements (granule cell bodies or proximal dendrites [postsynaptic]). Asterisk indicates that the GABA values in terminals at 55 mM K⁺ were significantly different from those in terminals at 2.5 mM K⁺ (P < 0.05, unpaired t-test, two tails, GraphPad). In contrast, NAAG labeling in GABA terminals and postsynaptic element was unaffected by depolarization (P > 0.05, unpaired t-test, two tails, GraphPad). The total numbers of tissue profiles included in the quantifications were as follows: Basal GABA terminal, 95; depolarized GABA terminal, 138; basal postsynaptic elements, 64; depolarized postsynaptic element, 99.

Discussion

The present study shows that NAAG is predominantly localized to postsynaptic compartments, with background levels in other neuronal compartments, including in presynaptic terminals.
NAAG immunogold particles were spatially related to vesicular structures in these profiles; such vesicles have previously been described in postsynaptic spines in the hippocampus (Spacek and Harris 1997). In acute hippocampal slices, we demonstrate that NAAG is depleted from the spines in response to neuronal stimulation. The finding of high postsynaptic NAAG-content is very robust, with comparable post-to-presynaptic ratios in rat hippocampus fixed in vivo and acute hippocampal slices fixed after 2 h in vitro. The same goes for the depletion of postsynaptic NAAG content in response to depolarization. The depletion was dependent on Ca\(^{2+}\) and sensitive to BoNT/B, strongly suggesting that NAAG undergoes vesicular release from the postsynaptic spine and acts as a retrograde transmitter in the brain. The observation that inhibited exocytosis by BoNT/B during depolarization gave postsynaptic NAAG levels above what was found in the resting condition suggesting a basal release of NAAG. The NAAG content in stem dendrites is not altered in the experimental conditions tested in this study.

The presence of BoNT/B, which cleaves the vesicular SNARE (v-SNARE) VAMP2, completely blocked depletion of NAAG from postsynaptic spines/thorns. This is in line with a recent report showing that VAMP3 is important for regulated postsynaptic exocytosis (Jurado et al. 2013) and strongly suggests that NAAG undergoes VAMP-mediated exocytosis from postsynaptic elements. How BoNT/B is internalized in the postsynaptic element is not totally clear, but gangliosides and other proteins in the postsynaptic membrane have recently been shown to mediate internalization of BoNTs (Rummel 2013). For instance, the ganglioside GD1a seems to be especially important for binding BoNT/B (Ochanda et al. 1986). Furthermore, translocation of the BoNT/B light chain requires a pH of 4.5–6 (Pirazzini et al. 2013). The lumen of recycled synaptic vesicles, with an estimated pH of ∼5.8, (Miesenbock et al. 1998), therefore provides an appropriate environment where light chain translocation can be induced. Therefore, a postsynaptic effect of BoNT/B after extracellular application of the drug could be expected. Concentration curves for the extracellular concentration of BoNT/B that effectively inhibits Ca\(^{2+}\)-dependent exocytosis from postsynaptic dendritic spines are not available. The concentration used in the present study (133 nM) is one-third higher than the concentration previously shown to effectively inhibit Ca\(^{2+}\)-dependent glutamate exocytosis from synaptosomes (100 nM) (McMahon et al. 1992).

Indeed, a postsynaptic localization of VAMP2 has been reported (Hussain and Davanger 2015) and that SNARE proteins are located in dendritic “synaptic-like vesicles” (Hussain and Davanger 2015; Hussain et al. 2017; 2019). The dendritic vesicles observed by us are morphologically similar to synaptic vesicles (Fig. 1D,E) and similar to those presented by Hussain and Davanger (2015) and Hussain et al. (2017), (2019). Supporting vesicular accumulation of NAAG, a vesicular transport mechanism for NAAG through the sialic acid transporter sialin has been suggested (Lodder-Gadaczek 2013). This supports the idea that NAAG can act as a retrograde transmitter through binding to one or both of these receptors. One possibility is that NAAG provides the postsynaptic neuron with the ability to control its own inputs both at high and low activation levels. When the spine/thorn receives very low glutamate signals, low levels of NAAG are released, consistent with the increased NAAG levels found in the spines/thorns after exposure to BoNT/B during GluR depolarization. When the glutamate signal is low, NAAG has been shown to activate the synaptic, GluN2A-containing NMDARs, while inhibiting the extrasynaptic GluN2B-containing NMDARs (Khacho et al. 2015), thereby enhancing the synaptic glutamate signal. In contrast, in situations of extensive glutamatergic signaling and a subsequent lowering of tissue pH, NAAG released from the postsynapse could reduce excitotoxicity by inhibiting both intra- and extrasynaptic NMDARs. The activation of mGluR3 would further strengthen this neuroprotective effect by reducing glutamate release from the nerve terminal. In support of this hypothesis, knock-out or pharmacological inhibition of glutamate carboxypeptidase II, which breaks down synthetically released NAAG (Robinson and Coyle 1987; Stauch et al. 1988; Berger et al. 1999; Bergeron et al. 2007), is protective in animal models of ischemia (Slusher et al. 1999; Tortella et al. 2000; Jackson et al. 2001; Williams et al. 2001; Bacich et al. 2005; Long et al. 2005; Zhang et al. 2016) and epilepsy (Witkin et al. 2002). With this said, we should emphasize that our means of inducing membrane depolarization is not physiologic. Thus, the exact effects of NAAG release from postsynaptic dendrites during physiological conditions must await further studies.

A neurotransmitter role of NAAG has long been suggested, despite lack of direct evidence for presynaptic localization and release. Such evidence could only be obtained by immune-electron microscopic detection of NAAG in stimulated and unstimulated brain tissue. The commonly used aldehyde fixatives preserve good morphology for ultrastructural studies. NAAG, however, lacking free amine groups, is not fixed by aldehydes (Moffett et al. 1993). For fixation of NAAG, the less common, water soluble carbodiimide fixation agent EDC must be used. It acts by forming amine bonds between carboxyl groups or primary amine groups (Bauminger and Wilchek 1980). Despite this, a previous study (Renno et al. 1997) investigated the ultrastructural localization of NAAG in the
cerebellum, using only aldehyde fixation. This strongly suggests that their NAAG labeling was not selective.

Using EDC fixation, selective detection of NAAG is possible, but EDC fixation produces inferior morphology compared to tissue fixed with aldehydes. Further adding to a suboptimal morphologic result is that we used tissue embedding in an acrylic resin (Lowicryl HM20) over an epoxy resin. Low temperature embedding with Lowicryl HM20 preserves the antigens to a large extent and results in a superior immunogold sensitivity, but the ultrastructure is inferior compared with epoxy-embedded tissue (van Lookeren Campagne et al. 1991; Newman and Hobot 1999). Our group has successfully used Lowicryl HM20 in immunogold cytochemistry for many years (Bergersen et al. 2008 Nat Prot). In the present study, the morphology of the perfusion-fixed tissue (Fig. 1D,E) was better compared to the morphology of the immersion-fixed tissue from acute brain slices (Figs 2A–CE–GI–K, 3B,C and 48–I). Perfusion fixation gives a rapid and uniform penetration of the fixative into the tissue, reducing the risk of ischemic damage. This method is therefore the preferred method for morphology preservation (Gonzalezaguilar and Derobertis 1963). For in vitro experiments, like acute brain slices, immersion fixation must be performed, which gives a slower tissue fixation. Thus, some morphological changes are known to happen during slice incubation (Schurr et al. 1984; Fiala et al. 2003), although all precautions were made to preserve the morphology (Schurr et al. 1984; Fiala et al. 2003). However, in the present material, pre- and postsynaptic elements were clearly visible, making the identification of nerve terminals and postsynaptic dendritic spines/thorns reliable. To ensure that we could distinguish between pre- and postsynaptic elements, we double-labeled some tissue for NAAG and glutamate and could distinguish between pre- and postsynaptic elements, postsynaptic dendritic spines/thorns reliable. To ensure that clearly visible, making the identification of nerve terminals and the morphology (Schurr et al. 1984; Fiala et al. 2003), although all precautions were made to preserve the risk of ischemic damage. This method is therefore the preferred method for morphology preservation (Gonzalezaguilar and Derobertis 1963). For in vitro experiments, like acute brain slices, immersion fixation must be performed, which gives a slower tissue fixation. Thus, some morphological changes are known to happen during slice incubation (Schurr et al. 1984; Fiala et al. 2003), although all precautions were made to preserve the morphology (Schurr et al. 1984; Fiala et al. 2003). However, in the present material, pre- and postsynaptic elements were clearly visible, making the identification of nerve terminals and postsynaptic dendritic spines/thorns reliable. To ensure that we could distinguish between pre- and postsynaptic elements, we double-labeled some tissue for NAAG and glutamate and consistently found that NAAG was localized in the postsynaptic compartment low in glutamate (unpublished results; KN, CM, VG).

Conclusion
The present study shows evidence for exocytotic postsynaptic release of NAAG, implying that fine-tuning of the glutamate signal occurs through retrograde rather than anterograde NAAG signaling. To our knowledge, this is the first demonstration of vesicular release of a transmitter exclusively from dendrites.

Supplementary material
Supplementary material is available at Cerebral Cortex online.

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