Seeding and transgenic overexpression of alpha-synuclein triggers dendritic spine pathology in the neocortex

Sonja Blumenstock¹²³, Eva F Rodrigues², Finn Peters², Lidia Blazquez-Llorca⁴, Felix Schmidt¹⁵, Armin Giese¹ and Jochen Herms¹²³,*

Abstract

Although misfolded and aggregated α-synuclein (α-syn) is recognized in the disease progression of synucleinopathies, its role in the impairment of cortical circuitries and synaptic plasticity remains incompletely understood. We investigated how α-synuclein accumulation affects synaptic plasticity in the mouse somatosensory cortex using two distinct approaches. Long-term in vivo imaging of apical dendrites was performed in mice overexpressing wild-type human α-synuclein. Additionally, intracranial injection of preformed α-synuclein fibrils was performed to induce cortical α-syn pathology. We find that α-synuclein overexpressing mice show decreased spine density and abnormalities in spine dynamics in an age-dependent manner. We also provide evidence for the detrimental effects of seeded α-synuclein aggregates on dendritic architecture. We observed spine loss as well as dystrophic deformation of dendritic shafts in layer V pyramidal neurons. Our results provide a link to the pathophysiology underlying dementia associated with synucleinopathies and may enable the evaluation of potential drug candidates on dendritic spine pathology in vivo.

Keywords alpha-synuclein; dendritic spines; in vivo imaging; seeding; synucleinopathies

Subject Category Neuroscience

DOI 10.15252/emmm.201607305 | Received 7 November 2016 | Revised 27 February 2017 | Accepted 2 March 2017 | Published online 28 March 2017

EMBO Mol Med (2017) 9: 716–731

Introduction

The mammalian brain contains a complex network of billions of neurons communicating through different types of synapses that can be severely and irreversibly disturbed by neurodegeneration. In synucleinopathies like Parkinson’s disease (PD) or dementia with Lewy bodies (DLB), progressive neurodegeneration is linked to misfolding and intracellular aggregation of the synaptic protein α-synuclein (α-syn; Clayton & George, 1999; Lücking & Brice, 2000) and leads to motor as well as cognitive deficits and ultimately to dementia (Spillantini et al., 1997; Burn, 2004; Aarsland et al., 2008). Familial cases of PD or DLB can be caused by mutations in the α-syn gene (Polymeropoulos et al., 1997; Krüger et al., 1998; Zarranz et al., 2004) or overexpression of α-syn (Simón-Sánchez et al., 2009) and strengthen the association between protein load/aggregation and the onset of disease. Although the filamentous, cytosolic α-syn inclusions called Lewy bodies (Spillantini et al., 1998) are a prerequisite for the histopathological diagnosis of PD and DLB, only an imperfect correlation between the Lewy body load and severity of cognitive impairment is known (Hughes et al., 1992; Parkkinnen et al., 2005). Several studies rather support synaptic failure as the predominant pathophysiological mechanism. Presynaptic neurotransmitter release has been proven to be unbalanced in PD (Nikolaus et al., 2009). Moreover, presynaptic α-syn oligomers and micro-aggregates are thought to be responsible for neurodegeneration as well as for dendritic spine loss observed in postmortem DLB brains (Kramer & Schulz-Schaeffer, 2007).

Over the past few years, an increasing body of evidence supports the concept of the propagation of misfolded α-syn species from neuron to neuron, where they are transported along axons and trigger the conversion of endogenous α-syn into a pathologic form in culture and in vivo (Luk et al., 2012a,b). Aspects of progressive α-syn aggregation—namely the spread along synthetically connected brain regions and the maturing of Lewy-like inclusions—can be recapitulated in model systems by exogenous introduction of in vitro-generated, preformed fibrils (PFFs) of α-syn, which under certain conditions is even possible in non-transgenic animals (Luk et al., 2012a,b; Masuda-Suzukake et al., 2013; Osterberg et al., 2015). Dendritic spines are small protrusions from the dendritic shaft which are highly regulated and have been shown to be altered in...
several neurological disorders including neurodegenerative diseases (Fiala et al., 2002; Penzes et al., 2011; Murmu et al., 2013). Various studies in the field of α-syn-related disorders have addressed the impairment of dendritic spines in brain areas like the striatum (McNeill et al., 1988; Day et al., 2006; Zaja-Milatovic et al., 2006; Finkelstein et al., 2016), the hippocampus (Winner et al., 2012), the olfactory bulb (Neuner et al., 2014), and naturally the substantia nigra (Patt et al., 1991). However, chronic effects of cortical α-syn have not been studied extensively, even though it is becoming more and more evident that the cortical involvement in the pathophysiology of PD and DLB cannot be neglected.

This study aims to extend the understanding of the structural consequences of neocortical α-syn accumulation on dendritic spines in a time-resolved manner. To address this question, we used chronic two-photon in vivo imaging to monitor spine dynamics over time, in young and aged transgenic mice overexpressing human wild-type α-synuclein (PDGF-h-α-syn) (Masliah, 2000). Furthermore, we used PFFs as a tool to induce cortical α-syn accumulation in mice expressing only the reporter transgene eGFP in order to study the structural consequences of α-syn seeding on dendritic architecture and spines.

Here, we report detrimental effects on dendrites and dendritic spines due to α-syn overexpression as well as seeding. Accumulation of α-syn in the neocortex triggers the loss of dendritic spines, interferes with spine dynamics, and leads to dystrophic malformation of dendritic branches. Our combination of a transgenic and a PFF-seeding mouse model provides new insights on the α-syn-dependent mechanisms that lead to dendritic spine decay in the cortex and therefore might create a better understanding of cognitive decline in synucleinopathies.

Results

Overexpression of wild-type human α-synuclein leads to changes in dendritic spine density and dynamics in aged mice

In vivo two-photon imaging was performed in three age groups of mice (3, 6, and 12 months), which were chosen according to previously published data on the progression of α-syn accumulation and its corresponding behavioral phenotype (Masliah, 2000; Rockenstein et al., 2002; Amschil et al., 2013). We consider the 3-month-old cohort as “presymptomatic”, whereas the 6- and 12-month-old cohorts show a developing and fully fledged phenotype, respectively.

Previous long-term in vivo imaging studies have shown that spines are highly dynamic structural elements and can be classified accordingly: Persistent spines are long lasting and stable for weeks up to the entire life span, whereas transient spines are newly formed and lost within days (Gruzdendorf et al., 2002; Trachtenberg et al., 2002; Holtmaat et al., 2005). In both h-α-syn transgenic and control mice, dendritic images were analyzed for spine density and dynamics. Newly appeared and disappeared spines relative to the previous imaging session were marked as “gained” or “lost”, respectively. Spines that were gained at a certain time point and persisted for <7 days were considered as being “transient” (Fig 1A).

Our image analysis in the 6- and 12-month-old cohorts revealed that the number of spines per μm was significantly lower in mice overexpressing h-α-syn. The ANOVA genotype main factor was significantly changed (6 M: F(1,7) = 27.11; P = 0.0012 and 12 M: F(1,7) = 55.78; P = 0.0001), denoting that h-α-syn mice exhibit considerably less (∼30%) dendritic spines on apical tuft dendrites but show no increase or decrease in spine density over the imaging time period relative to controls (Fig 1B).

Concerning spine dynamics, the fractions of both gained and lost spines were found to be elevated; however, only the comparison of the lost fractions reached statistical significance (6 M: F(1,7) = 5.85; P = 0.0461 and 12 M: F(1,7) = 8.86; P = 0.0206). As newly gained spines were disproportionally more often lost again, the fraction of transient spines showed a significant increase in both 6- and 12-month-old h-α-syn mice (6 M: F(1,7) = 7.19; P = 0.0315 and 12 M: F(1,7) = 24.25; P = 0.0017; Fig 1C).

As a consequence of the change in spine dynamics, the daily turnover ratios were found to be significantly elevated in h-α-syn mice (6 M: F(1,7) = 5.65; P = 0.0491 and 12 M: F(1,7) = 6.93; P = 0.0338; Fig 1D). In summary, we could show that spine density is significantly reduced and does not further change over the imaging interval of 35 days. Moreover, parameters of spine dynamics were profoundly changed in mice overexpressing α-syn compared to controls. The significant increase in transient spine fraction both in 6 and 12 months of age indicates that compensatory mechanisms may take place. Since there are no significant differences between 6 and 12 months of age, spine pathology does not further progress after 6 months of age but has reached a stable equilibrium.

Young adult α-synuclein overexpressing mice exhibit a different dynamic phenotype than aged mice

In order to reveal when the spine loss starts leading to reduced spine density in 6- and 12-month-old h-α-syn mice, we analyzed a 3-month-old cohort over 6 weeks. Different to the older cohort, we observed a progressive decrease in spine density, with a significant interaction between genotype and time in Bonferroni’s post hoc test (F(5,30) = 15.18; P < 0.001). Compared to controls, spine density in h-α-syn mice was reduced 12% on average at the first imaging time point (age of mice: 3 months) and 30% at the last imaging time point (age of mice: 4.5 months; Fig 2A).

Moreover, the dynamic fractions of spines show a somewhat contradictory picture compared to the 6- and 12-month-old mice. Our data revealed a significant impairment in the formation of new spines in h-α-syn mice (F(1,6) = 8.71; P = 0.0256), while the fraction of lost spines was unaffected, which accounts for the progressive decrease in spine density. By implication, stable spines represent a significantly larger fraction in these mice (F(1,6) = 8.71; P = 0.0256), while the fraction of transient spines is not significantly different from controls (Fig 2B). We also observed no significant change of the daily turnover ratio (TOR) between the groups at 3 months of age (Fig 2C).

Our results demonstrate that overexpression of human α-syn leads to structural alterations of dendritic spines in the cerebral cortex. The alterations develop already in young adults at the age of 3–4.5 months, significantly before behavioral or motoric phenotypes become apparent (Amschil et al., 2013), but reach a state of dynamic equilibrium later and do not progress further until 12 months of age.
Alterations in spine density and morphology in young adult and aged PDGF-h-α-syn mice

The progressive decrease in spine density seen in young adult mice could be monitored in vivo between 3 and 4.5 months of age. Extrapolation of the data curve (Fig 2A) of α-syn overexpressing animals to an earlier age would yield a time point at which no difference in synaptic spine densities between the groups is present. As chronic in vivo imaging in younger mice is technically not possible (bone growth in mice until 6–8 weeks of age, 4 weeks postsurgery necessary for chronic window clearing/healing), we addressed this question with a single time point ex vivo approach. Confocal

Figure 1. α-Synuclein overexpression alters spine density and dynamics in vivo.
A. Representative in vivo two-photon recordings of eGFP-labeled apical tuft dendrites in the somatosensory cortex in h-α-syn and control animals. Arrowheads mark representative spines that were stable (white, present > 7 days), newly formed (green), or lost (magenta). Gained spines that do not stabilize (yellow/green, present < 7 days) are defined as transient. Scale bar, 5 μm.
B. Spine density is reduced in both 6- (**P = 0.0012) and 12-month-old (***P = 0.0001) h-α-syn animals.
C. The fractions of both gained (*P6 months = 0.0527, P12 months = 0.0678) and lost spines (*P6 months = 0.0461, *P12 months = 0.0206) are elevated in h-α-syn mice compared to controls; the fraction of transient spines is significantly higher (*P6 months = 0.0335, **P12 months = 0.0017).
D. Consequently, the daily turnover ratio (TOR) is significantly increased in both 6- and 12-month-old h-α-syn mice (*P6 months = 0.0491, *P12 months = 0.0338).
Data information: n = 5 (h-α-syn), n = 4 (control) animals, mean ± s.e.m.; two-way ANOVA genotype main factor, *P < 0.05, **P < 0.01, ***P < 0.001.
Figure 2.

A 3 months

spine density [µm]

control h-α-syn

time [d]

B gained spines lost spines

fraction [%]

time [d]

C daily turnover

daily TOR

time [d]

D young adult PDGF-h-α-syn

spine density [µm]

2 mo 2 mo 4.5 mo 4.5 mo

ctrl h-α-syn ctrl h-α-syn

fraction [%]

mushroom stubby thin filopodia

E aged PDGF-h-α-syn

spine density [µm]

7.5 mo 7.5 mo 13.5 mo 13.5 mo

ctrl h-α-syn ctrl h-α-syn

fraction [%]

mushroom stubby thin filopodia
Alpha-synuclein mediates spine pathology  Sonja Blumenstock et al

**Figure 2.** α-Synuclein overexpression alters spine density, dynamics, and morphology differently depending on age.

A. In young, 3-month-old h-α-syn mice, progressive decrease in spine density is observed in vivo.
B. The loss of spines in young h-α-syn animals is driven by a reduced fraction of newly gained spines, while the fraction of lost spines remains unchanged and the fraction of stable spines is increased ($P_{\text{gain/total}} = 0.0256$).
C. The daily turnover of spines shows no significant difference between groups ($P = 0.4062$).
D. In vivo confocal data in young mice confirm synapse loss in h-α-syn mice between 2 and 4.5 months of age ($P_{\text{syn 2-4.5 months}} = 0.0064$; $P_{\text{conf 4.5 months}} = 0.0012$) and show a shift in spine morphology toward relatively more stubby ($P_{\text{stubby}} = 0.0418$) and less thin spines ($P_{\text{thin 4.5 months}} = 0.0044$; $P_{\text{thinn 4.5 months}} = 0.0155$).
E. In aged h-α-syn mice show a decrease in total spine density ($P_{\text{total 5 months}} = 0.0005$; $P_{\text{thinn 5 months}} = 0.0039$) as well as in the fraction of mushroom spines ($P_{\text{mushroom 5 months}} = 0.0004$; $P_{\text{stubby 5 months}} = 0.0298$), whereas the fraction of thin spines is increased ($P_{\text{thin 5 months}} = 0.0097$; $P_{\text{stubby 5 months}} = 0.0155$).

Data information: (A–C) $n = 4$ animals per group; (D, E) $n = 3–4$ animals per group as illustrated, mean with s.e.m., Bonferroni’s post hoc test (A): $^{*}P < 0.05$, $^{**}P < 0.01$; two-way ANOVA genotype main factor (B, C). $^{*}P < 0.05$, $^{**}P < 0.01$, $^{***}P < 0.001$.

imaging of apical tuft dendrites (layer I) was performed in brain slices of control as well as h-α-syn mice that were 2 and 4.5 months old. Note that the higher resolution of confocal microscopy in brain slices leads to an overall increased detection of spine density and a better ability to judge spine morphology. Two remarkable differences concerning the age and the genotype of the animal were observed. Firstly, at 2 months of age, there was no difference in spine density between control and transgenic animals, whereas at 4.5 months, spine density in h-α-syn mice showed a ~40% decrease ($t_{(5)} = 5.770$; $P = 0.0012$) relative to age-matched controls. Secondly, there was a significant difference in the spine densities of the h-α-syn mice ($t_{(5)} = 4.097$; $P = 0.0064$; ~30% reduction at 4.5 months) depending on their age, whereas this had no impact on the spine densities of control animals (Fig 2D).

In order to clarify whether the progressive spine loss in h-α-syn mice is reflected by changes in spine morphology, we furthermore analyzed spine shape for which three morphological subtypes have been defined (Rochefort & Konnerth, 2012). Thin spines have an elongated appearance, with a long neck and a small spine head. Mushroom spines have the largest, round head, while stubby spines are devoid of a neck and appear as round elevation from the dendritic shaft. In addition, filopodia are very long and thin structures that lack a spine head and are considered the most dynamic of dendritic structures, which appear and disappear rapidly and only sometimes develop into one of the other spine types. In the morphological analysis of our confocal data, the absolute numbers (Appendix Fig S1A) as well as the fractions of mushroom, stubby and thin spines as well as filopodia, were comparable in 2-month-old mice independent of the expression of human α-syn (Fig 2D).

In 4.5-month-old h-α-syn mice, the density of all morphological spine types is decreased (Appendix Fig S1A); however, not all types are lost to the same extent. The normalized fractions reveal a slight increase in the fraction of stubby spines ($t_{(5)} = 2.579$; $P = 0.0418$), whereas thin spines clearly represent a smaller fraction compared to both age-matched controls ($t_{(5)} = 3.346$; $P = 0.0155$) and to younger h-α-syn mice ($t_{(5)} = 4.443$; $P = 0.0044$) (Fig 2D).

The confocal imaging data from aged mice confirmed the reduction in spine density of ~30% in both 7.5- and 13.5-month-old h-α-syn brains. Compared to controls, the fractions of mushroom spines were decreased, while the fractions of thin spines were increased (Fig 2E and Appendix Fig S1B).

Taken together, our data confirmed our hypothesis that structural changes of dendritic spines develop within a certain time frame in young α-syn overexpressing mice. This time frame was determined to be approximately between 2 and 4.5 months of age. Before, the dendritic appearance in these mice is indistinguishable from healthy controls. Later, α-syn overexpression causes changes in dendritic spines toward an overall lower spine density and a shift in spine morphology.

**Intrastriatal injection of preformed α-synuclein fibrils triggers a progressive spread of endogenous protein aggregation**

α-syn pathology can be induced in wild-type mice by injecting preformed fibrils (PFFs) (Luk et al, 2012a). Using this as a tool to investigate whether seeded pathology of disease-associated α-syn will lead to structural alterations in synapses, we injected healthy 2-month-old GFP-M mice with PFFs prepared in vitro. Prior to injection, the quality of the injection material was controlled. The amyloid nature of PFFs was confirmed in a thioflavin T (ThT) assay, as ThT changes its fluorescent properties upon binding to amyloid β-sheet structures (Fig 3A). Differential centrifugation through a sucrose gradient confirmed the strongly increased sedimentation coefficient and thus the increased molecular density of PFFs compared to α-syn monomers (Fig 3B). Finally, an electron micrograph visualized the fibrillary structure of PFFs (Fig 3C).

Sonicated PFFs were stereotactically injected into the striatum (Fig 3D), an area which contains extensive efferent and afferent projections to other brain regions and is affected in PD (Shepherd, 2013). As cortical neurons are the targets of interest in this study, the time-course and extent of α-syn pathology progression into different layers of the neocortex was monitored at different days postinjection (dpi). The phosphorylation of α-syn at S129 (pS129) is widely considered as a pathologic postranslational modification (Tenreiro et al, 2014; Oueslati, 2016) and was used in this study to identify pathologic inclusions.

Striatal injection of PFFs resulted in intraneuronal aggregation of phosphorylated α-syn (pS129-immunopositive) in the neocortex, at first confined to layer IV and V neurons (30–150 dpi), developing later to spread over all cortical layers (9 months postinjection; Fig 3E). pS129-α-syn inclusions appeared as cytoplasmic fibrillar forms (Fig 3F), which could also be stained by anti-ubiquitin (Fig 3G) and thioflavin S (Fig 3H and I), thereby presenting Lewy-body-like traits. Inclusions were also observed in the contralateral cortex relative to the injection site and the amygdala (Appendix Fig S2B). Due to a reported cross-reactivity between the pS129 antibody and phosphorylated neurofilament subunit L (NFL; Sacino et al, 2014b), we confirmed the presence of phosphorylated α-synuclein inclusions with a costaining for NFL (Appendix Fig S2B and C).
Figure 3.

Sonja Blumenstock et al.  
Alpha-synuclein mediates spine pathology

EMBO Molecular Medicine  
Vol 9 | No 5 | 2017
presence of pS129-α-syn inclusions across different brain regions is shown in Appendix Fig S2D.

Control brains injected with PBS did not show comparable immunopositive inclusions (Fig 3E). Importantly, also monomeric α-syn injected into GFP-M mice as well as PFFs injected into α-syn knock-out mice (SNCA KO) failed to induce the phenotype as observed after injection of PFFs (Appendix Fig S2E).

In order to further examine whether the effect can be attributed to templated conversion of endogenous mouse α-syn into misfolded and aggregated species rather than just the uptake of PFFs by cells, control injections of fluorescently labeled PFFs into the striatum were performed. Twenty-four hours after the injection, the injection material was prominently detectable in the striatum, corpus callosum, and the injection canal (Appendix Fig S3B and C). At 6 dpi, however, the fluorescent material was removed from the injection site (Appendix Fig S3D) and later showed very little colocalization with developing pS129-α-syn-positive aggregates at 30 dpi (Appendix Fig S3E and F). We therefore conclude that the spreading of intracellular α-syn aggregation in our model is dependent on the introduction of fibrillar α-syn as a molecular template on top of endogenous α-syn expression and not simply due to a cellular uptake of injected PFFs.

Considering inflammation as a contributor in neurodegenerative diseases (Perry et al., 2010) and synaptic plasticity (Morris et al., 2013) and as PFF-induced α-syn accumulations have been associated with microglia (Sacino et al., 2014a), we assessed the presence and activation of microglia in the neocortex with two markers. Anti-ILb-1 marks all microglial cells including their processes, CD68 is a marker associated with active phagocytosis. We find that at the time point of spine analysis (5 months postinjection), the cortical area covered by microglia shows a tendency toward higher coverage, especially in layer IV (Appendix Fig S4A and B). We also observed a tendency for an increase in the coverage with activated microglia (Appendix Fig S4C and D) as well as occasional colocalization of CD8-positive microglia and pS129-positive aggregates (Appendix Fig S4E). However, variation between samples is high and the effect is not significant.

**Seed α-synuclein aggregates have detrimental effects on dendrites and spines**

Advantageous to other mouse models, the seeding model does not require a disease-related transgene to induce a pathologic phenotype. In this way, we wanted to investigate structural alterations that might be the consequence of dose- and time-controlled molecular templating of α-syn aggregates. Five months after the injection of PFFs or PBS (control) into the dorsal striatum, brain slices were stained for pS129-positive α-syn aggregates, which were most abundant in layer IV and upper layer V of the cortex, including pyramidal eGFP-expressing neurons (Appendix Fig S3A). In order to study whether structural changes might be solely attributed to a local effect of intracellular aggregates on neighboring dendrites or due to a more global disturbance in the synaptic network, we imaged apical dendrites located in two different layers (Fig 4A). Dendrites imaged in layer IV or V were spatially close to cells bearing an intraneuronal α-syn accumulation, while tuft dendrites of layer I were distal to these cells (Fig 4B).

Indeed, we found alterations in spine density irrespective of the cortical location of the dendrites. In both layer IV/V (t(11) = 3.628; P = 0.004) and layer I (t(11) = 3.838; P = 0.0028), spine density was found to be lowered in brains which had been injected with PFFs and contained α-syn aggregates (Fig 4C).

We next investigated how vulnerable the morphological spine types are in response to seeded α-syn. In the layer IV/V dendrites, all types were lost to the same extent, leaving the fractions of mushroom, stubby, and thin spines comparable in controls and seeded animals (Appendix Fig S5B and Fig 4C). Contrarily, in apical tuft dendrites (layer I), thin spines proved to be more and stubby spines less vulnerable to seeded α-syn. The fraction of thin spines was therefore decreased (t(11) = 2.350; P = 0.0385), while the fraction of stubby spines was increased (t(11) = 2.599; P = 0.0247) (Appendix Fig S5A and Fig 4C). This phenotype was already observed in our ex vivo spine data of young transgenic h-α-syn mice (see Fig 2) and could be reproduced here. On the presynaptic side, no overt change in the density of excitatory boutons could be detected in both the transgenic and the seeding mouse model (Appendix Fig S6L–L).

The irregular form of the tuft dendritic shafts in PFF-seeded brains was an additional observation. In controls, injected only with PBS, the diameter of dendritic shafts was found to be relatively uniform. In contrast, apical tuft dendrites of PFF-seeded mice showed an irregular appearance of the dendritic shaft. Over the length of the dendrites, large swellings could be observed as well as segments with an extremely narrow diameter (Fig 4D). In a histogram plotting the frequency of diameter values, dendritic shaft diameters follow a normal distribution ($R^2 = 0.9263$) for the control animals. For the seeded mice, the variety of values is broader and the values are best described with two distributions: one at very low dendritic diameter ($R^2 = 0.9250$) and one comparable to the control distribution ($R^2 = 0.9664$), with the exception of the presence of abnormally high values (Fig 4E). As a consequence of the high variation, the mean standard deviation of the dendritic shaft diameter was found to be significantly increased (t(11) = 3.054; P = 0.011) in mice that were seeded with α-syn PFFs (Fig 4F). Note that this phenotype was only observed in our seeding, but not in the transgenic model.

With our approach, we could show that the introduction of α-synuclein seeds and the consecutive accumulation in the cerebral
cortex triggers structural changes both at the level of the dendritic spine and the dendritic shaft.

**Discussion**

This study presents converging evidence from two distinct approaches for the adverse effects of α-syn accumulation for the density and structural plasticity of dendritic spines in the cerebral cortex. Dendritic spines are excitatory postsynapses, and their loss has been described repeatedly as a structural correlate for cognitive impairment and dementia (Terry *et al.*, 1991; Dickson *et al.*, 1995; Bellucci *et al.*, 2012; Picconi *et al.*, 2012). Moreover, the effects of mutated and accumulated α-syn have been studied in several brain regions. Striatal spine loss has been reported in various animal models of PD and in PD and DLB patients (McNeill *et al.*, 1988; Zaja-Milatovic *et al.*, 2006). In a mouse model overexpressing the human A53T α-syn mutation, spine loss has been observed in the cerebral cortex. In layer I, pS129-positive structures are much less dense. Lines exemplarily mark dendrites used for analysis. In layer IV/V, spine density is reduced as well (**P = 0.004), without a significant effect on spine morphology.

Figure 4. The presence of accumulated α-synuclein induced by seeding 5 months prior to analysis causes spine loss and malformation in layer V apical dendrites.

A α-Synuclein aggregates occur as intrasomal (arrowheads) and neuritic (arrows) accumulations and are present predominantly in upper layer V and layer IV of the cortex. In layer I, pS129-positive structures are much less dense. Lines exemplarily mark dendrites used for analysis.

B Spine analysis was performed on apical dendrites located in the cortical layers IV/V and I.

C In layer I, spine density is reduced relative to PBS-injected controls (**P = 0.0028), with the fraction of stubby spines being increased (P = 0.0247) and the fraction of thin spines being decreased (P = 0.0385). In layer IV/V, spine density is reduced as well (**P = 0.004), without a significant effect on spine morphology.

D In apical tuft dendrites of PFF-injected mice, dendrites display dystrophic swellings and parts of very small diameter; white lines: measurement positions.

E Histograms of the dendritic shaft diameter.

F Variation in the diameter of single dendrites in seeded mice compared to controls (P = 0.011).

Data information: n = 6 (control), n = 7 (PFF) animals, mean with s.e.m.; Student’s t-test: *P < 0.05, **P < 0.01. Scale bars: 20 μm (A), 2 μm (B, D).
striatum and in the hippocampus (Finkelstein et al., 2016). Some authors report that D2R striatopallidal neurons in reserpin and 6-OHDA-treated mice selectively lose spines (Day et al., 2006), while others found spine loss on both direct and indirect pathway neurons in 6-OHDA- or MPTP-treated models (Suárez et al., 2014). In contrast, another study failed to detect striatal spine loss in A53T mice, despite a decrease in the number of Snpc neurons (Oaks et al., 2013). α-syn was also demonstrated to have negative impact on newly generated neurons of the hippocampus (Winner et al., 2012) and the olfactory bulb (Neuner et al., 2014), particularly on their dendrite outgrowth and spine development.

In our current study, we aimed to extend this knowledge with a chronic examination of spines in the cortex. As the chronic effects of α-syn on cortical spine plasticity remain incompletely understood, we used two different mouse models which exhibit α-syn accumulation in the cortex. In our transgenic model, wild-type human α-syn was expressed under the PDGF promoter, leading to neocortical accumulation of mostly soluble protein located as inclusions in neuronal cell bodies as well as synaptic terminals (Masliah, 2000; Rockenstein et al., 2002; Amschel et al., 2013). Our seeding model was based on the striatal injection of preformed wild-type mouse α-syn fibrils and caused a slowly progressing phenotype of phosphorylated α-syn-positive neuronal inclusions, which first developed in the deeper layers of the neocortex and condensed over time to mature into fibrillary Lewy-like inclusions.

Our study also provides in vivo evidence that accumulation of wild-type α-syn interferes with cortical spine plasticity and spine density which declined in young adult mice starting from about 3 months of age and stabilized at a reduced overall density in older mice (≈30%). This spine loss occurs well before behavioral impairments become apparent in these mice (Masliah, 2000; Amschel et al., 2013) and may be the result of a presymptomatic change in pre- and postsynaptic function.

While numerous studies have made great effort to unravel the molecular mechanism behind α-syn toxicity, it remains to be clarified whether a toxic gain-of-function or a loss-of-function or both is responsible for synapse and cell loss. Among the toxic gain-of-function effects of α-syn are the impairment of lysosomal and proteasomal protein degradation, induction of endoplasmic reticulum (ER) stress, Golgi fragmentation, and active formation of pores on cellular membranes (reviewed by Cookson & van der Brug, 2008). For instance, it was shown that the acute application of α-synuclein oligomers onto autaptic hippocampal cultures as well as onto hippocampal slices significantly enhances basal NMDA receptor activation and the amplitude of AMPA-receptor-mediated synaptic currents (Hüls et al, 2011; Diogenes et al., 2012). This augmented excitatory transmission could be explained by (i) α-syn oligomer-mediated pore formation in the postsynaptic membrane (Volles et al., 2001; Furukawa et al., 2006; Schmidt et al., 2012), (ii) enhancement of voltage-operated Ca\(^{2+}\) channel activity (Hettiarachchi et al., 2009), or (iii) inefficient membrane repolarization (Shrivastava et al., 2015), all causing increased calcium influx which in turn results in AMPA-receptor recruitment to the dendritic spine membrane (Byth, 2014). The resulting postsynaptic is saturated with receptors and therefore unable to recruit extra AMPA receptors upon theta-burst stimulation which causes a long-term potentiation (LTP) decline and overall excitotoxicity (Yuste & Bonhoeffer, 2001; Diogenes et al., 2012). On the long term—as we analyzed the structural impact of transgenic and seeded α-synuclein during and after several months—this proposed mechanism based on a slightly but chronically enhanced calcium influx may disturb the tightly controlled, calcium-dependent machinery of synaptic plasticity. Likewise, a primary presynaptic pathology might alter synaptic vesicle release which as a long-term consequence may alter dendritic spine plasticity. Excessive α-synuclein has been shown to inhibit neurotransmitter release via specifically reducing the size of the synaptic vesicle recycling pool (Nemani et al., 2010) as well as to regulate SNARE-driven membrane fusion (García-Reitbock et al., 2010; Burre et al., 2015). A study in DLB brain tissue showed massive deposits of diffuse aggregates in the cortical and subcortical gray matter, which were located at synaptic terminals and were linked to dendritic spine loss. This suggests that the subcellular presynaptic location of α-syn is important for its neurodegenerative effect (Kramer & Schulz-Schaeffer, 2007) and although our models show no overt presynaptic loss, an alteration in function seems likely. The functional roles of α-synuclein, in conjunction with regulating actin dynamics and chaperoning the polymerization of microtubule-associated proteins (Sousa et al., 2009), would certainly impair spine stability and dendritic plasticity (Tsaneva-Atanasova et al., 2009).

The α-syn loss-of-function hypothesis comprises that with progressing aggregation, endogenous α-syn becomes sequestered into inclusions; be they oligomers, protofibrils, or mature LBs and LNs, leaving less functional protein to perform normal cellular functions (reviewed by Benskey et al., 2016). Particularly the changes seen after seeding with PFFs are supportive of this hypothesis. As previously shown and confirmed in this study through seeding of fluorescently tagged α-syn fibrils, the aggregates formed are chiefly composed of endogenous α-syn, with very little of the initial PFF seed being part of the developed inclusions (Luk et al., 2009). The concept of sequestration is further reinforced by the complete absence of inclusions and toxicity in α-syn null mice seeded with PFFs (Volpicelli-Daley et al., 2011; Luk et al., 2012a). To complicate things even further, recent evidence suggests that microglial activity can contribute to both detrimental and protective mechanisms in neurodegenerative diseases (Morris et al., 2013; Miyamoto et al., 2016; Tang & Le, 2016). In our model, the infusion of PFFs did not cause overt and chronic microglial activation at the observed time point, but additional experiments might be necessary to gain a broader picture about the glial involvement in α-syn-affected synapse dynamics.

From our imaging data, we cannot pin down which of these described mechanisms exactly are involved in the loss of spines, which emphasizes the need for further studies for clarification. Furthermore, as our in vivo data suggest, the mechanism underlying synapse loss and the shift in morphology may not even be uniform throughout the course of disease. Spine dynamics were found to be changed remarkably depending on the age of the animal upon observation. The decrease in gained spines observed in 3-month-old animals caused a progressive spine loss and is morphologically reflected in the diminished fraction of thin spines, which are known to correspond to non-synaptic transient precursors of the larger stabilized spines (Holtmaat et al., 2005; Arellano et al., 2007). Contrarily, in the two older groups (6 and 12 months), further spine loss was restrained by an elevation of the daily turnover ratios. This was in turn reflected by an increase in the fraction of thin spines.
Thus, we propose that the observed spine remodeling in these animals represents a compensatory mechanism for the loss of absolute spine number, thereby strengthening the remaining synaptic contacts.

In both our seeding and transgenic model, proteinaceous aggregates are present in the neocortex, albeit in two histopathological variances: while the PDGF promoter in the transgenic model drives a more or less uniform accumulation of non-fibrillary (Masliah, 2000) α-syn in cortical neurons and synapses, seeding of PFFs leads to the formation and maturation of fibrillary Lewy-like structures in neurons connected to the dorsal striatum. Our results from our in vivo study in young transgenic mice are in line with the dendritic spine phenotype which can be induced through inoculation with PFFs. Dendritic spine loss in layer V pyramidal neurons as well as a redistribution of spine morphologies toward less thin spines can be observed 5 months after inoculation with PFFs. The observed spine loss was independent of whether the respective dendrite was spatially close to the cortical layer V, in which neuronal α-syn inclusions were most pronounced. Interestingly, dystrophic deformation of apical tuft dendrites was present in PFF-seeded brains. Dendritic varicosities have been described in other progressive neurodegenerative diseases and have been linked to dendritic spine loss in scrapie-injected rodents (Fuhrmann et al., 2007) and an Alzheimer’s disease model (Bittner et al., 2010). The presence of axonal varicosities caused by α-synuclein seeding has been reported (Rockenstein et al., 2005); furthermore, it has been shown that inclusions mature and condense over time and that aggregate bearing neurons selectively degenerate (Osterberg et al., 2015). Therefore, it seems likely that the degeneration of the respective cell body and the accompanying transport deficits are the cause for dystrophies and contribute to spine loss in our PFF-seeded brains.

For answering scientific questions, animal models in principle comprise advantages as well as drawbacks (Bezard et al., 2013). For example, all available genetic mouse models of synucleinopathies are able to model only a subset of symptoms known in the human disease cases. Promoter-driven protein expression is

---

**Figure 5. Cortical α-synuclein accumulation and long-term in vivo imaging of PDGF-α-syn × GFP-M mice.**

A Immunostaining with 13G7 antibody shows cortical layer V overexpression of α-synuclein (magenta) including accumulation in cell bodies (arrows).

B Experimental timeline: 4 weeks after window implantation (white arrowhead), imaging was performed over 6 weeks, once every 7 days (gray arrowheads) and finally followed by perfusion and tissue fixation directly after the last imaging session (black arrowhead).

C Overview and detailed projections of eGFP-labeled layer V apical dendrites in the somatosensory cortex, imaged through a chronic cranial window.

Data information: Scale bars: 20 μm (A), 50 μm (C), 5 μm (C, inset).
limited to a cell population in which the promoter is active and is even complicated through the fact that background strain might affect the expression pattern (Chesselet et al., 2008). Apart from that, α-synuclein pathology induced by seeding of PFFs can be induced in wild-type animals (Luk et al., 2012a) in a dose- and region-specific manner. Here, however, the unphysiologically large protein amount used for “planting the seed” should be considered. One strong point of our combined approach was to apply and compare both models, in order to best break down the actual structural impact of α-syn accumulation on cortical neurons. Collectively, our results support the idea that α-syn homeostasis is constantly balanced on a knife-edge and that perturbations that change its expression produces neurotoxicity with a common mechanism in which the critical events are protein misfolding, aggregate deposition, aggregate propagation, and synaptic dysfunction (Jucker & Walker, 2011; Guo & Lee, 2014; Luna & Luk, 2015). Our data provide evidence for the harmful structural consequences of excessive wild-type α-synuclein on cortical dendrites throughout different pathological stages and may pave the way toward new therapeutic strategies for PD and DLB that focus on the maintenance of synaptic function.

Materials and Methods

Animals

PDGF-h-α-syn animals were obtained from QPS Austria Neuropharmacology (Grazbach, Austria) and bred on a C57Bl/6 background. These transgenic mice overexpress human α-synuclein (h-α-syn) under the regulatory control of the human platelet-derived growth factor-β (PDGF-β) promoter (Masliah, 2000). Expression is strongest in the neocortex, hippocampus, olfactory bulb, and limbic system, leading to intraneuronal and synaptic α-syn accumulation as early as 3 months of age (Fig 5A and Appendix Fig S6A–D) as well as progressive motor deficits (Masliah, 2000; Amschl et al., 2013). Furthermore, C57Bl/6 wild-type, Thy1-eGFP transgenic mice (GFP-M) (Jackson Laboratory, Bar Harbor, ME, USA), and α-syn knockout (SNCA KO) mice (Abeliovich et al., 2000) were used. For in vivo imaging, the heterozygous PDGF-h-α-syn × GFP-M (termed h-α-syn in this paper) line was created by interbreeding and the experiment involved three age cohorts of both male and female mice, in which mice were 3, 6 or 12 months old at the first imaging time point. All animals were housed in groups under pathogen-free conditions and bled from purified E. coli BL21(DE3) (Invitrogen, MA, USA) were transformed with the plasmid, and expression was induced with isopropyl β-D-1-thiogalactopyranoside (IPTG, Pefab, Erlangen, Germany). Cells were lysed by boiling after heat-inactivation of proteases. After centrifugation, the supernatant was filtered through Filterpore 0.2 filters (Sartorius, Göttingen, Germany) and eluted with a linear 25 mM to 500 mM NaCl gradient. Synuclein containing fractions were concentrated using VivaSpin 2 columns (Sartorius, Göttingen, Germany). Protein concentration was assessed to 5 mg/ml in 50 mM Tris–HCl, pH = 7.0. After freezing in liquid nitrogen, the protein was stored at −80°C. Appendix Fig S2A shows plots from the monomer preparation. Preformed fibrils (PFFs) were assembled from purified α-synuclein monomer (5 mg/ml) by incubation at 37°C with constant agitation (1,400 rpm) in an orbital mixer (Eppendorf, Hamburg, Germany) for 96 h and stored at −80°C (Conway et al., 2000; Deeg et al., 2015). For fluorescent labeling, purified α-synuclein monomer (3.75 mg/ml, containing 100 mM NaHCO3) was incubated with 0.34 mg/ml Alexa Fluor® 488 NHS Ester (Life Technologies, Darmstadt, Germany) and stereotactically injected with 5 μl (25 μg) of PFFs into the dorsal striatum (coordinates relative to the bregma: +0.2 mm anterior, +2.0 mm from midline, +2.6 mm beneath the dura) of the right hemisphere using a 5-μl Hamilton syringe. Injections were performed at 400 nl/min with the needle in place after injection for at least 5 min. Control animals received sterile PBS or 5 μl of 5 mg/ml monomeric α-synuclein. Animals were monitored regularly after the surgery and sacrificed at different predetermined time points (30 days, 60 days, 90 days, 5 months, and 9 months after injections) for perfusion and tissue fixation with 4% paraformaldehyde.

Thioflavin T binding and sucrose gradient

For in vitro measurement of thioflavin T (ThT) binding, 10 μM ThT was added to 100 μg/ml monomeric α-synuclein or PFFs and incubated at 30°C for 10 min. Fluorescence signal was excited at 420–460 nm and emission detected at 480–530 nm with a spectrofluorometer (FluoStar Optima, BMG Lab Tech, Jena, Germany). Continuous sucrose-gradient assay was performed as described previously (Friedlander et al., 2002; Wagner et al., 2013). Briefly, six layers of solutions with decreasing sucrose concentration (50 mM Tris pH 7.5, 0.1% NP-40, 10–60% D(+)-sucrose, respectively) were filled into a 4 ml 11 × 60 mm polylamellar tube (Beckman Coulter, CA, USA). Finally, 200 μl of 5 μM protein in 1× TBS (pH 7.5) containing 0.1% NP-40 was loaded on the top of the gradient. Ultracentrifugation with 100,000 g at 4°C for 1 h was performed using a SW60Ti rotor (Beckman Coulter, USA). Resulting continuous gradients were
Electron microscopy

α-Synuclein fibrils diluted to 5 μM in H2O were adsorbed onto a Formvar-coated, carbon-stabilized copper grid. The grid was then rinsed briefly with H2O and stained with uranyl acetate and lead citrate. Digital images were captured at different magnifications on a Jeol JEM-1011 TEM (JEOL Inc., MA, USA) equipped with an 11 megapixel Orios CCD digital camera (Gatan Inc. CA, USA).

Cranial window

A cranial window was implanted over the right cortical hemisphere as previously reported (Fuhrmann et al., 2007). In short, the mice were given an intraperitoneal injection of ketamine/xylazine to reach surgical anesthesia. Additionally, dexamethasone (0.01 mg/g body weight; CP Pharma, Burgdorf, Germany) was intraperitoneally administered immediately before surgery (Holtmaat et al., 2009). A circular piece of the skull (4 mm in diameter) over the right hemisphere (centered over the parietal bone, approx. 5.5 mm caudal from the bregma and 5.5 mm lateral from midline) was removed using a dental drill (Schick-Technikmaster C1; Pluraden; Offenbach, Germany). The craniotomy was closed immediately with a round coverslip (4 mm in diameter), held with dental acrylic. A small z-shaped titan bar was glued next to the coverslip to allow repositioning of the mouse during subsequent imaging sessions. After surgery, mice received subcutaneous analgesic treatment with cefotaxime (0.06 mg/kg body weight; Sanofi-Aventis, Frankfurt, Germany). The craniotomy was closed immediately with a round coverslip (4 mm in diameter), held with dental acrylic. A small z-shaped titan bar was glued next to the coverslip to allow repositioning of the mouse during subsequent imaging sessions. After surgery, mice received subcutaneous analgesic treatment with cefotaxime (0.06 mg/kg body weight; Sanofi-Aventis, Frankfurt, Germany).

Long-term in vivo imaging

Imaging started 4 weeks after the cranial window preparation to allow the animals to recover from surgery. After 6 weekly 2-photon imaging sessions, mice were sacrificed and their brains were fractionated in volumes of 200 μl. 20 μl per fraction was analyzed by denaturing Western blot using a monoclonal antibody against mouse-α-synuclein (New England Biolabs, Frankfurt, Germany).

Image processing and analysis of in vivo and ex vivo data

For image analysis, only microscope image data with sufficient eGFP expression and good signal-to-noise ratio over time were included and the experimenter was blinded to the genotype of the mice by assigning a random number to each animal. From each mouse (male or female, n = 4–5 in vivo, n = 6–7 seeding experiment, without randomization), 9–10 dendrites from at least three different positions underneath the cranial window or in slices were analyzed. The length of individual dendrites was measured in ZEN 2012 (Zeiss). Spines identified along the dendrite were marked as gained, lost, or stable. The mean density of dendritic spines was estimated for each time point and expressed over 1 μm of dendrite length. The stability of spines was calculated based on the amount of spines that remained unaltered for at least two subsequent imaging sessions. The spine turnover rate (TOR) was assessed based on gain and loss of spines over each day of imaging, calculated as follows: TOR = (N\textsubscript{gained} + N\textsubscript{lost})/(2 × N\textsubscript{present} × t), where N\textsubscript{gained}, N\textsubscript{lost}, and N\textsubscript{present} represent the number of gained, lost, or total spines at time points of interest, respectively, while t\textsubscript{i} is the number of days between consecutive imaging sessions. Ex vivo image analysis measurements were performed manually from maximal projection images of deconvoluted (AutoQuantX3, Media Cybernetics) confocal stacks. All spines along the dendrite were marked and categorized into three morphologically different classes, according to established criteria (Jung et al., 2011). For analysis of the dendritic diameter, length measurements were taken every 40 pixels along the dendritic shaft (control: 756 measurements, n = 6, PFF-seeded: 840 measurements, n = 7) using ImageJ. To quantify the density of glutamatergic presynaptic boutons VGLUT1-positive Imaris 7.7.2 software was applied to detect VGLUT1-positive puncta. In detail, the spots detection algorithm was applied using an estimated diameter of 0.5 μm in xy and an 1.5 μm in z. Background subtraction was enabled, and region-growing type was set to local contrast with a manual threshold of 30 defined for all datasets. Only spots with a xy diameter greater than 0.4 μm were counted as boutons. The data were compiled in MATLAB using ImarisXT interface.

Statistical analysis

The sample size of animals and imaged dendrites per animal were chosen according to our previous experience in long-term imaging (Neuner et al., 2014; Filser et al., 2015). Graphs were created, and statistics were calculated in Prism v 5.04 (GraphPad Software, San Diego, CA, USA). For in vivo time series data, two-way ANOVA followed by the Bonferroni’s post hoc test was used to compare the variance of spine parameters assessed over time in control and h-α-syn animals. For assessment of inter-group differences at single time points, Student’s t-test (unpaired, two-sided) was applied. Normal distribution was assumed according to the central limit theorem, as spine densities were calculated as the means of means for every mouse. For t-tests, the variance between groups was tested (F-test) and not found to be significantly different. Data are expressed as mean ± SEM unless otherwise indicated, with P < 0.05 defining differences as statistically significant (**P < 0.05; ***P < 0.01; ****P < 0.001; for post hoc test: aP < 0.05; bP < 0.01).
The paper explained

Problem
Parkinson’s disease (PD) and is the second most common neurodegenerative disease with a prominent loss of nigrostriatal dopaminergic neurons. The resultant dopamine deficiency underlies the onset of typical motor symptoms. Apart from this, PD and other neurodegenerative conditions featuring misfolded and aggregated forms of the synaptic protein α-synuclein (α-syn) are frequently accompanied by cognitive decline and dementia, arising from the structural and functional changes in the brain cortical synapses. The role of α-syn in the impairment of cortical circuitries and synaptic plasticity remains incompletely understood. We therefore investigated how α-synuclein accumulation affects the plasticity of dendritic spines as the loss of these small protrusions from the neuronal dendritic shaft is widely considered a structural correlate for cognitive decline.

Results
Our study of the cortex involved two distinct in vivo mouse models. Long-term in vivo two-photon imaging of apical dendrites was performed in mice overexpressing wild-type human α-syn. Additionally, intracranial injection of preformed α-syn fibrils was performed to seed cortical α-syn pathology in mice without an additional disease-related transgene. We find that α-synuclein overexpressing mice show decreased spine density and abnormalities in spine dynamics in an age-dependent manner. We also provide evidence for the detrimental effects of seeded α-syn aggregates on dendritic architecture, as we observed spine loss as well as dystrophic deformation of dendritic shafts in layer V pyramidal neurons of the cortex.

Impact
Our results demonstrate the impact of cortical α-syn in a time-resolved manner and provide a link to the pathophysiology underlying dementia associated with α-synucleinopathies. They may also enable the evaluation of potential drug candidates on dendritic spine pathology in vivo.

Immunohistochemistry and confocal microscopy
Animals in deep ketamine/xylazine anesthesia were killed by transcardiac perfusion with 1× phosphate-buffered saline (PBS) followed by 4% paraformaldehyde (w/v). The brains were postfixed in PBS containing 4% paraformaldehyde overnight before cutting followed by 4% paraformaldehyde (w/v). The brains were postfixed × cardiac perfusion with 1

Appendix Fig S6E–H illustrates the automatic bouton detection applying the algorithm.

Expanded View for this article is available online.

Acknowledgements
We thank Katharina Bayer, Sarah Hanselka, and Eric Grießinger for their excellent technical support and animal care. We like to give special thanks to Prof. Virginia Lee and Dr. Kelvin Luk for providing us with the plasmid for α-syn expression and to S. Tahirovic for advice and kindly providing the CD68 antibody. We also thank MM. Doroštar, S. Crux, S. Fiser, and C. Sgobio for scientific support and advice on the manuscript. This work was funded the Munich Cluster for Systems Neurology SyNergy (EXC 1010).

Author contributions
SB performed design, data collection, analysis and interpretation of experiments and wrote the manuscript. EFR contributed to the data collection in in vivo experiments. FP analyzed presynaptic terminals. LB-L provided electron micrographs from α-synuclein PFFs. AG and FS provided experimental material and expertise in α-synuclein purification in PFF assembly. EFR and FS helped with manuscript preparation. JH supervised the study; contributed to conception, design, and manuscript writing; and provided financial support and final approval of the manuscript.

Conflict of interest
The authors declare that they have no conflict of interest.

References
Aarsland D, Beyer MK, Kurz MW (2008) Dementia in Parkinson’s disease. Curr Opin Neurol 21: 676–682
Abeliovich A, Schmitz Y, Fariñas I, Choi-Lundberg D, Ho W-H, Castillo PE, Shinsky N, Verdugo JMC, Armanini M, Ryan A et al (2000) Mice lacking α-synuclein display functional deficits in the nigrostriatal dopamine system. Neuron 25: 239–252
Amschel D, Neddens J, Havas D, Flunkert S, Römer H, Rockenstein E, Masliah E, Windisch M, Hutter-Paier B (2013) Time course and progression of wild type α-Synuclein accumulation in a transgenic mouse model. BMC Neurosci 14: 6
Areliano JI, Espinosa A, Fairén A, Yuste R, DeFelipe J (2007) Non-synaptic dendritic spines in neocortex. Neuroscience 145: 464–469
Bellucci A, Zaltieri M, Navarria L, Grigolatto J, Missale C, Spano P (2012) From α-synuclein to synaptic dysfunctions: new insights into the pathophysiology of Parkinson’s disease. Brain Res 1476: 183–202
Benskey MJ, Perez RG, Manfredsson FP (2016) The contribution of alpha synuclein to neuronal survival and function – Implications for Parkinson’s disease. J Neurochem 137: 331 – 359
Bezard E, Yue Z, Kirk D, Spillantini MG (2013) Animal models of Parkinson’s disease: limits and relevance to neuroprotection studies. Mov Disord 28: 61 – 70
Bittner T, Fuhrmann M, Burgold S, Ochs SM, Hoffmann N, Mittregger G, Kretzschmar H, LaFerla FM, Herms J (2010) Multiple events lead to dendritic spine loss in triple transgenic Alzheimer’s disease mice. PLoS One 5: e15477
Burn DJ (2004) Cortical Lewy body disease. J Neurol Neurosurg Psychiatry 75: 175 – 178
Burre J, Sharma M, Sudhof TC (2015) Definition of a molecular pathway mediating α-synuclein neurotoxicity. J Neurosci 35: 5221 – 5232
Byth LA (2014) Ca²⁺- and CaMKII-mediated processes in early LTP. Ann Neurosci 21: 151 – 153
Chesselet M-F, Fleming S, Mortazavi F, Meurers B (2008) Strengths and limitations of genetic mouse models of Parkinson’s disease. Parkinsonism Relat Disord 14: 584 – 587
Clayton DF, George M (1999) Synucleins in synaptic plasticity and neurodegenerative disorders. J Neurosci Res 58: 120 – 129
Conway KA, Harper JD, Lansbury PT (2000) Fibrils formed in vitro from α-Synuclein and two mutant forms linked to Parkinson’s disease are typical amyloid. Biochemistry 39: 2552 – 2563
Cookson MR, van der Brug M (2008) Cell systems and the toxic mechanism(s) of α-synuclein. Exp Neurol 209: 5 – 11
Day M, Wang Z, Ding J, An X, Ingham CA, Shering AF, Wokosin D, Ilijic E, Sun Z, Sampson AR et al (2006) Selective elimination of glutamatergic synapses on striatopallidal neurons in Parkinson disease models. Nat Neurosci 9: 251 – 259
Deeg AA, Reiner AM, Schmidt F, Schueder F, Ryazanov S, Ruf VC, Giller K, Becker S, Leonov A, Griesinger C et al (2015) Anle138b and related compounds are aggregation specific fluorescence markers and reveal high affinity binding to α-synuclein aggregates. Biochim Biophys Acta 1850: 1884 – 1890
Dickson DW, Crystal HA, Bevona C, Honer W, Vincent I, Davies P (1995) Correlations of synaptic and pathological markers with cognition of the elderly. Neurobiol Aging 16: 285 – 298
Diogenes MJ, Dias RB, Rombo DM, Vicente Miranda H, Maiolino F, Guerreiro P, Nasstrom T, Franquelim HG, Oliveira LMA, Castanho MARB et al (2008) Extracellular α-synuclein oligomers modulate synaptic transmission and impair LTP via NMDA-receptor activation. J Neurosci 32: 11750 – 11762
Fiala JC, Spacek J, Harris KM (2002) Dendritic spine pathology: cause or consequence of neurological disorders? Brain Res Rev 39: 29 – 54
Filser S, Osvesian SV, Masana M, Blazquez-Llora L, Brandt Elang A, Volbracht C, Müller MB, Jung CKE, Herms J (2015) Pharmacological inhibition of BACE1 impairs synaptic plasticity and cognitive functions. Biol Psychiatry 77: 729 – 739
Finkelstein Di, Hare DJ, Billings JL, Sedjadera A, Nurjono M, Arthofer E, George S, Culvenor JC, Bush AI, Adlard PA (2016) Clioquinol improves cognitive, motor function, and microanatomy of the alpha-synuclein hA53T transgenic mice. ACS Chem Neurosci 7: 119 – 129
Friedlander G, Schönberger O, Horonchik L, Yedida Y, Shaked G, Gabizon R, Taraboulos A (2002) Protease-sensitive scrapie prion protein in aggregates of heterogeneous sizes. Biochemistry 41: 12868 – 12875
Fuhrmann M, Mitteregger G, Kretschmar H, Herms J (2007) Dendritic pathology in prion disease starts at the synaptic spine. J Neurosci 27: 6224 – 6233
Furukawa K, Matsuzaki-Kobayashi M, Hasegawa T, Kikuchi A, Sugeno N, Itouyama Y, Wang Y, Yao PJ, Bushlin I, Takeda A (2006) Plasma membrane ion permeability induced by mutant alpha-synuclein contributes to the degeneration of neural cells. J Neurochem 97: 1071 – 1077
Garcia-Reitbock P, Anichtchik O, Bellucci A, Lovino M, Ballini C, Fineberg E, Ghetti B, Della Corte L, Spano P, Tofaris GK et al (2010) SNARE protein redistribution and synaptic failure in a transgenic mouse model of Parkinson’s disease. Brain 133: 2032 – 2044
Gruetzendler J, Narayanan K, Can W-B (2002) Long-term dendritic spine stability in the adult cortex. Nature 420: 810 – 812
Guo JL, Lee VM-Y (2014) Cell-to-cell transmission of pathogenic proteins in neurodegenerative diseases. Nat Med 20: 130 – 138
Helmchen F, Denk W (2005) Deep tissue two-photon microscopy. Nat Methods 2: 932 – 940
Hettiarachchi NT, Parker A, Dallas ML, Pennington K, Hung C-C, Pearson HA, Boyle JP, Robinson P, Peers C (2009) α-Synuclein modulation of Ca²⁺ signaling in human neuroblastoma (SH-SYSY) cells. J Neurochem 111: 1192 – 1201
Holtmaat AJG, Trachtenberg JT, Wilbrecht L, Shepherd GM, Zhang X, Knott GW, Svoboda K (2005) Transient and persistent dendritic spines in the neocortex in vivo. Neuron 45: 279 – 291
Holtmaat A, Bonhoeffer T, Chow DK, Chuckowree J, De Paola V, Hofer SB, Hübener M, Keck T, Knott G, Lee W-CA et al (2009) Long-term, high-resolution imaging in the mouse neocortex through a chronic cranial window. Nat Protoc 4: 1128 – 1144
Hughes AJ, Daniel SE, Kilford L, Lees AJ (1992) Accuracy of clinical diagnosis of idiopathic Parkinson’s disease: a clinico-pathological study of 100 cases. J Neurol Neurosurg Psychiatry 55: 181 – 184
Hüls S, Högen T, Vassallo N, Danzer KM, Hengerer B, Giese A, Herms J (2011) AMPA-receptor-mediated excitatory synaptic transmission is enhanced by iron-induced α-synuclein oligomers: α-synuclein oligomers alter synaptic transmission. J Neurochem 117: 868 – 878
Jucker M, Walker LC (2011) Pathogenic protein seeding in Alzheimer disease and other neurodegenerative disorders. Ann Neurol 70: 532 – 540
Jung CKE, Fuhrmann M, Honarejadj K, Van Leuven F, Herms J (2011) Role of presenilin1 in structural plasticity of cortical dendritic spines in vivo: effect of PS1 on structural plasticity. J Neurochem 119: 1064 – 1073
Kostka M, Högen T, Danzer KM, Levin J, Habeck M, Wirth A, Wagner R, Glabe CG, Finger S, Heinzelmann U et al (2008) Single particle characterization of iron-induced pore-forming α-synuclein oligomers. J Biol Chem 283: 10992 – 11003
Kramer ML, Schulz-Schaeffer WJ (2007) Presynaptic α-synuclein aggregates, not lewy bodies, cause neurodegeneration in dementia with Lewy bodies. J Neurosci 27: 1405 – 1410
Kruger R, Kuhn W, Müller T, Woitalla D, Graeber M, Küsel S, Przuntek H, Epplen JT, Schöls L, Riess O (1998) Ala30Pro mutation in the gene encoding alpha-synuclein in Parkinson’s disease. Nat Genet 18: 106 – 108
Lücking CB, Brice A (2000) Alpha-synuclein and Parkinson’s disease. Cell Mol Life Sci 57: 1894 – 1908
Luk KC, Song C, O’Brien P, Stieber A, Branch J, Brunden KR, Trojanowski JQ, Lee VM-Y (2009) Exogenous α-synuclein fibrils seed the formation of Lewy body-like intracellular inclusions in cultured cells. Proc Natl Acad Sci USA 106: 20051 – 20056
Luk KC, Kehm V, Carroll J, Zhang B, O’Brien P, Trojanowski JQ, Lee VM-Y (2012a) Pathological α-synuclein transmission initiates Parkinson-like neurodegeneration in nontransgenic mice. Science 338: 949 – 953
Luk KC, Kehm VM, Zhang B, O’Brien P, Trojanowski JQ, Lee VM-Y (2012b) Intracellular inoculation of pathological α-synuclein initiates a rapidly progressive neurodegenerative α-synucleinopathy in mice. J Exp Med 209: 975 – 986
Luna E, Luk KC (2015) Bent out of shape: α-synuclein misfolding and the convergence of pathogenic pathways in Parkinson’s disease. FEBS Lett 589: 3749 – 3759
Masliah E (2000) Dopaminergic loss and inclusion body formation in α-synuclein mice: implications for neurodegenerative disorders. Science 287: 1265 – 1269
Masuda-Suzukake M, Nonaka T, Hosokawa M, Oikawa T, Arai T, Akiyama H, Mann DMA, Hasegawa M (2013) Prior-like spreading of pathological α-synuclein in brain. *Brain* 136: 1128–1138

McNeill TH, Brown SA, Rafols JA, Shoulson I (1988) Atrophy of medium spiny striatal dendrites in advanced Parkinson’s disease. *Brain Res* 455: 148–152

Miyamoto A, Wake H, Ishikawa AW, Eto K, Shibata K, Murakoshi H, Koizumi S, Moorman AJ, Yoshimura Y, Nabekura J (2016) Microglia contact induces synapse formation in developing somatosensory cortex. *Nat Commun* 7: 12540

Morris GP, Clark IA, Zinn R, Vissel B (2013) Microglia: a new frontier for synaptic plasticity, learning and memory, and neurodegenerative disease research. *Neurobiol Learn Mem* 105: 40–53

Murmu RP, Li W, Holtmaat A, Li Y (2013) Dendritic spine instability leads to progressive neocortical spine loss in a mouse model of Huntington’s disease. *J Neurosci* 33: 12997–13009

Nemani VM, Lu W, Berge V, Nakamura K, Onoa B, Lee MK, Chaudhry FA, Nicoll RA, Edwards RH (2010) Increased expression of α-synuclein reduces neurotransmitter release by inhibiting synaptic vesicle reclustering after endocytosis. *Neuron* 65: 66–79

Neuner J, Ovsepian SV, Dorostkar M, Filser S, Gupta A, Michalakis S, Biel M, Morris GP, Clark IA, Zinn R, Vissel B (2012) Amyloidogenic α-synuclein seeds do not invariably induce rapid, widespread pathology in mice. *Acta Neuropathol* 127: 645–665

Schmidt F, Levin J, Kamps F, Kretzschmar H, Giese A, Bötzler K (2012) Single-channel electrophysiology reveals a distinct and uniform pore complex formed by α-synuclein oligomers in lipid membranes. *PLoS One* 7: e42545

Shepherd CMG (2013) Corticostriatal connectivity and its role in disease. *Nat Rev Neurosci* 14: 278–291

Shrivastava AN, Redeker V, Fritz N, Pieri L, Almeida LG, Spolidoro M, Liebmann T, Bousset L, Renner M, Len A et al (2015) α-Synuclein assemblies sequester neuronal 3-Na⁺/K⁺-ATPase and impair Na⁺ gradient. *EMBO J* 34: 2408–2423

Oaks AW, Franklum F, Finkelstein DJ, Sidhu A (2013) Age-dependent effects of A5TF α-synuclein on behavior and dopaminergic function. *PLoS One* 8: e60378

Osterberg VR, Spinelli KJ, Weston LJ, Luk KC, Wolter R, Unni VK (2015) Progressive aggregation of alpha-synuclein and selective degeneration of Lewy inclusion-bearing neurons in a mouse model of Parkinsonism. *Cell Rep* 10: 1252–1260

Oueslati A (2016) Implication of alpha-synuclein phosphorylation at S129 in synucleinopathies: what have we learned in the last decade? *J Parkinsons Dis* 6: 39–51

Parkkinen L, Kauppinen T, Pirrittilä T, Autere JM, Alafuzzof I (2005) α-Synuclein pathology does not predict extrapyramidal symptoms or dementia. *Ann Neurol* 57: 82–91

Patt S, Gertz HJ, Gerhard L, Cervós-Navarro J (1991) Pathological changes in dendrites of substantia nigra neurons in Parkinson’s disease: a Golgi study. *Histo Histopathol* 6: 373–380

Penzes P, Cahiill ME, Jones KA, VanLeeuwen J-E, Woolfrey KM (2011) Dendritic spine pathology in neuropsychiatric disorders. *Nat Neurosci* 14: 285–293

Perry VH, Nicoll JAR, Holmes C (2010) Microglia in neurodegenerative disease. *Nat Rev Neurol* 6: 193–201

Picconi B, Piccoli G, Calabresi P (2012) Synaptic dysfunction in Parkinson’s disease. In *Synaptic plasticity: dynamics, development and disease*, Kreutz RM, Sala C (eds), pp 553–572. Vienna: Springer

Polymeropoulos MH, Lavedan C, Leroy E, Ide SE, Deheja A, Dutra A, Pike B, Root H, Rubenstein J, Boyer R et al (1997) Mutation in the alpha-synuclein gene identified in families with Parkinson’s disease. *Science* 276: 2045–2047

Rochefort NL, Konnerth A (2012) Dendritic spines: from structure to in vivo function. *EMBO Rep* 13: 699–708

Rockenstein E, Mallory M, Hashimoto M, Song D, Shults CW, Lang I, Masliah E (2002) Differential neuropathological alterations in transgenic mice expressing alpha-synuclein from the platelet-derived growth factor and Thy-1 promoters. *J Neurosci Res* 68: 568–578

Rockenstein E, Schwach G, Ingel O, Adame A, Crews L, Mante M, Pfragner R, Schreiner E, Windsisch M, Masliah E (2005) Lysosomal pathology associated with alpha-synuclein accumulation in transgenic models using an eGFP fusion protein. *J Neurosci Res* 80: 247–259

Sacino AN, Brooks M, McKinney AB, Thomas MA, Shaw G, Golde TE, Giasson BI (2014a) Brain injection of α-synuclein induces multiple proteinopathies, glosis, and a neuronal injury marker. *J Neurosci* 34: 12368–12378

Sacino AN, Brooks M, Thomas MA, McKinney AB, McCarvy NH, Rutherford NJ, Ceballos-Diaz C, Robertson J, Golde TE, Giasson BI (2014b) Amyloidogenic α-synuclein seeds do not invariably induce rapid, widespread pathology in mice. *Acta Neuropathol* 127: 645–665

Spillantini MG, Crowther RA, Jakes R, Goedert M (1997) Alpha-synuclein in Lewy bodies. *Nature* 388: 839–840

Spillantini MG, Crowther RA, Jakes R, Hasegawa M, Goedert M (1998) α-Synuclein in filamentous inclusions of Lewy bodies from Parkinson’s disease and dementia with Lewy bodies. *Proc Natl Acad Sci USA* 95: 6469–6473

Suárez LM, Solis O, Caramés JM, Taravini IR, Solís M, Murer MG, Morallora R (2014) L-DOPA treatment selectively restores spine density in dopamine receptor D2–expressing projection neurons in dyskinetic mice. *Biol Psychiatry* 75: 711–722

Tang Y, Le W (2016) Differential roles of M1 and M2 microglia in neurodegenerative diseases. *Mol Neurobiol* 53: 1181–1194

Tenreiro S, Reimão-Pinto MM, Antas P, Rino J, Wawrzycka D, Macedo D, Rosado-Ramos R, Amen T, Waiss M, Magalhães F et al (2014) Phosphorylation modulates clearance of alpha-synuclein inclusions in a yeast model of parkinson’s disease. *PLoS Genet* 10: e1004302

Terry RD, Masliah E, Salmon DP, Butters N, DeTeresa R, Hill R, Hansen LA, Katzman R (1991) Physical basis of cognitive alterations in Alzheimer’s disease: synapse loss is the major correlate of cognitive impairment. *Ann Neurol* 30: 572–580

Trachtenberg JT, Chen BE, Knot GW, Feng G, Sanes JR, Welker E, Svoboda K (2002) Long-term in vivo imaging of experience-dependent synaptic plasticity in adult cortex. *Nature* 420: 788–794

Tsaneva-Atanasova K, Burgos A, Galli T, Holcman D (2009) Quantifying neurite growth mediated by interactions among secretory vesicles, microtubules, and actin networks. *Biophys J* 96: 840–857

Alpha-synuclein mediates spine pathology

Sonja Blumenstock et al

---

**EMBO Molecular Medicine** Vol 9 | No 5 | 2017

© 2017 The Authors
Volles MJ, Lee S-J, Rochet J-C, Sholerman MD, Ding TT, Kessler JC, Lansbury PT (2001) Vesicle permeabilization by protofibrillar α-synuclein: implications for the pathogenesis and treatment of Parkinson's disease. *Biochemistry* 40: 7812 – 7819

Volpicelli-Daley LA, Luk KC, Patel TP, Tanik SA, Riddle DM, Stieber A, Meaney DF, Trojanowski jQ, Lee VM-Y (2011) Exogenous α-synuclein fibrils induce Lewy body pathology leading to synaptic dysfunction and neuron death. *Neuron* 72: 57 – 71

Wagner J, Ryazanov S, Leonov A, Levin J, Shi S, Schmidt F, Prix C, Pan-Montojo F, Bertsch U, Mitteregger-Kretzschmar G et al (2013) A novel oligomer modulator for disease-modifying therapy of neurodegenerative diseases such as prion and Parkinson's disease. *Acta Neuropathol* 125: 795 – 813

Winner B, Regensburger M, Schreglimann S, Boyer L, Prots I, Rockenstein E, Mante M, Zhao C, Winkler J, Masliah E et al (2012) Role of α-synuclein in adult neurogenesis and neuronal maturation in the dentate gyrus. *J Neurosci* 32: 16906 – 16916

Yuste R, Bonhoeffer T (2001) Morphological changes in dendritic spines associated with long-term synaptic plasticity. *Annu Rev Neurosci* 24: 1071 – 1089

Zaja-Milatovic S, Keene CD, Montine KS, Leverenz JB, Tsuang D, Montine TJ (2006) Selective dendritic degeneration of medium spiny neurons in dementia with Lewy bodies. *Neurology* 66: 1591 – 1593

Zarranz JJ, Alegre J, Gómez-Esteban JC, Lezcano E, Ros R, Ampuero I, Vidal L, Hoenicka J, Rodriguez O, Atarés B et al (2004) The new mutation, E46K, of α-synuclein causes parkinson and Lewy body dementia. *Ann Neurol* 55: 164 – 173

License: This is an open access article under the terms of the Creative Commons Attribution 4.0 License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited.