A30P α-Synuclein interferes with the stable integration of adult-born neurons into the olfactory network

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Impaired olfaction is an early symptom in Parkinson disease (PD), although the exact cause is as yet unknown. Here, we investigated the link between PD-related mutant α-Synuclein (α-SYN) pathology and olfactory deficit, by examining the integration of adult-born neurons in the olfactory bulb (OB) of A30P α-SYN overexpressing mice. To this end, we chose to label one well-known vulnerable subpopulation of adult-born cells, the dopaminergic neurons. Using in vivo two-photon imaging, we followed the dynamic process of neuronal turnover in transgenic A30P α-SYN and wild-type mice over a period of 2.5 months. Our results reveal no difference in the number of cells that reach, and possibly integrate into, the glomerular layer in the OB. However, in mutant transgenic mice these new neurons have a significantly shortened survival, resulting in an overall reduction in the addition of neurons to the glomerular layer over time. We therefore propose unstable integration and impaired homeostasis of functional new neurons as a likely contributor to odour discrimination deficits in mutant α-SYN mice.

Hyposmia is one of the most frequent non-motor symptoms of Parkinson disease (PD)1–3, being detected in the majority of PD patients several years before the first appearance of motor symptoms4. Moreover, deposition of fibrillar α-synuclein (α-SYN) aggregates5–6, the main pathological hallmark of the disease - occurs inside olfactory bulb (OB) neurons of PD patients5,4. Despite evidence of olfactory deficits in α-SYN overexpressing mice4, no satisfactory explanation has yet been put forward which correlates the neuropathological findings with the behavioural deficit. Interestingly, a deficit of neuronal precursor cells was discovered in the OB of PD patients compared with healthy controls10. Studies in mice further linked α-SYN overexpression to reduced survival of adult-born OB neurons11–14, including both periglomerular neurons (PGNs) and granule cells (GCs). Despite the evidence gained from these studies, we still lack a thorough investigation of the long-term effects of α-SYN aggregates on this highly dynamic process of neuronal turnover which occurs in the OB15. Undoubtedly, continuous incorporation of ‘virgin’ neurons into pre-existing networks poses a considerable physiological challenge, since their survival depends very much on the activity of local circuits and extrinsic influences16–20. Functional synaptic plasticity and integration requires communication with network neurons at the integration site and is a prerequisite for their contribution to olfactory circuits21–23. For instance, by synapsing with mitral cells (MCs), adult-born neurons sharpen odour tuning and provide contrast enhancement, important determinants for odour discrimination24,25. It is thus not surprising that this complex, fine-tuned, turnover of gained-and-lost neurons is closely coupled with olfactory stimulation22. Thus, odour deprivation decreases the number of new neurons in the OB25, whereas odour enrichment promotes their survival20.

In our study, we aimed at correlating PD-related α-SYN pathology in the OB network with olfactory deficits and the fate of adult-born neurons. To this end, we investigated in detail the α-SYN pathology in the OB of A30P α-SYN mice26 and its impact on olfaction. Using chronic intravital two-photon imaging, we followed the turnover of adult-born neurons in the OB of transgenic α-SYN and wild-type mice over a period of 2.5 months. We focused on dopaminergic PGNs as a subset of adult-born neurons, because (i) these are known to be primarily affected in PD27,28, (ii) they can be selectively labeled, and (iii) they have a superficial location thus making them accessible to two-photon imaging.
Herein, we report that aggregation-prone A30P α-SYN interferes with the stable integration and longer-term survival of adult-born neurons, with possible consequences on odour discrimination.

**Results**

**A30P α-SYN mice are impaired in fine odour discrimination.** For the initial characterization of PD-related pathology, we examined α-SYN aggregation in the OB of 6-month-old mice overexpressing human A30P α-SYN under the Thy1-promoter. The A30P mutation is known to cause severe, early-onset familial PD, accelerating the formation of toxic protein protofibrils. Immunofluorescent staining with human α-SYN-specific antibody revealed broad protein expression throughout the OB network including its three main cell layers: glomerular, MC and GC layer (Fig. 1a). In comparison to mice with elevated α-SYN levels in neuroblasts and immature newborn cells (PDGFβ-promoter), protein expression under neuron-specific Thy1 starts relatively late, thus providing a suitable model for the study of neuronal integration and fate in α-SYN-loaded networks in early-stage PD.

In PD, α-SYN is post-translationally modified (glycated, nitrated, oxidized or phosphorylated) and thereby more prone to forming aggregates and fibrils. Thus, we characterized one form of pathological protein alteration by staining against phosphorylated α-SYN, a major component of Lewy bodies (LB) in the human brain. In transgenic, but not wild-type, mice phosphorylated α-SYN was detected in the wide majority of MCs (Fig. 1b,d), while the glomerular (Fig. 1c) and GC layers (Fig. 1e) were largely spared. In addition, α-SYN was characterized by its resistance to proteinase K (PK) digestion, a property shared in common with fibrillar amyloid-β, the major component of Alzheimer disease-related amyloid plaques, and with the disease-associated form of prion protein. α-SYN clusters were detected in some MCs of the transgenic mice (Fig. 1f,h) but were absent from wild-type mice. Rarely, we detected subcellular α-SYN accumulations in the glomerular (Fig. 1g) and GC layer (Fig. 1i) of mutant α-SYN mice. These findings in Thy1-A30P α-

![Figure 1](https://www.nature.com/scientificreports/)

**Figure 1 | Olfactory bulb pathology in A30P α-SYN mice.** (a) Immunofluorescent micrographs showing the overexpression pattern of human A30P α-SYN (15G7, red) under control of the Thy1-promoter in the OB of 6-month-old transgenic mice: z-stack projections of confocal series. Scale bar - 50 μm. (b–e) Representative OB sections immunostained for phosphorylated α-SYN (pSer129, DAB), a marker for aberrant protein modification. Scale bars - 50 μm. (b) Distribution of phosphorylated α-SYN in (c) PGNs, (d) MCs and (e) GCs. Note that only MCs are positive for pSer129. (f–j) Representative OB sections with immunostaining for fibrillar protein, detected by Proteinase K digestion and human α-SYN specific antibody (15G7, DAB). Scale bars - 50 μm. Images with higher magnification of the (g) glomerular, (h) MC and (i) GC layer. Some α-SYN aggregates were detected in MCs of transgenic, but not wild-type, mice. (j) Olfactory discrimination task. Six-month-old Control (wild-type) (n = 8) and A30P α-SYN mice (n = 7) were exposed to different binary odour mixtures (e.g. 55% S1 and 45% S2 vs. 45% S1 and 55% S2). Mice were trained to dig in a bowl containing more of S1 odour. Correct choices for S1 were plotted against the different mixture pairs. Note that with increasing similarity of the mixtures, the performance of A30P α-SYN mice dropped compared to Control. (k) Olfactory memory task. **p < 0.01; ***p < 0.005; two-way ANOVA/Bonferroni post-hoc test.
SYN mice thus reflect in part the neuropathology in the human PD brain, showing LBs in MCs and the inner plexiform layer.

We further tested whether α-SYN accumulation in the OB is associated with olfactory deficits in the transgenic mice. To this end, we used a simple battery test7,16, which allows the detection of subtle differences in olfactory discrimination. Mice had to choose between increasingly similar solutions consisting of a mixture of two different odours (conditioned S⁻; S⁺), down to concentration differences of only 2% (Fig. 1j). Up to a 10% difference (55% S⁻ and 45% S⁺; vice versa), A30P α-SYN mice performed just as wild-type Control. However, at mixtures with 6% difference (53% S⁻ and 47% S⁺; vice versa) the performance of PD-like mice dropped significantly when compared to Control (75 ± 8%; n = 7 mice vs. 89 ± 8%; n = 8 mice, respectively; p < 0.01). This discrepancy between the two genotypes grew markedly with the next difficulty step. Whereas wild-type mice managed to discriminate 2% different odorant mixtures (51% S⁻ and 49% S⁺; vice versa) in 93 ± 8% of the trials, α-SYN mice succeeded in only 79 ± 9% (p < 0.001). As expected, 50% binary mixtures could not be distinguished by either group, with mice succeeded in only 79% S⁻ vs. 12% S⁺, respectively; 60% of all usable windows for high-quality microscopy.

Adult-born dopaminergic neurons are vulnerable to A30P α-SYN deposition. As an important aim of our study, we investigated the impact of A30P α-SYN accumulation on the dynamic integration process of adult-born neurons. Due to their superficial location in the OB and their particular role in PD27,28, we chose one subset of adult-born neurons: dopaminergic PGNs, which constitute ~20–40% of all GABAergic PGNs29–41. In the glomerular layer of the transgenic α-SYN mice, A30P α-SYN was enriched in both tyrosine hydroxylase (TH)-positive dopaminergic neurons and other PGN subtypes (Fig. 2a). Notably, A30P α-SYN had a significant impact on the density of TH-positive PGNs in both 6- (30%; 16,000 ± 1,700 vs. 23,000 ± 1,700 neurons per mm; n = 4 mice per group; p < 0.05) and 12-month-old transgenic mice (~47%; 16,100 ± 1,900 vs. 30,200 ± 2,600; n = 3 mice per group) when compared to Control (p < 0.05; Fig. 2b). While the dopaminergic neuron density remained on a similar level in transgenic α-SYN mice of both ages (p > 0.05), a clear trend towards an increase in these neurons was apparent in wild-type mice from 6 to 12 months (p = 0.06), consistent with previous reports15. Collectively, these results demonstrate a sensitivity of dopaminergic PGNs to toxic A30P α-SYN in the PD mouse model.

To confirm the negative effect of A30P α-SYN deposition on the survival of newborn dopaminergic PGNs, we performed BrdU-injections (Fig. 2c–e). Of note, A30P α-SYN mice showed ~50% reduction of the total new PGNs after 1 month (0.9 ± 0.1%; n = 6 mice vs. 1.8 ± 0.3% in Control; n = 5 mice; p < 0.05) (Fig. 2d), whereas cell proliferation was not affected (p > 0.05; not shown). Similar analysis in the GC layer revealed a comparable result: ~45% less adult-born GCs in the PD-like mice (2.0 ± 0.1%; n = 6 vs. 3.6 ± 0.5% in Control; n = 5) (p < 0.01; not shown). The fraction of newborn dopaminergic PGNs, characterized by co-expressing of TH and BrdU, was decreased to ~33% in the PD-like mice (27.7 ± 3.8%; n = 4 mice/97 cells vs. 41.6 ± 3.4%; n = 3 mice/91 cells, respectively; p < 0.05) (Fig. 2e). As expected, the loss of dopaminergic adult-born PGNs was less than in PDGFβ-human A53T α-SYN mice12 (70%), PDGFβ-human wild-type α-SYN mice11 (59%), and CaMK-human A30P α-SYN mice11 (41%). Together, these results confirm the reduced survival of adult-born PGNs and GCs in the OB of A30P α-SYN mice, with the dopaminergic PGNs as one vulnerable subpopulation.

Genetic labelling of dopaminergic neurons in A30P α-SYN mice. To specifically target dopaminergic PGNs for the in vivo study in α-SYN mice, we established a Cre/lox mouse model of PD (Fig. 3a). We first crossed knock-in driver lines expressing Cre recombinase from the TH locus (TH::IRESCre)39 with tdTomato reporter mice14. These double-transgenic wild-type mice were referred to as TH-Control. Further, crossing to Thy1-human A30P α-SYN mice26 enables the analysis of dopaminergic neurons with PD-like phenotype, cited as TH-A30P α-SYN. Previous studies reported a non-specific leak in this Cre/lox system34. To test the accuracy of genetic labelling of dopaminergic neurons within the glomerular layer of TH-Control and TH-A30P α-SYN mice (9-month-old), we stained OB sections with antibodies against TH and two other PGN markers39, Calbindin (CB) and Calretinin (CR) (Fig. 3b–d). In accordance with Adam and Mizrahi43, we detected tdTomato expression in the GC and MC layer (Fig. 3b), regions in which neurons normally do not express TH. Around the glomeruli, quantification of single and double-labelled neuronal somata expressing tdTomato with or without marker (Fig. 3c,d), revealed a majority of 71.0 ± 4.8% tdTomato-positive cells expressing simultaneously TH (n = 3 mice/1,698 cells). A very small proportion of tdTomato cells expressed CR (5.0 ± 7.5%; n = 4 mice/2,752 cells), while nearly none expressed CB (0.3 ± 0.2%; n = 4 mice/2,313 cells). To confirm stability of this labelling system within the time window for two-photon imaging, we compared these results to 12-month-old mice, revealing similar levels of tdTomato-positive cells expressing TH (69.9 ± 2.6%; n = 2 mice/226 cells; p > 0.05). Overall, these findings are similar to previous reports15 using heterozygous Z/EG GFP Cre reporter mice for the labelling of dopaminergic PGNs. Two months after transduction of newborn neurons with lentiviral (LV) eGFP constructs41,44 in the rostral migratory stream (RMS), we observed a small fraction of cells being positive for both eGFP and tdTomato (Fig. 3e), confirming that the TH-A30P α-SYN mouse labels adult-born dopaminergic PGNs. In conclusion, we consider this Cre/lox system appropriate for the genetic targeting of dopaminergic PGNs in the OB of transgenic α-SYN mice.

A30P α-SYN interferes with the stable integration of adult-born dopaminergic neurons in vivo. To monitor the dynamic integration and fate of the adult-born PGNs in vivo, we implanted an open-skull cranial window over both OBs (Fig. 4a–d). In line with previous reports for the OB13 and cortex15, we achieved with our preparation a success rate of ~60% usable windows for high-quality microscopy. Optical clarity for intravital two-photon imaging remained stable throughout the test period of 2.5 months. As open-skull surgery is associated with activation of glial cells in the OB12 and cortex45 within the first three weeks after operation, we started imaging 4 weeks post operatively to avoid the potential impact of gliosis on the analysis (Fig. 4e). Chronic imaging was performed in 9-month-old TH-A30P α-SYN and aged-matched TH-Control mice. Assuming a continuous neuronal turnover, we chose this time frame to elucidate in more detail the lack of dopaminergic PGNs observed in 6- and 12-month-old transgenic α-SYN mice (Fig. 2b). Because differentiation into the final fate of adult-born neurons requires synaptic contacts23,47, the appearance of tdTomato-labelled cell bodies indicated synaptic integration of newborn dopaminergic PGN, whereas disappearance of red somata clearly displayed the death of these neurons within the interneuron network35,48 (Fig. 4f,g). Accordingly, all the stable, lost or gained PGNs were tracked in a 2-week interval as summarized in Fig. 4h. In both genotypes, the neuron turnover rates were similar (0.8 ± 0.1; n = 4 mice/1,454 cells and 0.6 ± 0.2; n = 5 mice/1,437
cells; PD-like and Control, respectively) \(p < 0.05\); not shown). Of note, in TH-Control mice more PGNs gradually added to the network as assessed relative to the first imaging time point (Fig. 4i), resulting in a steady increase of these neurons up to 16.1 ± 5.5% after 2.5 months \(p < 0.01\). This continuous enlargement of the dopaminergic PGN population in wild-type mice both confirms our results shown in Fig. 2b and accords with a previous study of dopaminergic neuron turnover. In contrast, in TH-A30P α-SYN mice the number of PGNs marginally decreased over time to 23.8 ± 2.4%, but statistically remained on the same level (Fig. 4i; \(p < 0.05\)). Comparison of the fractional changes in the neuron density of both genotypes revealed significant differences at the last two time points \(p < 0.05\) and \(p < 0.01\), respectively. In addition, we determined the percentage of stable, lost and gained PGNs per imaging interval in both groups (Fig. 4k–m). Interestingly, both genotypes showed similar fractions of gained PGNs (Fig. 4l; 10.2 ± 1.4% vs. 10.7 ± 0.9%; PD-like and Control, respectively), while a significant increase in lost PGNs was recorded in the transgenic α-SYN mice (Fig. 4m; 11.4 ± 1.1% vs. 7.6 ± 1.0%; \(p < 0.05\)). To determine the timing of cell loss, we further quantified the fraction of cells that appeared at one imaging time point, but quickly disappeared in the next, as a parameter for integration stability (Fig. 4n). This readout further provides novel insight into the temporal definition of the toxic effect of A30P α-SYN. Notably, TH-A30P α-SYN mice showed more than double the amount of ‘short-lived’ cells per imaging interval compared to TH-Control (28.2 ± 1.2% vs. 12.7 ± 3.9%; \(p < 0.05\)). The higher neuron loss in PD-like mice was also reflected in a lower fraction of stable neurons (86.6 ± 2.7% vs. 92.2 ± 0.8%) \(p = 0.06\); Fig. 4k) and a 23% decrease in the overall survival rate over time (Fig. 4j). Indeed,
while 70.9 ± 5.5% PGNs, counted in TH-Control at the first time point, were still preserved after 2.5 months, this number was reduced to 54.7 ± 6.8% in TH-A30P α-SYN mice (p, 0.05). In conclusion, this detailed turnover analysis reveals deleterious consequences of A30P α-SYN on the stable incorporation of new PGNs, which reach the glomerular layer and appear to integrate into the abnormal ‘diseased’ network but are eliminated shortly afterwards.

Discussion

Our study provides the first in vivo evidence that A30P α-SYN accumulation interferes with the stable integration of the adult-born neurons in the OB, with significant effects on their survival. As summarized in Fig. 5, we describe a progressive increase in cell loss in the transgenic mutant α-SYN mice (~12%) as compared to wild-type (~8%), while at the same time the gain in new neurons is similar in both groups (~10%). Of note, this neuronal loss specifically targets those cells which have been recently integrated in the network. Such data demonstrates that early stages of incorporation and differentiation into dopaminergic neurons are not affected in transgenic Thy1-A30P α-SYN mice. Rather, it is their longer-term consistency and survival within the OB network which is defective, resulting in an overall reduction in the net addition of neurons to the OB over time. Consequently, PD-like mice lack the successful substitution of pre-existing adult-born neurons with new functional cells. Maintaining a stable, and precise, equilibrium between the influx of new neurons and the elimination of old neurons represents a key determinant of olfactory discrimination. We therefore propose a disturbed neuronal plasticity resulting from unstable integration of adult-born PGNs and GCs in A30P α-SYN mice, as a probable contributing factor to the observed olfactory discrimination deficits.

Furthermore, with this chronic, intravital two-photon imaging study of the integration kinetics of new neurons in transgenic α-SYN mice, we have also developed a potentially useful platform for the testing and evaluation of drugs that promote the integration and survival of neurons against α-SYN-induced toxicity. For this purpose, the OB is ideal since it is located superficially and thus easy to access, whilst representing at the same time the second main source of dopaminergic neurons in the brain. We studied this subset of adult-born neurons as representative of a vulnerable population of interneurons. Their susceptibility to increased amounts of α-SYN

Figure 3 | Genetic labelling of dopaminergic neurons in the olfactory bulb of A30P α-SYN mice. (a) Crossing scheme for targeting dopaminergic neurons in PD-like (top) and Control (wild-type; bottom) mice. Mice with Cre recombinase knocked into the TH locus were crossed to Ai9 tdTomato Cre reporter (TH-Control); and further to human A30P α-SYN mice (TH-A30P α-SYN). (b) Immunofluorescent micrographs showing tdTomato (red) and TH (green) expression in various OB layers in TH-A30P α-SYN mice, including ‘Merge’ view: z-stack projections of confocal series. Scale bar - 50 μm. (c) Representative high magnification images of one glomerulus with tdTomato-positive PGNs, stained for various subtype markers (c1) TH, (c2) calbindin (CB) or (c3) calretinin (CR) (green); z-stack projections of confocal series. White arrows point to double-labelled cells (Merge, yellow). Scale bar - 20 μm. (d) Summary plots showing the percentage of single/double-labelled PGN somata in TH-A30P α-SYN mice, expressing tdTomato with or without TH, CR and CB. TH from tdTomato, 71.0 ± 4.8% of tdTomato-positive cells expressed TH; tdTomato from TH, 98.0 ± 0.4% of TH cells expressed tdTomato (n = 1,698 cells from 3 mice); CB from tdTomato, 0.3 ± 0.2% of tdTomato-positive cells expressed CB (n = 2,313 cells from 4 mice); CR from tdTomato, 5.0 ± 7.5% of tdTomato-positive cells expressed CR (n = 2,752 cells from 4 mice). (e) High-resolution micrograph of a 2-month-old lentivirally-labelled (LV-eGFP; green) dopaminergic (tdTomato; red) adult-born PGN in TH-A30P α-SYN mice: z-stack projections of in vivo two-photon series. Scale bar - 20 μm.
had been demonstrated by several other studies, analysing variously proliferation, differentiation, survival and signalling of dopaminergic PGNs. A possible explanation for the vulnerability of dopaminergic neurons involves increased cellular toxicity in the presence of dopamine. Other groups report on the role of α-SYN in regulating dopamine biosynthesis, which is disturbed upon α-SYN aggregation. We further showed that increased cell death is not a unique feature of dopaminergic adult-born PGNs but appears also in other integrating neurons, because adult-born GCs were affected similarly. Our study specifically reveals that the critical step between early and stable integration/survival of the adult-born neurons, which depends on neuronal input and activity, is affected by A30P α-SYN aggregation. In favour of the altered network hypothesis, we show that transgenic A30P α-SYN mice are characterized by broad α-SYN accumulation in the OB, while only MCs - the main synaptic partners of adult-born PGNs and GCs - have pathology-associated α-SYN modifications. Interestingly, Thy1 expression in the OB stops in MCs, and transgene expression under Thy1 increases with neuron maturation. We thus attribute the specific location of the α-SYN pathology in MCs to their stability within the OB network, allowing gradual accumulation and deposition of the protein, whereas dynamic populations of PGNs and GCs are constantly replaced during adult neurogenesis. This study thus contributes to a better understanding of the role of α-SYN in regulating dopamine biosynthesis and the development of novel strategies for the treatment of neurological disorders related to α-SYN pathology.

**Figure 4** | Unstable integration of dopaminergic neurons in TH-A30P α-SYN mice in vivo. (a, b) Intravital two-photon microscopy of dopaminergic neuron turnover through a 3 mm round open-skull cranial window implanted over both OBs. Scale bar - 1 mm (a). (c) Blood vessel map of the region of interest (black box), as defined in (a). Scale bar - 100 µm. (d) Representative micrograph of the position (white box) marked in (c) with tdTomato-labeled dopaminergic PGNs: z-stack projection of two-photon image series. Scale bar - 10 µm. (e) Experimental time scheme of window surgery (black arrow) followed by in vivo two-photon imaging (grey arrow heads) in a two-week interval for a period of 2.5 months. Mice were 8 month old at the time point of surgery. Tracking of stable (red), gained (green) and lost (blue) PGNs in the (f1) first, (f2) second, and (f3) third imaging session, respectively: z-stack projection of two-photon image series. Scale bar - 10 µm. (g1, g2) High-magnification images of tdTomato-labelled PGNs: (g2) one lost cell (blue) in between a majority of stable somata (red) in two consecutive imaging sessions. Scale bar - 10 µm. (h) Table with absolute cell counts for stable (Nstable), lost (Nlost), gained (Ngained) and present (Npresent) cells at various imaging time points in both genotypes. (i) Quantification of the change (in %) in the PGN population over time relative to the first imaging session in TH-Control (n = 5) and TH-A30P α-SYN mice (n = 4). (j) Quantification of the neuronal survival rate (in %) relative to the first imaging session in TH-Control (n = 5) and TH-A30P α-SYN mice (n = 4). Summary plots showing the fraction of (k) stable, (l) gained, (m) lost, and (n) 'short-lived' neurons per imaging interval (%) in TH-Control (n = 5) and TH-α-SYN mice (n = 4). Note the high fraction of 'short-lived' neurons in TH-A30P α-SYN mice, namely PGNs which appeared in one session but quickly disappeared in the following. *p < 0.05, **p < 0.01; one way ANOVA/Tukey-Kramer post-hoc test (i); two-way ANOVA/Bonferroni post-hoc test (ij); Student’s t-test (k-n).
synaptic integration. Moreover, extracellular effects on the new neurons by secreted protein are possible. Definitely, the total dosage of \( \alpha \)-SYN is a significant factor to determine its toxicity in the cause of PD.

In conclusion, we demonstrate the pathological effects of aggregation-prone A30P \( \alpha \)-SYN on the homeostatic turnover of adult-born neurons, and on their stability and longer-term survival during integration into the OB network. Therefore, our transgenic mouse model with late Thy-1-A30P \( \alpha \)-SYN expression reflects, in part, the pathology observed in brains of PD patients with Lewy bodies in MCs of the OB, the human olfactory deficit at an early stage of the disease process, and a lack of progenitor cells. Importantly, our results support the previously reported association between adult neurogenesis and fine odour discrimination; although we still lack the molecular and cellular mechanism to explain olfactory deficits in these mice. Finally, a detailed investigation of the dendritic and synaptic maturation of these neurons along with an analysis of their synaptic connectivity to MCs will better our understanding of the effects of toxic/mutant/aggregated \( \alpha \)-SYN on odour discrimination deficiency in transgenic \( \alpha \)-SYN mice.

**Methods**

**Animals.** We used adult mice (5–12-month-old) that were bred and maintained in a specified pathogen-free animal facility: transgenic mice (C57Bl6 background) overexpressing human A30P \( \alpha \)-SYN under control of the Thy1-promoter and age-matched wild-type mice with same genetic background. For specific labeling of dopaminergic PGNs in A30P \( \alpha \)-SYN (1) and wild-type mice (2), the following two lines were crossed: (1) TH::IRE-S-Cre driver line \( \times \) A9 tdTomato Cre reporter line with floxed stop codon \( \times \) A30P \( \alpha \)-SYN line, cited as TH-A30P \( \alpha \)-SYN; (2) TH::IRE-S-Cre driver line \( \times \) A9 tdTomato Cre reporter line, referred to as TH-Control. In both lines, the floxed stop sequence is excised by the Cre enzyme and tdTomato expression is activated only in TH-positive neurons of the F1 offspring. All mice were kept in normal 12 h-light-dark cycle, had free access to food and water and showed no signs of inflammation or metabolic compromise. Genotypes were determined from mouse-tail DNA samples by PCR for tdTomato (forward primer: CGTTCTCCGTATGGGATGG; reverse primer: GGCATTAAAGCAGGATATCC) and human A30P \( \alpha \)-SYN (forward primer: TTAGCACTGCAGGGATAG; reverse primer: TTACCTTCTTCTCATGAG). Together, 58 animals were used in this study. All procedures were in accordance with an animal protocol approved by the University of Munich and the government of Upper Bavaria (Az. 55.2-1-54-2531-184-09).

**Immunohistochemistry and confocal microscopy.** Mice were perfused transcardially with 4% paraformaldehyde. For immunohistochemistry, the following primary antibodies were applied to 50 nm free-floating coronal OB sections: mouse monoclonal anti-tyrosine hydroxylase (TH) (1: 500), anti-calretinin (CR) (1: 1000) and anti-NeuN (1: 200) from Millipore, mouse monoclonal anti-calbindin (CB) (1: 500) from Sigma, rat monoclonal anti-Brdu (1: 200) from Dako, rabit monoclonal pSer129 \( \alpha \)-SYN (1: 500) from Abcam, and rat monoclonal antibody 15G7 against human \( \alpha \)-SYN (116–131) peptide MPVDPDIAYEMPSEE (1: 4000) from Connex. As secondary antibodies, we used: goat anti-mouse Alexa 488/647 (1: 500), rabbit anti-rat Alexa 488/647 (1: 500) from Invitrogen and biotinylated rabbit anti-rat (1: 300) from Dako. Enzymatic digestion using Proteinase K (PK; Roche) was employed for 8–10 min. Peroxidase detection was applied for 8–10 min (DAB detection kit, DCS Diagnostics). Sections were imaged at 0.75 pm pixels resolution in xy dimension and 1 pm in z dimension on a confocal microscope (Zeiss LSM 510) equipped with a 10X, 25X or 40X oil objective (Zeiss). Counting of somata and nuclei was performed manually from confocal image stacks, using Zeiss AIM software. For consistent analysis, every tenth OB slice was used and imaged at predefined positions within the glomerular layer.

**BrdU injections.** To study the survival of adult-born neurons, 5-month-old mice received intraperitoneal (i.p.) injections of bromodeoxouridine (BrdU; 50 mg/kg) for 5 consecutive days, and were perfused transcardially at 32 days after the 1st injection. To test possible effects of A30P \( \alpha \)-SYN on the proliferation of new cells, the mice received one single BrdU injection (100 mg/kg, i.p.), and were perfused after 24 hrs.

**Viral injection and open-skull craniotomy.** Mice were deeply anaesthetized with ketamine/xylazine (0.13:0.01 mg/g body weight: i.p.). Additionally, dexamethasone (6 mg/g body weight; i.p.; Sigma) was administered before surgery. Stereotaxic injection of LV particles for the transduction of adult-born neurons was performed, using pressure via a micropipette. Mouse received 300 nl injection on both left and right RMS (coordinates relative to Bregma: anterior –3.3 mm, lateral –0.8 mm, depth 2.9 mm), where new dopaminergic neurons are predominantly generated. Lentivirus, containing the gene for eGFP, controlled by the CAG promoter (plasmids from Dr. Stylianos Michalakis) was used at a titer of >10^8 infecting units ml^-1. Implantation of open-skull cranial windows over both OBs was performed in eight-month-old mice as described elsewhere. Briefly, the skull over both OBs was cleaned and dried. A round 3 mm craniotomy was opened over both OB hemispheres. Bone was carefully removed and a 3 mm round cover slip (VWR) was gently placed on the cavity and glued to the bone with thin layers of Histoacryl glue (B. Braun) and dental cement (Cyano Veneer fast; H. Schein). For repeated imaging in a custom-made stereotaxic stage, a 0.1 g metal bar was glued to the skull. After surgery, mice received a subcutaneous analgesic dose of carprofen (5 mg/g body weight Rimadyl; Pfizer) and the antibiotic cefotaxime (0.25 mg/g body weight, Phamore) for three days. Imaging began after a resting period of four weeks. In ~60% of the animals (9/15 mice) the brain region under the window was clear for imaging. If new bone grew...
under the window and impeded high-quality imaging, mice were excluded from the study.

**In vivo two-photon imaging.** Imaging was performed under light anaesthesia (ketamine/xylazine, 0.1/0.0075 mg/kg body weigh, ip). Mice were placed under the microscope in a custom-made holder with fixed position relative to the objective lens. Two-photon imaging data presented here were from 6 consecutive imaging sessions from 9 mice in 6 different litters. Mice took place over an imaging period of up to 2.5 months in a two-week interval. Imaging was performed using an upright two-photon microscope (Zeiss), equipped with a 2X water immersion objective (1.0 NA; Zeiss). TdTomato was excited with a femtosecond laser (MaiTai DeepSee, Spectra-Pics) at 915 nm and emission was collected at 527–582 nm, respectively (LSM 7MP, Zeiss). Images of cell bodies were acquired at 0.41 μm pixels resolution in xy dimension and 2 μm in z dimension (1024 × 1024 pixels). Same imaging volumes were precisely aligned over the imaging period. The laser intensity was adjusted to keep the emitted tDTomato fluorescence stable.

**Olfaction tests.** To detect differences in olfactory performance, 6-month-old mice were tested in a small discrimination task. Briefly, before the start of the experiment, mice were placed on a restricted diet to maintain a 10% reduced body weight during training and testing phase. Odorants were presented on fresh shavings (1 ml per 3 g shavings) in 2 round plastic dishes, mounted horizontally on a carrier, and separated by a vertical barrier between them. During training, the mice learned to dig in a dish with unscented shavings to find a small piece of chocolate. For the discrimination tasks, two identical dishes were presented simultaneously on the carrier. In the beginning, one dish was scented with apple odorant (designated as Sα; 10% Phenylthacacetate; Sigma) and the other with the same amount of solvent (DMSO; Thermo Fisher). Digging in the S+ dish was consistently rewarded with chocolate. To train the mice to associate the smell of Sα (and not of chocolate) with the reward, the chocolate was now buried in both scented shavings (15 successful trials). Mice had to discriminate between Sα and strawberry odorant (Sβ; 1% Methyl trans-cinnamate; Sigma). To increase the level of difficulty, the mice had to discriminate between different binary mixtures of Sα and Sβ with increasing similarity between the odors, [Sα : Sβ] vs [Sα : S0]: [70:30] vs [30:70]; [55:45] vs [45:55]; [53:47] vs [47:53]; [51:49] vs [49:51] and [50:50] vs [50:50]. A correct choice was defined as digging in the dish with the higher amount of Sα (8 trials/day). Successful choices were rewarded with a small piece of chocolate, whereas failed discrimination (choice of Sβ) immediately terminated the trial. To assess differences in olfactory long-term memory, the mice were confronted with the same odours again after 2 months, and the success in choosing the Sα odour was evaluated (12 trials).

**Data analysis and statistics.** Quantitative analysis of neuron turnover was performed manually from two-photon image stacks. Using Zeiss AIM software, from each stack in consecutive imaging sessions, several regions of interest were identified and cropped. Stacks were z-projected and each region of interest was compared with the same region in the previous session. Thereby, each cell was manually scored as stable (Nstable), lost (Nlost), or new (Nmade). Turnover ratios represent the gain and loss of neurons from day to day were calculated as DTOR = (Nstable + Nlost)/2 × Nmade/I, where Nmade is the number of all apparent neurons at a time point and I, is the number of days between subsequent imaging sessions. Fractional changes in the neuronal population were quantified for every time point relative to the first imaging session. Mean fractions of stable, lost and gained neurons were calculated for each time point and mean in addition, the neuronal cell numbers which both alpha-synuclein mutations linked to early-onset Parkinson’s disease: implications for pathogenesis and therapy. Proc. Natl. Acad. Sci. 97, 571–576 (2000).

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J.N., S.F. and J.H. designed the experiments. S.M. and M.B. provided relevant materials. All authors discussed the results and commented on the manuscript.

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