This is an Open Access document downloaded from ORCA, Cardiff University's institutional repository: https://orca.cardiff.ac.uk/id/eprint/3332/

This is the author's version of a work that was submitted to / accepted for publication.

Citation for final published version:


Publishers page: http://www.jneurosci.org/content/26/15/3942

Please note:
Changes made as a result of publishing processes such as copy-editing, formatting and page numbers may not be reflected in this version. For the definitive version of this publication, please refer to the published source. You are advised to consult the publisher’s version if you wish to cite this paper.

This version is being made available in accordance with publisher policies. See http://orca.cf.ac.uk/policies.html for usage policies. Copyright and moral rights for publications made available in ORCA are retained by the copyright holders.
Neurobiology of Disease

Pathological Changes in Dopaminergic Nerve Cells of the Substantia Nigra and Olfactory Bulb in Mice Transgenic for Truncated Human α-Synuclein(1–120): Implications for Lewy Body Disorders

George K. Tofariss,1 Pablo Garcia Reitböck,1 Trevor Humby,2 Sarah L. Lambourne,2 Mark O’Connell,1 Bernardino Ghetti,3 Helen Gossage,1 Piers C. Emson,2 Lawrence S. Wilkinson,3 Michel Goedert,4 and Maria Grazia Spillantinis1

1Centre for Brain Repair and Department of Clinical Neurosciences, University of Cambridge, Cambridge CB2 2PY, United Kingdom, 2The Babraham Institute, Cambridge CB2 4AT, United Kingdom, 3Department of Pathology and Laboratory Medicine, Indiana University School of Medicine, Indianapolis, Indiana 46202, and 4MRC Laboratory of Molecular Biology, Cambridge CB2 2QH, United Kingdom

Dysfunction of the 140 aa protein α-synuclein plays a central role in Lewy body disorders, including Parkinson’s disease, as well as in multiple system atrophy. Here, we show that the expression of truncated human α-synuclein(1–120), driven by the rat tyrosine hydroxylase promoter on a mouse α-synuclein null background, leads to the formation of pathological inclusions in the substantia nigra and olfactory bulb and to a reduction in striatal dopamine levels. At the behavioral level, the transgenic mice showed a progressive reduction in spontaneous locomotion and an increased response to amphetamine. These findings suggest that the C-terminal of α-synuclein is an important regulator of aggregation in vivo and will help to understand the mechanisms underlying the pathogenesis of Lewy body disorders and multiple system atrophy.

Key words: aggregation; behavior; tyrosine hydroxylase; fibril; dopamine; nigrostriatal; Parkinson; α-synuclein

Introduction

Parkinson’s disease (PD) is the most common movement disorder. Neuropathologically, it is defined by nerve cell loss in several brain regions, including the substantia nigra, and by the presence of Lewy bodies and Lewy neurites (Goedert, 2001; Braak et al., 2003). Abundant Lewy bodies and Lewy neurites in cerebral cortex are also the defining neuropathological characteristics of dementia with Lewy bodies (DLB), a common late-life dementia. Ultrastructurally, Lewy bodies and Lewy neurites are composed of filamentous and granular material (Forno, 1996). Missense mutations (A30P, E46K, and A53T) and multiplications of the α-synuclein gene cause dominantly inherited forms of PD and DLB (Polymeropoulos et al., 1997; Krüger et al., 1998; Singleton et al., 2002; Chartier-Harlin et al., 2004; Ibanez et al., 2004; Zarrañez et al., 2004), and α-synuclein is the major component of the filamentous inclusions of Lewy bodies and Lewy neurites (Spillantini et al., 1997, 1998a). The correlation between Lewy body formation and neurodegeneration suggests that the aggregation of α-synuclein is an important event during disease pathogenesis. Therefore, dissection of the mechanisms underlying inclusion formation in vivo is essential for understanding disease pathogenesis. In this respect, several mouse lines have been generated that express wild-type or mutant human α-synuclein from several different promoters (Kahle et al., 2000; Masliah et al., 2000; van derPutten et al., 2000; Matsuoka et al., 2001; Rathke-Hartlieb et al., 2001; Giasson et al., 2002; Lee et al., 2002; Neumann et al., 2002; Richfield et al., 2002).

Overexpression of wild-type human α-synuclein was found to lead to the formation of granular deposits and subtle biochemical abnormalities (Masliah et al., 2000). Subsequent work showed that mice overexpressing A30P and A53T human α-synuclein developed accumulation of α-synuclein in cell bodies and dystrophic neurites in several brain regions (van der Putten et al., 2000; Giasson et al., 2002; Lee et al., 2002; Neumann et al., 2002), with one study (Giasson et al., 2002) documenting the presence of filamentous α-synuclein deposits in cerebellum and spinal cord. However, none of these studies, which used either the Thy1 or the prion protein promoter, resulted in the accumulation of α-synuclein in dopaminergic nerve cells of the substantia nigra. Furthermore, expression of wild-type or mutant human α-synuclein from the tyrosine hydroxylase (TH) promoter did not result in the formation of pathological inclusions (Matsuoka et al., 2001; Rathke-Hartlieb et al., 2001; Richfield et al., 2002).
Previous work has shown that Lewy body extracts are enriched in C-terminally truncated α-synuclein (Baba et al., 1998; Tofarisi et al., 2003) and that human α-synuclein lacking the C-terminal 20 amino acids α-synuclein(1–120) (α-Syn120) assembles into filaments in vitro faster than either wild-type or mutant protein (Crowther et al., 1998; Serpell et al., 2000; Murray et al., 2003). It is therefore possible that truncation of α-synuclein could accelerate aggregation in a similar manner in vivo, resulting in neuropathological changes at an earlier age. Here, we investigated the effects of this modification by generating transgenic mice expressing human α-Syn120 from the rat TH promoter.

Materials and Methods

Generation of transgenic mouse lines. Transgene constructs were generated using the rat TH promoter (see Fig. 1A). The TH cassette (a kind gift from Dr. H. van der Putten, Novartis, Basel, Switzerland) (Min et al., 1994) directs high expression in dopaminergic and other TH-positive neurons. Human α-synuclein(1–120) DNA was generated from the wild-type cDNA (Jakes et al., 1994) by PCR amplification, and a Kozak consensus sequence was introduced upstream of the initiation codon. The expression construct was generated by subcloning human α-synuclein(1–120) DNA into the unique EcoRI site of the expression vector. Before microinjection, vector sequences were removed by digestion with NotI. Founders and transgenic progeny were identified by PCR analysis of lyses from tail biopsies using two sets of primers. Transgenic lines were established as C57BL/6 × CBA/ca hybrids and subsequently backcrossed to C57BL/6, which lacks α-synuclein (Specht and Schoepfer, 2001).

Immunoblotting. Total protein concentrations in tissue lyses were measured using the biocinchoninic acid assay (BCA; Pierce, Rockford, IL), and equal amounts were run on 12 or 15% SDS-PAGE. Proteins were measured using the biocinchoninic acid assay (BCA; Pierce, Rockford, IL). The membranes were blocked with 4% milk and incubated with the appropriate primary antibodies. Bound antibodies were then visualized using peroxidase-conjugated secondary antibodies (1:2000) and enhanced chemiluminescence (PerkinElmer Life Sciences, Boston, MA).

Immunohistochemistry. Mice were anesthetized with an intraperitoneal injection of xylazine and ketamine or pentobarbital, followed by intracardial perfusion with TBS and 4% ice-cold (w/v) paraformaldehyde. The brain was dissected and postfixed in 4% paraformaldehyde (PFA) in distilled water and placed in TBS, pH 7.6. Sections (50 μm thick) were cut using a vibratome. Immunogold labeling with antibody Syn-1 (1:100) was performed using 10 nm gold particles (Goldmark Biologicals, Phillipsburg, NJ) conjugated to goat anti-mouse IgG (diluted 1:10). Sections were postfixed in 2.5% glutaraldehyde, followed by 1% osmium tetroxide, dehydrated through a graded alcohol series, and embedded in epoxy resin. Semithin sections were stained with toluidine blue. Ultrathin sections were contrasted with lead citrate and uranyl acetate and scanned with electron microscopy.

Sequential extraction of α-synuclein. Extraction of α-synuclein was performed as described previously (Tofarisi et al., 2003). Tissues were homogenized on ice in 5 vol of TBS+ [50 mM Tris-HCl, pH 7.4, 175 mM NaCl, 5 mM EDTA, 0.1 mM PMSF, 1 mM N-ethylmaleimide, plus complete proteasome inhibitor cocktail (Roche Diagnostics, Mannheim, Germany)] and spun for 30 min at 120,000 × g at 4°C. The resulting supernatants were homogenized in TBS+ containing 1% Triton X-100, followed by centrifugation. The pellets were then homogenized in TBS+ containing 1% Triton X-100, followed by centrifugation. The pellets were then homogenized in TBS+ containing 1% Triton X-100, followed by centrifugation. The pellets were then homogenized in TBS+ containing 1% Triton X-100, followed by centrifugation. The pellets were then homogenized in TBS+ containing 1% Triton X-100, followed by centrifugation. The pellets were then homogenized in TBS+ containing 1% Triton X-100, followed by centrifugation. The pellets were then homogenized in TBS+ containing 1% Triton X-100, followed by centrifugation. The pellets were then homogenized in TBS+ containing 1% Triton X-100, followed by centrifugation. The pellets were then homogenized in TBS+ containing 1% Triton X-100, followed by centrifugation.

HPLC. Striatal monoamine levels were compared between transgenic and littermate control mice at 1, 3, 6, 9, and 12 months of age. The anterior striatum was dissected on ice, weighed, and homogenized in 0.1 M cold perchloric acid. Homogenates were centrifuged twice at 15,000 rpm for 5 min at 4°C, and the supernatants were diluted 1:10 in 0.1 M perchloric acid. They were stored at −80°C for 2 d. On the day of analysis, the supernatants were thawed and mixed with 3,4-dihydroxybenzylamine as an internal standard. Levels of dopamine, homovanillic acid, and a-amphetamine to modulate LMA

Behavioral testing. Twenty-three α-Syn120 transgenic and 25 wild-type control mice were assessed for spontaneous locomotor activity (LMA) and the effectiveness of a-amphetamine sulfate to modulate LMA at 6 and 18 months of age (n = 12 and 13 for wild-type mice at 6 and 18 months of age and n = 12 and 11 for transgenic mice at 6 and 18 months of age, respectively). Mice were individually placed into test chambers [for a description of the apparatus, see Isles et al. (2004)] for 30 min before administration of 1 mg/kg (i.p.) a-amphetamine sulfate or 0.7% saline vehicle and LMA, and then recorded for an additional 120 min. The mice were given two sessions, 7 d apart, and drug treatments were counterbalanced across groups and ages. All procedures were conducted in accordance with the requirements of the United Kingdom Animals
Results
Expression of human α-Syn120
Five founders were obtained, three of which transmitted the transgene in a Mendelian manner. By immunoblotting using antibody Syn-1, two lines expressed high levels of α-Syn120 in the nigrostriatal pathway and the olfactory bulb, brain regions expressing high levels of TH. One of these lines was used for additional characterization (Fig. 1B). Relative to immunoreactivity for TH, the expression of α-Syn120 was higher in the olfactory bulb than in the substantia nigra, and high levels of expression were detected in both regions at all ages examined (Fig. 1D,E).

When compared with the substantia nigra and the olfactory bulb of wild-type C57BL/6 mice with endogenous α-synuclein, transgenic α-Syn120 was present at lower levels than the endogenous protein (Fig. 1C). However, it has to be considered that although most neurons in the wild-type mouse express α-synuclein, transgenic α-Syn120 is only expressed in TH neurons. Antibodies PER4 and LS509, which are specific for the last 20 amino acids of α-synuclein (Jakes et al., 1999), failed to recognize the transgenic protein. As expected (Specht and Schoepfer, 2001), no mouse α-synuclein was detected in either transgenic lines or littermate controls. Heterozygous transgenic mice were used for this study.

Immunohistochemical distribution of human α-Syn120
The cellular localization of α-Syn120 was studied by single- and double-labeling immunohistochemistry. Strong immunoreactivity was present in the somatodendritic compartment of the vast majority of TH-positive neurons from 6 weeks of age onwards. Within the nigrostriatal pathway of young mice, staining for α-Syn120 closely followed TH staining throughout the cell soma and around the nucleus, extending into dendritic processes (Fig. 2). Colocalization of the two proteins was observed in 90% of cells. α-Syn120 was also present in the corresponding terminal fields, as indicated by punctate staining in the striatum (Fig. 3a). However, when stained with an antibody against TH, axons of transgenic mice appeared with focal varicose swellings in contrast to littermate controls. Using antiphosphorylated neurofilament antibodies, axons appeared to have a larger diameter in transgenic mice compared with wild-type α-synuclein in olfactory bulb from C57BL/6 mice. D, E, Immunoblotting (antibody Syn-1) of extracts from olfactory bulb (D) and substantia nigra (E) at 6, 12, and 24 months showed stable expression of the transgene. rS, Recombinant full-length human α-synuclein.

Figure 1. Expression of α-Syn120 in transgenic mouse brain. A, Schematic diagram of the construct: human α-synuclein(1–120) DNA was cloned downstream of the rat TH promoter. B, Immunoblot probed for α-synuclein (antibody Syn-1) and TH. The tissues were obtained from 6-week-old transgenic mice and littermate controls. Cx, Cerebral cortex; Ch, cerebellum; OB, olfactory bulb; SN, substantia nigra. No endogenous α-synuclein was detected at 19 kDa. C, Expression levels of α-Syn120 in olfactory bulb of transgenic mice compared with full-length α-synuclein in olfactory bulb from C57BL/6 mice. D, E, Immunoblotting (antibody Syn-1) of extracts from olfactory bulb (D) and substantia nigra (E) at 6, 12, and 24 months showed stable expression of the transgene. rS, Recombinant full-length human α-synuclein.
Transgenic mice were studied at 3, 6, and 12–14 months of age for the function of age-related changes in the striatum of transgenic (endogenous protein (Fig. 3). Granular deposits and vacuolation of the cytoplasm were also detected in older mice (Fig. 4j). Furthermore, there was an increase in microglial cells associated with affected brain regions, indicative of an inflammatory response (Fig. 5A, B). There was no significant cell loss by cell counting of TH-positive neurons in the substantia nigra using stereology (data not shown). Staining was observed with α-synuclein antibodies Syn-1, Syn h119, and PER7, but not with PER4 or LB509. Inclusions were not stained with anti-neurofilament or anti-ubiquitin antibodies (data not shown).

**Electron microscopy**
Pathological profiles were investigated by electron microscopy and immunogold labeling of olfactory bulb using antibody Syn-1. The α-Syn120 aggregates had a mixed granular and fibrillar morphology (Fig. 6a). A subset of inclusions in olfactory bulb and substantia nigra was thioflavin S-positive, consistent with the electron microscopic findings (Fig. 6b,c). Occasional inclusions were present in 3-month-old mice, and their numbers increased with age.

**Sequential extraction of human α-Syn120**
α-Synuclein was sequentially extracted from olfactory bulb and substantia nigra of 6-month-old transgenic mice using TBS, 1% Triton X-100, RIPA buffer, and 8 M urea/5% SDS (Fig. 7). A protein band with an apparent molecular mass of 12 kDa was detected with antibody Syn-1 in the TBS and Triton X-100 fractions. It corresponds to monomeric α-Syn120. When the RIPA-insoluble material was treated with urea, in addition to a high-molecular weight smear, a band corresponding to monomeric truncated α-Syn120 was found in substantia nigra and olfactory bulb. The smear and the monomeric α-Syn120 band present in the urea fraction were stained by antibodies Syn-1, Syn204, and PER7 (Fig. 7). The α-Syn120 aggregate in a case of PD was extracted in parallel and the urea-soluble material was stained with anti-neurofilament or anti-ubiquitin antibodies (data not shown).

**Staining for human α-Syn120 in substantia nigra as a function of age**
Transgenic mice were studied at 3, 6, and 12–14 months of age (n = 3 per time point). At 3 months (Fig. 4a), α-Syn120 accumulated in the somatodendritic compartment, where its staining pattern resembled that of TH. By 12–14 months of age, pathological changes were detected (Fig. 4b–j). They included dense, shrunken perikarya (Fig. 4b), beaded or dystrophic processes (Fig. 4c,e), as well as inclusions that were either perinuclear (Fig. 4dj, arrows) or very localized (Fig. 4f,g, arrowheads). Double immunostaining for both TH and α-synuclein confirmed the localization of dense inclusions within the cytoplasm of dopaminergic neurons (Fig. 4g). Granular deposits and vacuolation of the cytoplasm were also detected in older mice (Fig. 4h–j). Furthermore, there was an increase in microglial cells associated with affected brain regions, indicative of an inflammatory response (Fig. 5A, B). There was no significant cell loss by cell counting of TH-positive neurons in the substantia nigra using stereology (data not shown). Staining was observed with α-synuclein antibodies Syn-1, Syn h119, and PER7, but not with PER4 or LB509. Inclusions were not stained with anti-neurofilament or anti-ubiquitin antibodies (data not shown).

**Neurochemical changes**
The impact of human α-Syn120 expression on dopamine synthesis and turnover was assessed in the anterior striatum of transgenic mice at 1, 3, 6, 9, and 12 months of age (n = 4 per group).
When compared with age-matched littermate controls, although striatal dopamine levels were the same at 1 month, a statistically significant 30% reduction ($p < 0.0001, F = 32.8, R^2 = 0.653$) was detected at later time points in the transgenic mice (Fig. 8). This was paralleled by a 30% reduction in the dopamine metabolite homovanillic acid (Fig. 8). Measurement of 5-HT levels showed no significant difference between transgenic and littermate control mice (Fig. 8).

**Spontaneous locomotor activity**

The effect of α-Syn120 expression on spontaneous activity of 6- and 18-month-old transgenic mice is shown in Figure 9, A and B. At 6 months, there was no significant difference in total activity between wild-type and transgenic mice, whereas at 18 months, transgenic mice made significantly fewer beam breaks during the 30 min test than age-matched controls (AGE × GENOTYPE interaction, $F_{(1,44)} = 6.6, p < 0.02$); this was observed against a general reduction in locomotor activity by the older group of mice (effect of AGE, $F_{(1,44)} = 15.2, p < 0.001$). The pattern of reduced activity in 18-month-old transgenic mice was consistent throughout the session [Fig. 9B, no AGE × TIME × GENOTYPE interaction, $F_{(5,220)} = 0.3$, not significant (n.s.)]. However, all mice showed equivalent levels of habituation to the environment (effect of TIME, $F_{(5,220)} = 28.8, p < 0.001$), and the older mice made consistently fewer beam breaks in each 5 min time bin (AGE × TIME interaction, $F_{(5,220)} = 5.9, p < 0.001$).

**Effects of amphetamine**

After the 30 min habituation sessions, control and α-Syn120 transgenic mice were given an injection of either saline vehicle or amphetamine (Fig. 9C). When the drug treatment effects were calculated as a percentage of the final 10 min of the preinjection habituation session, 18-month-old transgenic mice exhibited an enhanced response to amphetamine relative to age-matched controls and 6-month-old transgenic mice and controls (effect of AGE × TREATMENT × GENOTYPE, $F_{(1,88)} = 6.8, p < 0.02$). Overall, amphetamine yielded a significant increase in locomotor activity (effect of TREATMENT, $F_{(1,88)} = 58.3, p < 0.001$) that peaked 60 min after administration (TIME × DRUG interaction, $F_{(3,264)} = 15.3, p < 0.001$). There were no genotype-related differences in the total number of beam breaks made in the 2 h after drug or vehicle injection (data not shown, effect of GENOTYPE, $F_{(1,88)} = 1.8$, n.s.), although 18-month-old mice made significantly fewer beam breaks than 6-month-old mice (effect of AGE, $F_{(1,88)} = 12.5, p < 0.001$). There was no interaction between genotype, age, and drug treatment ($F_{(1,88)} = 0.1$, n.s.).

**Discussion**

α-Synuclein is a presynaptic protein, which gives a characteristic punctate staining in rodent and human brain (Jakes et al., 1994). Similar findings were obtained for α-Syn120 in the striatum and olfactory bulb, indicating that it was transported like the full-length protein. However, the progressive accumulation of α-Syn120 in the somatodendritic compartment was pathological and associated with the accumulation of microglia and inflammation. In 3-month-old transgenic mice, α-synuclein showed a diffuse somatodendritic distribution; by 12–14 months, perikarya in the substantia nigra were shrunk or contained cytoplasmic inclusions, and axons in the striatum contained vacuole swellings and had an enlarged diameter, resembling the pathological profiles in PD (Duda et al., 2002; Neumann et al., 2004).

In the olfactory bulb, α-Syn120-positive inclusions were already present by 3 months of age. This is reminiscent of human brain, where Lewy bodies and Lewy neurites appear in the olfactory bulb before the substantia nigra (Braak et al., 2003). Olfactory dysfunction precedes clinical motor signs in PD (Katzenschlager and Lees, 2004). Electron microscopy and immunogold labeling of α-Syn120 deposits revealed the presence of granular and fibrillar structures in the olfactory bulb. Furthermore, some inclusions in substantia nigra and olfactory bulb were trichofilin S-positive, consistent with the presence of aggregated α-Syn120. By sequential extraction, we detected insoluble α-Syn120 aggregates in the olfactory bulb and the substantia nigra of transgenic mice. Their limited solubility was similar to that of α-synuclein extracted from PD and DLB brains (Campbell et al., 2000; Tofaris et al., 2003), although there was less insoluble material than in the human conditions. Furthermore, although in other reports the insoluble material appeared as a smear containing distinct bands as well as monomeric α-synuclein (Kahle et al., 2001), this was not the case here, where only one band corresponding to mono-
meric α-Syn 120 was present in the insoluble fraction. This difference could be attributable to the different methods of extraction, the nature of the transgenic protein (truncated versus full-length) or the amount of insoluble aggregates and their distribution, which was limited to substantia nigra and olfactory bulb in the α-Syn120 transgenic mice (Kahle et al., 2001; Giasson et al., 2002). The α-Syn120 inclusions were usually ubiquitin negative, consistent with the extensive evidence indicating that the ubiquitination of α-synuclein in Lewy bodies and Lewy neurites is a late event (Spillantini et al., 1998a; Tofaris et al., 2001, Sampathu et al., 2003; Tofaris et al., 2003). To our knowledge, this is the first transgenic mouse model with mixed filamentous and granular α-synuclein aggregates in dopaminergic neurons. It shows that truncated human protein expressed on a endogenous α-synuclein null background can induce inclusion formation in vivo.

The neurochemical effects of human α-Syn120 expression in the striatum were assessed in mice up to 12 months of age. Although at 1 month there was no difference in dopamine levels between transgenic mice and littermate controls, a significant reduction in dopamine and homovanillic acid, but not 5-HT, was detected from 3 months onwards. These findings indicate a specific dysfunction of the dopaminergic system, in agreement with the differential expression of α-Syn120, and are consistent with
neurochemical changes in PD, which is characterized by a deficiency in striatal dopamine and homovanillic acid (Hornykiewicz, 1998). Unlike in PD, there was no evidence of significant nerve cell death in our model. It is conceivable that the reduction in dopamine levels is best explained by a metabolic dysfunction resulting from the early accumulation of α-Syn120 in presynaptic nerve terminals in the striatum rather than by changes at the cell body level in the substantia nigra. In support of the latter is the temporal correlation between the presence of dilated striatal terminals and dopamine deficiency and the evidence from studies in human brain, which showed that, in PD, although striatal dopamine is decreased by 80%, nigral neurons are only reduced by 50% (Hornykiewicz, 1998).

The behavioral analysis, which revealed a progressive reduction in spontaneous locomotor activity and an increased response to amphetamine, was consistent with transgene-induced effects on dopamine-mediated functions. Similar findings have previously been reported in animal models of PD after chemical and genetic lesions, which reduce presynaptic dopamine (Cai et al., 2002; Marazziti et al., 2004). Furthermore, expression of full-length mutant, but not wild-type, α-synuclein under the control of the TH promoter led to a significant decrease in locomotor activity with aging (Thiruchelvam et al., 2004), suggesting that C-terminal truncation of the protein can have a pathological phenotype similar to that of a disease-causing missense mutation. The enhanced response to amphetamine is reminiscent of dopamine “supersensitivity,” a postsynaptic adaptation related to the chronic lowering of basal dopamine levels in terminal regions, especially the striatum. The synaptic mechanisms underlying this phenomenon can involve increased dopamine receptor numbers or be related to changes in binding efficiencies and/or downstream signal transduction pathways (Cai et al., 2002; Gerfen et al., 2002). Although several other transgenic mouse models of α-synuclein expression have been produced, they showed accumulation of α-synuclein in brain regions other than the substantia nigra (Giasson et al., 2002), or the inclusions were granular and nuclear (Masliah et al., 2000), or there were no inclusions (Thiruchelvam et al., 2004). Our model is the first to demonstrate a direct link between granular and filamentous α-synuclein cytolytic pathologies confined to the dopaminergic system and a progressive behavioral deficit without involvement of areas of motor function (i.e., corticospinal tract and anterior horn cells), which were affected in other models (Masliah et al., 2000; van der Putten et al., 2000; Giasson et al., 2002; Neumann et al., 2002).

These data, together with our immunohistochemical and neurochemical findings, raise the possibility that behavioral deficits may predate overt neurodegeneration and point to nerve terminals as possible anatomical substrates for the primary insult in α-synucleinopathies.

The above findings show that mice expressing human α-Syn120 in dopaminergic neurons recapitulate many changes characteristic of human Lewy body diseases. The major limitations of our model are the small number of fibrils, the absence of significant cell death, and the lack of endogenous full-length α-synuclein. One of our aims was to produce a model whereby the effect of aggregated human α-synuclein could be investigated without interference from endogenous mouse protein. A previous in vitro study has reported that the fibrillization of human α-synuclein is inhibited in the presence of the mouse protein (Rochet et al., 2000). However, truncated human α-synuclein can cross-seed the fibrillation of full-length protein in vitro (Murray et al., 2003). It will therefore be important to determine whether the presence of full-length protein influences the pathological and behavioral phenotype detected in α-Syn120 mice. C-terminal truncation of α-synuclein is considered by some to play a role in the pathogenesis of Lewy body diseases and the degeneration of dopaminergic nerve cells (Tofaris et al., 2003; Li et al., 2005; Liu et al., 2005). C-terminally truncated α-synuclein forms filaments at a faster rate than the full-length protein (Crowther et al., 1998; Serpell et al., 2000; Murray et al., 2003). Furthermore, C-terminally truncated α-synuclein has been detected in Lewy bodies in human diseases and in the brains of transgenic mice expressing mutant human α-synuclein (Giasson et al., 2002; Lee et al., 2002; Tofaris et al., 2003). How truncated α-synuclein can be generated in cells, under pathological conditions, is currently unclear; the proteosome could be involved (Tofaris et al., 2001; Liu et al., 2003). Isolated α-synuclein filaments from human brain are made of the full-length protein, suggesting that truncation may occur after assembly (Spillantini et al., 1998a,b; Crowther et al., 2000; Takao et al., 2004). This notwithstanding, it is clear that the C-terminal region of α-synuclein, which binds dopamine derivatives (Norris et al., 2005), is a negative regulator of self-assembly. Therefore, modifications in this region, such as oxidation, nitration, and phosphorylation (Hashimoto et al., 1999; Giasson et al., 2000; Fujiwara et al., 2002), may well influence the propensity of α-synuclein to aggregate in vivo in a way similar to truncation. The same is true of molecules that bind to the monomeric protein. Thus, polyamines have been shown to promote the aggregation of α-synuclein through binding to its C-terminal region (Antony et al., 2003; Fernandez et al., 2004). Other positively charged molecules may act in a similar way (Goers et al., 2003).
fical inclusions in substantia nigra and olfactory bulb, as revealed by morphological and biochemical studies. This was accompanied by a reduction in dopamine levels in the striatum, a progressive reduction in spontaneous locomotor activity, and an increased response to amphetamine. Our findings suggest that the C-terminal region of α-synuclein is an important regulator of aggregation.

References


