Dopamine induces soluble α-synuclein oligomers and nigrostriatal degeneration

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Parkinson’s disease (PD) is defined by the loss of dopaminergic neurons in the substantia nigra and the formation of Lewy body inclusions containing aggregated α-synuclein. Efforts to explain dopamine neuron vulnerability are hindered by the lack of dopaminergic cell death in α-synuclein transgenic mice. To address this, we manipulated both dopamine levels and α-synuclein expression. Nigrally targeted expression of mutant tyrosine hydroxylase with enhanced catalytic activity increased dopamine levels without damaging neurons in non-transgenic mice. In contrast, raising dopamine levels in mice expressing human A53T mutant α-synuclein induced progressive nigrostriatal degeneration and reduced locomotion. Dopamine elevation in A53T mice increased levels of potentially toxic α-synuclein oligomers, resulting in conformationally and functionally modified species. Moreover, in genetically tractable Caenorhabditis elegans models, expression of α-synuclein mutated at the site of interaction with dopamine prevented dopamine-induced toxicity. These data suggest that a unique mechanism links two cardinal features of PD: dopaminergic cell death and α-synuclein aggregation.

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RESULTS
Expression of TH-RREE elevates dopamine levels and causes hyperactivity in NonTg mice
To increase dopamine levels in vivo, we used a lentiviral vector containing the TH gene encoding human tyrosine hydroxylase (TH). TH is the rate-limiting enzyme in dopamine biosynthesis converting tyrosine to L-DOPA, which is then metabolized by aromatic amino acid decarboxylase to generate dopamine. A mutated TH enzyme that has R37R38 replaced with E37E38 (TH-RREE) is insensitive to feedback inhibition by dopamine and can be used to achieve increased catalytic activity of TH. We first tested the function of the vector on human neuroblastoma SH-SY5Y cells. This cell line has undetectable levels of endogenous TH, despite the expression of other enzymes required for synthesis and processing of dopamine. We transduced the cells after 5 d of differentiation with 20 µM retinoic acid and confirmed TH expression by western blot and immunocytochemistry compared with empty vector (CtrlVect)-transduced controls (Supplementary Fig. 1a–c). The TH-RREE transduced cells had measurable intracellular levels of L-DOPA, dopamine and DOPAC in the range of 25–200 fmol µg⁻¹, whereas catecholamines were not detected in CtrlVect-transduced cells (Supplementary Fig. 1d), indicating that TH-RREE vector enhances catecholamine production.

We performed bilateral injections of TH-RREE vector into the SN of aged (10 month old) nontransgenic (NonTg) mice. This age was selected because the mice expressing A53T mutant human α-synuclein are asymptomatic and do not exhibit the characteristic motor phenotype derived from spinal cord degeneration. The typical age of onset of this phenotype in our colony was approximately 18 months. At 5 months post-injection (mpi), TH expression was increased in both SN and striatum in NonTg TH-RREE mice compared withagematched NonTg CtrlVect mice (Fig. 1a,b). TH-RREE significantly elevated striatal concentrations of L-DOPA and dopamine by 36% (P = 0.0235) and 52% (P = 0.0137), respectively, whereas DOPAC was not affected. Notably, 5-HT levels did not change, indicating that the effect was specific for the catecholamine neurotransmitter system (Fig. 1c). Consistent with elevated striatal dopamine, NonTg TH-RREE mice exhibited hyperactivity in open field-testing compared with NonTg CtrlVect mice (Fig. 1d). There was no difference in rotarod performance between the NonTg injection groups (Fig. 1e). These data indicate that the TH-RREE vector substantially enhanced dopamine levels fully 5 months following injection.

Dopamine-induced neurodegeneration and motor deficit is dependent on α-synuclein expression
Bilateral injection of the TH-RREE vector into the SN of 10-month-old A53T mutant α-synuclein mice also resulted in TH overexpression (Supplementary Fig. 2). However, in contrast with NonTg mice, A53T TH-RREE mice exhibited striatal degeneration and neuronal loss in the SN at 5 mpi. To circumvent detection of vector-encoded TH, we used another marker of dopaminergic neurons, VMAT2, to evaluate neuronal loss. A53T TH-RREE mice had fewer VMAT2-positive neuronal loss in the SN at 5 mpi. To circumvent detection of vector-encoded TH, we used another marker of dopaminergic neurons, VMAT2, to evaluate neuronal loss. A53T TH-RREE mice had fewer VMAT2-positive neuronal loss in the SN at 5 mpi. To circumvent detection of vector-encoded TH, we used another marker of dopaminergic neurons, VMAT2, to evaluate neuronal loss. A53T TH-RREE mice had fewer VMAT2-positive neuronal loss in the SN at 5 mpi.

Figure 1 TH-RREE lentiviral vector increases dopamine levels and causes hyperactivity in NonTg mice. (a) TH protein levels were increased in the striatum of TH-RREE-vector-injected mice relative to empty-vector-injected controls (CtrlVect) at 5 mpi. Densitometric analysis was conducted by normalizing TH to the GAPDH loading control. Blots are cropped; for full-length blots see Supplementary Figure 11 (n = 3 mice per group, P = 0.0201, t = 3.742, d.f. = 4; two-tailed Student’s t test). (b) Increased TH expression was confirmed by immunohistochemistry in the SN (left) and striatum (right). Scale bars represent 200 µm (n = 4 mice per group). (c) TH overexpression significantly increased the steady state concentrations of striatal catecholamines L-DOPA and dopamine (DA), but did not alter DOPAC or 5-HT (L-DOPA, n = 4 mice per group, P = 0.0235, t = 3.017, d.f. = 6; DA, n = 4 mice per group, P = 0.0137, t = 3.445, d.f. = 6; DOPAC, n = 5 mice for CtrlVect and n = 4 mice for TH-RREE, P = 0.2713, t = 1.194, d.f. = 7; 5-HT, n = 4 mice for CtrlVect and n = 3 mice for TH-RREE, P = 0.658, t = 0.4703, d.f. = 5; two-tailed unpaired Student’s t test). (d) TH-RREE injected NonTg mice exhibited greater locomotion as measured by open field activity (n = 5 mice per group, P = 0.0399, t = 2.451, d.f. = 8; two-tailed unpaired Student’s t test). (e) There was no change in rotarod performance between the injection groups (n = 3 mice per group, P = 0.9959, t = 0.005455, d.f. = 4; two-tailed unpaired Student’s t test). Box plots show median, 25th and 75th percentiles, and minimum and maximum values. *P < 0.05.
cells in the SN than any of the other injection groups (Fig. 2a). We quantified the loss of neurons using unbiased stereological counting of Nissl cells, which revealed a significant 25% reduction in A53T TH-RREE mice compared with controls (Fig. 2b). The degeneration of cell bodies was accompanied by a significant 62% decrease in VMAT2 staining in the striatum of A53T TH-RREE mice as compared to A53T CtrlVct. Blots are cropped; for full-length blots see Supplementary Figure 11 (n = 3 mice per group except n = 4 mice for A53T TH-RREE, P_{A53T TH-RREE versus A53T CtrlVct} = 0.0049, P_{A53T TH-RREE versus NonTg CtrlVct} = 0.0316, P_{A53T TH-RREE versus NonTg TH-RREE} = 0.0253, F_{(3,9)} = 8.810; one-way ANOVA with Tukey's correction for multiple comparisons). (c,d) Histological analysis of VMAT2 staining in the striatum, with subtraction of background staining in the cortex, revealed severe dopaminergic denervation in A53T TH-RREE mice. Scale bar represents 200 \mu m (n = 3 mice per group). Consistent with progressive degeneration of synapses, we noted a time-dependent decline of striatal dopamine content in A53T TH-RREE mice as compared to A53T CtrlVct. Blots are cropped; for full-length blots see Supplementary Figure 11 (n = 3 mice per group except n = 4 mice for NonTg CtrlVct, P_{A53T TH-RREE versus A53T CtrlVct} = 0.0043, P_{A53T TH-RREE versus NonTg CtrlVct} = 0.0163, P_{A53T TH-RREE versus NonTg TH-RREE} = 0.0163, F_{(3,8)} = 10.02; one-way ANOVA with Tukey's correction for multiple comparisons). (e,f) Striatal DAT levels normalized to the actin loading control were increased in NonTg TH-RREE mice relative to NonTg CtrlVct, whereas DAT levels were decreased in A53T TH-RREE mice as compared to A53T CtrlVct. Box plots show median, 25th and 75th percentiles, and minimum and maximum values. *P < 0.05, **P < 0.01.

Figure 2 Dopamine-induced neurodegeneration of the SN is dependent on \( \alpha \)-synuclein. (a) At 5 mpi, fewer VMAT2-positive cells (arrows) were present in the SN of A53T TH-RREE mice than in any other injection groups. Scale bar represents 100 \mu m (n = 3 mice per group). (b) Unbiased stereological counting of Nissl-positive neurons in the SN revealed a significant 25% loss of cells in A53T TH-RREE mice. The data are presented as mean ± s.e.m. (n = 3 mice per group except n = 4 mice for A53T TH-RREE, P_{A53T TH-RREE versus A53T CtrlVct} = 0.0049, P_{A53T TH-RREE versus NonTg CtrlVct} = 0.0316, P_{A53T TH-RREE versus NonTg TH-RREE} = 0.0253, F_{(3,9)} = 8.810; one-way ANOVA with Tukey's correction for multiple comparisons). (c,d) Histological analysis of VMAT2 staining in the striatum, with subtraction of background staining in the cortex, revealed severe dopaminergic denervation in A53T TH-RREE mice. Scale bar represents 200 \mu m (n = 3 mice per group). Consistent with progressive degeneration of synapses, we noted a time-dependent decline of striatal dopamine content in A53T TH-RREE mice as compared to A53T CtrlVct. Blots are cropped; for full-length blots see Supplementary Figure 11 (n = 3 mice per group except n = 4 mice for NonTg CtrlVct, P_{A53T TH-RREE versus A53T CtrlVct} = 0.0043, P_{A53T TH-RREE versus NonTg CtrlVct} = 0.0163, P_{A53T TH-RREE versus NonTg TH-RREE} = 0.0163, F_{(3,8)} = 10.02; one-way ANOVA with Tukey's correction for multiple comparisons). (e,f) Striatal DAT levels normalized to the actin loading control were increased in NonTg TH-RREE mice relative to NonTg CtrlVct, whereas DAT levels were decreased in A53T TH-RREE mice as compared to A53T CtrlVct. Box plots show median, 25th and 75th percentiles, and minimum and maximum values. *P < 0.05, **P < 0.01.

receptors remained unchanged (Supplementary Fig. 3a,b). Levels of dopamine- and cAMP-regulated phosphoprotein-32 (DARPP-32), which serves as an important signaling molecule in spiny projection neurons, were also unaffected (Supplementary Fig. 3a,c).

To further investigate the progressive degenerative phenotype in A53T TH-RREE mice, we quantified the total number of SN neurons at the earlier time point of 2.5 mpi. No difference was observed between A53T CtrlVct and TH-RREE mice, indicating that cell loss had not yet occurred (Fig. 3a). Despite the maintenance of neuronal cell bodies, striatal VMAT2 levels were decreased by 25% (Fig. 3b), revealing a milder loss of nerve terminals than at the later time point of 5 mpi (Fig. 2c–d). Consistent with progressive degeneration of synapses, we noted a time-dependent decline of striatal dopamine content in A53T TH-RREE mice. At 2.5 mpi, dopamine was increased significantly by 71% compared with age-matched A53T CtrlVct mice as a result of mutant TH expression (Fig. 3c). However, A53T TH-RREE mice subsequently exhibited a 37% drop in dopamine concentration between 2.5 and 5 mpi, whereas A53T CtrlVct mice showed no change during the same period (Fig. 3c).
The reduction in dopamine levels cannot be explained by increased conversion to DOPAC, as levels of this metabolite remained unchanged from 2.5 to 5 mpi (Fig. 3d). We also did not detect an increase in dopamine-protein adducts by near infrared fluorescence (Supplementary Fig. 4). Rather, the loss of dopamine is likely a result of the ongoing nigrostriatal degeneration documented in these mice. Concomitant with the decrease in dopamine levels, A53T TH-RREE mice exhibited a significant impairment in ambulatory activity that was not observed in TH-RREE mice at 2.5 mpi, which had not yet exhibited loss of neuronal cell bodies in the SN. The data are presented as mean ± s.e.m. (n = 3 mice per group for CtrlVect and n = 3 mice for TH-RREE, P = 0.8453, t = 0.2055, d.f. = 5; two-tailed unpaired Student’s t test). (b) VMAT2 staining in the striatum revealed a modest loss of terminals, suggesting that dopaminergic synapses had degenerated before overt cell death. Scale bar represents 200 μm (n = 4 mice per group, P = 0.0424, t = 2.569, d.f. = 6; two-tailed unpaired Student’s t test). (c) A53T TH-RREE mice exhibited an elevation of dopamine (DA) levels only transiently, with an initial increase of 71% at 2.5 mpi compared with age-matched A53T CtrlVect mice. Subsequently, however, A53T TH-RREE mice underwent a significant 37% drop in striatal dopamine between 2.5 and 5 mpi, which was not observed in A53T CtrlVect mice (n = 3 mice per group except n = 5 mice for TH-RREE 2.5 mpi, P\textsubscript{TH-RREE 2.5 mpi versus CtrlVect 2.5 mpi} = 0.0092, F\textsubscript{TH-RREE 2.5 mpi versus TH-RREE 5 mpi} = 0.0187, F\textsubscript{(3,10)} = 7.460; one-way ANOVA with Tukey’s correction for multiple comparisons). (d) Levels of DOPAC in the striatum remained unchanged over time regardless of lentiviral treatment. (CtrlVect, n = 3 mice per group; TH-RREE, n = 6 mice for 2.5 mpi and n = 4 mice for 5 mpi, F\textsubscript{(3,12)} = 0.5481; one-way ANOVA with Tukey’s correction for multiple comparisons). (e) Consistent with a late-onset depletion of dopamine in the striatum, from 2.5 to 5 mpi, A53T TH-RREE mice developed a reduction in locomotor activity that was not observed in CtrlVect mice (n = 3 mice per group, P\textsubscript{TH-RREE 5 mpi versus CtrlVect 5 mpi} = 0.1512, P\textsubscript{TH-RREE 5 mpi versus TH-RREE 2.5 mpi} = 0.0488, F\textsubscript{TH-RREE 5 mpi versus CtrlVect 5 mpi} = 0.0213, F\textsubscript{(3,8)} = 7.107; one-way ANOVA with Tukey’s correction for multiple comparisons). (f) The motor deficit in A53T TH-RREE mice was not severe enough to affect coordination or balance, as rotarod performance remained intact (n = 4 mice per group except n = 3 mice for TH-RREE 2.5 mpi and n = 3 mice for CtrlVect 5 mpi, F\textsubscript{(3,10)} = 0.07172; one-way ANOVA with Tukey’s correction for multiple comparisons). Box plots show median, 25th and 75th percentiles, and minimum and maximum values. *P < 0.05, **P < 0.01.

Dopamine modifies α-synuclein oligomer conformations in A53T mice

These data indicate that increased steady state levels of dopamine result in neurotoxicity in an α-synuclein-dependent manner. To investigate possible mechanisms, we examined the influence of dopamine on α-synuclein aggregation and, in particular, the kinetic stabilization of potentially toxic α-synuclein oligomers. The presence of Lewy-body-like inclusions in circumscribed regions, including the brainstem of aged A53T mice, has been well-documented\textsuperscript{32}. α-synuclein-positive inclusions were indeed detected in the brainstem of A53T TH-RREE mice at 5 mpi (age 15 months), similar to age-matched A53T CtrlVect mice (Supplementary Fig. 5a). In contrast, the SN is known to remain devoid of inclusions during the lifespan of A53T mice\textsuperscript{32}. Expression of TH-RREE vector did not induce α-synuclein inclusion formation or alter levels or conformations of detergent-insoluble α-synuclein in the SN at 5 mpi (Supplementary Fig. 5).

Examination of detergent-soluble α-synuclein extracted from SN of A53T mice at 5 mpi revealed substantial alterations in the quantities and conformations of α-synuclein species as a result of dopamine elevation. A53T TH-RREE mice had a significant 22% reduction in monomeric human α-synuclein compared with A53T CtrlVect mice (Fig. 4a,b). Following fractionation of the soluble SN extract by native size exclusion chromatography (SEC), we detected the presence of

Figure 3 Dopaminergic neurodegeneration in A53T TH-RREE mice is progressive and ultimately leads to locomotor deficit. (a) At the early time point of 2.5 mpi, A53T TH-RREE mice did not yet exhibit loss of neuronal cell bodies in the SN. The data are presented as mean ± s.e.m. (n = 4 mice for CtrlVect and n = 3 mice for TH-RREE, P = 0.8453, t = 0.2055, d.f. = 5; two-tailed unpaired Student’s t test). (b) VMAT2 staining in the striatum revealed a modest loss of terminals, suggesting that dopaminergic synapses had degenerated before overt cell death. Scale bar represents 200 μm (n = 4 mice per group, P = 0.0424, t = 2.569, d.f. = 6; two-tailed unpaired Student’s t test). (c) A53T TH-RREE mice exhibited an elevation of dopamine (DA) levels only transiently, with an initial increase of 71% at 2.5 mpi compared with age-matched A53T CtrlVect mice. Subsequently, however, A53T TH-RREE mice underwent a significant 37% drop in striatal dopamine between 2.5 and 5 mpi, which was not observed in A53T CtrlVect mice (n = 3 mice per group except n = 5 mice for TH-RREE 2.5 mpi, P\textsubscript{TH-RREE 2.5 mpi versus CtrlVect 2.5 mpi} = 0.0092, P\textsubscript{TH-RREE 2.5 mpi versus TH-RREE 5 mpi} = 0.0187, F\textsubscript{(3,10)} = 7.460; one-way ANOVA with Tukey’s correction for multiple comparisons). (d) Levels of DOPAC in the striatum remained unchanged over time regardless of lentiviral treatment. (CtrlVect, n = 3 mice per group; TH-RREE, n = 6 mice for 2.5 mpi and n = 4 mice for 5 mpi, F\textsubscript{(3,12)} = 0.5481; one-way ANOVA with Tukey’s correction for multiple comparisons). (e) Consistent with a late-onset depletion of dopamine in the striatum, from 2.5 to 5 mpi, A53T TH-RREE mice developed a reduction in locomotor activity that was not observed in CtrlVect mice (n = 3 mice per group, P\textsubscript{TH-RREE 5 mpi versus CtrlVect 5 mpi} = 0.1512, P\textsubscript{TH-RREE 5 mpi versus TH-RREE 2.5 mpi} = 0.0488, F\textsubscript{TH-RREE 5 mpi versus CtrlVect 5 mpi} = 0.0213, F\textsubscript{(3,8)} = 7.107; one-way ANOVA with Tukey’s correction for multiple comparisons). (f) The motor deficit in A53T TH-RREE mice was not severe enough to affect coordination or balance, as rotarod performance remained intact (n = 4 mice per group except n = 3 mice for TH-RREE 2.5 mpi and n = 3 mice for CtrlVect 5 mpi, F\textsubscript{(3,10)} = 0.07172; one-way ANOVA with Tukey’s correction for multiple comparisons). Box plots show median, 25th and 75th percentiles, and minimum and maximum values. *P < 0.05, **P < 0.01.
various oligomeric α-synuclein species of different molecular weights (Fig. 4c). Quantification of total oligomeric species revealed a significant increase in A53T TH-RREE mice, greater than twofold above that in A53T CtrlVect mice (Fig. 4d). To characterize the oligomers, we used multiple α-synuclein antibodies and a combination of SDS-PAGE, immunoelectron microscopy and biochemical assays. In A53T CtrlVect mice, oligomers had Stokes radii of up to 65 Å and ranged in molecular weight from 36 to 80 kDa (Fig. 4c and Supplementary Fig. 6a,b), consistent with previous observations. However, in addition to these low-angstrom species, larger species of up to 122 Å were detected in A53T TH-RREE mice. On 12% SDS-PAGE, these species migrated as dimers, trimers and high-molecular-weight polymers stable to SDS and heat, consistent with the presence of oxidized and crosslinked species. The appearance of unique, high-angstrom species in A53T TH-RREE mice was observed with multiple α-synuclein antibodies that recognize N- or C-terminal epitopes (Fig. 4c and Supplementary Fig. 6a,b). Moreover, analysis of quinone-associated protein by near-infrared fluorescence revealed that only the high-angstrom oligomeric fractions from A53T TH-RREE mice were positive, supporting a direct interaction of dopamine and α-synuclein oligomers in these mice (Supplementary Fig. 6c).

Further structural analysis of oligomers isolated from high- (72–122) or low-angstrom (41–65) SEC fractions was undertaken by immunoelectron microscopy. Oligomers typically appeared as clusters with >2 gold particles in both high- and low-angstrom fractions from A53T TH-RREE mice, as well as low-angstrom fractions from A53T CtrlVect mice. The species were labeled with α-synuclein antibodies directed at either the N- or C terminus, indicating that both ends were exposed (Fig. 4e). Clusters of >2 gold particles were not observed in high-angstrom fractions from A53T CtrlVect mice or fractions from A53T TH-RREE mice that were immunodepleted of α-synuclein before imaging. Similar to mouse-derived oligomers, oligomers generated from incubating purified recombinant human wild-type α-synuclein with dopamine in vitro migrated as 36- to >98-kDa bands on SDS-PAGE and were recognized by both N- and C-terminally directed α-synuclein antibodies in western blots and by immunoelectron microscopy (Supplementary Fig. 7c,d). Electron microscopy imaging of the recombinant oligomers revealed clusters of >2 gold particles as observed in oligomer fractions from mouse SN (Supplementary Fig. 7d).

Given that there is growing evidence for prion-like spreading of α-synuclein pathology in disease, we investigated whether dopamine...
30 worms per genotype per replicate, 3 independent replicates, resistance to CAT-2 induced toxicity. Representative images from day 5 the site of interaction with dopamine (A53T(125-9m)) showed significant resistance to CAT-2 induced toxicity. Overexpression of dopamine neurons of C. elegans resulted in 62% of worms with wild-type dopamine neurons at day 5 post-hatching, and 50% at day 6 post-hatching. Overexpression of CAT-2 in A53T worms significantly exacerbated toxicity at both time points, whereas coexpression of CAT-2 with A53T α-synuclein mutated at the site of interaction with dopamine (A53T(125-9m)) showed significant resistance to CAT-2 induced toxicity. Representative images from day 5 post-hatching are shown in a, with intact neurons marked by arrowheads and degenerating or missing neurons by arrows. Scale bar represents 10 μm, ns, not significant. The data are presented as mean ± s.e.m. (n = 30 worms per genotype per replicate, 3 independent replicates, P(A53T + CAT-2 Day 5 versus A53T Day 5) = 0.0026, P(A53T + CAT-2 Day 5 versus A53T(125-9m) + CAT-2 Day 5) = 0.0013, F(3,8) = 15.19; P(A53T + CAT-2 Day 6 versus A53T Day 6) = 0.029, P(A53T + CAT-2 Day 6 versus A53T(125-9m) Day 6) = 0.0232, P(A53T + CAT-2 Day 6 versus A53T(125-9m) + CAT-2 Day 6) = 0.0365, F(3,8) = 6.488; one-way ANOVA on each time point with Tukey’s correction for multiple comparisons). **P < 0.01.

These findings indicate that dopamine promotes the generation of conformationally and functionally modified α-synuclein oligomeric species in the mouse brain.

To further establish the link between dopamine-induced oligomers and toxicity, we exposed primary neuronal cultures to recombinant α-synuclein oligomers generated in the presence of dopamine in vitro. Exogenous α-synuclein from oligomer preparations had become internalized in neurons and localized to neurites by 2 weeks post-treatment (Supplementary Fig. 9a). At this time point, we evaluated cells for viability using Calcein AM and propidium iodide (PI) dyes. Exposure of neurons to 1 μM dopamine-incubated α-synuclein induced a significant 44% reduction in Calcein-positive, PI-negative cells relative to phosphate-buffered saline (PBS)-treated controls (Supplementary Fig. 9b,c). Treatment with equivalent doses of monomeric α-synuclein, or dopamine that had been incubated under aggregation conditions without α-synuclein, did not reduce cell viability (Supplementary Fig. 9b,c). The observed neurotoxicity can therefore be attributed specifically to α-synuclein oligomers, consistent with our findings in vivo linking dopamine-modified species with severe nigrostriatal degeneration.

**Disrupting the interaction of dopamine and α-synuclein mitigates neurotoxicity in C. elegans**

To gain further mechanistic insight into the synergistic toxicity of dopamine and α-synuclein, we took advantage of the genetic tractability of the nematode model system Caenorhabditis elegans. Previous studies have shown that expressing human wild-type or PD-linked mutants of α-synuclein in dopaminergic neurons of C. elegans causes progressive age-dependent neurodegeneration. Overexpression of the TH homolog CAT-2 also induces dopamine neuron loss in worms. To test the combined effect of these factors on neuronal health, we first generated transgenic worms expressing human A53T α-synuclein under the control of the worm dopamine transporter (dat-1) promoter. C. elegans hermaphrodites have precisely eight pair of neuronal cells relative to phosphate-buffered saline (PBS)-treated controls (Supplementary Fig. 10a). Consistent with previous C. elegans models of A53T α-synuclein toxicity, we documented that 62% and 50% of A53T worms maintained normal dopamine neurons on days 5 and 6 post-hatching, respectively (Fig. 5). Notably, CAT-2 overexpression in A53T worms significantly exacerbated the degeneration of dopamine neurons, with only 33% and 26% of worms exhibiting normal neurons at days 5 and 6, respectively (Fig. 5). These data are consistent with our observations in the A53T TH-RREE mouse model and suggest that neuronal susceptibility to dopamine/α-synuclein toxicity is conserved.

We next investigated whether mutating α-synuclein at the site of interaction with dopamine would rescue neurons from degeneration. Mutation of the Y125EMP29 motif in the C terminus of α-synuclein to F125AFA29 abolishes the interaction with dopamine and inhibits dopamine-mediated stabilization of oligomers. Transgenic worms expressing A53T α-synuclein with the F125AFA29 mutation (A53T(125-9m)) at similar levels as A53T α-synuclein showed no difference in dopamine neuron degeneration compared with A53T worms (Fig. 5 and Supplementary Fig. 10). However, A53T(125-9m) worms displayed complete resistance to CAT-2-induced neurodegeneration, with the percentage of worms exhibiting normal dopamine neurons restored to 66% and 49% on days 5 and 6, respectively (Fig. 5). These findings suggest that the specific interaction of dopamine with the C terminus of α-synuclein is a critical contributor to the degeneration of dopamine neurons in vivo. The demonstration of this phenomenon in two in vivo platforms (worms and mice) strongly supports a role for dopamine-induced oligomers in the mechanism of neurotoxicity.
DISCUSSION

We used a new approach to enhance nigrostriatal dopamine levels in A53T mutant human α-synuclein transgenic mice. In this well-characterized mouse model, the SN does not develop α-synuclein inclusions and does not undergo neuronal cell loss. However, elevation of dopamine levels in the A53T mice induced substantial nigrostriatal degeneration and a previously undescribed locomotor impairment. Neurodegeneration was not observed in NonTg mice receiving the same treatment, indicating that dopamine-induced toxicity is dependent on α-synuclein. To the best of our knowledge, this is also the first demonstration that dopamine promotes α-synuclein oligomerization in vivo and that disrupting the ability of dopamine to stabilize/modify oligomers rescues neurons from dopamine toxicity. Taken together, these findings suggest that dopamine-induced α-synuclein oligomers may be a promising new target for PD treatment.

Dopamine has long been considered a contributor to the death of dopaminergic neurons in disease. At cytosolic pH, dopamine auto-oxidizes to form reactive quinone species, hydrogen peroxide and other electrophiles. The notion that dopamine may be neurotoxic has resulted in the widespread practice of 'DOPA sparing', which delays administration of the highly effective symptomatic therapy L-DOPA until more advanced stages of the disease. Although there is presently no conclusive evidence indicating that L-DOPA accelerates PD progression, clinical trials have provided mixed results and the issue of L-DOPA toxicity remains, unfortunately, unresolved. Despite the importance of investigating catecholamine toxicity, there are few studies in animal models. Single high-dose injections of dopamine into the striatum of rats have been shown to be acutely toxic in the short timeframe of 1 week. Other reports have shown that redistribution of endogenous dopamine to the cytosol is associated with toxicity in VMAT2 mutant mice. However, the long-term effects of increasing nigrostriatal dopamine levels remain unknown.

Our experiments represent the first investigation of dopamine toxicity by chronic enhancement of dopamine synthesis in mice. By expressing mutant TH that is insensitive to feedback inhibition by dopamine, we found that striatal dopamine in NonTg mice was increased by over 50%. To our surprise, this substantial perturbation of dopamine homeostasis was insufficient to induce degeneration. NonTg TH-RREE mice maintained a normal number of SN neurons, intact synaptic contacts with the striatum and robust motor coordination/balance. The excess dopamine was likely released into the synapse, leading to the observed hyperactivity, which is consistent with other studies. It is possible the excess dopamine was well tolerated as a result of compensatory upregulation of metabolic enzymes and transporters; indeed, there was a substantial increase in DAT levels in these mice.

α-synuclein has been independently linked to PD through dominantly inherited mutations and multiplications of the SNCA gene, and by the presence of aggregated α-synuclein in Lewy bodies and Lewy neurite inclusions. However, efforts to model PD by overexpression of α-synuclein in transgenic mice have often resulted in a lack of inclusion pathology and neurodegenerative changes in the SN. Dopaminergic neurons are entirely spared in the A53T transgenic mouse model that we used, and we documented a normal complement of neuronal cell bodies and terminals and a lack of intraneuronal α-synuclein inclusions in the SN in A53T CtrlVect mice. This may be related to several factors, including the fact that mouse α-synuclein naturally has a threonine at position 53. In vitro, mouse α-synuclein fibrillizes more rapidly than human wild-type or A53T α-synuclein, and yet aged mice do not spontaneously develop α-synuclein inclusions or a PD-like disorder. It is likely that mice have evolved protective mechanisms against α-synuclein aggregation, such as an enhanced ability to maintain α-synuclein in a physiological lipid- or protein-bound state, and that these mechanisms may counteract efforts to initiate disease by transgenic α-synuclein expression.

Our study offers another possible explanation for the challenges of producing SN neurodegeneration in α-synuclein transgenic mice: lack of dopamine dysregulation. In stark contrast with NonTg mice, elevation of dopamine levels in A53T mice resulted in progressive nigrostriatal degeneration with locomotor impairment, markedly improving this mouse as a model of PD. Similarly, coexpression of CAT-2 and A53T α-synuclein in C. elegans induced more severe dopamine neuron degeneration than A53T α-synuclein expression alone. These findings are consistent with a limited number of studies reporting toxicity associated with the convergence of dopamine and α-synuclein. In the case of VMAT2-deficient mice, loss of SN neurons was only observed in animals on a normal α-synuclein background, whereas identical reduction of VMAT2 in animals with Snca gene locus deletion did not result in cell loss. In C. elegans expressing wild-type human α-synuclein, increasing cytosolic dopamine by mutation of the worm homolog of VMAT2 accelerated neurodegeneration. Conversely, protection from neuronal loss has been reported for worms expressing wild-type human α-synuclein and a nonfunctional mutant of CAT-2 that prevents dopamine synthesis. In addition, in rodent midbrain cultures, elevating cytosolic dopamine concentration reduces dopaminergic cell survival, and this effect is dependent on α-synuclein.

The synergistic toxicity of dopamine and α-synuclein may be mediated by α-synuclein oligomers, which are thought to be neurotoxic species. Lentiviral vector delivery of artificial α-synuclein variants with enhanced oligomerization into the rat SN resulted in the selective loss of dopaminergic cells. Similarly, artificial mutants with impaired β-sheet structure and increased propensity to form oligomers induced degeneration of dopaminergic nerve terminals in worms and overt dopaminergic cell loss in flies. In addition, oligomer-containing extracts from A53T transgenic mice were toxic when applied to primary mouse cortical cultures. Although it is known that oxidized dopamine kinetically stabilizes soluble α-synuclein oligomers in cell-free systems and in SH-SY5Y cultures, the toxicity of these species is largely unexplored. Evidence from studies conducted in vitro suggests that dopamine-induced oligomers can block their own degradation and that of other substrates by chaperone-mediated autophagy, and can reduce neurotransmitter release by inhibition of SNARE complex formation. In human fetal dopaminergic cultures, reduced viability was linked to endogenous dopamine and the accumulation of soluble α-synuclein protein complexes.

Notably, the effects of dopamine on α-synuclein oligomerization in vivo have not been investigated. Thus, we sought to determine whether A53T TH-RREE mice had alterations in oligomer biochemical and/or functional properties. Soluble α-synuclein oligomers were extracted from the SN and enriched by SEC under non-denaturing conditions to preserve native conformations. Increasing dopamine resulted in greater total levels of α-synuclein oligomers detected by western blot. The oligomers included species with Stokes radii of up to 65 Å that were also detected in A53T CtrlVect mice, consistent with previous characterization of oligomers in aged A53T mice. Larger oligomers of up to 122 Å were uniquely observed in A53T TH-RREE mice, suggesting that dopamine is capable of modifying oligomer conformations. These species may reflect dopamine-induced remodeling of existing oligomers and/or formation of oligomers from α-synuclein monomers de novo. In addition, oligomers with larger Stokes radii may contain a greater number of monomer units than lower angstrom species, and/or occupy less compacted arrangements.
Multiple biochemical and imaging approaches were employed to characterize the oligomers extracted from the SN. In addition to well-established methods such as western blotting and SEC, immunoelectron microscopy was performed on SEC fractions to visualize mouse-derived α-synuclein oligomers. Currently, there is no accepted method of specifically imaging α-synuclein oligomers in brain tissue in situ. However, with our approach, we were able to identify and image oligomers of known Stokes radii. Oligomer species from both A53T TH-RREE and A53T CtrlVect mice were labeled with antibodies directed at the N- or C terminus of α-synuclein, suggesting that epitopes at both ends of the protein were exposed. These epitopes were also exposed in soluble α-synuclein oligomers generated with dopamine in vitro, consistent with previous imaging by immunoelectron microscopy. Furthermore, LB509 and Syn505 antibodies are known to react robustly with α-synuclein in pathological inclusions in human disease brain. The reactivity of both mouse-derived and recombinant oligomers with these antibodies suggests that they may share disease-associated conformations.

Unlike the α-synuclein species extracted from A53T CtrlVect mice, oligomers from A53T TH-RREE mice were unable to act as seeds for α-synuclein fibrillization in vitro. In addition, we did not detect dopamine-modified oligomers in cortical tissue from A53T TH-RREE mice. These data argue against both intracellular and intercellular prion-like propagation of these species. Although α-synuclein aggregates that exhibit prion-like behavior have been shown in many contexts to be cytotoxic, seeding ability is neither necessary nor sufficient for toxicity. In a comparison of different oligomer species generated in vitro, α-synuclein oligomers that were toxic to SH-SY5Y cultures were also incapable of seeding intracellular aggregation of α-synuclein. Conversely, seeding-competent oligomers were found to be non-toxic when applied to SH-SY5Y cells. Although the spread of α-synuclein pathology in PD may primarily be driven by self-replicating seeding and cell-to-cell transmission, toxicity may depend on the interaction of aggregates with cell-type-specific factors. In this regard, dopamine may act as an enhancer of oligomer toxicity. Moreover, dopamine-modified species may resist sequestration into non-toxic fibrils or inclusions, and may therefore be available to participate in pathological interactions that ultimately lead to neuron death.

Although the mouse data clearly demonstrate an association between the neurodegenerative/locomotor disease phenotype and the presence of modified and more abundant α-synuclein oligomers, the functional evidence from our transgenic C. elegans models further establishes this link. It is well-documented that the Y123EMPS129 motif in α-synuclein is required for the stabilization of α-synuclein oligomers by dopamine in vitro. Mutation or deletion of this motif restores the ability of α-synuclein to fibrillize in the presence of dopamine, and the increase in oligomeric species in SH-SY5Y cells with elevated dopamine is prevented by expression of α-synuclein that lacks this region. Worms expressing A53T α-synuclein specifically mutated in the Y123EMPS129 domain were entirely resistant to the neurotoxic effects of increasing dopamine levels. Thus, preventing the interaction of dopamine and α-synuclein and thereby inhibiting its effects on α-synuclein oligomers is neuroprotective. Although we were not able to biochemically characterize α-synuclein oligomer species in these worms as a result of expression in only eight cells in the entire animal, our findings nevertheless support a key role of dopamine in neurodegeneration, specifically through its influence on α-synuclein.

Collectively, our findings underscore the potential critical importance of the interaction of dopamine and α-synuclein in driving disease. In PD, dopaminergic terminals are thought to degenerate before cell bodies, suggesting that the disease may arise at the synapse.

Increasing dopamine levels in A53T mice recapitulated the substantial loss of terminals that precedes overt nigral cell death, offering a new model of disease progression in PD. These mice also underwent an eventual decline in dopamine levels and developed an associated motor deficit, mimicking the depletion of striatal dopamine and the resulting hypokinesia that occurs in PD. Our findings also demonstrate that dopamine modifies α-synuclein aggregation in vivo, resulting in oligomer conformations that are biochemically and structurally similar to neurotoxic oligomers induced by dopamine in vitro. Finally, we were able to rescue dopaminergic neurons from dopamine toxicity by specifically inhibiting its interaction with α-synuclein in C. elegans. Dopamine-modified α-synuclein species may mediate neurodegeneration by disrupting cellular membranes, as previously proposed for α-synuclein oligomers. Moreover, damage to synaptic vesicle membranes could lead to dopamine leakage and further induction of pathogenic α-synuclein species.

METHODS

Methods, including statements of data availability and any associated accession codes and references, are available in the online version of the paper.

Note: Any Supplementary Information and Source Data files are available in the online version of the paper.

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AUTHOR CONTRIBUTIONS

D.E.M., E.T., J.R.M., R.G.K. and H.I. conceived and designed the experiments. D.E.M., E.T., H.K., N.S.G., M.J.D., S.D., P.G., J.L.G. and V.X.T. performed the experiments and analyzed the data. D.E.M. wrote the paper, with important contributions from J.R.M., E.T., I.H.W., R.G.K., K.A.C., G.A.C. and H.I.

COMPETING FINANCIAL INTERESTS

The authors declare no competing financial interests.

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1. Ehringer, H. & Hornykiewicz, O. Verteilung von noradrenalin und dopamin (3-hydroxytyramin) im Gehirn des menschen und ihr verhalten bei erkrankungen des extrapyramidalen systems. (Distribution of noradrenaline and dopamine (3-hydroxytyramine) in the human brain and their behavior in diseases of the extrapyramidal system). Klin. Wochenschr. 38, 1236–1239 (1960).

2. Graham, D.G. Oxidative pathways for catecholamines in the genesis of neuromelanin and cytotoxic quinones. Mol. Pharmacol. 14, 633–643 (1978).

3. Jenner, P. & Olanow, C.W. Oxidative stress and the pathogenesis of Parkinson's disease. Neurology 47 Suppl 3: S161–S170 (1996).

4. Dexter, D.T. et al. Basal lipid peroxidation in substantia nigra is increased in Parkinson's disease. J. Neurochem. 52, 381–389 (1989).

5. Zhang, J. et al. Parkinson's disease is associated with oxidative damage to cytoplasmic DNA and RNA in substantia nigra neurons. Am. J. Pathol. 154, 1423–1429 (1999).

6. Giasson, B.I. et al. Oxidative damage linked to neurodegeneration by selective alpha-synuclein nitration in synucleinopathy lesions. Science 290, 989–989 (2000).

7. Zhang, J. et al. Parkinson's disease is associated with oxidative damage to cytoplasmic DNA and RNA in substantia nigra neurons. Am. J. Pathol. 154, 1423–1429 (1999).

8. Giasson, B.I. et al. Oxidative damage linked to neurodegeneration by selective alpha-synuclein nitration in synucleinopathy lesions. Science 290, 989–989 (2000).
7. Hastings, T.G., Lewis, D.A. & Zigmond, M.J. Role of oxidation in the neurotoxic effects of intrastriatal dopamine injections. Proc. Natl. Acad. Sci. USA 93, 1956–1961 (1996).

8. Colebrooke, R.E. et al. Age-related decline in striatal dopamine content and motor performance occurs in the absence of nigral cell loss in a genetic mouse model of Parkinson’s disease. Eur. J. Neurosci. 24, 2622–2630 (2006).

9. Caudle, W.M. et al. Reduced vesicular storage of dopamine causes progressive nigrostriatal neurodegeneration. J. Neurosci. 27, 8138–8148 (2007).

10. Polymeropoulos, M.H. et al. Mutation in the alpha-synuclein gene identified in families with Parkinson’s disease. Science 276, 2045–2047 (1997).

11. Krüger, R. et al. Ala30Pro mutation in the gene encoding alpha-synuclein in multiple system atrophy and Parkinson’s disease-type pathology. Neurology 68, 4087–4092 (2013).

12. Singleton, A.B. et al. alpha-Synuclein locus triplication causes Parkinson’s disease. Science 302, 841 (2003).

13. Chartier-Harlin, M.-C. et al. Alpha-synuclein locus duplication as a cause of familial Parkinson’s disease. Nat. Genet. 18, 106–108 (1998).

14. Iwai, A. Synucleins are a presynaptic protein of the central nervous system. J. Biol. Chem. 275, 13344–13349 (2000).

15. Krüger, R. et al. Alpha-synucleinopathies. Adv. Neurol. 80, 1062–1064 (2013).

16. Lesage, S. et al. French Parkinson’s Disease Genetics Study Group. G51D alpha-synuclein mutation causes a novel parkinsonian-pyramidal syndrome. Ann. Neurol. 73, 459–471 (2013).

17. Pasanen, P. et al. Novel alpha-synuclein mutation A53E associated with atypical multiple system atrophy and Parkinson’s disease-type pathology. Neurobiol. Aging 35, 2180.e1–2180.e5 (2014).

18. Spillantini, M.G. et al. Alpha-synuclein in Lewy bodies. Nature 388, 839–840 (1997).

19. Baba, M. et al. Aggregation of alpha-synuclein in Lewy bodies of sporadic Parkinson’s disease and dementia with Lewy bodies. Am. J. Pathol. 152, 879–884 (1998).

20. Iwai, A. et al. The precursor protein of non-A beta component of Alzheimer’s disease amyloid is a presynaptic protein of the central nervous system. Neuron 14, 467–475 (1995).

21. George, J.M., Jin, H., Woods, W.S. & Clayton, D.F. Characterization of a novel protein regulated during the critical period for song learning in the zebra finch. Neuron 15, 361–372 (1995).

22. Abeliovich, A. et al. Mice lacking alpha-synuclein display functional deficits in the nigrostriatal dopamine system. Neurobiol. Dis. 25, 239–252 (2000).

23. Murphy, D.D., Rueder, S.M., Trojanowski, J.Q. & Lee, V.M. Synucleins are developmentally expressed, and alpha-synuclein regulates the size of the presynaptic vesicular pool in primary hippocampal neurons. J. Neurosci. 20, 3214–3220 (2000).

24. Conway, K.A., Rochet, J.C., Bieganski, R.M. & Lansbury, P.T. Jr. Kinetic stabilization of the alpha-synuclein protofibril by a dopamine-alpha-synuclein adduct. Science 294, 1346–1349 (2001).

25. Norris, E.H. et al. Reversible inhibition of alpha-synuclein fibrillation by dopaminomelatonin-mediated conformational alterations. J. Biol. Chem. 280, 21212–21219 (2005).

26. Mazzulli, J.R. et al. Cytosolic catecholins inhibit alpha-synuclein aggregation and facilitate the formation of intracellular soluble oligomeric intermediates. J. Neurosci. 26, 10068–10078 (2006).

27. Mazzulli, J.R., Armstrong, M., Dumolin, M., Parastatidis, I. & Ischiropoulos, H. Cellular oligomerization of alpha-synuclein is determined by the interaction of oxidized catechol with a C-terminal sequence. J. Biol. Chem. 282, 31621–31630 (2007).

28. Herrera, F.E. et al. Inhibition of alpha-synuclein fibrillation by dopamine is mediated by interactions with furan and with furan and with furan. J. Neurosci. 31, 3682–3691 (2011).

29. Martinez-Vicente, M. et al. Dopamine-modified alpha-synuclein blocks chaperone-mediated autophagy. J. Clin. Investig. 118, 777–788 (2008).

30. Choi, B.K. et al. Large alpha-synuclein oligomers inhibit neuronal SNARE-mediated vesicle docking. Proc. Natl. Acad. Sci. USA 110, 4087–4092 (2013).

31. Nakashima, A. et al. The mutation of two amino acid residues in the N-terminus of tyrosine hydroxylase (TH) dramatically enhances the catalytic activity in neuroendocrine AIT-20 cells. J. Neurochem. 82, 202–206 (2002).

32. Giasson, B.I. et al. Neuronal alpha-synucleinopathy with severe movement disorder in mice expressing A53T human alpha-synuclein. Neuron 34, 521–533 (2002).

33. Teu, K. et al. D-beta-hydroxybutyrate rescues mitochondrial respiration and mitigates features of Parkinson disease. J. Clin. Investig. 112, 892–901 (2003).

34. Tsika, E. et al. Distinct region-specific alpha-synuclein oligomers in A30T transgenic mice: implications for neurodegeneration. J. Neurosci. 30, 3409–3418 (2010).

35. Souza, J.M., Giasson, B.I., Chen, Q., Lee, V.M. & Ischiropoulos, H. Dityrosine cross-linking promotes formation of stable alpha-synuclein polymers. Implication of nitrative and oxidative stress in the pathogenesis of neurodegenerative synucleinopathies. J. Biol. Chem. 275, 18344–18349 (2000).

36. Luk, K.C. et al. Pathological alpha-synuclein transmission initiates Parkinson-like neurodegeneration in nontransgenic mice. Science 338, 949–953 (2012).

37. Lakos, M. et al. Dopaminergic neuronal loss and motor deficits in Caenorhabditis elegans overexpressing human alpha-synuclein. J. Neurochem. 86, 165–172 (2003).

38. Cao, S., Gelwix, C.C., Caldwell, K.A. & Caldwell, G.A. Torsin-mediated protection for selective neurodegeneration in Parkinson disease. Proc. Natl. Acad. Sci. USA 102, 2622–2630 (2005).

39. Karpinar, D.P. et al. Pre-fibrillar alpha-synuclein variants with impaired beta-structure increase neurotoxicity in Parkinson’s disease models. EMBO J. 28, 3256–3268 (2009).

40. Olanow, C.W. Levodopa: effect on cell death and the natural history of Parkinson’s disease. Mov. Disord. 30, 37–44 (2015).

41. Loth, K.M. et al. Increased vesicular monoamine transporter enhances dopamine release and opposes Parkinson disease-related neurodegeneration in vivo. Proc. Natl. Acad. Sci. USA 111, 9977–9982 (2014).

42. Roche, J.C., Conway, K.A. & Lansbury, P.T. Jr. Inhibition of fibrillation and accumulation of prefibrillar oligomers in mixtures of human and mouse alpha-synuclein. Biochemistry 39, 10619–10626 (2000).

43. Specht, C.G. & Schoepfer, R. Deletion of the alpha-synuclein locus in a subgroup of C57BL/6J inbred mice. BMC Neurosci. 2, 11 (2001).

44. Mosharov, E.V. et al. Interplay between cytosolic dopamine, calcium and alpha-synuclein causes selective death of substantia nigra neurons. Neurology 62, 218–229 (2009).

45. Winner, B. et al. In vivo demonstration that alpha-synuclein oligomers are toxic. Proc. Natl. Acad. Sci. USA 108, 4194–4199 (2011).

46. Xu, J. et al. Dopamine-dependent neurotoxicity of alpha-synuclein: a mechanism for selective neurodegeneration in Parkinson disease. Nat. Med. 8, 600–606 (2002).

47. Danzer, K.M. et al. Different species of alpha-synuclein oligomers induce calcium influx and seeding. J. Neurosci. 27, 9220–9232 (2007).

48. Cheng, H.-C., Ulane, C.M. & Burke, R.E. Clinical progression in Parkinson disease and the neurobiology of axons. Ann. Neurol. 67, 715–725 (2010).
ONLINE METHODS

Animals. The mice (Mus musculus) used in this study were homozygous for expression of human A53T α-synuclein under the mouse PrP promoter (line M83). These mice undergo spinal cord degeneration at approximately 18 months of age in our colony, and the phenotype has been described previously32. Any animals showing symptoms of spinal cord degeneration, that is, hunched back, altered gait, or hindlimb paralysis, were excluded from the study. NonTg littermates were also used for experiments. No care was taken as to the sex of the animals. Mice were randomly assigned to the lentiviral vector injection groups and subsequent analyses, and the organization of experimental conditions was also random. Data collection and analyses were not performed blind to the conditions of the experiments except where otherwise indicated. All animal work was conducted according to National Institute of Health guide for the care and use of laboratory animals and in compliance with procedures approved by the Children’s Hospital of Philadelphia Institutional Animal Care and Use Committee.

Production of lentiviral vectors and determination of titer. Generation of human tyrosine hydroxylase-1 with mutation of R72P to R72E (TH-RREE) has been previously described26. The coding sequence was removed from the pcDNA3.1 vector and was subcloned into the self-inactivating pTY-linker lentiviral expression vector at the PmeI site downstream of the CMV promoter. Generation of replication-deficient pseudotyped HIV-derived lentiviral vectors was achieved by transient co-transfection of HEK293T cells with the expression vector along with the vectors carrying the additional transcripts required for encapsulation, packaging and envelope proteins. 24 h after the cells were seeded into 150-mm dishes coated with poly-D-lysine they were transiently transfected with a mixture containing 550 μg of the pTY-TH-RREE expression vector, 357.5 μg of CMVΔD82 packaging vector, and 192.5 μg of pVSV-G envelope vectors using the calcium phosphate transfection protocol. The culture medium was changed every day and the media containing the virus was collected 72 h post-transfection, filtered through 0.45 μm membranes, and concentrated by ultracentrifugation at 50,000 × g for 2 h at 4 °C. The viral pellet was subsequently re-suspended in 800 μl of DMEM medium to make a concentrated viral stock. An empty vector control virus containing all elements of V. M. Lee, University of Pennsylvania32. Incubation with biotin-conjugated secondary antibodies was followed by avidin-biotin-peroxidase complex, each

Immunohistochemistry and stereological cell counts. For histological analysis, the mice were deeply anesthetized and perfused transcardially with saline followed by 4% paraformaldehyde (PFA) in 0.1 M phosphate buffer (pH 7.4). The brains were post-fixed overnight at 4 °C in 4% PFA and then cryoprotected in 30% sucrose solution for 2 d at 4 °C. The brains were submerged in dry ice-chilled isopentane for 30 s and then stored at −80 °C until sectioning. On the day of sectioning, brains were mounted on chucks with OCT (Tissue-Tek) and the brainstem, SN and striatum were each sectioned using a cryostat (Leica, Jung Frigocut 2800N). Free-floating coronal sections of alternating 30- and 30-μm thickness were collected from each region and stored at 4 °C in 0.1 M phosphate buffer containing 0.1% sodium azide.

To assess motor coordination and balance, mice were tested on the Rotarod (Ugo Basile, model 7650). The mice were acclimated to the apparatus with four training sessions of 5 min each at 4 rpm, followed by 5 min of rest in the home cage. Testing was performed 1 h later with two trials in which speed was accelerated from 4 to 40 rpm in 300 s. For each mouse, the latency to fall off the rotator within this time period was recorded for the two trials and averaged.

Catechol quantification by HPLC with electrochemical detection. Striatal tissues were homogenized by sonication in ten volumes of 0.1 M perchloric acid containing 1 μM 3,4 dihydroxybenzylamine as an internal standard. The samples were centrifuged at 16,000 g for 10 min at 4 °C and the supernatants were filtered through 0.22-μm filters. 50 μl of each sample was injected onto an Agilent 1100 series HPLC controlled by ChemStation software version 1.04 (Agilent). Catechols were resolved on a reverse-phase C18 Luna column (150 × 4.6 mm, 5 μm; Phenomenex) and detected using a Coullarray detector (ESA Biosciences) as previously described26. Protein pellets were solubilized in 50 mM Tris pH 7.4 containing 2% SDS and protein concentration was determined using the BCA microassay kit (Pierce). Monoamine levels were normalized to protein concentration and expressed as femtomoles analyte per μg of protein.

Sequential extraction and native SEC. Striatal or SN tissue from individual mice was homogenized in ten volumes of lysis buffer: 1% Triton X-100 in 20 mM HEPES pH 7.4, 150 mM NaCl, 10% glycerol, 1 mM EGTA, and protease inhibitor cocktail (P2714, Sigma). The tissue was grinded with a mechanical homogenizer and centrifuged at 16,000 g for 10 min at 4 °C. The pellet was further extracted by sonication in 2% SDS, 50 mM Tris pH 7.4 with protease inhibitor cocktail, boiling at 95 °C for 10 min, and centrifugation at 16,000 g for 10 min. The resulting supernatant was designated the Triton-insoluble fraction. Protein concentration was determined using the BCA assay (Pierce). For SEC, 500 μg of Triton-soluble SN tissue in a total volume of 270 μl was loaded onto a Superdex 200 HR10/30 column (GE Healthcare) connected to an
Agilent 1100 series HPLC system controlled by ChemStation software version 1.04 (Agilent). Mobile phase consisted of 25 mM HEPES and 150 mM NaCl, pH 7.25 and the flow rate was set to 0.3 ml/min. Fractions corresponding to 122–94, 94–81, 81–72, 72–65, 65–59, 59–54, 54–50, 45–41, 41–38, 38–34, and 34–32 Å were each pooled and concentrated with 3,000 NWML Ultracel Microcon filters (Millipore). Fractions 122–72, 65–41, and 38–32 were further combined to enhance the α-synuclein signal on SDS-PAGE. The SEC column was calibrated using globular protein standards (GE Healthcare).

Western blotting. Proteins sequentially extracted or further fractionated by SEC were run on 10% or 12% SDS-PAGE gels and transferred to PVDF membranes. The membranes were blocked for 1 h at 20–25 °C in 5% (w/v) milk in 20 mM Tris pH 7.4, 150 mM NaCl and 0.1% Tween. Incubation with primary antibodies diluted in blocking buffer was overnight at 4 °C. Antibodies used against synaptic proteins were TH (rabbit, 1:4,000, Calbiochem 657012)32, DAT (rat, 1:1,000, Millipore MAB369)36, D1 receptor (rat, 1:1,000, Sigma D2944)36, DARPP-32 (rabbit, 1:1,000, Cell Signaling 23065)37. Antibodies directed against α-synuclein were LBS09 (mouse, 1:1,000)32,34, Syn505 (mouse, 1:1,000)34, and SNL-4 (rabbit, 1:1,000)37, all courtesy of V.M. Lee, University of Pennsylvania. Loading controls for Triton-soluble proteins were GAPDH (mouse, 1:10,000, Abcam ab8245)38, NSE (rabbit, 1:4,000, Abcam ab53023)39, and Actin (rabbit, 1:1,000, Sigma A2066)34. The loading control for Triton-insoluble proteins was Vimentin (mouse, 1:1,000, BD Pharmingen 550513)34. Membranes were incubated for 1 h at 20–25 °C with secondary antibodies conjugated to IRDye 680 or 800 (1:5,000, Rockland)34 and then scanned using an Odyssey Infrared Imaging System (Li-Cor). Quantification of protein levels was performed using ImageStudio software version 4.0.21 (Li-Cor) and was normalized to loading control levels. For α-synuclein oligomer bands, all individual immunoblots > 19 kD that were detected were quantified and the intensities summed, and the total was divided by the total NSE.

Near-infrared (nIR) scanning of oxidized catechols was performed as described previously59. Briefly, before transfer the SDS-PAGE gels were scanned in the 700-nm channel at intensity 10 on an Odyssey Infrared Imaging System (Li-Cor).

Immunoelectron microscopy of SEC fractions. SEC fractions from TN tissue corresponding to low (41–65) and high (72–122) angstrom Stokes radii or oligomeric SEC fractions (41–122 Å) for each mouse with 425 µg fresh recombinant α-synuclein (see below) before grid preparation or had primary antibody omitted.

In vitro aggregation and seeding assays. Recombinant human wild-type α-synuclein was expressed and purified as described previously34. Purified α-synuclein was incubated at 6 mg/ml (415 µM) with or without equimolar dopamine at 37 °C and shaking at 1,400 rpm for up to 6 d. At indicated time points, fibrillar content of the reaction mixture was assayed by the addition of Thioflavin T (Sigma) to a final concentration of 25 µM in 0.05 M KH2PO4 pH 7.5, 150 mM NaCl. The α-synuclein oligomers were labeled with the Syn505 antibody against the N terminus or LB509 antibody against the C terminus, followed by 10 nm gold conjugated secondary (Electron Microscopy Sciences) as previously described25. Grids were negatively stained with 1% uranyl acetate and imaged at the University of Pennsylvania Electron Microscopy Resource Laboratory. Control samples were immunodepleted of α-synuclein (see below) before grid preparation or had primary antibody omitted.

Seedling assays were performed by incubating 5 µg total protein from pooled oligomeric SEC fractions (41–122 Å) for each mouse with 425 µg fresh recombinant α-synuclein (300 µM final concentration) at 37 °C and shaking at 1,400 rpm. Aliquots at indicated time points were analyzed by Thioflavin T. To immunodeplete α-synuclein for control samples, pooled SEC fractions were incubated overnight at 4 °C with LB509 antibody at a 1:5 ratio of antibody to total µg protein. Protein G-conjugated beads (Sigma) were equilibrated in 25 mM HEPES, 150 mM NaCl, pH 7.4 and incubated with the immunocomplexes for 1 h at 4 °C. α-Synuclein oligomers were then pulled down by centrifugation at 2,000 g for 2 min, and the resulting supernatant was used for experiments.

C. elegans models. α-Synuclein plasmids for microinjection into worms were generated using Gateway technology (Invitrogen). Plasmids containing sequences for human A53T α-synuclein or A53T α-synuclein with Y125EMPS129 mutated to Y125AFA130 [A53T(125-9m)] were generated previously77. The coding regions were PCR-amplified with addition of flanking attB1 and attB2 sites, and the resulting fragments were used to generate Gateway entry vectors by BP reaction with pDONR221. LR reactions were then performed with each entry vector and the previously constructed pDEST-DAT-1 destination vector containing the dat-1 promoter38. All primer sequences are available upon request.

Nematodes were maintained using well-established methods69. Constructs were injected into worms to create transgenic animals as previously described61. Strains UA287 (baEx168 a,b,c [Pdat::A53T α-synuclein, Punc-54::tdTomato];[Y125 EMPS129::GFP]) and UA288 (baEx169 a,b,c [Pdat::A53T(125-9m) α-synuclein, Punc-54::tdTomato];[Y125 EMPS129::GFP]) were generated by injecting a solution of 50 ng/µl of either Pdat::A53T α-synuclein or Pdat::A53T(125-9m) α-synuclein into strain BY230 ([572 EMPS129::GFP]; with a phenotypic marker (Punc-54::tdTomato, 50 ng/µl) for body wall muscle expression). Three independent stable lines were created for each group (a, b, c). To analyze CAT-2 overexpression in these same backgrounds, we removed the Pdat::GFP transgene from the UA287 and UA288 transgenic backgrounds by outcrossing to the N2 background. These two strains, which still contained the Punc-54::tdTomato phenotypic marker, were both crossed into UA57 ([baEx135 a,b,c CAT-2, Pdat::GFP])

Strains UA296 (baEx168 a,b,c [Pdat::A53T α-synuclein, Punc-54::tdTomato];[Y125 EMPS129::GFP]) and UA297 ([baEx169 a,b,c [Pdat::A53T(125-9m) α-synuclein, Punc-54::tdTomato];[Y125 EMPS129::GFP]) and UA297 ([baEx169 a,b,c [Pdat::A53T(125-9m) α-synuclein, Punc-54::tdTomato];[baEx415 a,b,c CAT-2, Pdat::GFP]) for dopaminergic neurodegeneration analyses, transgenic hermaphrodite animals were scored as described previously62. Briefly, on the day of analysis, the six anterior dopaminergic neurons (four CEP (cephalic) and two ADE (anterior dorsal)) were examined in 30 randomly selected worms that also express the tdTomato marker in the body wall muscle cells. Each animal was considered normal when there was a full complement of six anterior DA neurons. However, if a worm exhibited any degenerative phenotype, such as a missing dendritic process, cell body loss, or a bleeding neuronal process, it was scored as degenerating. Three independent transgenic worm lines were analyzed per genetic background and an average of the total percentage of worms with normal neurons was reported in the study.

For label-free quantification of α-synuclein expression by mass spectrometry, protein from GFP and tdTomato positive day 1 adult nematodes was extracted in a buffer containing 50 mM Tris, pH 7.5 with 8 M urea and 2% SDS. 50 µg of soluble protein underwent buffer exchange to PBS and concentrated using 10 kDa cutoff spin filters (Amicon). The concentrated protein was combined with 6x loading buffer and run on a 10% NuPAGE gel with MES running buffer (Invitrogen). The gel was fixed and stained using Novex Colloidal Blue staining kit (Invitrogen). Purified recombinant α-synuclein was used as a marker for sample bands of interest. Sample bands of interest with apparent molecular weight of 14 kDa were excised from the gel and digested with trypsin overnight. Trypsin digests were then analyzed by LC-MS/MS on a hybrid LTQ Orbitrap Elite mass spectrometer (ThermoFisher) coupled with a nanoLC Ultra (Eksigent). The mass spectrometer was set for parallel reaction monitoring for human α-synuclein sequence with the A53T mutation.

Primary neuronal cultures. Hippocampal neurons were provided by the University of Pennsylvania Neuron Culture Service Center. After dissection from C57BL/6 mouse embryos at day 18–19, cells were mechanically dissociated, trypsinized, and seeded at 100,000 cells per well into 24-well plates freshly coated with 50 µg/ml poly-D-lysine (Sigma). After plating, neurons were cultured for 2 h in culture media containing 5% heat-inactivated FBS, 1% Glutamax, 2% B-27 supplement, 100 U/ml penicillin, 100 µg/ml streptomycin in Neurobasal media (all from Invitrogen). After 2 h, the media was changed to culture media without FBS to discourage survival of glial cells. Twice per week, half of the media was replaced with fresh culture media.
Following 1 week in culture, neurons were treated with dopamine-incubated α-synuclein (from day 4–6 of in vitro aggregation) at a final concentration of 1 µM. Control conditions were equivalent doses of PBS, monomeric α-synuclein, or dopamine that had been incubated in parallel under aggregation conditions but without α-synuclein. 2 weeks post-treatment, cell viability was assayed using calcein AM (Sigma) and propidium iodide (PI) (Sigma) incubated for 20 min at 20–25 °C at final concentrations of 3 µM in PBS. Fluorescence images were obtained using MetaMorph software version 7.7 (Molecular Devices) and an inverted Olympus IX70 microscope equipped with an IX-FLA fluorescence observation attachment (Olympus Optical). For quantification of viable cells (calcein positive and PI negative), at least ten random fields of view were blindly counted per treatment condition and averaged to obtain one replicate value. Independent culture experiments performed with separate plating/treatments were considered independent replicates and were reported as the n value.

To image human α-synuclein in treated neurons, cells were labeled in a two-stage protocol as previously described61. 2 weeks following treatment, cells were live-incubated in media containing Syn204 antibody (mouse IgG2a, 1:500) for 1 h at 4 °C to label extracellular α-synuclein. The cells were then fixed with 4% PFA for 20 min at 20–25 °C, and permeabilized with 0.1% Triton X-100 in 5% NGS, 3% BSA in PBS for 1 h at 20–25 °C. Cells were incubated with LB509 (mouse IgG1, 1:500) overnight at 4 °C to label both extracellular and intracellular α-synuclein. Secondary antibodies conjugated to Alexa Fluor (anti-mouse IgG1 488, 1:500, Thermo Fisher A-21135) and Alexa Fluor (anti-mouse IgG2a 594, 1:500, Thermo Fisher A21121; anti-mouse IgG2a 594, 1:500, Thermo Fisher A-21135) were incubated for 1 h at 20–25 °C. The staining was imaged by laser-scanning confocal microscopy (Olympus Fluoview) and the n value equals the number of independent culture experiments.

Statistics. All statistical analysis was done using Prism 6 software (GraphPad). Two-tailed unpaired Student’s t test was used for all comparisons between two groups. For comparisons with multiple groups, one-way ANOVA using Tukey’s or Bonferroni’s correction for multiple comparisons was used. No statistical methods were used to pre-determine sample sizes but our sample sizes are similar to those reported previously34,36. All experiments were performed at least twice, with measurements from the same samples treated as technical replicates. In the case of representative images, several images per sample were obtained. Readers are referred to the Life Sciences Reporting Summary available online for additional information.

Data Availability Statement. The data that support the findings of this study are available from the corresponding author upon reasonable request.
Life Sciences Reporting Summary

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### Experimental design

| Sample size | No statistical methods were used to pre-determine sample sizes but our sample sizes are similar to those reported in previous publications (ref. 34, 36, 62). |
| Data exclusions | The mice used in this study undergo spinal cord degeneration at approximately 18 months of age in our colony, and the phenotype has been described previously (ref. 32). Any animals showing symptoms of spinal cord degeneration, i.e. hunched back, altered gait, or hindlimb paralysis, were excluded from the study. |
| Replication | All attempts at replication were successful. |
| Randomization | Mice were randomly assigned to the lentiviral vector injection groups and subsequent analyses. Transgenic worms were randomly chosen for analysis. |
| Blinding | Data collection and analyses were not performed blind to the conditions of the experiments except for stereological quantification of neurons, and counting of cells in culture. |

Note: all studies involving animals and/or human research participants must disclose whether blinding and randomization were used.

### Statistical parameters

For all figures and tables that use statistical methods, confirm that the following items are present in relevant figure legends (or in the Methods section if additional space is needed).

- **Confirmed**
  - The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement (animals, litters, cultures, etc.)
  - A description of how samples were collected, noting whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
  - A statement indicating how many times each experiment was replicated
  - The statistical test(s) used and whether they are one- or two-sided (note: only common tests should be described solely by name; more complex techniques should be described in the Methods section)
  - A description of any assumptions or corrections, such as an adjustment for multiple comparisons
  - The test results (e.g. P values) given as exact values whenever possible and with confidence intervals noted
  - A clear description of statistics including central tendency (e.g. median, mean) and variation (e.g. standard deviation, interquartile range)
  - Clearly defined error bars

See the web collection on statistics for biologists for further resources and guidance.
7. Software

Describe the software used to analyze the data in this study.

All statistical analysis was done using Prism 6 (GraphPad). Open field testing used Multi Device Interface version 1.3 (Columbus Instruments). Images of tissue staining were accessed using Aperio Imagescope version 11.2 (Leica Biosystems) and quantification of staining was performed using ImageJ version 1.48 (National Institutes of Health). Stereological cell counts were done using StereoInvestigator version 11.02.1 (MBF Bioscience). HPLC analyses were done using ChemStation version 1.04 (Agilent), and catechol quantification additionally used ESA CoulArray version 3.10. Fluorescence images were obtained using MetaMorph version 7.7 (Molecular Devices). Western blot band intensities were quantified using Image Studio software version 4.0.21 (Li-Cor).

For manuscripts utilizing custom algorithms or software that are central to the paper but not yet described in the published literature, software must be made available to editors and reviewers upon request. We strongly encourage code deposition in a community repository (e.g. GitHub). Nature Methods guidance for providing algorithms and software for publication provides further information on this topic.

8. Materials and reagents

Materials availability

Indicate whether there are restrictions on availability of unique materials or if these materials are only available for distribution by a for-profit company.

There are no restrictions on availability of unique materials, such as TH-RREE and CtrlVect plasmids.

Antibodies

Describe the antibodies used and how they were validated for use in the system under study (i.e. assay and species).

The following antibodies were used: VMAT2 (Covance Custom Immunology Services, ref. 43), TH (Calbiochem, ref. 51, 52), SynSOS (ref. 32, 54), DAT (Millipore, ref. 36), D1R (Sigma, ref. 36), DARPP-32 (Cell Signaling, ref. 53), LB509 (ref. 32, 34), SNL-4 (ref. 55), GAPDH (Abcam, ref. 56), NSE (Abcam, ref. 57), Actin (Sigma, ref. 58), Vimentin (BD Pharmingen, ref. 34), Syn204 (ref. 63).

Eukaryotic cell lines

a. State the source of each eukaryotic cell line used.

b. Describe the method of cell line authentication used.

c. Report whether the cell lines were tested for mycoplasma contamination.

d. If any of the cell lines used are listed in the database of commonly misidentified cell lines maintained by ICLAC, provide a scientific rationale for their use.

SH-SYSY human neuroblastoma cells were obtained from American Type Culture Collection.

No cell line authentication was performed.

No test for mycoplasma contamination was performed.

No commonly misidentified cell lines were used.

Animals and human research participants

Policy information about studies involving animals; when reporting animal research, follow the ARRIVE guidelines

Provide details on animals and/or animal-derived materials used in the study.

The mice (Mus musculus) used in this study were homozygous for expression of human A53T α-synuclein under the mouse PrP promoter (line M83) (ref. 32), and non-transgenic littermates. No care was taken as to the sex of the animals. Mice were 10 months old at viral injection and either 12.5 or 15 months old at analysis. Four transgenic C. elegans strains were generated and hermaphrodites were analyzed at 3, 5, and 6 days post-hatching. Primary neuronal cultures were derived from C57BL/6 mouse embryos at day 18-19.

Policy information about studies involving human research participants

Provide details on human research participants.

The study did not involve human research participants.