Effects of innate immune receptor stimulation on extracellular α-synuclein uptake and degradation by brain resident cells

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Abstract

Synucleinopathies are age-related neurological disorders characterized by the progressive deposition of α-synuclein (α-syn) aggregates and include Parkinson’s disease (PD) and dementia with Lewy bodies (DLB). Although cell-to-cell α-syn transmission is thought to play a key role in the spread of α-syn pathology, the detailed mechanism is still unknown. Neuroinflammation is another key pathological feature of synucleinopathies. Previous studies have identified several immune receptors that mediate neuroinflammation in synucleinopathies, such as Toll-like receptor 2 (TLR2). However, the species of α-syn aggregates varies from study to study, and how different α-syn aggregate species interact with innate immune receptors has yet to be addressed. Therefore, we investigated whether innate immune receptors can facilitate the uptake of different species of α-syn aggregates. Here, we examined whether stimulation of TLRs could modulate the cellular uptake and degradation of α-syn fibrils despite a lack of direct interaction. We observed that stimulation of TLR2 in vitro accelerated α-syn fibril uptake in neurons and glia while delaying the degradation of α-syn in neurons and astrocytes. Internalized α-syn was rapidly degraded in microglia regardless of whether TLR2 was stimulated. However, cellular α-syn uptake and degradation kinetics were not altered by TLR4 stimulation. In addition, upregulation of TLR2 expression in a synucleinopathy mouse model increased the density of Lewy-body-like inclusions and induced morphological changes in microglia. Together, these results suggest that cell type-specific modulation of TLR2 may be a multifaceted and promising therapeutic strategy for synucleinopathies; inhibition of neuronal and astroglial TLR2 decreases pathogenic α-syn transmission, but activation of microglial TLR2 enhances microglial extracellular α-syn clearance.

Introduction

Synucleinopathies such as Parkinson’s disease (PD) and dementia with Lewy bodies (DLB) are the second most common age-related neurodegenerative disorders and are characterized by abnormal deposits of α-synuclein (α-syn) called Lewy bodies (LBs) and Lewy neurites (LN)s. α-Syn pathology spreads through various brain regions during the progression of the disease. Although the underlying mechanisms of this spread are not fully understood, cell-to-cell transmission of α-syn has been strongly implicated. Under physiological conditions, α-syn is a neuronal cytosolic protein involved in processes related to synaptic homeostasis, such as vesicle recycling. In the diseased brain, however, a small amount of monomeric and oligomeric α-syn can be released from neurons into the extracellular space and can be taken up by neighboring neurons and glial cells, thereby inducing neurotoxicity. If the neuron-released α-syn interacts with glial cells, the resulting cellular stress can promote inflammatory responses. Thus, chronic inflammation of the brain is a...
key feature of synucleinopathies and is thought to exacerbate neuronal death.

Several immune receptors have been shown to interact with extracellular α-syn to activate processes strongly associated with neuroinflammation. We previously demonstrated that the pattern recognition receptor Toll-like receptor 2 (TLR2) can directly interact with β-sheet-enriched oligomeric forms of neuron-released α-syn to induce proinflammatory microglial activation. Another innate immune receptor, TLR4, is also known to play a role in α-syn clearance, and studies have shown that genetic depletion of TLR4 ameliorates neurodegeneration, neuroinflammation, and behavioral deficits in toxin-based mouse models of PD. Additional studies employing recombinant α-syn aggregates of various forms have identified several other α-syn receptors, such as lymphocyte-activation gene 3 (LAG3), cluster of differentiation 36 (CD36), macrophage-1 antigen (MAC-1), nicotinamide adenine dinucleotide phosphate (NADPH) oxidase, β1-integrin, and heparan sulfate. α-Syn release by neurons can take on various conformations, including monomeric, oligomeric, and fibrillar forms. We have previously shown that microglial uptake of oligomeric α-syn is dependent on TLR2 signaling. Monomeric and low concentrations of oligomeric and fibrillar α-syn, however, are internalized by TLR2-deficient microglia. These observations suggest that cellular α-syn internalization is reliant on multiple receptors, each specific to a different conformation of α-syn. Moreover, most studies that identified novel α-syn receptors used a heterogeneous mix of various α-syn aggregate species; thus, cross-interactions among aggregate species with identified receptors have not been addressed. For example, while TLR2 has been shown to directly interact with oligomeric neuron-released α-syn but not α-syn fibrils, its activity may influence fibril uptake and clearance. Therefore, we investigated whether TLR2 activation can modulate the cellular internalization of recombinant α-syn fibrils despite a lack of direct interaction.

Materials and methods

Antibodies and chemicals

Protease inhibitor cocktail, phosphatase inhibitors, and lipopolysaccharide (LPS) were purchased from Sigma-Aldrich (St. Louis, MO). Pam3CSK4 was purchased from InvivoGen (San Diego, CA). The following antibodies were used for western blot analysis and immunostaining: α-syn (Syn-1; BD Bioscience, San Diego, CA), Iba-1 (Wako, Richmond, VA), α-syn (Syn 211), p62SQSTM1, and β-actin (Sigma-Aldrich).

Cell culture

Human SH-SY5Y neuroblastoma cells, human primary astrocytes, mouse primary astrocytes, and mouse BV2 microglia were cultured as previously described. Differentiated SH-SY5Y (dSY5Y) cells were prepared as described elsewhere. Preparation of recombinant α-syn fibrils and neuron-released α-syn

α-Syn fibrils were prepared as described elsewhere. Briefly, endotoxin-free recombinant α-syn was dissolved in phosphate-buffered saline (PBS) and filtered with a syringe filter (0.2-µm pore size). The concentration of monomeric α-syn was determined by the bicinchoninic acid protein assay. A total of 1 ml of dissolved α-syn (200 µM) was incubated at 37 °C with agitation (280 rpm). After a 14-day incubation, the samples were centrifuged at 10,000×g for 10 min. The supernatants were transferred to new tubes to determine the amount of soluble α-syn. The pellets were washed with PBS, and the amount of insoluble α-syn fibrils was calculated based on the amount of remaining soluble α-syn in the supernatants. The pellets were resuspended in PBS (final concentration: 2 mg/ml) and briefly sonicated before use.

Neuron-released α-syn (α-syn-conditioned medium, αSCM) was prepared as described elsewhere. In brief, dSY5Y cells were infected with human wild-type α-syn expressing adenoviral vectors (serotype Ad5) at a multiplicity of infection of 100. Two days post-infection, the cells were cultured with serum-free Dulbecco’s modified Eagle’s medium (DMEM) for an additional 18 h. Cell debris was removed from the culture medium, and culture medium was concentrated using 10 K cutoff centrifugal filters. Concentrated neuron-released α-syn contained 1.06 ± 0.371 mg/ml α-syn. In the current study, dSY5Y cells were treated with ~1 µg/ml α-syn.

Cellular α-syn uptake and degradation assays

The α-syn fibril uptake and degradation assays were performed as described elsewhere. For the uptake assay, dSY5Y cells, human primary astrocytes, mouse primary astrocytes, and BV2 microglia were preincubated in the presence or absence of Pam3CSK4 (10 µg/ml) or LPS (0.1 µg/ml). After 1 h of incubation, fibrils (200 nM) were added to the cells and incubated for the indicated duration (Fig. 1a and Supplementary Fig. 1a). Whole-cell lysates of fibril-treated cells were harvested at the indicated time points.

For the degradation assay, cells were pretreated with fibrils (200 nM) for the indicated duration, washed with PBS four times, and incubated with fresh media in the presence or absence of Pam3CSK4 (10 µg/ml) or LPS (0.1 µg/ml) (Fig. 2a and Supplementary Fig. 1c). At the indicated time points, whole-cell lysates were harvested. For the neuron-released α-syn uptake assay, dSY5Y cells were preincubated in the presence or absence of Pam3CSK4 (10 µg/ml). After a 1-h incubation, αSCM was added to the cells. Following incubation for the indicated...
Fig. 1 Stimulation of TLR2 and kinetics of α-synuclein fibril uptake by neurons, astrocytes, and microglia. 

**a** Experimental scheme of the α-syn uptake assay. 

**b-d** hSY5Y cells (b), human primary astrocytes (c), and BV2 microglia (d) were treated with α-syn fibrils (200 nM) in the presence or absence of Pam3CSK4 (10 μg/ml) for the indicated duration. Whole-cell lysates were probed for α-synuclein, p62, and β-actin. The levels of α-synuclein and p62 were determined by densitometric quantification. The data are the mean ± SEM. *p < 0.05 and **p < 0.001.
Fig. 2 Stimulation of TLR2 and kinetics of α-synuclein fibril degradation by neurons, astrocytes, and microglia. 

a) Experimental scheme of the α-syn degradation assay. 

b-d) dSY5Y cells (b), human primary astrocytes (c), and BV2 microglia (d) were pretreated with α-syn fibrils (200 nM) for the indicated duration. After washing with PBS, the cells were incubated in the presence or absence of Pam3CSK4 (10 μg/ml) and harvested at the indicated time points. Whole-cell lysates were probed for α-synuclein, p62, and β-actin. The levels of α-synuclein and p62 were determined by densitometric quantification. The data are the mean ± SEM. *p < 0.05 and **p < 0.01.
duration, whole-cell lysates were harvested (Supplementary Fig. 2a).

Western blot analysis
Western blot analysis was performed as described elsewhere\(^{21}\). Briefly, cells were rinsed with PBS and then lysed with NuPAGE LDS sample buffer (Thermo Fisher Scientific) in the presence of protease and phosphatase inhibitors. Whole-cell lysates were briefly sonicated, separated by electrophoresis and transferred to PVDF membranes using a semidyry Trans-Blot Turbo Transfer System (Bio-Rad, Hercules, CA). The membranes were blocked with Odyssey blocking buffer (LI-COR Biosciences, Lincoln, NE) and probed with primary antibody followed by fluorescence-tagged secondary antibody. Fluorescence signal detection and densitometric analysis were performed using ODYSSEY CLx (LI-COR Biosciences) and Image Studio (LI-COR Biosciences).

Delivery of lentiviral vectors to the mouse brain
To investigate the effects of innate immune receptor stimulation on \(\alpha\)-syn neuropathology and microglial activation, we delivered LV-control or LV-TLR2 to non-tg and \(\alpha\)-syn-tg mice expressing human wild-type \(\alpha\)-syn under the PDGF-\(\beta\) promoter (D line) (Fig. 3a). This transgenic model is known to replicate the neuropathological features of PD, such as mild \(\alpha\)-syn pathology and microgliosis\(^{24}\). The lentiviral vectors (LV-control and LV-TLR2) were constructed and prepared as previously described\(^{21,25}\).

Briefly, either LV-control or LV-TLR2 (2 \(\mu\)l, 2.2 \(\times\) 10\(^7\) infection units) was stereotaxically injected into the bilateral hippocampi of the mice (anterior–posterior [AP], −2.0 mm; medial–lateral [ML], 1.5 mm; and dorsal–ventral [DV], −1.3 mm). At 5 weeks post-injection, the brains were fixed in 4% paraformaldehyde (PFA) at 4 °C for neuropathological analysis. All animal procedures were approved by the Institutional Animal Care and Use Committee of University of California, San Diego under protocol S02221.

Immunohistochemistry, double immunolabeling, and neuropathological analysis
Immunohistochemical, double immunolabeling, and neuropathological analysis were performed as described elsewhere\(^{26}\). Briefly, sagittal brain (blind-coded) sections were incubated with primary antibodies at 4 °C overnight. The next day, the sections were incubated with...
biotinylated-, FITC-conjugated, or Texas red-conjugated secondary antibodies. The biotinylated secondary antibodies were detected with an avidin D-HRP detection system (ABC elite, Vector Laboratories, Burlingame, CA). The immunostained sections were imaged with an Olympus BX41 microscope. α-Syn immunoreactivity was determined by measuring the optical density using Image Quant 1.43 software (NIH). The number of ameboid and fusiform Iba-1-positive cells was determined per field (230 μm × 184 μm) for each animal based on cell body recognition via Image Quant 1.43 software (NIH).

**Statistical analysis**

InStat (GraphPad Software, San Diego, CA) was used for all statistical analyses. All data were analyzed for statistical significance by unpaired *t* test or one-way ANOVA. All data are presented as the means ± SEMs.

**Results**

**Stimulation of innate immune receptors and cellular α-syn fibril uptake**

To address the role of innate immune receptors in the kinetics of α-syn fibril uptake, we performed an uptake assay with brain resident cells, such as neurons, astrocytes, and microglia (Fig. 1a–d and Supplementary Fig. 1a–d). dSY5Y neuronal cells, human and mouse primary astrocytes, and BV2 microglia were stimulated with either a TLR2 agonist (pam3CSK4, 10 μg/ml) or a TLR4 agonist (LPS, 0.1 μg/ml) for 1 h and then incubated with α-syn fibrils (200 nM) (Fig. 1a and Supplementary Fig. 1a). The resulting whole-cell lysates were harvested at various time points and analyzed by western blot analysis (Fig. 1b–d and Supplementary Fig. 1b–d).

In nonstimulated control cells of all three types, α-syn fibril uptake increased with prolonged exposure (Fig. 1b–d and Supplementary Fig. 1b–d). Interestingly, stimulation of TLR2 for 8, 24, and 2 h resulted in significantly higher levels of α-syn fibril uptake in neurons, astrocytes, and microglial cells (Fig. 1b–d); however the levels of α-syn fibril uptake were not altered by TLR4 stimulation (Supplementary Fig. 1b–d). We next treated dSY5Y cells with neuron-released α-syn obtained from the culture media of neuronal cells overexpressing human α-syn (αSCM) in the presence or absence of pam3CSK4 (Supplementary Fig. 2a). The uptake of high-molecular-weight neuron-released α-syn was slightly accelerated by TLR2 stimulation (Supplementary Fig. 2b).

Expression of the autophagy indicator p62/SQSTM1 (p62) by nonstimulated cells increased as the duration of exposure to α-syn fibrils increased (Fig. 1b–d). Surprisingly, while stimulation of TLR2 further enhanced p62 expression in dSY5Y cells and astrocytes (Fig. 1b, c), it did not affect p62 expression in microglia (Fig. 1d). Taken together, these results suggest that stimulation of TLR2, but not TLR4, enhances α-syn fibril uptake by brain-resident cells.

**Innate immune receptor stimulation and cellular α-syn degradation**

Once internalized by the cell, α-syn aggregates can be cleared by protein degradation processes, such as autophagy. To investigate whether degradation of internalized α-syn is also modulated by innate immune receptor stimulation, dSY5Y cells, human and mouse primary astrocytes and BV2 microglia were pretreated with α-syn fibrils for 24 or 18 h and then incubated in fresh media with or without TLR agonists, such as pam3CSK4 (10 μg/ml) and LPS (0.1 μg/ml) (Fig. 2a and Supplementary Fig. 1e). Whole-cell lysates were collected at various time points and analyzed by western blot analysis (Fig. 2b–d and Supplementary Fig. 1f, g).

In dSY5Y cells and astrocytes, internalized α-syn fibrils were gradually degraded in a time-dependent manner (Fig. 2b, c and Supplementary Fig. 1f, g). Surprisingly, this degradation was significantly delayed by TLR2 stimulation (Fig. 2b, c) but was not affected by TLR4 stimulation (Supplementary Fig. 1f, g). In microglia, which are known to be the most efficient brain-resident cell at removing extracellular α-syn, most of the internalized α-syn was degraded within 6 h regardless of whether TLRs were stimulated (Fig. 2d and Supplementary Fig. 1h).

Whereas p62 levels increased in dSY5Y cells and astrocytes as the duration of exposure to α-syn fibrils increased (Fig. 1b–d), in the absence of α-syn fibrils, p62 expression progressively declined (Fig. 2b, c). However, the reduction in p62 expression was inhibited and delayed by TLR2 stimulation in neurons and astrocytes, respectively (Fig. 2b, c). Interestingly, p62 expression in microglia was immediately suppressed following the removal of α-syn fibrils and then completely restored; TLR2 stimulation had no effect on p62 expression (Fig. 2d). In summary, degradation of internalized α-syn fibrils in neurons and astrocytes was delayed by TLR2 stimulation but not by TLR4 stimulation. However, microglia efficiently degraded internalized α-syn fibrils regardless of whether TLRs were stimulated.

**Genetic stimulation of innate immune receptors exacerbated α-syn neuropathology and inflammation in the α-syn transgenic mouse model**

To verify our in vitro findings in an animal model, we stimulated TLR2 in α-syn transgenic mice overexpressing human wild-type α-syn (α-syn-tg mice) (Fig. 3a). To stimulate TLR2 in the mice, we delivered a TLR2-overexpressing lentiviral vector (LV-TLR2) to the hippocampi of nontransgenic (non-tg) and α-syn-tg mice (Fig. 3a). A control lentiviral vector (LV-control) was
injected into the mice as a control (Fig. 3a). A viral vector approach was used in favor of pharmacological agents to prevent unwanted peripheral activation of TLR2. This α-syn-tg mouse model is known to exhibit the hallmark pathologies of synucleinopathies, such as LB-like intracellular α-syn accumulation and neuroinflammation. Studies have also shown that compared to delivery of LV-control, delivery of LV-TLR2 significantly increases TLR2 expression in neurons and glia in these mice. TLR2 overexpression subsequently accelerates neurodegeneration, astrogliosis, and microgliosis in α-syn-tg mice. Consistent with previous findings, we observed widespread α-syn pathology, most notably LB-like α-syn inclusions in the neocortex and hippocampus, in α-syn-tg but not non-tg mice (Fig. 3b). Interestingly, compared to injection of LV-control, overexpression of TLR2 by LV-TLR2 injection significantly increased the density of LB-like α-syn inclusions in α-syn-tg mice (Fig. 3b–d).

Microgliosis is an indicator of neuroinflammation and can be assessed by determining the number of activated microglia in the pathogen-affected brain region via immunohistochemistry. We observed a mild increase in the number of total microglia in the hippocampus and neocortex in LV-control-injected α-syn-tg mice compared to non-tg mice (Fig. 4a–c). In addition, TLR2 overexpression significantly increased the number of microglia in the hippocampi of both non-tg and α-syn-tg mice injected with LV-TLR2 compared to those administered LV-control (Fig. 4a–c). Most remarkably, however, the proportion of microglia displaying an ameboid and reactive morphology was significantly increased and the proportion of fusiform, resting microglia was decreased in the...
neocortices and hippocampi of α-syn-tg mice injected with LV-TLR2 (Fig. 4a–c). Double immunolabeling corroborated these morphological changes (Fig. 4d, e). In LV control-injected non-tg mice, most of the microglia displayed a fusiform phenotype with few branches (Fig. 4d, e). Interestingly, overexpression of TLR2 in the mice was associated with slightly more swollen microglia with thicker branches, indicative of an ameboid phenotype (Fig. 4d, e). The number of ameboid microglia was increased in α-syn-tg mice compared to non-tg mice and significantly increased by TLR2 overexpression (Fig. 4d, e). Based on α-syn immunoreactivity, LB-like inclusions were detected only in α-syn-tg mice, and the density of these inclusions was significantly increased in mice injected with LV-TLR2 (Fig. 4d, e). Furthermore, TLR2 overexpression increased the colocalization of α-syn and microglia in α-syn-tg mice (Fig. 4d, e). Taken together, these results suggest that genetic stimulation of TLR2 exacerbates α-syn neuropathology and induces morphological changes in microglia in α-syn-tg mice.

**Discussion**

In the current study, we demonstrated that stimulating the innate immune receptor TLR2 on neurons, astrocytes, and microglia increased the uptake of α-syn fibrils. Interestingly, innate immune receptor stimulation delayed the degradation of internalized α-syn and promoted p62 expression in neurons and astrocytes but had no effect on the rapid degradation of α-syn in microglia. Furthermore, upregulation of TLR2 expression in a mouse model of synucleinopathy aggravated α-syn neuropathology and promoted ameboid morphology of microglia (Fig. 5).

TLR2 is expressed in neurons and glial cells and is elevated in the brains of synucleinopathy model mice and patients. We previously demonstrated that TLR2 interacts with β-sheet-enriched oligomeric forms of neuron-released α-syn, thereby inducing activation of downstream signaling cascades. In microglia and astrocytes, activation of TLR2 induces α-syn internalization and neurotoxic inflammatory responses, such as cytokine/chemokine gene expression, nitric oxide induction, intracellular reactive oxygen species production, and morphological changes, via the nuclear factor kappa-light-chain-enhancer of activated B cells (NF-κB) and leucine-rich repeat kinase 2/nuclear factor of activated T-cells and cytoplasmic 2 (LRRK2/NFATc2) signaling pathways.

In neurons, TLR2 activation not only induces neuronal α-syn internalization but also increases neurotoxic α-syn accumulation through autophagy inhibition via mammalian target of rapamycin (mTOR). Therefore, genetic...
and pharmacological inhibition of TLR2 ameliorates neuroinflammation, neurodegeneration, α-syn abnormal deposition, and memory impairment in a synucleinopathy mouse model. In this study, we selectively enhanced TLR2 expression rather than depleting gene expression in a mouse model of synucleinopathy, and TLR2 gene induction increased the number of ameboid microglia and the density of LB-like α-syn inclusions in α-syn-tg mice. Studies have identified several other receptors for extracellular α-syn in addition to TLR2, such as TLR4, LAG3, CD36, MAC-1, NADPH oxidase, β1-integrin, and heparan sulfate. While each study employed different types of α-syn aggregates, extracellular α-syn may exist as a heterogeneous collection of proteins. For example, we previously demonstrated that TLR2 interacts with oligomeric neuron-released α-syn but not monomeric, fibrillar, 4-hydroxy-2-nonenal (HNE)-mediated, nitrilation-mediated α-syn aggregates. These findings support the existence of multiple mechanisms of extracellular α-syn cellular internalization based on the molecular structure of each aggregate type. In this study, we examined the effect of two innate immune receptors, TLR2 and TLR4, on cellular α-syn fibril uptake and degradation. While TLR2 is known to interact with neuron-released oligomeric forms of α-syn, TLR4 has been reported to interact with recombinant α-syn fibrils. Stimulation of TLR2 significantly increased and delayed α-syn fibril uptake and degradation, respectively; however, TLR4 stimulation did not affect the kinetics of α-syn fibril uptake and degradation in any type of cell. Given these findings, we speculate that both LPS and α-syn fibrils target the same receptor, TLR4; therefore, stimulation of TLR4 did not induce a further increase in α-syn fibril uptake or further delay in α-syn degradation. Therefore, understanding the cross-interactions between innate immune receptors and α-syn species may be helpful for clarifying the disease mechanism.

α-Syn propagation by cell-to-cell transmission is strongly associated with disease progression in synucleinopathies. Although a comprehensive mechanism remains elusive, we have previously shown the importance of receptors with α-syn-binding ability and α-syn aggregation driven by mass action, including genetic defects, impairment of protein homeostasis, and oxidative stress. Recent studies have also demonstrated the existence of α-syn oligomers and seeds in the extracellular space, such as the cerebral fluid, in PD/DLB patients. In addition, cryo-EM and injection studies with preformed fibrillar and fibrillar α-syn species isolated from the human brain have demonstrated the ability of α-syn seeding in the pathogenesis of synucleinopathies. Remarkably, a recent study suggested that disease type and neuropathology might be determined by the strain of fibrillar α-syn species.

Although neurons, astrocytes, and microglia all showed increased α-syn fibril uptake following innate immune receptor stimulation, the underlying mechanisms of degradation in these cells appear to be different. p62 expression was increased and sustained in stimulated neurons and astrocytes compared to nonstimulated controls. In addition, we recently demonstrated that TLR2 activation impaired autophagic flux, resulting in the accumulation of p62 followed by an increase in abnormal neurotoxic α-syn deposition. Therefore, we hypothesized that TLR2 stimulation may impair autophagy and subsequently inhibit the degradation of internalized α-syn fibrils. In microglia, however, p62 levels and the rapid degradation of internalized α-syn were unaffected by innate immune receptor activation. It is possible that the regulatory machinery for autophagic activity in microglia is independent of the innate immune receptor in question. Further study is needed to investigate whether TLR2 stimulation enhances microglial uptake kinetics via other immune receptors given that autophagic flux is not affected.

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Conflict of interest
S.-J.L. is a founder and CEO of Neuamedy Co., Ltd.

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