Parkinson’s disease associated mutation E46K of α-synuclein triggers the formation of a distinct fibril structure

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Amyloid aggregation of α-synuclein (α-syn) is closely associated with Parkinson’s disease (PD) and other synucleinopathies. Several single amino-acid mutations (e.g. E46K) of α-syn have been identified causative to the early onset of familial PD. Here, we report the cryo-EM structure of an α-syn fibril formed by N-terminally acetylated E46K mutant α-syn (Ac-E46K). The fibril structure represents a distinct fold of α-syn, which demonstrates that the E46K mutation breaks the electrostatic interactions in the wild type (WT) α-syn fibril and thus triggers the rearrangement of the overall structure. Furthermore, we show that the Ac-E46K fibril is less resistant to harsh conditions and protease cleavage, and more prone to be fragmented with an enhanced seeding capability than that of the WT fibril. Our work provides a structural view to the severe pathology of the PD familial mutation E46K of α-syn and highlights the importance of electrostatic interactions in defining the fibril polymorphs.

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Deposition of α-syn amyloid fibrils in Lewy bodies (LB) and Lewy neuritis (LN) is a common histological hallmark of Parkinson’s disease (PD) and synucleinopathies such as dementia with Lewy bodies (DLB) and multiple system atrophy (MSA)1–3. Accumulating evidence shows that α-syn amyloid fibrils serve as prion-like seeds for the propagation and cell-to-cell transmission of the pathological entity of α-syn4,5. The spread of pathological α-syn fibril is closely associated with the disease progression6,7. Several single point mutations and genomic duplication or triplication of SNCA, the gene encoding α-syn, have been identified to be causative to the familial forms of the diseases with a broad spectrum of distinctive clinical symptoms8–12. Moreover, the hereditary mutations exhibit exacerbated pathology in various cellular and PD-like animal models13,14. Intriguingly, the cryo-EM structure of full-length wild-type (WT) α-syn fibril demonstrates that in the five mutation sites found in familial PD, four of them, i.e. E46K, A53T/E, G51D, and H50Q, are located at the protofilament interface of the WT fibril15,16, which indicates that these mutations may alter the fibril structure and consequently influence α-syn amyloid aggregation and PD pathology.

α-Syn hereditary mutation E46K was originally identified from a Spanish family with autosomal dominant parkinsonism11. The clinical phenotype of E46K patient features rapid and severe disease progression with early onset of the parkinsonism of DLB11. Previous studies have shown that the E46K mutation can enhance α-syn fibril formation and α-syn pathology in vitro and in cultured cells13,17,18. Solid-state NMR study has shown that the E46K mutation causes large conformational changes of the fibril structure19.

In this work, we determine the cryo-EM structure of the N-terminally acetylated E46K α-syn (Ac-E46K) fibril at an overall resolution of 3.37 Å. The structure reveals a distinct fold of α-syn, in which the electrostatic interactions in the WT fibril are broken and reconfigured due to the E46K mutation. We further show that the Ac-E46K fibril is less stable than the WT fibril under harsh conditions and protease cleavage, while the mutant fibril is more efficient in seeding amyloid fibril formation. This work provides a structural mechanism for the influence of E46K on the amyloid fibril formation of α-syn, and suggests that electrostatic interactions may serve as one of the driving force for the polymeric fibril formation of α-syn.

Results

E46K mutation alters the morphology and stability of α-syn fibril. To investigate the influence of E46K on α-syn fibril formation, we prepared recombinant full-length α-syn with E46K mutation. Since in vivo α-syn is generally acetylated at the N terminus20,21, we modified the recombinant α-syn with N-terminal acetylation (Supplementary Fig. 1). The recombinant N-acetylated E46K α-syn (termed as Ac-E46K) formed amyloid fibrils after incubation in the buffer containing 50 mM Tris, 150 mM KCl, pH 7.5, at 37 °C with agitation for 7 days. We first used the atomic force microscopy (AFM) to characterize the fibril morphology. The result showed that the Ac-E46K fibril features a different fibril structure from the N-acetylated WT (Ac-WT) fibril formed under the same condition22 (Supplementary Fig. 2). The Ac-E46K fibril is approximately two times more twisted than the Ac-WT fibril with a half pitch of ~64 nm in comparison with ~120 nm of the Ac-WT. Moreover, the Ac-E46K fibril features a right-handed helical twist (Supplementary Fig. 2), which is distinct from the left-handed twist commonly found in the WT and other PD-familial mutant α-syn fibrils15,16,22–24. The different fibril morphologies suggest different properties of the Ac-E46K and Ac-WT fibrils, which may associate with their different pathologies.

To identify the different properties of the Ac-E46K and Ac-WT fibrils, we first sought to test the stability of the fibrils by cold denaturation25. The fibrils were cooled down and incubated at 0 °C for 48 h. The result of circular dichroism (CD) spectroscopy showed that the Ac-E46K fibril lost its β-sheet structure significantly faster than that of the Ac-WT fibril (Fig. 1a). Consistently, negative-staining transmission microscopic (TEM) imaging showed a faster disassembly of the Ac-E46K fibril than that of the Ac-WT fibril (Supplementary Fig. 3a). Intriguingly, we noticed that the Ac-E46K fibril stored at ~80 °C underwent apparent fragmentation after thawing; in contrast, the Ac-WT fibril were generally stable, and fragmentation was only observed after several cycles of freeze-thaw (Supplementary Fig. 3b). CD spectra further showed that in eight cycles of freeze-thaw, the structure of Ac-WT fibril was well maintained with a consistent content of β-sheet structure; in contrast, the Ac-E46K fibril gradually lost its β-sheet structure as the freeze-thaw cycles increased (Fig. 1b). In consistence with the freeze-thaw experiment, as we sonicated the preformed fibrils (PFFs), we noticed that under the same sonication condition, the Ac-E46K fibril broke into significantly shorter fragments than that of the Ac-WT fibril visualized by TEM and AFM (Fig. 1c, d, and Supplementary Fig. 4). It has been shown that fibril fragmentation is a key factor in nucleation-depended fibrillation26. Thus, these results indicate that the Ac-E46K fibril is less stable than the Ac-WT fibril, and may gain an enhanced pathology.

Since α-syn fibrils propagate by seeding the amyloid fibril formation in the transfected neurons, we attempted to compare the seeding property of the Ac-E46K and Ac-WT PFFs. We sonicated the PFFs and confirmed a comparable amount of Ac-E46K and Ac-WT PFF seeds by SDS-PAGE (Supplementary Fig. 5a). Of note, although in the absence of PFF seeds, the fibrillation of both Ac-WT and Ac-E46K were weak in our assay (Supplementary Fig. 5b and c), we observed a faster aggregation of Ac-WT than that of the Ac-E46K (Supplementary Fig. 5d). On the contrary, in the presence of PFF seeds, Ac-E46K aggregated faster than Ac-WT (Fig. 1e). Consistently, the results of TEM and SDS-PAGE showed more amyloid fibrils formed in the seeded Ac-E46K samples than that of the Ac-WT (Supplementary Fig. 5b and c). These results indicate that the Ac-E46K PFF seeds are more potent than that of the Ac-WT in seeding α-syn fibril formation.

Furthermore, we compared the stability of the fibrils under proteinase K (PK) digestion. The result showed that the Ac-E46K fibril is digested significantly faster than that of the Ac-WT fibril (Fig. 1f). Together, these data indicate that the Ac-E46K fibril is less stable than the Ac-WT fibril in both physical and chemical treatments and more efficient in fragmentation and propagation.

Structure determination of the Ac-E46K α-syn fibril by cryo-EM. To understand the structural basis underlying the different properties between the Ac-E46K and Ac-WT fibrils, we sought to determine the atomic structure of Ac-E46K fibril by cryo-EM. The Ac-E46K fibril sample was fixed on the carbon grid and frozen in liquid ethane. The cryo-EM micrographs were acquired at ×105,000 magnification on the 300 keV Titan Krios microscope equipped with the K2 Summit camera. 13,064 micrographs picked from 754 micrographs were used for the reconstruction of the Ac-E46K fibril (Table 1). The Ac-E46K fibril sample is morphologically homogeneous with one dominant species in the 2D classification of the fibrils (Supplementary Fig. 6). After helical reconstruction of the dominant fibril species by Relion, we obtained a 3D density map of the Ac-E46K fibril to an overall resolution of 3.37 Å (Supplementary Fig. 7). The density map showed a right-handed helix with a width of ~10 nm and a half helical pitch of ~68 nm.
(Fig. 2), which is consistent with the AFM measurement (Supplementary Fig. 2). The fibril contains two protofilaments that intertwine along an approximate 2-fold screw axis (Fig. 2). The helical twist between α-syn subunits is $-179.37^\circ$ and the helical rise is 2.38 Å (Fig. 2b, c, Table 1).

The Ac-E46K fibril exhibits a distinct fold of α-syn fibril. The high quality cryo-EM density map allowed us to unambiguously build an atomic structure model for the Ac-E46K fibril (Fig. 3a). This structure represents the fibril core (FC) of the Ac-E46K fibril consisting of residues 45–99 out of a total of 140 amino acids of α-syn (Fig. 3a), which is slightly smaller than the Ac-WT FC consisting of residues 37–99 formed under the same condition\textsuperscript{15}. While, similar to that of the WT fibril, the N- and C-termini of Ac-E46K α-syn remain flexible and are not visible by cryo-EM.

The Ac-E46K FC exhibits a serpentine fold, in which the topology of the β strands is different from any reported α-syn...
Aside from the large conformational change of the FC-N, the α-syn molecule (termed as FC-C) via interactions with T75, K45, and K80 forms inter-molecular salt bridge with E61 of the opposing FC-C. The E46K mutation also leads to the disruption of the WT α-syn subunit to involve in the FC-N from the other two essential electrostatic interactions (K58-E61 and E61-K58), locks the FC-N with FC-C side-chain interactions within the same proto-hairpin and extends from the β-hairpin conformation (Fig. 3d). In the Ac-WT fibril, E46 forms a salt bridge with K80, which together with the β-hairpin conformation in the adjacent layer (Fig. 3d).

Thus, the single mutation of E46K entirely changes the α-syn fibril structure.15,16,22-24,27,28 (Fig. 3b). Compared to the structure of Ac-WT α-syn fibril, the major difference lies in the N-terminal region of the FC (termed as FC-N), which covers residues 37–59 in the Ac-WT fibril or 45–59 in the Ac-E46K fibril. In the Ac-WT structure, the FC-N region stretches and packs around the rest of the α-syn molecule (termed as FC-C) via interactions with T75, A76, A78, and K80 of FC-C (Fig. 3c). In contrast, in the Ac-E46K structure, this segment forms a β-hairpin and extends from the FC-C with interactions to the side chains of Q62, T64, and V66 (Fig. 3c).

Aside from the large conformational change of the FC-N, the FC-C regions of the Ac-E46K and Ac-WT fibrils adopt a similar topology with a Greek key-like fold (Fig. 3c); however, the two FC-C structures are very different with an r.m.s.d. (Ca atoms) of 5.558 Å (Supplementary Fig. 8). In the Ac-WT fibril, α-syn subunits fold in a flat layer; in contrast, in the Ac-E46K fibril, β6 and β7 of the FC-C swap to the next layer to form inter-molecular side-chain interactions within the same protofilament (Fig. 3d). Thus, the single mutation of E46K entirely changes the α-syn fibril structure.

E46K rearranges the electrostatic interactions in the α-syn fibril. Since the overall structural rearrangement of α-syn fibril is initiated by one single mutation of E46K, we looked close to residue 46 in the Ac-WT and Ac-E46K fibrils. In the Ac-WT fibril, E46 forms a salt bridge with K80, which together with the salt bridge formed by E61 and K58, locks the FC-N with FC-C (Fig. 4a, b). In contrast, in the Ac-E46K fibril, the E46K mutation breaks the salt bridge of E46-K80, which results in the breaking of the other two essential electrostatic interactions (K58-E61 and K45-H50-E57) (Fig. 4a, b). Alternatively, in the Ac-E46K fibril, K80 forms inter-molecular salt bridge with E61 of the opposing α-syn subunit to involve in the fibril interface, and K45 directly forms salt bridge with E57 to stabilize the β-hairpin conformation of the FC-N (Fig. 4a, b).

The E46K mutation also leads to the disruption of the WT protofilament interface (Fig. 4c, d). Upon the flipping away of the FC-N from the fibril interface, a distinct fibril interface is formed. The exposed segments 74–79 of opposing α-syn subunits form...
typical class I parallel in-register homo-steric zippers to mediate the inter-protofilamental interactions (Fig. 4d). The steric-zipper interface is stabilized by the hydrophobic interactions between the side chains of V74, A76, and A78 and flanked by the intermolecular salt bridges formed by K80-E61, which further lock the two protofilaments (Fig. 4a).

The release of FC-N from the fibril interface in the Ac-E46K fibril increases the flexibility of this segment, which exhibits higher B-factors and poor density for tracing the N-terminal residues 37–44 that are visible in the Ac-WT FC (Fig. 4c). Thus, the relatively loose packing of the Ac-E46K fibril in comparison with that of the Ac-WT fibril may explain the decreased stability and increased sensitivity of the Ac-E46K fibril to the environment, which may associate with the pathology of this PD familial mutation.

**Discussion**

Structural polymorphism is a common characteristic of amyloid fibrils formed by different amyloid proteins such as α-syn, tau, and Aβ. Different fibril polymorphs may represent different pathological entities that are associated with different subsets of neurodegenerative diseases. Determination of polymorphic amyloid fibril structures is useful for the mechanistic understanding of amyloid pathology. In this work, we report the cryo-EM structure of an α-syn fibril with N-terminal acetylation and familial mutation E46K found in parkinsonisms. The Ac-E46K α-syn fibril exhibits a distinct structure that is distinct from the reported α-syn polymorphs. Compared to the Ac-WT fibril prepared under the same condition, the E46K mutation breaks an important electrostatic interaction of E46-K80 and thus induces the reconfiguration of the overall structure. In the mutant fibril, the FC-N region that used to form the protofibrillar interface in the Ac-WT fibril, flips away from the fibril interface and becomes less stable. In line with the structural information, we find that the Ac-E46K fibril is less stable upon both physical and chemical treatments. Furthermore, seeding experiments indicate that the instability and fragmentation-prone property of Ac-E46K fibril may lead to an increased capability of propagation. Of note,
the Ac-E46K fibril reported in this work represents the only right-handed α-syn fibril identified so far. Intriguingly, a recent study on Aβ structure showed that brain-derived Aβ fibrils are right-hand twisted, rather than left-hand twisted as seen in the in vitro formed Aβ fibrils. What defines the helical chirality of the amyloid fibrils? Whether and how does it determine fibril pathology? These are interesting questions to be answered.

Protein structures are determined by their primary sequences. While, regarding amyloid proteins, this dogma becomes more complicated. The reported α-syn fibril structures usually contain the central region of α-syn including residues ~37–99. This ~60 a.a. long sequence can fold into various structures to construct polymorphic α-syn fibrils. Several WT α-syn polymorphs have been observed in the in vitro prepared fibrils (Supplementary Fig. 9). Furthermore, a recent study on brain-derived α-syn fibrils showed that in vivo cofactors can participate into α-syn fibril structure and introduce more polymorphs. Recently, several fibril structures formed by inherited α-syn mutants have been reported including Ac-A53T, H50Q, and E46K. The Ac-A53T and H50Q fibrils exhibit similar morphology, and α-syn monomer in these structures folds similarly to that in the Ac-WT fibril (Supplementary Fig. 9). Two E46K α-syn fibril structures have been reported. These two structures closely resemble the so-called polymorph 2a WT α-syn fibril.
**Fig. 5 Electrostatic interactions in α-syn fibril polymorphs.** Primary sequence of WT α-syn fibril core (FC) is shown on top. K residues are colored in blue; H is light blue; E/D is red. Interacting charged residues in the structures of different α-syn polymorphic fibrils are connected with solid lines. One layer of different polymorphic structures of α-syn fibrils are shown below. The dimeric α-syn molecules are colored in different levels of gray. Electrostatic interactions are highlighted with K colored in blue, H in light blue and E in red. PDB IDs of the structures are provided in parentheses.

(Supplementary Fig. 9). Distinctive from the known polymorphs, the Ac-E46K fibril structure in the current work represents a distinct fold of α-syn (Supplementary Fig. 9), which demonstrates how the E46K mutation might rearrange the fibril structure. The polymorphic structures of E46K fibrils may derive from the different fibrillation conditions, and N-terminal acetylation may also play a role by kinetically affecting aggregation.

Then, how many fibril structures can α-syn potentially form? What determines the possibilities? We notice that different α-syn structures contain different electrostatic interactions, which play an important role in defining and stabilizing the overall structure (Fig. 5). In the Ac-WT fibril, intra-molecular electrostatic interactions of K58-E61 and E46-K80 lock the Greek key-like fold of α-syn, and the inter-molecular interactions of E45-H50-E57 are involved in stabilizing the fibril interface, which pairs α-syn protofilaments to form mature fibrils (Fig. 5). In contrast, in the polymorph 2 a WT fibril, electrostatic interaction forms between K45 and E57, which mediates the dimerization of the protofilaments (Fig. 5). Alternatively, the Ac-E46K fibril contains an intra-molecular electrostatic interaction between K45 and E57, which locks the β-hairpin fold of the FC-N region, and an inter-molecular interaction between E61 and K80′, which stabilizes the interface of the mutant fibril (Fig. 5). Different patterns of electrostatic interactions are also observed in other amyloid fibrils. For instance, tau fibrils derived from Pick’s disease exhibit different electrostatic interactions from those from AZ and chronic traumatic encephalopathy (CTE) [40-42] (Supplementary Fig. 10). These structural information suggest that different combinations of charged residues may serve as one of the key factors for the selection of fibril polymorph. In addition, mutations, post-translational modifications and cofactors can further enrich the potential variations of α-syn fibril structures.

**Methods**

**Preparation of Ac-WT and Ac-E46K α-syn.** Protein expression and purification of full-length α-syn Ac-WT and Ac-E46K mutation follows the same protocol described previously[39]. Briefly, the WT and E46K α-syn gene were inserted into pET22 vector and transformed into E. coli expression cell line BL21(DE3) (TransGen Biotech, Cat. # CD601-02). Primer sequences used in this study are listed in Supplementary Table 1. Fission yeast N-acetyltransferase complex B was co-expressed with α-syn to obtain the N-terminally acetylated α-syn[43], 1 mM isopropyl-1-thio-D-galactopyranoside (IPTG) was used to induce protein expression at 37 °C for 4 h. Cells were harvested in 50 mM Tris-HCL, pH 8.0, 1 mM phenylmethylsulfonyl fluoride, 1 mM EDTA and lysed by sonication. After centrifugation at 16,000×g for 30 min, the supernatant was boiled at 100 °C for 10 min. After another centrifugation, streptomycin (20 mg ml⁻¹) was added to remove nucleic acid in the supernatant. pH was adjusted to 3.5 using 2 M HCl to precipitate other unwanted components. 50 mM Tris-HCL (pH 8.0) was used to dialyze the supernatants overnight. Q column (GE Healthcare, 17-5156-01) and size-exclusion chromatography Superdex 75 (GE Healthcare, 28-9893-33) were used for protein purification. Protein samples were loaded to Q column with buffer A (50 mM Tris-HCL, pH 8.0), and eluted with a gradient (20–60%) of buffer B (50 mM Tris-HCL, 1 M NaCl, pH 8.0). For Superdex75, the buffer containing 50 mM Tris-HCL, 150 mM KCl pH8.0 was used. After purification, an on-line EASY-nL-LC 1000 coupled with an Orbitrap Q-Exactive HF mass spectrometer was used to verify that the Ac-E46K α-syn protein sample is indeed acetylation (Supplementary Fig. 1).

**Fibrillation of Ac-WT and Ac-E46K α-syn.** 100 μM recombinant Ac-WT and Ac-E46K α-syn in 50 mM Tris, pH 7.5, 150 mM KCl were incubated at 37 °C with constant agitation (900 rpm) in ThermoMixter (Eppendorf) for a week, respectively. The preformed fibrils were sonicated at 20% power for 15 times (1 s per time, 1 s interval) on ice by JY92-IIN sonicator (Xinyi Sonication Equipment Company, Ningbo, China). Then, the α-syn PFF seeds (0.5 mol%) were added to 100 μM α-syn monomer, and further incubated at 37 °C with agitation (900 rpm) for a week to obtain mature fibrils.

**CD spectroscopy.** For cold incubation assay, 20 μM Ac-WT or Ac-E46K α-syn fibrils were incubated at 0 °C. Fibril samples were picked up at different incubation time for CD measurement. For freeze-thaw assay, fibril samples were incubated in liquid nitrogen for 10 min and then warmed up in water bath at room temperature for 5 min. The secondary structure of the fibril sample was measured by a Chirascan CD spectrometer (Applied Photophysics, UK). Spectra were recorded from 200 nm to 260 nm at 0 °C or room temperature using a cell with a path length of 0.1 cm. All spectra were collected more than 3 times with a background-corrected against buffer blank.

**Atomic force microscopy (AFM).** As for characterizing fibril fragmentation, 20 μM Ac-WT or Ac-E46K α-syn fibrils were sonicated for 4 times (1 s per time, 1 s interval) on ice by JY92-IIN sonicator. 8 μl of diluted samples (10 μM) were mounted on a freshly cleaved mica for 3 min and gently rinsed with Milli-Q water to remove unbound fibrils. The samples were dried by nitrogen flow. Images were acquired with ScanAsyst air mode by using NanoScope V Multimode 8 (Bruker). SNL-10 probe was taken with a constant of 0.35 N m⁻¹ for scanning. Measurements were recorded at 512 x 512 pixels at a line rate at 1.5 Hz. The percentage for different length of fibrils were processed and analyzed by using NanoScope Analysis software (version 1.5).
As for characterizing the morphology of mature α-syn fibrils, samples were acquired with 2 μm α-syn fibrils. The image and height were analyzed on the Nanoscope software and image.

Proteolytic digestion of α-syn PFFs. α-Syn fibrils were concentrated by centrifugation (14,462 × g, 25 °C, 45 min), washed and resuspended with PBS, followed by sonication at 20% power for 15 times (1 s per 1 s interval) on ice by T92-IIN sonicator. The concentration of sonicated PFF seeds was calculated as equal to that of the PFFs before sonication. The concentration of PFFs before sonication was calculated as the amount of α-syn monomer subtracting the amount of soluble α-syn after fibril formation. 0.3 mol% or 0.5 mol% α-syn AcWT or Ac-E46K PFF seeds were added into 50 μM α-syn monomer (50 mM Tris, pH 7.5, 150 mM KCl, 0.05% NaN3). Final concentration of 10 μM ThT was added to the reaction mixture. Reactions were performed in triplicates in a 384 well optical plate (Thermo Scientific Nunc). The plate was detected by a Varian Flash plate reader (Thermo Scientific). Fluorescent intensities of each reaction were acquired at 440 nm (excitation) and 485 nm (emission) wavelengths, with a bottom read. Graphing was performed with GraphPad Prism 6.

Negative-staining electron microscopy. Five microtiters of aliquot of fibril sample was applied to a glow-discharged 200 mesh carbon support film (Beijing Zhongkingkey Technology Co., Ltd.) for 45 s. Then the grid was washed with 5 μl double-distilled water and followed by another wash of 5 μl 3% w/v uranyl acetate. Another 5 μl 3% w/v uranyl acetate was added to the grid to stain the sample for 45 s. After removing the excess buffer by filter paper, the grid was dried by infrared lamp. The sample imaging was applied by Tecnai T12 microscope (FEI Company) operated at 120 KV.

Cryo-electron microscopy. An aliquot of 4 μl of 6 μM Ac-E46K α-syn fibril solution was applied to a glow-discharged holey carbon grid (Quantifoil R1/2/1.3, 300 mesh), blotted for 6.5 s, and plunge-frozen in liquid ethane using FEI Vitrobot Mark IV with 95% humidity at 16 °C. The grids were loaded onto a FEI Titan Krios microscope operated at 300 kV with magnification of ×105,000, in which a GIF Quantum energy filter (slit width 20 eV) were equipped. A Gatan K2 Summit camera recorded the cryo-EM micrographs using super-resolution mode with pixel size 0.66 Å. Defocus values were set from −2.4 to −1.5 μm. All of the micrographs were dose-thresholded to 32 frames and the electron dose rate was set to ~8.3 counts per physical pixel per second (~6.25 e−/Å² per s), total exposure time is 8 s, and 0.25 s per frame. Hence, the total dose is ~50 e−/Å². AutoEMOTION2 written by Dr. Lei was used for all data collection 45.

Image pre-processing. All 32 frames were aligned, summed and dose-weighted by MotionCor2 and further binned to a pixel size of 1.33 Å. The defocus values of dose-weighted micrographs were estimated by CTFFIND4.1,5, and micrographs with the highest-resolution Thon ring >6 Å, were kept for the subsequent processing in RELION3.0 47. 13,064 filaments were picked manually from 754 micrographs. The extracted segments (box size: 160 × 160, inter-box distance of 2.5 μm) were used to reconstruct a 3D model of the helical polymer.

Helical parameters determination. Twist angle and rise value of helical polymer was calculated before reconstruction. Micrographs, which were involved in the selected 2D classes, were used to estimate the rise value by a C++ program that searches the strongest diffraction position around the 1/2θ Å−1 in Fourier space. The average of these positions was converted to the rise value in real space. The way to estimate the twist angle was the same as that described in our previous structural determination of Ac-WT α-syn fibril 31, in which helical pitch length of each filament was estimated and the average of these helical pitch lengths was used to calculate the twist angle.

Helical reconstruction. Several iterations of 2D classification, 3D classification and 3D refinement were performed to filter the segments belonging to the same conformation. Segments of the selected 2D classes were used to reconstruct a reference with more feature in 3D refinement from the initial featureless cylinder model created by the relion_helix_toolbox program. 3D classification (k = 2) then built a reasonable map based on the improved reference model. Auto-refine procedure was used using single class using resolution from the result of 3D classification. No explicit τau2_fudge value was specified, and no mask was used. The final map was convergence with the rise of 2.38 Å and the twist angle of ~179.37°. The map was sharpened with a B-factor of ~97.8051 Å2 using post-processing subroutine in RELION. The overall resolution was reported at 3.37 Å by the standard FSC = 0.143 criteria. Local resolution estimation calculated by ResMap 48.

Polymorphs of Ac-E46K α-syn fibril. Two polymorphs at least existed in micrographs. One is twisted filament, and the another one is non-twisted filament which currently cannot be solved since this polymorph has no twist. All filament including twisted and non-twisted filaments in 754 micrographs were counted as 18891. Among them, 13064 was twisted filaments which was used for reconstruction. The percentage of twisted filament is 69.15% and was dominated in the two fibril polymorphs.

Model building and refinement. A homology model based on the NMR structure (PDB entry code 2N0A) was built and modified by COOT 28,49. The model with the three adjacent layers (six promoters) was refined using the real-space refinement program in PHENIX 50. The subunit dimers in the middle of three layers was extracted and used as the final model.

Reporting summary. Further information on research design is available in the Nature Research Reporting Summary linked to this article.

Data availability. Density maps of the Ac-E46K α-syn fibril are available through EMDB with entry code: EMD-0833. The structural model was deposited in the Protein Data Bank with entry code: 6L4S. Other structural models used in this study are available in the Protein Data Bank with entry codes: 6A6B (Ac-WT), 6RT0 (WT polymorph 2a), 6RT1 (WT polymorph 2b), 6CU8 (WT), 6LRQ (Ac-A53T), 6PES (H50Q), 6UFR (E46K), and 2N0A (solid-state NMR structure of WT). The source data underlying Figs. 1a, 1b, 1e, 1f and Supplementary Fig. 3d are provided as a Source Data file. Other data that support the findings of this study are available from the corresponding authors upon reasonable request.

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Author contributions

C.L., X.L., and D.L. designed the project. K.Z., H.L., C.Z., and F.L. prepared the Ac-WT and Ac-E46K monomer and fibril samples. Z.L., H.L., K.Z., and Y.T. performed the biochemical assays. Z.L. and Y.S. performed the AFM experiments. Y.L, K.Z. and X-D.S. performed the cryo-EM experiments. Y.L. built and refined the structure model. All of the authors are involved in analyzing the data and contributed to manuscript discussion and editing. D.L. and C.L. wrote the manuscript.

Competing interests

The authors declare no competing interests.

Additional information

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