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Title:

(-)-Epigallocatechin gallate prevents the aggregation of tau protein into toxic oligomers at substoichiometric ratios

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Keywords

Tau protein, tau oligomers, aggregation inhibitors, EGCG, Alzheimer's disease, polyphenol

Abbreviations:

AD, Alzheimer's disease; EGCG, (-)-epigallocatechin gallate; ThT, Thioflavin T; PCR, polymerase chain reaction; NBT, nitroblue tetrazolium; MTT, 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide; AFM, atomic force microscopy

Total: 2972

Abstract

The accumulation of amyloid-beta ($A\beta$) and tau aggregates is a pathological hallmark of Alzheimer's disease. Both polypeptides form fibrillar deposits, but several lines of evidence indicate that $A\beta$ and tau form toxic oligomeric aggregation intermediates. Depleting such structures could thus be a powerful therapeutic strategy. We generated a fragment of tau (His-K18 Δ K280) that forms stable, toxic, oligomeric tau aggregates *in vitro*. We show that (-)-epigallocatechin gallate (EGCG), a green tea polyphenol which previously was found to reduce $A\beta$ aggregation, inhibits the aggregation of tau K18 Δ K280 into toxic oligomers at ten- to hundred-fold substoichiometric concentrations, rescuing toxicity in neuronal model cells.

Introduction

A variety of peptides and proteins are known to assemble into highly ordered aggregate structures with a cross-beta sheet structure, termed amyloid. The brains of patients suffering from Alzheimer's disease (AD) exhibit two types of aggregated protein deposits: extracellular plaques mainly composed of amyloid-beta (A β) peptide and intracellular neurofibrillary tangles composed of the hyperphosphorylated protein tau, in which the microtubule-binding repeat domain adopts a cross-beta structure¹⁻⁴. Tau exists in multiple splicing variants and is a highly hydrophilic and thus soluble protein with little propensity to aggregate⁵. The K18 Δ K280 tau protein fragment spans the four repeat domains of the microtubule-binding domain of the longest human isoform tau40; the deletion of lysine 280 has been described in FTDP-17⁶ (Figure 1 A). Filaments of wild-type (wt) tau only form *in vitro* in the presence of polyanions such as heparin^{7,8}, however K18 Δ K280 aggregates into β -sheet rich structures without heparin⁹.

Therapeutic efforts into slowing or reversing the progression of neurodegenerative diseases have concentrated on clearing insoluble protein aggregates or inhibiting the aggregation of amyloidogenic proteins. In recent years, new therapeutic strategies have emerged, which include the stabilization of mature fibrils to deplete toxic oligomers or the redirection of the aggregation cascade to increase the formation of non-toxic, off-pathway aggregates¹⁰.

Since two types of fibrillar aggregates are formed from A β and tau in AD, small molecules targeting both polypeptides could be a valuable therapeutic strategy. Inhibitors of heparin-induced tau paired helical filament formation were found in the classes of anthraquinones, polyphenols, porphyrins and phenothiazines^{11,12}. The green tea polyphenol (-)-epigallocatechin gallate (EGCG) interferes with aggregation of numerous amyloidogenic proteins and small polypeptides and ameliorates their detrimental effects¹³⁻¹⁸. It stimulates the assembly of non-toxic, unstructured, off-pathway oligomers from A β , α -synuclein and IAPP *in vitro*^{19,20}. Despite challenges in its variable oral bioavailability, short pharmacokinetic half-life, and limited partitioning across the blood-brain barrier, this may make it a promising model drug for AD^{21,22}. However, the effect of EGCG on tau aggregation has not yet been characterized. Here, we developed a cell-free aggregation assay using the tau fragment His-K18 Δ K280 to probe the effect of EGCG under physiological conditions in the absence of heparin. We demonstrate that EGCG prevents the formation of β -sheet rich aggregates at substoichiometric concentrations and reduces their toxicity in cell-based assays, suggesting a specific interaction of EGCG with a crucial early aggregation intermediate.

Materials and Methods

Protein expression and purification

The construct encoding the tau fragment K18 (K18ΔK280, Figure 1 A), comprising the mutant four-repeat microtubule-binding domain, was kindly provided by M. Holzer. Expression clones for K18ΔK280 with an N-terminal His₆-tag were generated using the Gateway® technology (Life Technologies). *AttB* sites were added to the construct by PCR (forward primer: 5'-GGGGACAAGTTTGTACAAAAAAGCAGGCTGGatgcagacagccccgtgcccatgc-3'; reverse primer: 5'-GGGGACCACTTTGTACAAGAAAGCTGGGTGTCATTCAATCTTTTATTTCTCC-3'). pDONR221 and pDESTco were chosen as entry and expression vectors, respectively. Ni-NTA matrix columns were used for purification of recombinant protein (QIAGEN). Columns were washed with native wash buffer or with wash buffers of decreasing urea concentrations, and the His-tagged protein was eluted under native conditions following the manufacturer's specifications (QIAGEN).

The protein was subjected to reversed-phase purification using Poros 50 R1 resin (Applied Biosciences), washed using 0.1% trifluoroacetic acid (TFA), and eluted with 60% acetonitrile/0.1% TFA, lyophilized and the aliquots stored at -20°C. Protein concentration was determined by OD₂₈₀ using an extinction coefficient of 8480 M⁻¹cm⁻¹.

Protein aggregation

Lyophilized His-K18ΔK280 protein was reconstituted in PBS and sonicated at 4°C for 5 min. The protein solution was cleared by centrifugation at 200,000 x g and 4°C for 20 min. His-K18ΔK280 was incubated at 37°C in 96-well plates at 12.5 μM, with 5 mM dithiothreitol (DTT) added daily to prevent formation of intramolecular disulfide bonds. EGCG (Sigma Aldrich) solutions were freshly prepared with ultrapure H₂O.

Thioflavin T assay

To measure aggregation formation, 20 μM Thioflavin T (ThT) was added to each reaction and fluorescence measured in triplicates in black non-binding 96-well plates with clear, flat bottoms (#3651, Corning) using a microplate reader (Infinite M200, Tecan), at excitation and emission wavelengths of 440 and 485 nm, respectively. Results were calculated as means ± s.d.

SDS-PAGE and dot blot assays

Protein samples were either denatured by boiling at 95°C for 10 min with 2% SDS + 50 mM DTT for denaturing SDS-PAGE and denaturing dot blots or diluted with an equal volume of H₂O for non-

denaturing dot blots. 20 μ l were applied to nitrocellulose membranes. Membranes were blocked for 1 h in 3% nonfat milk and incubated overnight with the following primary antibodies at 4°C: polyclonal anti-tau A0024 (aa 243-441, DAKO; 1:2000), anti-amyloid oligomer A11 (1:2000, Life Technologies). Proteins were visualized by chemiluminescence (Chemicon) or the AttoPhos reagent (Promega). NBT staining of nitrocellulose membranes was performed as previously described^{19,23}.

Centrifugal sedimentation

Samples were collected every 24 h and subjected to ultracentrifugation for 20 min at 200,000 x g (70,000 rpm) and 4°C in a TL-100 centrifuge and TLA120.2 rotor (Beckman). Pellets were resuspended in PBS. Unfractionated samples, supernatants and resuspended pellets were boiled in SDS sample buffer, resolved by SDS-PAGE and visualized by silver staining as previously described²⁴.

Atomic force microscopy (AFM)

20 μ l of His-K18 Δ K280 samples were pipetted onto freshly cleaved mica fixed onto a glass slide and incubated for 10 min at room temperature, washed 3x with ddH₂O, and dried overnight. Samples were analyzed using the Nanowizard AFM (JPK, Berlin) in intermittent contact mode.

Tryptophan fluorescence spectroscopy

Protein solutions were diluted 1:10 in PBS. Fluorescence emission was measured at 280 nm absorption wavelength using a fluorescence spectrometer (LS 50, Perkin Elmer).

Cell viability assay

PC-12 cells (American Type Culture Collection) were grown in DMEM F12-K medium supplemented with 14.5% horse serum, 4% fetal calf serum, 100 U/mL penicillin and 100 μ g/mL streptomycin at 37°C with 5% CO₂ in a humidified environment. Prior to viability assays, DTT was removed from K18 Δ K280 aggregate reactions by buffer exchange using a 10 kDa centrifugal filter (Amicon) or by pelleting aggregates by ultracentrifugation (200,000 x g, 20 min), washing and resuspending in PBS. Cell viability was measured using the MTT assay as previously described¹⁹. Student's t-test was used to calculate p values.

Results

EGCG substoichiometrically prevents the formation of SDS-stable tau oligomers

First, we assessed the aggregation of tau K18ΔK280 (Figure 1 A) in the absence or presence of EGCG (Figure 1 B) by measuring Thioflavin T (ThT) fluorescence emission at 485 nm under reducing conditions (Figure 1 C). ThT increases fluorescence on binding to β -sheet rich amyloid-like structures²⁵. In the absence of EGCG, ThT fluorescence increased after an initial lag phase of ca 24 - 30 h. In contrast, incubation of K18ΔK280 with a substoichiometric concentration of EGCG (0.01x tau protein) elongated the lag phase. At higher concentrations of EGCG (0.1x, 1x and 5x), no increase in ThT fluorescence was observed (Figure 1 C), indicating that EGCG blocks the formation of amyloid-like, β -sheet rich tau aggregates. In contrast, EGCG did not prevent the rapid formation of β -sheet rich K18ΔK280 aggregates that were induced by heparin (Supplementary Figures S1 and S2 A), but changed aggregate morphology from large fibrillar assemblies into smaller spherical structures (Supplementary Figure S2 B). Likewise, EGCG reduced ThT binding to heparin-induced full length tau aggregates both under reducing (wt tau) and oxidizing conditions (Δ C tau), but did not delay aggregation or alter the fibrillar morphology of tau aggregates under these conditions (Supplementary Figure S3).

We next analyzed the effects of EGCG on tau aggregation by centrifugal fractionation of samples followed by denaturing SDS-PAGE and silver staining (Figure 1 D). In the absence of EGCG, we found a time-dependent increase of tau in the pellet fractions, indicating the formation of insoluble, high molecular weight aggregates (Figure 1 D). In addition, SDS-stable multimers appeared from 48 hours onwards. In contrast, incubation of the tau fragment with equimolar EGCG eliminated the formation of SDS-stable aggregates (Figure 1 D). Thus, our centrifugation assays confirm that EGCG treatment potently blocks K18ΔK280 aggregation *in vitro*.

To visualize K18ΔK280 aggregate species, we performed atomic force microscopy (AFM) on samples incubated with or without equimolar concentrations of EGCG. In the absence of EGCG, we observed the time-dependent formation of non-fibrillar, oligomeric K18ΔK280 aggregate species (Figure 1 E). In contrast, only a few small tau aggregate species were observed after incubation with EGCG.

Finally, we performed dot blot assays with the anti-oligomer antibody A11 in order to examine whether the spontaneously formed K18ΔK280 aggregates are toxic, disease relevant structures. Previous studies have demonstrated that A11 detects proteotoxic, oligomeric aggregate species of A β , tau and other polypeptides but not fibrils or monomers^{26,27}. In the absence of EGCG, dot blot analysis revealed strong A11 signals at 72 and 96 hours, supporting the results by AFM that predominantly oligomeric K18ΔK280 aggregates are spontaneously formed *in vitro* (Figure 1 E). Interestingly, at 120 hours, the A11 signal decreased, which could be due to the formation of the

larger aggregates that are less efficiently recognized by the antibody. In samples of K18ΔK280 co-incubated with EGCG, we observed no time-dependent increase in the A11 signal, suggesting that EGCG inhibits the formation of potentially proteotoxic oligomeric tau species.

Taken together, these data indicate that EGCG is a potent inhibitor of K18ΔK280 aggregation *in vitro*, which blocks the formation of insoluble, high-molecular weight, SDS-stable tau oligomers at substoichiometric concentrations.

EGCG prevents tau conformational changes and rescues tau toxicity in vitro

Next, we scrutinized the effect of EGCG on the structure of tau K18ΔK280. First, we tested if EGCG bound to the monomer or to some other tau species. After incubation with EGCG (5x molar ratio) we performed SDS-PAGE and detected tau by silver staining and EGCG by electroblotting and subsequent staining with the redox-sensitive dye nitroblue tetrazolium (NBT), which detects EGCG-bound polypeptides^{19,28}. As before, treatment with EGCG prevented the formation of SDS-resistant tau dimers, oligomers, or aggregates (Figure 2 A). However, a second (18 kDa) band appeared after 24 h with slightly lower mobility than the tau monomer (16 kDa). This band reacted weakly with silver staining, but strongly with the EGCG-sensitive dye NBT. This suggests that the bulk of the protein exists as a monomer that is not bound to EGCG, but that a subpopulation of EGCG-bound tau molecules exists. These molecules are either monomers that have a conformation that is distinct from the unbound tau monomers, or they are present in the form of SDS-labile oligomers.

To further assess the effects of EGCG on K18ΔK280 misfolding and aggregation, we performed dot blot assays under native conditions using the anti-tau antibody A0024 (Figure 2 B and C). In the absence of EGCG, antibody binding decreased from 48 hours onwards. After 96 hours of incubation, no tau signal could be detected (Figure 2 B). In contrast, the protein could be detected with the A0024 antibody after 96 h or 120 h in samples incubated with EGCG at 1:100 or 1:10 ratios, respectively. Tau was detected in all samples when re-probed with the same antibody under denaturing conditions (Figure 2 C), confirming that the loss of antibody binding indicated a change in secondary and/or tertiary structure during tau aggregation, which was prevented by EGCG treatment.

We speculated that EGCG might prevent early conformational changes in the tau fragment K18ΔK280 that could trigger subsequent aggregation. To investigate this hypothesis, we performed tryptophan fluorescence and circular dichroism spectroscopy (Figure 2 D and E). In the absence of EGCG, the tryptophan fluorescence emission spectrum of monomeric K18ΔK280 (incubation time 0 hours) showed a maximum emission at approximately 350 nm. After 120 hours of incubation, a

hypsochromic shift in the emission spectrum was observed, with maximum emission at 335 nm. This shift in emission was abrogated by co-incubation of K18 Δ K280 with EGCG, suggesting that EGCG suppresses the conformational changes observed in untreated protein samples. Correspondingly, circular dichroism spectra before and after incubation with EGCG both indicated that the protein was largely unstructured and did not adopt a β -sheet conformation (Figure 2 E).

Finally, we investigated whether EGCG-treated or -untreated tau aggregates are toxic to mammalian cells. To this end, we incubated K18 Δ K280 for 144 h in the presence or absence of EGCG and then filtered K18 Δ K280 aggregate samples using a 10 kDa filter to remove all small molecules while retaining all tau species. Solutions were then applied to PC-12 cells for 72 hours, and cell viability was measured by MTT reduction (Figure 2 F). Our results showed a significant cellular toxicity of aggregated K18 Δ K280, which was rescued by co-incubation of the protein with EGCG. To confirm that tau toxicity and its rescue by EGCG was linked to the aggregation of tau, we pelleted aggregated tau by ultracentrifugation and washed the pellet to remove DTT, EGCG, and soluble tau protein (Figure 2 G). We found that the resuspended aggregates were toxic to PC-12 cells as assessed by the MTT assay, whereas pellets from EGCG-treated samples were neither toxic nor contained any tau protein. This confirmed that tau toxicity was linked to its aggregation and that EGCG prevented the formation of toxic tau aggregates.

Discussion

Alzheimer's disease is characterized by the two aggregated protein species A β and tau^{1,2}. While much effort has been put into elucidating the mechanisms of A β polymerization, and a large body of evidence exists for a toxic role of A β oligomers in AD pathology²⁹⁻³¹, studies investigating the aggregation mechanism of the microtubule-associated protein tau have been less numerous. Recently, oligomeric forms of tau isolated from AD brains were found to be potently toxic^{32,33} and have been discussed as potential targets for therapeutic intervention³⁴⁻³⁶. The aim of this study was therefore to establish a model for tau oligomerization *in vitro*, and to investigate the effects of the green tea polyphenol EGCG on tau oligomerization without the help of polyanions.

EGCG has been implicated in several studies as a drug candidate that modulates the aggregation of proteins implicated in neurodegenerative diseases, such as huntingtin, amyloid-beta and α -synuclein^{19,23}. We found that EGCG strongly inhibits tau aggregation even at highly substoichiometric concentrations, suggesting that the compound may target an aggregation intermediate that appears early in the amyloid formation cascade. However, loss of ThT fluorescence by itself is not a reliable marker for a loss of amyloid-like structures³⁷. Several hypotheses could explain the effect of EGCG on ThT fluorescence: (1) EGCG binds to K18 Δ K280 without altering aggregation, but displaces ThT; (2) in the presence of EGCG, K18 Δ K280 aggregation is redirected into a distinct aggregate species that has no affinity for ThT; (3) EGCG prevents the polymerization of K18 Δ K280, so that the protein largely remains in its natively unfolded monomeric form. We will discuss these possible mechanisms in the following paragraphs.

Sedimentation assays indicated that EGCG indeed inhibits polymerization of tau into high molecular weight aggregates; however, they do not exclude the formation of SDS-labile, low-molecular aggregates in the presence of EGCG. We performed atomic force microscopy to investigate whether such aggregates might be formed in the presence of EGCG. Spherical and amorphous aggregates of K18 Δ K280 were detectable from 48 hour onwards in the absence of EGCG, but not in its presence, indicating that if such aggregates form in the presence of EGCG, they are rare. The hypsochromic shift in tryptophan fluorescence of tau oligomers and the masking of antibody binding epitopes compared to monomeric protein indicated a structural change during aggregation that was prevented by EGCG. Dot blot analysis using anti-oligomer antibody A11 and circular dichroism spectroscopy further corroborated that EGCG prevents the formation of β -sheet-rich oligomeric tau species.

In summary, these data strongly support the hypothesis that EGCG prevents the formation of toxic tau oligomers and keeps the bulk of the tau fragment in a monomeric, unfolded state. In contrast, previous data on A β , α -synuclein, and IAPP indicated that EGCG induces a specific

conformational change and redirects these polypeptides into off-pathway aggregation^{14,19}. Since EGCG inhibited tau aggregation at highly substoichiometric concentrations, stable, quantitative binding to the tau monomer cannot explain its effect. Rather, we conclude that EGCG binds to a rare partially misfolded monomeric or oligomeric tau species. Quinoid substances such as EGCG can covalently bind proteins²⁸. While the role of covalent modification in the mechanism of EGCG is still debated^{20,37}, the appearance of a distinct population of EGCG-bound tau molecules in SDS gels (Figure 2 A) indicates tight, and possibly irreversible, binding of EGCG to a subpopulation of the tau protein. This suggests a mechanism in which EGCG binds to a rare aggregation nucleus and converts it into an inactive conformation, thus removing its seeding activity.

The apparent difference in mechanism of EGCG between K18ΔK280 polymerization on one hand and Aβ, IAPP and α-synuclein polymerization on the other hand, would then result from the rare occurrence of nucleus formation in tau, when compared to nucleus formation of other polypeptides^{19,20} (Figure 3). Correspondingly, the effect of EGCG on heparin induced tau aggregation more resembled its effect on Aβ. Here, the lag phase was very short and nucleus formation was not a rare rate-limiting step. Consequently, EGCG had little effect on aggregation kinetics even though it did change the structure of heparin-induced K18ΔK280 aggregates.

Alternatively, EGCG could transiently bind rare oligomers and catalytically return tau to its unfolded monomeric state, acting as a chemical chaperone^{38,39}. Chemical chaperones have been shown to both stabilize the native conformation of amyloidogenic proteins and to destabilize partially folded states³⁹⁻⁴¹. The presence of tightly bound EGCG-tau complexes and the absence of kinetic inhibition in heparin-induced aggregation both argue against such a mechanism. However, the methods used in our study are not sensitive to conformational changes in small subpopulations of the tau protein, so more refined experiments would be needed to conclusively rule out a catalytic mechanism.

In conclusion, our study indicates that EGCG is a potent inhibitor of tau aggregation and toxicity. Its effect on K18ΔK280 polymerization is distinct from its effects on Aβ and α-synuclein aggregation. Its dual inhibition of both Aβ and tau aggregation into β-sheet rich, toxic structures, might be of synergistic benefit for the treatment of AD.

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Figures

Figure 1 EGCG prevents the formation of tau oligomers and SDS-stable aggregates at substoichiometric concentrations. A) Structure of the K18Δ280 tau fragment comprising all four repeat domains (T1-4) of the microtubule-binding domain found in the longest human tau isoform Tau-441 and with a deletion of lysine residue 280. B) Structure of (-)-epigallocatechin gallate (EGCG). C) Effect of EGCG on tau K18ΔK280 aggregation (12.5 μM) as assessed by ThT fluorescence assay. Results represent mean fluorescence signal ± s.d. (n = 3). D) Effects of EGCG (equimolar concentration) on the formation of SDS-stable, high molecular weight tau aggregates. T = total, unfractionated sample, S = supernatant, P = pellet. Fractionation by centrifugal sedimentation at 200,000 x g. E) Analysis of untreated and EGCG-treated (equimolar concentration) K18ΔK280 by atomic force microscopy after 0 h, 48 h and 120 h incubation at 37°C. Scale bar = 500 nm. F) Effects of EGCG on the formation of tau oligomers as assessed by A11 dot blot analysis.

Figure 2 EGCG prevents tau conformational changes and rescues tau toxicity. A) EGCG binding to K18ΔK280 visualized by NBT staining (right) compared to silver stained protein (left). Asterisk indicates monomer band, arrowhead band stained by NBT. B) Dot blot analysis of non-denatured K18ΔK280 protein incubated in the absence (0x) or presence of substoichiometric (0.01x and 0.1x) concentrations of EGCG for 0 - 120 hours. C) Dot blot analysis of denatured samples of K18ΔK280 incubated under the same conditions. D) Tryptophan fluorescence spectra of K18ΔK280 (12.5 μM) incubated for 0 h or 120 h in the absence or presence of equimolar concentrations of EGCG (tryptophan absorption wavelength 280 nm). E) Circular dichroism spectra of K18ΔK280 (12.5 μM) incubated with EGCG. F), G) Assessment of tau K18ΔK280 toxicity using the MTT metabolic assay. PC-12 cells were incubated with protein incubated in the absence or presence of EGCG for 144 h. E) Protein samples were filtered using a 10 kDa size exclusion filter. F) Samples were pelleted by ultracentrifugation, washed and resuspended in PBS. Results represent means ± s.d. (n = 3). * p < 0.05, ** p < 0.01, *** p < 0.001 (Student's t-test).

Figure 3 Working model for the effects of (-)-epigallocatechin gallate on the aggregation of tau. In this model, EGCG preferentially binds to a misfolded monomer or a rare transient nucleus (triangle), not to the unfolded monomer (circle). Binding of EGCG to the misfolded monomer or nucleus either catalytically converts it back to its unfolded monomeric form (dashed arrow (1)), or converts it to an inactive conformation and removes it from the aggregation cascade (dashed arrow (2)), thus inhibiting subsequent aggregation steps.

Supplementary Information

(-)-Epigallocatechin gallate prevents the aggregation of tau protein into toxic oligomers at substoichiometric ratios

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Supplementary methods

Expression of full length tau protein

The wild type 2N4R (P10636-8) tau expression plasmid (pRK172-Tau 2N4R) was a generous gift from Virginia Lee (University of Pennsylvania). Site-directed mutagenesis to obtain 2N4R C291A/C322A was accomplished using *Pfu* turbo polymerase and transforming into XL 10-Gold ultracompetent cells. The WT as well as mutant plasmid was transformed into BL21DE3-Gold competent cells for protein expression and cells grown in 1L of LB media at 37°C followed by induction with 1 mM IPTG at 37°C when OD 600 was 0.5. The cells were pelleted 2 hours post induction and stored at -80°C. The pelleted cells were re-suspended in ice cold 80 mM PIPES, pH 6.8, 1 mM MgSO₄ and 1 mM EGTA and lysed using a French pressure cell press. The cell lysate was then centrifuged at 5,000 rpm for 20 min at 4°C and the supernatant was collected. The supernatant was then boiled at 100°C for 10 min and further centrifuged at 5,000 rpm for 10 min. The supernatant was loaded on a phosphocellulose-packed column equilibrated with lysis buffer and eluted with lysis buffer containing increasing concentrations of NaCl(0.2 - 1.0 M). The column fractions were screened by gel electrophoresis and the peak tau fractions were pooled and dialyzed against buffer containing 10 mM NH₃HCO₃. The dialyzed sample was then aliquoted and lyophilized.

Full length tau aggregation

Lyophilized wt tau monomer was dissolved in water + DTT (25 mM) to a concentration of 40 μM and incubated at room temperature for 1 h. The protein was diluted in aggregation buffer (100 mM HEPES, 100 mM NaCl, 20 μM ThT) and 8 μM heparin (8 μM) were added. Tau DC was prepared under identical conditions, but in the absence of DTT. Protein samples were incubated in a 96 well microplate (Corning #3651), at 37°C in a fluorescence microplate reader (Infinite F200) with intermittent shaking (5 s every 5 min). ThT signals were recorded at 436 nm excitation and 482 nm emission. After 72 h, 10 μl of sample was adsorbed onto mica for 5 min, washed with 4 x 40 μl water and 40 μl ethanol, dried for 2 h and imaged on a NanoMan AFM (Veeco) in tapping mode using a FESP tip.

Supplementary Figures

Supplementary Figure S1

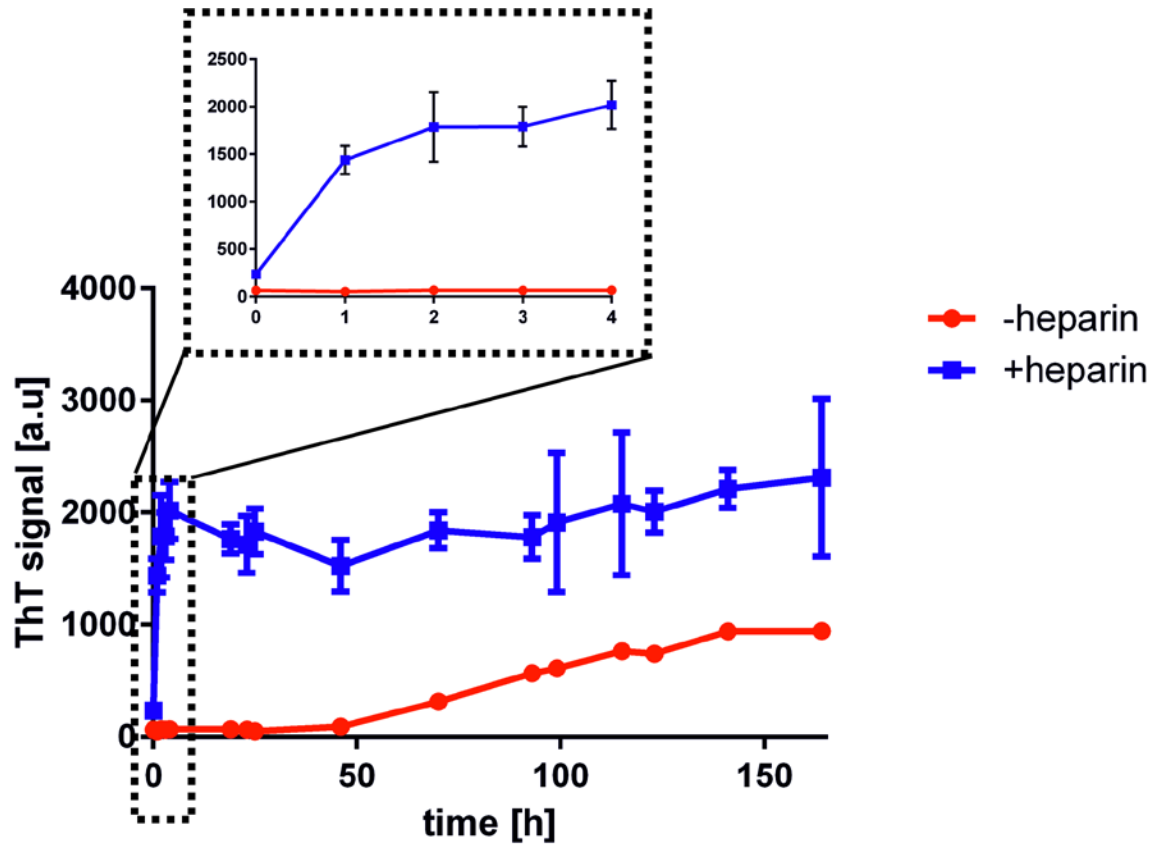
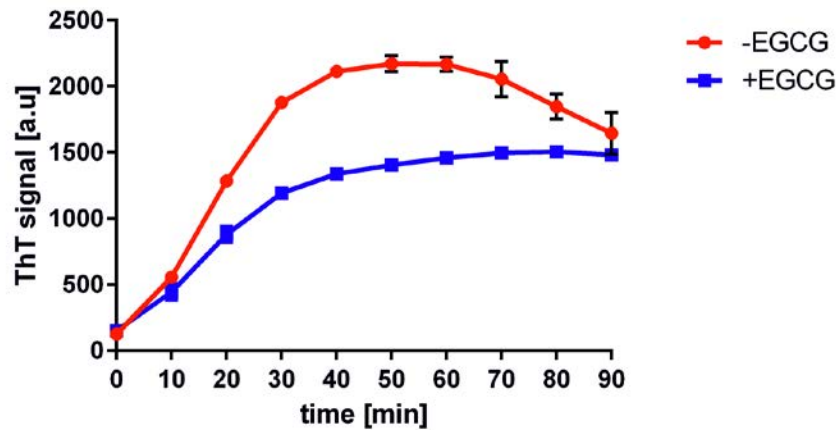


Figure S1 Heparin eliminates the lag phase of the tau aggregation reaction. Aggregation of K18 Δ K280 in the presence of heparin (Sigma) as assessed by ThT fluorescence assay (tau protein concentration: 12.5 μ M, heparin concentration 5 μ M) for 165 h (smaller diagram represents the first 4 hours of aggregation). Results represent mean fluorescence signal \pm s.d. (n = 3).

Supplementary Figure S2

A)



B)

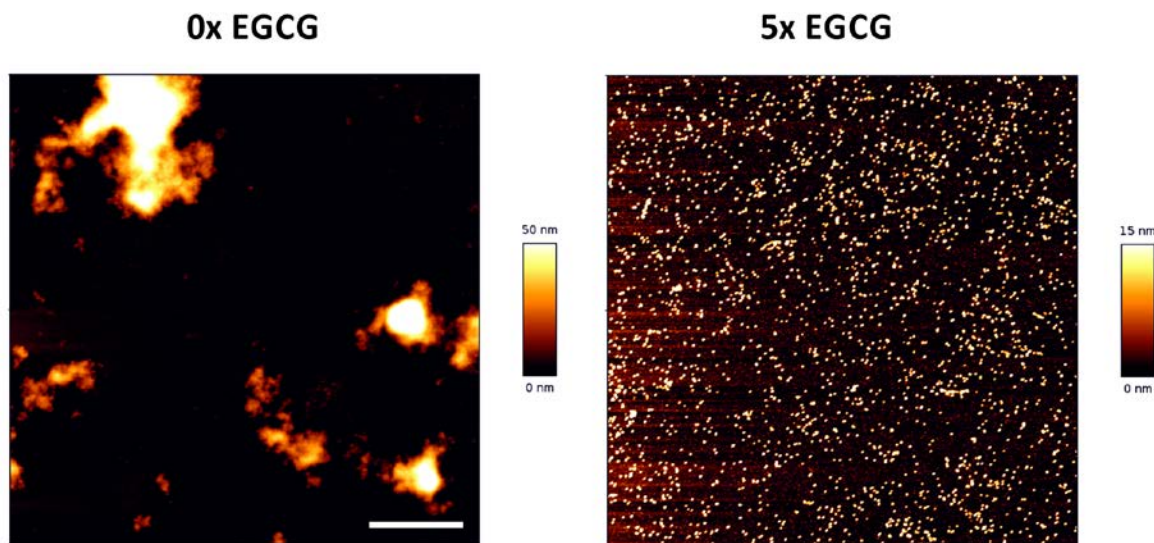


Figure S2 EGCG alters the aggregation kinetics of heparin-induced tau aggregation and alters the aggregate conformation. A) Aggregation of tau (12.5 μ M) in the presence of heparin (5 μ M) and EGCG (5x molar concentration) as assessed by ThT fluorescence. Results represent mean fluorescence signal \pm s.d. (n = 3). B) Analysis of EGCG-treated (5x molar concentration) K18 Δ K280 by atomic force microscopy after 3 h incubation. Samples were sonicated for 3 minutes in a water bath. Scale bar = 2 μ M.

Supplementary Figure S3

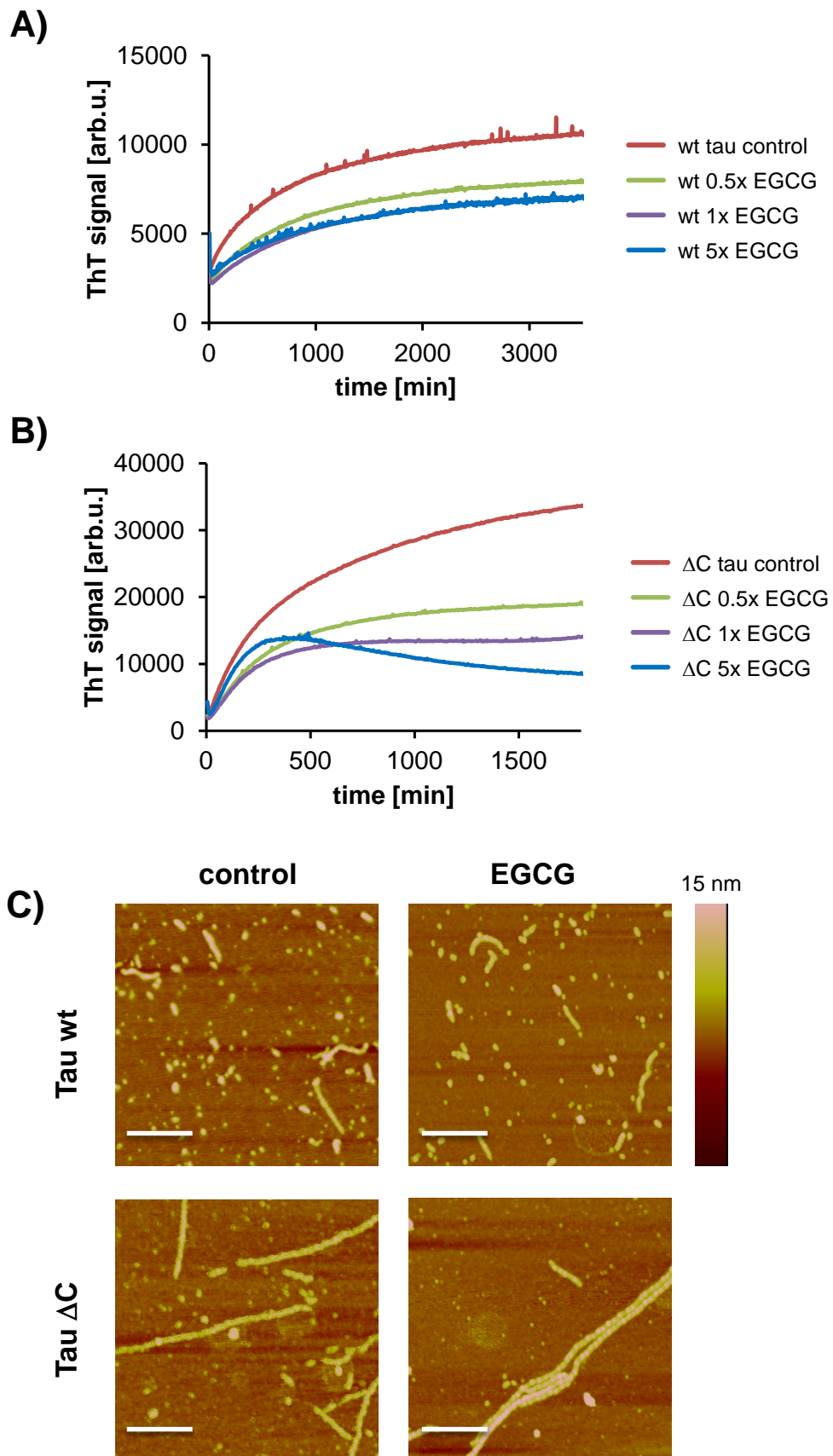


Figure S3 EGCG alters the aggregation kinetics of heparin-induced full length tau aggregation. A) Aggregation of wt full length tau (8 μ M) in the presence of heparin (8 μ M), DTT (5 mM) and EGCG (4 – 20 μ M, 0.5x - 5x tau concentration) under reducing conditions as assessed by ThT fluorescence. B) Aggregation of mutant C291A, C322A (Δ C) full length tau (8 μ M) in the presence of heparin (8 μ M) and EGCG (4 – 20 μ M, 0.5 - 5x tau concentration) as assessed by ThT fluorescence. Δ C tau aggregates under non-reducing conditions in the presence of heparin. C) Atomic force microscopy images of aggregation end points (72 h) of wt tau and DC tau aggregation assays shown in A), B) in the absence or presence of EGCG (40 μ M); scale bar 500 nm.