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Major differences between the self-assembly and seeding behavior of heparin-induced- and in-vitro-phosphorylated tau and their modulation by potential inhibitors

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ABSTRACT: Self-assembly of the microtubule-associated protein tau into neurotoxic oligomers, fibrils, and paired helical filaments, and cell-to-cell spreading of these pathological tau species are critical processes underlying the pathogenesis of Alzheimer's disease (AD) and other tauopathies. Modulating the self-assembly process and inhibiting formation and spreading of such toxic species are promising strategies for therapy development. A challenge in investigating tau self-assembly in vitro is that unlike most amyloidogenic proteins, tau does not aggregate in the absence of posttranslational modifications (PTM), aggregation inducers, or pre-formed seeds. The most common induction method is addition of polyanions, such as heparin, yet this artificial system may not represent adequately tau self-assembly in vivo, which is driven by aberrant phosphorylation and other PTMs, potentially leading to in-vitro data that do not reflect the behavior of tau and its interaction with modulators in vivo. To tackle these challenges, methods for in-vitro phosphorylation of tau to produce aggregation-competent forms recently have been introduced (*Despres et al., Identification of the Tau phosphorylation pattern that drives its aggregation, Proc. Natl. Acad. Sci. USA, 2017*). However, the oligomerization, seeding, and interaction with assembly modulators of the different forms of tau have not been studied to date. To address these knowledge gaps, we compared here side-by-side the self-assembly and seeding activity of heparin-induced tau with two forms of in-vitro phosphorylated tau and tested how the molecular tweezer CLR01, a negatively charged compound, affected these processes. Tau was phosphorylated by incubation either with activated extracellular signal-regulated kinase 2 or with a whole rat-brain extract. Seeding activity was measured using a FRET-based biosensor-cell method. We also used solution-state NMR to investigate the binding sites of CLR01 on tau and how they were impacted by phosphorylation. Our systematic structure-activity relationship study demonstrates that heparin-induced tau behaves differently from in-vitro phosphorylated tau. The aggregation rates of the different forms are distinct as is the intracellular localization of the induced aggregates, which resembles brain-derived tau strains suggesting that heparin-induced tau and in-vitro phosphorylated tau have different conformations, properties, and activities. CLR01 inhibits aggregation and seeding of both heparin-induced and in-vitro-phosphorylated tau dose-dependently, although heparin induction interferes with the interaction between CLR01 and tau.

Introduction

As one of the major molecular actors in Alzheimer's disease (AD) and other tauopathies, including frontotemporal dementia, progressive supranuclear palsy, corticobasal syndrome, Pick's disease, and chronic traumatic encephalopathy, the microtubule-associated protein tau has been investigated in the last decades as an attractive, yet challenging therapeutic target.¹⁻³ In humans, tau exists as six alternatively spliced isoforms, which normally are bound to microtubules and are thought to stabilize the microtubule structure. In tauopathies, tau detaches from the microtubules and self-assembles into neurotoxic oligomers and fibrillar aggregates, which are found in the brain as neurofibrillary tangles (NFTs) and other types of deposits.⁴

NFTs, one of the two characteristic proteinaceous lesions of AD,⁴ faithfully correlate with clinical progression of the disease, in contrast to the other lesion, amyloid plaques, which are poorly associated with cognitive decline.⁵⁻⁷ Tau pathology propagates through the brain in a reproducible temporal and regional manner.^{8,9} In fact, immunostaining using monoclonal antibody (mAb) AT8, which binds specifically to hyperphosphorylated tau, is used for AD staging.¹⁰ Nonetheless, as in other proteinopathies, metastable tau oligomers have been suggested to be a major neurotoxic agent and possibly the main culprit responsible for impairing and eventually killing susceptible neurons.¹¹

The main components of NFTs in AD are tau Paired Helical Filaments (PHF) consisting of two helically twisted ribbons of 10–20 nm width with an 80-nm periodicity. In these insoluble structures, tau transforms from a soluble, natively disordered conformation into cross- β structures, typical of amyloid, which encompass primarily its microtubule-binding domain (MTBD, residues 244–372).¹²⁻¹⁵ A recent study using cryo-electron microscopy on PHFs and straight filaments (SF) isolated from patients with AD has defined the structure of tau fibrils and highlighted a distinct structural arrangement in PHF and SF.¹⁶ In AD-brain fibrils, tau protofilaments adopt a cross- β / β -helix structure encompassing microtubule-binding repeats R3 and R4. The core of the fibrils is flanked by a “fuzzy coat” comprising the unfolded N- and C-terminal domains.^{17,18}

In AD, all six isoforms of tau are found abnormally hyperphosphorylated in PHFs¹⁹⁻²². The majority of the abnormal phosphorylation sites are distributed over ~40 positions clustered in the proline-rich region (PRR) and C-terminal domains flanking the MTBD.²²⁻²⁴ Hyperphosphorylation of tau in these regions is thought to cause its pathologic aggregation²⁵⁻²⁷ though the precise mechanism of this process remains elusive. Interestingly, certain phosphorylation sites, e.g., S262 in the MTBD and

S214 in the PRR, have been found to inhibit aggregation.²⁸⁻³⁰

Many studies have characterized the self-assembly process of tau in vitro. However, due to the practical difficulty of obtaining purified hyperphosphorylated or otherwise post-translationally modified tau, in-vitro aggregation of tau typically is studied using tau forms devoid of posttranslational modifications (PTMs), which do not aggregate spontaneously. Fragments of tau containing the MTBD or parts thereof are aggregation-prone and their fibrils may serve as a template to promote the aggregation of full-length tau,^{15,31-35} yet this strategy does not alleviate the concern about the lack of pathologically relevant PTMs. More commonly, in-vitro aggregation assays use external inducers, including polyanions, such as heparin,^{36,37} RNA,³⁸ fatty acids, such as arachidonic acid,^{39,40} or small anionic molecules⁴¹ that stimulate the aggregation process through mechanisms that are not fully understood.⁴²⁻⁴⁵ Although external aggregation inducers may lead to formation of tau fibrils that can be morphologically similar to PHFs,^{18,36} the aggregation process and structural details of these fibrils may be distinct from those of disease-relevant tau aggregates.^{16,46} Along these lines, recently, tau fibrils formed in vitro by addition of heparin or RNA were found to contain substantial amounts of the inducers and to disaggregate upon treatment with heparinase or RNase, respectively.⁴⁷ These findings are of particular concern in view of recent data demonstrating that conformationally distinct tau “strains” display substantially different biological and pathological behavior^{48,49} because they suggest that tau induced to aggregate by polyanions may not recapitulate adequately the structure of tau found in patient brains. In addition, the use of artificial aggregation inducers complicates the interpretation of experiments testing assembly modulators and inhibitors as potential therapeutic agents for tauopathies. An alternative approach is to produce in-vitro phosphorylated tau for studies of aggregation, structure, pathological effects, and modulation of all of these processes. However, to date, only a few studies have used this strategy and their focus was on the fibril formation and morphology.^{30,50,51} To our knowledge, tau oligomerization, the seeding activity of tau, and the inhibition of aggregation or seeding have not been studied using in-vitro phosphorylated recombinant tau to date.

As a test modulator, we chose in this study the molecular tweezer CLR01. Molecular tweezers are broad-spectrum modulators of the aberrant self-assembly process of amyloid proteins.^{52,53} They act by moderate, labile binding to Lys, and to a lower extent to Arg residues through inclusion of the side-chain in the cavity of the molecular tweezer, which leads to temporary masking of hydrophobic interactions in which these side chains participate and reversal of the positive charge of their end groups to a negative charge. This causes major perturbation of interactions involving Lys (or Arg), which play an important role in the abnormal self-assembly of amyloidogenic proteins. A lead molecular tweezer called CLR01, substituted by two negatively charged phosphate groups at the central benzene spacer-unit (Figure 4D), has

been shown to modulate the oligomerization and aggregation of over a dozen disease-associated amyloidogenic proteins and to be an effective inhibitor of the toxicity of these proteins in cell-culture and animal models of the associated diseases.^{52,53} However, relatively little is known about the interaction of molecular tweezers with tau.

Here, we conducted a systematic investigation of tau oligomerization, aggregation, and seeding, comparing heparin-induced tau with two forms of in-vitro phosphorylated tau. We also tested how CLR01 interacted with each tau form and characterized the binding sites of CLR01 on tau. Our data suggest that the oligomerization and aggregation kinetics of heparin-induced tau are different from those of in-vitro phosphorylated tau, and that these different forms lead to different tau strains that can be detected in biosensor cells, as described previously for brain-derived tau aggregates supporting the notion that these forms of tau fibrils are structurally distinct.⁵⁴ CLR01 interacts preferentially with Lys residues located in the MTBD and flanking regions and phosphorylation of tau has little effect on this interaction. CLR01 modulates the aggregation and seeding of the different forms of tau, yet in most experiments, the dose-response relationship of CLR01 with heparin-induced tau and in-vitro phosphorylated tau are different, supporting the notion that these forms of tau fibrils are distinct and suggesting that heparin induction interferes with the interaction of CLR01 with tau.

Materials and Methods

Heparin sodium salt from porcine intestinal mucosa was purchased from Sigma or Celsus Laboratories (Cincinnati, OH). Thioflavin T, okadaic acid, Pefabloc® and all other chemicals, unless indicated otherwise, were purchased from Sigma. For expression of isotopically labeled tau, bacterial culture media were reconstituted using ¹³C₆-glucose ¹⁵NH₄Cl, ¹⁵N/¹³C-ISOGRO®, and ¹⁵N₂,¹³C₆-L-lysine hydrochloride from Sigma-Aldrich (Isotec). Complete™ EDTA-free protease-inhibitor cocktail was purchased from Roche. BCA colorimetric assay was from Pierce. CLR01 was prepared as described previously⁵⁵ and purified as a sodium salt.

Production and purification of recombinant tau in isotopically labeled and unlabeled forms. The 2N4R tau isoform (residues 1-441 of human MAPT) in its wild-type or S262A form, and the fragment F4 (tau residues 208-324) carrying a C322S substitution were produced in the *E. coli* strain BL21(DE3) carrying a pET15b recombinant plasmid (Novagen). Residue numbering of F4 is based on

the 2N4R sequence. The wild-type mapt gene was inserted into NcoI/XhoI cloning sites. The S262A substitution was introduced by site-directed mutagenesis of the wild-type construct. For uniform ¹⁵N-labeling, cells were grown at 37 °C in M9 minimal medium containing 4 g/L glucose, 1 g/L ¹⁵N-NH₄Cl, 0.5 g/L ¹⁵N-Isogro® (Sigma), 1 mM MgSO₄, 100 μM CaCl₂, MEM vitamin cocktail, and 100 mg/L ampicillin. Expression was induced by addition of 0.5 mM isopropyl β-D-1-thiogalactopyranoside for 3 h at 37 °C. For selective ¹⁵N_α,¹³C-Lys and uniform ¹³C-labeling, the same growth medium was used except that it contained 1 g/L of unlabeled NH₄Cl and 2 g/L ¹³C₆-glucose, and 150 mg ¹⁵N₂,¹³C-Lys were added 20 min before induction. Cells were harvested by centrifugation at 5,000 × *g* for 30 min and the pellet was resuspended in 50 mM NaH₂PO₄/Na₂HPO₄, pH 6.2, 2.5 mM EDTA, 2 mM DTT and 0.5% Triton X-100 supplemented with a Complete™ protease-inhibitor cocktail. The lysate was obtained by homogenizing this suspension using a high-pressure homogenizer followed by centrifugation at 30,000 × *g* for 30 min. The extract was incubated at 75 °C for 15 min to precipitate most of the bacterial proteins from the soluble extract as a first purification step. The soluble proteins were separated from the precipitate by centrifugation at 4,000 × *g* for 20 min and the desired protein was purified by cation-exchange chromatography (HiTrap SP HP 5 ml, GE Healthcare). The column was equilibrated in 50 mM NaH₂PO₄/Na₂HPO₄, pH 6.4, 2 mM EDTA (buffer A). The bacterial extract was loaded onto the column and then the column was washed with buffer A. The protein was eluted with a gradient from 0% to 50% buffer B (buffer A supplemented with 2M NaCl) over 20 mL (i.e., 4 column volumes). Elution fractions were analyzed by SDS-PAGE and fractions containing the pure protein were pooled together for buffer-exchange in 50 mM ammonium bicarbonate (HiPrep 26/10 desalting, GE Healthcare) prior to lyophilization. Lyophilized proteins were stored at -20 °C or -80 °C until further use.⁵⁶ The recombinant tau proteins used in this study are listed in Table 1 with an indication of phosphorylation sites provided by the kinase activity of either activated ERK2 (ERK2*) or a rat-brain extract (RBE). These phosphorylation patterns are consistent with those described previously.^{29,30}

In vitro phosphorylation of tau by activated ERK2. Recombinant ERK2 and a constitutively active MEK3 variant, a mitogen-activated protein-kinase kinase that phosphorylates and activates ERK2, were produced as described previously.^{29,56,57} ¹⁵N- or ¹⁵N₂, ¹³C-Lys/¹³C-tau were dissolved at 100 μM with 1 μM of ERK2 and ~0.1 μM of MEK3 in 400 μl of buffer containing 50 mM HEPES-KOH, pH 8.0, 12.5 mM MgCl₂, 50 mM NaCl, 2 mM EDTA, 1 mM DTT, 1 mM EGTA, and 12.5 mM ATP. The mixtures were

Table 1. Tau variants used in this study.

Tau proteins	Residues	Phosphorylation sites
F4	208-324	
Tau	1-441	

Phospho-tau	1-441	Phosphorylated by ERK2* (p-tau ^{ERK})	S46, T50, T69, T153, T175, T181, S191, S199, S202, T205, T231, S235, S396, S404, S422
Phospho-tau, S262A	1-441, S262A	Phosphorylated by RBE ¹ (p-tau(S262A) ^{RBE})	S46, T50, T69, T153, T175, T181, S191, S199, S202, T205, <u>S208</u> , T231, S235, <u>S356</u> , S396, S404, S422

¹ The underlined residues correspond to phosphorylation sites uniquely found using the kinase activity in the rat brain extract.

incubated at 37 °C overnight. Enzymatic reactions were stopped by incubating the reaction mixtures at 75 °C for 15 min followed by centrifugation at 16,000 × *g* for 20 min. Then, the supernates were buffer-exchanged in 50 mM ammonium bicarbonate and lyophilized. Prior to further analyses, phosphorylation was checked qualitatively by a shift in the apparent mobility of the protein band on SDS-PAGE. Then, the phosphorylated proteins were analyzed by high-resolution NMR spectroscopy to determine the phosphorylation patterns and potential protein degradation in a quantitative manner.^{29,57}

In vitro phosphorylation of tau by rat-brain extract (RBE). RBE was prepared from adult Sprague-Dawley rats by homogenizing whole brains (~2 g) in 5 mL of homogenizing buffer, containing 10 mM Tris-HCl, pH 7.4, 5 mM EGTA, 2 mM DTT, 1 μM okadaic acid (OA), 20 μg/mL Leupeptin, and 40 mM Pefabloc®. Insoluble material was precipitated by ultracentrifugation at 100,000 × *g* and 4 °C for 1 h. The supernate was directly used for its kinase activity. Total protein concentration was estimated at 11 mg/mL by a BCA colorimetric assay. ¹⁵N,¹³C tau-S262A was dissolved at 10 μM in 2.5 mL of phosphorylation buffer, containing 40 mM HEPES, pH 7.3, 2 mM MgCl₂, 5 mM EGTA, 2 mM DTT, 2 mM ATP, and 1 μM OA, supplemented with a Complete™ protease-inhibitor cocktail. The protein was incubated with 500 μL of brain extract at 37 °C for 24 h. The enzymatic reaction was stopped by heating the mixture at 75 °C for 15 min followed by centrifugation at 16,000 × *g* for 20 min. Then, the supernate was buffer-exchanged in 50 mM ammonium bicarbonate and the protein was lyophilized. Qualitative evaluation of protein phosphorylation was performed by SDS-PAGE and then the phosphorylation sites were determined by NMR analyses, as described above.

In vitro aggregation assay and transmission electron microscopy (TEM). For aggregation assays, unphosphorylated tau was mixed with heparin at a ratio of 4:1, respectively, in a buffer containing 25 mM NaH₂PO₄/Na₂HPO₄, pH 6.6, 25 mM NaCl, 2.5 mM EDTA, and 0.33 mM DTT at a final protein concentration of 5 μM. Reactions carried out in the absence of heparin were used as a negative control. ThT was prepared freshly at 5 mM and added at a final concentration of 50 μM before measurements. ThT emission was detected at 490 nm after excitation at 440 nm in 96-well plates on a plate reader (PHERAStar, BMG LABTECH GmbH, Ortenberg, Germany). Kinetic curves were fitted to a plateau followed by a single exponential association equation (Equation 1).

$$Y = Y_0 + (Y_{\max} - Y_0)(1 - e^{-k(t - t_0)}) \quad \text{Equation (1)}$$

where *t*₀ is the lag-time when nucleation occurs, *k* the association rate constant, *Y*₀ the value of the plateau at the initial step when *t* ≤ *t*₀, *Y*_{max} the maximal value of the plateau at the end of the elongation step. Inhibition of aggregation by CLR01 was tested under the same conditions by diluting a 10 mM stock solution of the molecular tweezer in triply-distilled water into the reaction mixture at different concentrations. Concentrations of CLR01 ranging from 1.25 to 20 μM were evaluated by 20-min-step kinetics for a total of 92 h.

IC₅₀ values were derived by fitting the percentage of inhibition as a function of CLR01 concentration (Equation 2).

$$\% \text{ inhibition} = \frac{k_0 - k}{k_0} \quad \text{Equation (2)}$$

Where *k* is the association rate constants at a given concentration of CLR01 and *k*₀ is the rate constant in the absence of CLR01.

After completion of the aggregation, the morphology of the protein alone or of Tau:CLR01 mixtures was examined by TEM. For these experiments, 10-μL aliquots of the aggregation reactions were applied to 400-meshed hexagonal Formvar-coated copper grids for 90 s. The liquid was wicked off using a filter paper and the grids were washed thrice with ultrapure water, stained with 10 μL of 2% uranyl acetate for 60 s, and wicked off using a filter paper. TEM was performed using a HITACHI H7500 microscope at 80 kV.

Aggregation assays using tau phosphorylated by RBE (p-Tau^{RBE}, p-tau(S262A)^{RBE}) were performed at 30 μM of the protein in a buffer containing 100 mM 2-(*N*-morpholino)ethanesulfonic acid, pH 6.9, 2 mM EGTA, 1 mM MgCl₂, 20 mM NaF, 0.33 mM DTT and 50 μM ThT. p-tau(S262A)^{RBE} was incubated without agitation for 5 d at 37 °C in the absence or presence of 7.5, 15, 22.5, 30, 60, or 120 μM CLR01 and aggregation kinetics were followed by ThT fluorescence. At the end of incubation, samples were analyzed by TEM as described above.

Preparation of tau seeds. To evaluate the seeding capability of tau, the different variants either were freshly dissolved and tested immediately or incubated at 37 °C for 5–10 days to form fibrils before addition to biosensor cells. Unphosphorylated tau was incubated in the presence of heparin or in the absence of heparin as a negative control, whereas tau phosphorylated by ERK2 (p-Tau^{ERK}) or p-tau(S262A)^{RBE} were incubated in the absence of heparin.

Before preparing transduction complexes, the proteins were sonicated for 10 min in a bath sonicator. Transduction complexes were made by combining a solution of 8.75 μ L Opti-MEM (Gibco) and 1.25 μ L Lipofectamine 2000 (Invitrogen) with a solution/suspension of tau seeds in Opti-MEM at the desired concentration, determined using the BCA colorimetric assay for a total volume of 20 μ L per well. The concentration was calculated based on tau monomer. Transduction complexes were incubated at room temperature for 20 min before adding to cells.

Transduction of tau seeds in biosensor cells. A monoclonal, fluorescence-resonance energy transfer (FRET)-based biosensor cell line⁵⁸ was maintained as described in instructions from ATCC (CRL-3275). Briefly, the cells were maintained in Dulbecco's Modified Eagle's Medium (DMEM, ATCC-30-2002) supplemented with 2 mM L-alanyl-L-glutamine and 10% fetal bovine serum (FBS) under a 5% CO₂ atmosphere at 37 °C. These cells stably express the tau MTBD containing the disease-associated substitution P301S and fused to either cyan fluorescent protein (CFP) or yellow fluorescent protein (YFP).⁵⁸ The cells were plated at 35,000 cells per well in 96-well plates (Sigma-Aldrich). Eighteen hours later, at ~60% confluency, the cells were transduced with tau seeds and incubated with the seeds for 24–48 h prior to flow cytometry analysis.

FRET Flow Cytometry. Cells were incubated in 50 μ L TrypLE (Cat. 12605010, ThermoFisher) per well for 5 min at 37 °C followed by addition of 150 μ L DMEM supplemented with 2 mM L-alanyl-L-glutamine and 10% fetal bovine serum to stop the reaction. The cells were transferred to a round-bottom 96-well plate (Cat. 07-200-89, ThermoFisher), centrifuged at 400 *g* for 5 min, and then resuspended in 200- μ L flow-cytometry buffer comprising 1% FBS and 1-mM EDTA in Hank's Balanced Salt Solution (GIBCO). A SORP BD Biosciences HTLSRII Analytic Flow Cytometer was used for FRET flow-cytometry measurements. To measure CFP and FRET, cells were excited using a 405-nm laser, and fluorescence was captured using 405/50-nm and 525/50-nm filters, respectively. To measure YFP, cells were excited using a 488-nm laser and fluorescence was captured with a 525/50-nm filter. To quantify FRET, we used a gating strategy similar to that described previously.⁵⁸ Briefly, CFP bleed-through into the YFP and FRET channels was compensated for using FACSDiva 8.0.1 software from BD Biosciences. Because some YFP-only cells exhibit emission in the FRET channel, we introduced an additional gate to exclude from analysis cells that generate a false-positive signal in the FRET channel (i.e., false FRET gate). Subsequently, we created a final bivariate plot of FRET vs. CFP and introduced a triangular gate to assess the number of FRET-positive cells. This FRET gate was adjusted to biosensor cells treated with lipofectamine but no tau seeds, which are thus FRET-negative. This allows for direct visualization of sensitized acceptor emission arising from excitation of the CFP donor at 405 nm. The integrated FRET density, defined as the percentage of FRET-positive cells multiplied by the median fluorescence intensity of

FRET-positive cells, was used for all analyses. For each experiment, 20,000 cells per replicate were analyzed and each condition was analyzed in quadruplicate. Data analysis was performed using Flow Jo v10 (Treestar).

Native PAGE/western blots. Recombinant protein aliquots from seeding experiments were mixed with 4X NativePAGE sample buffer, containing 50 mM Bis-Tris, 6 N HCl, 50 mM NaCl, 10% v/v glycerol, and 0.001% w/v Ponceau S, pH 7.2. Ten- μ L sample per well were loaded onto Novex NativePAGE 3-12%-gradient Bis-Tris gels (Thermo Fisher). The proteins were fractionated for 90-115 min at 150 V constant voltage using a XCell SureLock Mini-Cell system (ThermoFisher). The proteins then were transferred to a PVDF membrane for 1 h at 25 V constant voltage using a XCell II™ Blot Module (ThermoFisher). Membranes then were incubated in 20 mL of 8% acetic acid for 15 min to fix the proteins. The blots were probed with anti-human tau mAb HT7 diluted 1:1,000 in 5% (v/v) non-fat dry milk in TBST. ECL Plus (GE Healthcare) was used to visualize the bands in a Syngene PXi multi-application gel imaging system and band intensity was quantified densitometrically using Image J.^{59,60}

Surface Plasmon Resonance. Biotinylated heparin was prepared by reaction with amine-PEG₃-Biotin (Pierce, Rockford, IL) in the presence of NaCNBH₃ as described previously.⁶¹ The biotinylated heparin was immobilized on a streptavidin (SA) chip (GE Healthcare) based on the manufacturer's protocol. Briefly, 20 μ L of 0.1 mg/mL of the biotinylated heparin in HBS-EP running buffer (0.01 M HEPES, 0.15 M NaCl, 3 mM EDTA, and 0.005% surfactant P20, pH 7.4) were injected over flow cells 2, 3 and 4 of the SA chip at a flow rate of 10 μ L/min. The successful immobilization of heparin was confirmed by the observation of a ~200-resonance-unit (RU) increase in the sensor chip. The control flow cell was prepared by a 1-min injection of saturated biotin. Tau was dissolved in HBS-EP buffer and different dilutions of the protein solution were injected at a flow rate of 30 μ L/min. At the end of the sample injection, the same buffer was flowed over the sensor surface to facilitate dissociation. After a 3-min dissociation period, the sensor surface was regenerated by injection of 30 μ L of 2 M NaCl. The response sensograms were monitored as a function of time at 25 °C. For testing the inhibition of the tau-heparin interaction by CLR01, 500 nM of tau were mixed with 0, 5, 12.5, 25, or 50 μ M CLR01. Then, each mixture was injected over the heparin chip at a flow-rate of 30 μ L/min. At the end of the sample injection, the same buffer was flowed over the sensor surface to facilitate dissociation and the sensor surface was regenerated as described above. The highest response was obtained for calculation of normalized binding inhibition.

NMR spectroscopy. Spectra were acquired at 293 K on a Bruker 900-MHz spectrometer equipped with a triple-resonance cryogenic probe head. ¹⁵N-Lys-labeled tau was dissolved at 200 μ M in NMR sample buffer, containing 50 mM NaH₂PO₄/Na₂HPO₄, pH 6.4, 25 mM NaCl, 2.5 mM EDTA, 1 mM DTT and 10% D₂O in 3-mm NMR tubes. The proton frequency was calibrated with 1 mM sodium 3-(trimethylsilyl)propionate-2,2,3,3-d₄ as a reference. ¹H-¹⁵N

HSQC spectra were recorded for ^{15}N - and ^{15}N , ^{13}C -labeled samples with 64 scans per increment, and 3,072 and 512 points in the ^1H and ^{15}N dimensions, respectively.

The phosphorylation patterns of p-tau^{ERK}, p-tau^{RBE}, and p-tau(S262A)^{RBE} were analyzed as described previously.^{56,57} The patterns (Table 1) were similar to those characterized previously by Despres et al.³⁰ and Qi et al.²⁹ For ^{15}N , ^{13}C -Lys, ^{13}C -labeled p-tau assignment, a three-dimensional HN(CO)CACB experiment was performed with 16 scans per increment. Spectral widths were 16.0, 23.0, and 61.0 ppm in the ^1H , ^{15}N and ^{13}C dimensions with 2,048, 64 and 120 sample points, respectively.

For the NMR mapping of protein resonance intensities upon CLR01 binding, the intensity of each resonance in the ^1H - ^{15}N HSQC spectrum in the presence of CLR01 (I) was determined and compared to the resonance intensity in the absence of CLR01 (I_0). The normalized intensities I/I_0 were plotted as a function of the protein sequence.

For aggregation assays, protein concentrations were normalized based on their 1D proton spectra (zgpgpw5, Bruker) acquired at 600 MHz with 64 scans, 16 dummy scans and 16,384 points for a spectral width of 13.97 ppm centered on 4.70 ppm.

Results

Comparison of heparin-induced and in-vitro phosphorylated tau self-assembly. To establish baseline conditions for comparison of in-vitro phosphorylated tau with heparin-induced tau, we incubated a mixture of 5 μM tau and 1.25 μM heparin at 37 $^\circ\text{C}$ and measured ThT fluorescence up to 100 h. As expected, we observed a typical sigmoidal increase in ThT fluorescence, beginning with a short lag phase (~ 30 min), followed by a sharp increase in fluorescence over the next 6 h and then a slower increase up to a plateau at ~ 12 h (Figure 1A). We derived the aggregation rate constant, k , by fitting the kinetic curves to a one-phase association equation (Equation 1). Under these conditions, k was found to be $0.29 \pm 0.02 \text{ h}^{-1}$. Examination of the morphology at the end of the reaction by TEM showed abundant, relatively short fibrils (Figure 1B).

Despite site-specific, high-level phosphorylation at several sites, p-tau^{ERK} has been reported previously not to produce a detectable increase of the ThT fluorescence signal.^{29,30} In agreement with these previous observations, solutions of 100 μM p-tau^{ERK} did not aggregate up to 100 h (Figure 1A). TEM examination showed only non-fibrillar, amorphous aggregates (Figure 1C) as observed previously.^{29,30}

The quantitative phosphorylation pattern of p-tau^{RBE}, as determined by NMR, has been reported to be similar to that of p-tau^{ERK}, but contains three additional phosphorylation sites—at S208 in the PRR, and at S262 and S356 in the MTBD (Table 1).²⁹ Because phosphorylation of S262 is known to inhibit tau aggregation,²⁸ S262 was substituted by Ala for generation

of p-tau(S262A)^{RBE} to prevent phosphorylation at this site. At 5–10 μM , the aggregation of p-tau(S262A)^{RBE} was too slow to compare side-by-side with heparin-induced tau. However, at 30 μM , p-tau(S262A)^{RBE} reactions showed a sigmoidal increase in ThT fluorescence, beginning with a short lag phase of ~ 1 h, followed by an increase in fluorescence over 45 h to reach a plateau. Despite using a higher concentration, the aggregation kinetics of p-tau(S262A)^{RBE} was substantially slower than that of heparin-induced unphosphorylated tau (Figure 1A). The rate constant, k , in this case was $0.066 \pm 0.001 \text{ h}^{-1}$. Examination of the fibril morphology at the end of the reaction by TEM showed abundant fibrils, which were thicker and longer than those of heparin-induced tau (Figure 1D, Supplementary Figure S1). As shown previously,³⁰ the fibrils had both straight and squiggly morphology but did not show a helical twist (Supplementary Figure S1).

We asked next whether intermediate oligomers forming during the aggregation process could be detected and how they compared among the three forms of tau. We addressed this question by analyzing aliquots from aggregation mixtures containing 10 μM of each tau form at 0, 10, and 20 h using native-PAGE/western blot probed with the anti-human-tau mAb HT7. Interestingly, we found a distinct behavior for each form (Figure 1E). At $t = 0$ h, i.e., just after addition of heparin to unphosphorylated tau, the protein appeared as a smear over nearly the entire length of the gel. A band at ~ 70 kDa, presumably a monomer, could be discerned, as well as a lower band, at ~ 60 kDa, likely representing a truncation/degradation product. At $t = 10$ h, most of the protein seemed to “disappear,” likely due to aggregation preventing it from entering the gel, and only relatively faint smears with apparent mobility corresponding to 140–1,000 kDa were observed. At $t = 20$ h relatively little protein was apparent suggesting further aggregation. At this time point, possibly due to the relatively low amount of apparent protein, faint smeary bands corresponding to putative monomer through tetramer were observed.

In agreement with the lack of observed fibril formation in the case of p-tau^{ERK}, this protein showed several bands, including putative monomer through tetramer, a higher molecular weight smear at 800–1,000 kDa, and a band at $\sim 1,100$ kDa, likely a high-molecular-weight oligomer. At 10 and 20 h, the putative trimer, tetramer, and high-molecular-weight smear at 800–1,000 kDa seemed to convert to the high-molecular-weight oligomer at $\sim 1,100$ kDa, whereas little or no change was observed in the putative monomer and dimer abundance.

The behavior of p-tau(S262A)^{RBE}, which unlike p-tau^{ERK} did form β -sheet-rich fibrils, was more similar to that of heparin-induced unphosphorylated tau than to p-tau^{ERK}. At $t = 0$ h, the appearance was smeary with discernable putative monomer and dimer bands that migrated slightly slower than their unphosphorylated counterparts, whereas at 10 and 20 h, most of the signal gradually disappeared and lower molecular weight species appeared more abundant at 20 h than at 10 h. In all cases, the

solutions/suspensions were mixed thoroughly before loading on the gel to ensure that potential fibrils/aggregates precipitated at the bottom of the tube would be included. However, these large species apparently were too large to enter the gel and therefore were not observed. Alternatively, they may have been present but did not react with HT7.

Comparison of heparin-induced and in-vitro phosphorylated tau seeding in biosensor cells. Toxic tau species are thought to pass from affected cells to synaptically connected unaffected cells, where the seeds serve as templates that corrupt normal tau proteins and cause them to aggregate.^{62,63} To facilitate characterization of the seeds, Diamond's group has developed biosensor cells that allow detecting and quantifying tau seeding using a sensitive FRET assay.⁶⁴ The assay has been shown to have a wide dynamic range, allowing comparison of samples with vastly different levels of tau seeding and distinguishing among samples showing subtle differences.⁶³

Here, we used this assay to compare the seeding capability of the three tau variants. We began with testing the seeding activity of fibrils formed by heparin-induced tau or p-tau(S262A)^{RBE}, which were sheered by sonication for 10

min and mixed with the transfection agent, lipofectamine, for effective transduction into the biosensor cells, as described previously.⁶⁴ We examined the cells qualitatively by fluorescence microscopy for the presence of observable puncta and quantitatively for the CFP-YFP FRET signal using flow cytometry.

In untreated cells, only diffuse fluorescence was observed (Figure 2A), whereas bright puncta were detected following addition of either type of fibrils (Figure 2B, C) indicating seeding of tau aggregation. Quantitative analysis by flow-cytometry showed that both forms induced a robust seeding response. The magnitude of the response was higher overall for heparin-induced tau, though the difference reached statistical significance only at the highest concentration (1 μ M, Figure 2D). Interestingly, high-magnification examination revealed distinct morphology and localization of the aggregates. Aggregates induced by heparin-induced tau were larger and mainly cytosolic or perinuclear (Figure 2E), whereas those induced by p-tau(S262A)^{RBE} were smaller and co-localized with nuclei. These observations suggest that although both forms of tau form ThT-positive fibrils, the arrangements of the tau molecules in these fibrils are different and represent distinct strains based on their seeding activity and aggregate morphology in the biosensor cells.

Figure 1. Self-assembly of tau induced by heparin, p-tau^{ERK} and p-tau(S262A)^{RBE}. (A) Time-dependent ThT fluorescence of 5 μ M heparin-induced tau, 100 μ M p-tau^{ERK}, or 30 μ M p-tau(S262A)^{RBE}. (B-D) Morphological examination by TEM at the end of the aggregation reaction (t = 100 h) of heparin-induced tau (B), p-tau^{ERK} (C) and p-tau(S262A)^{RBE} (D). E) Native-PAGE analysis of 10 μ M each of heparin (hep)-induced tau, p-tau^{ERK}, and p-tau(S262A)^{RBE} at t = 0, 10, or 20 h of incubation under the same conditions. For heparin-induced tau, t = 0 is just before addition of heparin. Molecular weight markers are shown on the left. The mobilities of putative monomer (M), dimer (D), trimer (Tr), and tetramer (Te) are indicated on the right.

Because previously, both tau fibrils and oligomers as small as a trimer were shown to act as seeds,⁶⁴ we tested also whether the three tau variants could seed tau aggregation immediately after dissolution, when presumably only monomers and putative oligomers are present. Cells treated with freshly dissolved unphosphorylated tau in the absence (Figure 3A) or presence of heparin (Figure 3B) did not show formation of puncta. The latter condition produced a minimal FRET signal (Figure 3E). In contrast, freshly dissolved 1- μ M p-tau^{ERK} showed abundant bright puncta (Figure 3C) and induced a strong, dose-dependent seeding response (Figure 3E). Similarly, freshly dissolved 1- μ M p-tau(S262A)^{RBE} promoted formation of bright puncta (Figure 3D) and caused dose-dependent tau aggregation in the biosensor cells, albeit slightly weaker than p-tau^{ERK} (Figure 3E). Overall, the seeding activity of the freshly dissolved phosphorylated-tau proteins was ~4-fold weaker than that of the fibrils (*cf.* Figure 2D and 3E). These data suggested that unlike unphosphorylated tau, p-tau^{ERK} and p-tau(S262A)^{RBE} contained seed-competent tau monomers or pre-formed seeds, or formed seeds, presumably oligomers, during the experiment.⁶⁵ High-resolution images showed that the aggregate morphology and localization of freshly dissolved p-tau^{ERK} (Figure 3F)

and p-tau(S262A)^{RBE} (Figure 3G) resembled those of fibrillar p-tau(S262A)^{RBE} (Figure 2F), i.e., were small relative to the cell body and co-localized partially with nuclei, but not of heparin-induced tau fibrils (Figure 2E).

Formation of a small population of aberrant, cross-linked tau dimers via an intermolecular disulfide bond has been reported previously to promote seeding by the FTD-linked P301L tau isoform.⁶⁶ To test whether the seeding in our experiments was due to formation of disulfide-crosslinked dimers (or other oligomers), we reduced the protein by addition of 6-fold excess DTT and repeated the seeding measurements. The final concentration of DTT in the cell culture media, 6 μ M, did not cause any apparent change to the cells' morphology or viability. The measurements did not show significant differences between the seeding of either tau variant in the presence of DTT (Figure 3E), suggesting that aberrant intermolecular disulfide-mediated oligomerization did not contribute to the observed seeding.

Taken together, the results suggest that unphosphorylated tau needs to be induced to form fibrils in order to become

Figure 2. Induction of tau seeding by aggregated tau in biosensor cells. A–C) Fluorescence microscopy images of tau-biosensor cells. A) Negative control cells incubated with media alone. B) Cells incubated with heparin-induced tau fibrils. C) Cells incubated with p-tau(S262A)^{RBE} fibrils. D) Dose-response analysis of the seeding activity (Integrated FRET Density) of each tau form. E,F) High-magnification fluorescence microscopy images of cells incubated with 1 μ M of either heparin-induced tau fibrils (E) or p-tau(S262A)^{RBE} fibrils (F). Bright YFP fluorescence indicates tau aggregates. Nuclei were stained with 4',6-diamidino-2-phenylindole (DAPI, blue).

Figure 3. Induction of tau seeding by freshly dissolved tau in biosensor cells. A–D) Fluorescence microscopy images of tau-biosensor cells incubated with 1 μ M freshly dissolved tau (A), tau + heparin (B), p-tau^{ERK} (C), or p-tau(S262A)^{RBE} (D). E) Dose-response analysis of the seeding activity (Integrated FRET Density) of each tau form in the absence or presence of DTT. F,G) High-magnification fluorescence microscopy images of cells incubated with 300 nM freshly dissolved p-tau^{ERK} (F) or p-tau(S262A)^{RBE} (G). Bright YFP fluorescence indicates tau aggregates. Nuclei were stained with DAPI (blue).

seeding-competent. In contrast, in-vitro phosphorylated tau is seeding-competent already when freshly dissolved, presumably due to the presence of oligomers (Figure 1E). However, upon fibril formation, the seeding activity of p-tau(S262A)^{RBE} increases by ~4-fold. p-tau^{ERK} and p-tau(S262A)^{RBE} appear to form a similar strain, which is different from the one formed by heparin-induced tau. The strain formed by p-tau(S262A)^{RBE} seems to be maintained in both the freshly dissolved and fibrillar states.

CLR01 inhibits the aggregation of heparin-induced and in-vitro phosphorylated tau with distinct dose-dependencies. Similar to anionic aggregation inducers, such as heparin or arachidonic acid, CLR01 binds to positively charged residues in tau. This competition for the same binding sites makes data interpretation difficult because CLR01 binding could inhibit tau aggregation by two independent mechanisms: 1) competition with the inducer for the Lys/Arg residues, thereby attenuating the induction of tau aggregation; and 2) direct interference with intermolecular interactions among tau molecules. Therefore, we sought to distinguish between these two mechanisms and to compare between this non-ideal system and a simpler system in which CLR01 interacts with phosphorylated tau.

To determine the extent to which CLR01 competes with heparin for tau binding, we used SPR. Heparin was immobilized on a SPR chip and tau binding to the immobilized heparin was found to occur with $K_D = 10.9$ nM (Figure 4A), in agreement with recent results obtained using ELISA.⁶⁷ Similar experiments using solutions in which tau was mixed with CLR01 at different ratios then were performed, with the expectation that when the heparin-binding sites in tau are occupied by CLR01, tau binding to the surface-immobilized heparin should decrease, resulting in a reduction in the magnitude of the SPR signal. This well-established method has been used previously for characterizing heparin-protein interaction.^{68,69} The analysis showed that CLR01 competed with heparin for binding to tau, but in agreement with its highly labile binding, was a relatively weak competitor. The IC_{50} for the inhibition of 0.5 μ M tau binding to immobilized heparin by CLR01 was found to be 20 ± 1 μ M

(Figure 4C), i.e., 40-fold excess CLR01 relative to tau was needed for half-maximal inhibition.

Next, to test the effect of CLR01 on heparin-induced aggregation of tau, the unphosphorylated protein was incubated at 5 μ M in the presence of 1.25 μ M heparin and CLR01 was added at concentrations ranging from 2.5 to 40 μ M. CLR01 caused a dose-dependent decrease in the aggregation rate constant, k , and inhibited the aggregation completely at 6-fold excess. (Figure 5A). A plot of percentage of inhibition versus [CLR01] showed that the molecular tweezer inhibited the aggregation of heparin-induced tau with a half-maximal inhibitory concentration (IC_{50}) of 11.3 ± 0.6 μ M calculated using equation 1 (Figure 5B). Considering that both primary and secondary nucleation events may affect tau aggregation, we also attempted to use the method of Cohen et al.,^{70,71} which takes into account both nucleation mechanisms. We found little or no contribution of secondary nucleation, minimal changes in the data fitting, and a nearly identical IC_{50} value – 11.8 ± 0.7 μ M, using this method. Because inhibition of aggregation does not always produce the same effect on the slope of the ThT curve and the final plateau value, we calculated an IC_{50} value also based on the final plateau and found it to be in the same range, 19 ± 0.6 μ M (Supplementary Figure S2A). In this experiment, only ~3-fold excess CLR01 were sufficient for half-maximal inhibition, as opposed to the 40-fold excess needed for half-maximal inhibition of tau binding to heparin. Therefore, we conclude that the main inhibitory effect of CLR01 on tau aggregation is mediated by direct interference of tau self-assembly rather than by inhibition of heparin binding.

Due to the different aggregation kinetics of heparin-induced tau and p-tau(S262A)^{RBE} (Figure 1A), side-by-side comparison at the same concentration was not practical. Thus, to measure the effect of CLR01 on the aggregation of p-tau(S262A)^{RBE}, the protein was incubated at 30 μ M in the presence of increasing CLR01 concentrations. Interestingly, at substoichiometric concentrations, CLR01 appeared to have a mixed effect on the aggregation kinetics of p-tau(S262A)^{RBE}—initially, the kinetics was accelerated whereas after ~24 h, the aggregation rate was reduced compared to reactions in the absence of CLR01

(Figure 5C,D and Supplementary Figure S2B). In contrast, at 2- or 4-fold excess, CLR01 inhibited the aggregation completely (Figure 5C,D). These results further demonstrate that heparin-induced tau aggregation is dissimilar to the aggregation of hyperphosphorylated tau and in particular may yield misleading results when analyzing aggregation modulators, such as CLR01.

ThT fluorescence measures the formation of the cross- β structure when amyloidogenic proteins aggregate but the increase in ThT fluorescence does not always correlate with fibril formation.^{72–74} To assess whether in this case the initial acceleration in β -sheet formation reflected formation of fibrils, we examined the morphology of each preparation at the end of the aggregation reaction by TEM. The morphological examination showed that addition of CLR01 resulted in a dose-dependent inhibition of fibril

formation by both heparin-induced tau and p-tau(S262A)^{RBE} (Figure 6, Supplementary Figure S3). In the case of heparin-induced tau, thin, straight or squiggly fibrils could still be observed up to a tau:CLR01 concentration ratio of 1:4, respectively, whereas at higher concentrations the morphology was amorphous (Figure 6A). Fibrils of p-tau(S262A)^{RBE} were substantially reduced at a p-tau(S262A)^{RBE}:CLR01 concentration ratio 1:0.25, and eliminated at concentration ratio 1:0.5 or higher (Figure 6B), demonstrating that the acceleration of the ThT fluorescence increase at substoichiometric CLR01 concentrations did not reflect fibril formation. These data are in agreement with an increase in ThT fluorescence without fibril formation by other proteins in the presence of CLR01,^{75,76} yet the nature of the early ThT-positive structures forming in the presence of CLR01 is not well-understood.

Figure 4. Inhibition of tau binding to heparin by CLR01. Binding of tau to immobilized heparin and its inhibition by CLR01 were measured using SPR. A) A schematic structure of CLR01. The compound is partially protonated at physiologic pH. B) SPR sensorgram of tau binding to heparin immobilized on a SPR chip. C) Normalized tau binding in the presence of increasing concentrations of CLR01. D) % Inhibition versus CLR01 concentration used for calculating the half-maximal inhibition (IC_{50}).

CLR01 inhibits tau seeding in biosensor cells. Next, we asked whether CLR01 could inhibit the seeding activity of tau fibrils. To answer the question, we incubated the cells with CLR01 concentrations ranging from 10 nM to 50 μ M for 16 h to allow uptake of the molecular tweezer into the cells. Then, we added 150 nM heparin-induced tau or p-tau(S262A)^{RBE} fibrils, which had been sonicated for 10 min prior to addition to the cells, and measured the seeding activity after additional 24 h of incubation.

The seeding activity of heparin-induced tau fibrils was again higher than that of p-tau(S262A)^{RBE} (Figure 7A). Confocal microscopy confirmed the formation of abundant seeds in cells treated only with heparin-induced tau fibrils and not with CLR01 (Figure 7B) whereas no seeds could be seen in cells incubated with CLR01 prior to addition of tau seeds (Figure 7C). The presence of CLR01 in cells treated with heparin-induced tau up to a 1:1 tau:CLR01 concentration ratio, respectively, increased the seeding activity compared to cells treated with the same seeds in the absence of CLR01, whereas at higher concentrations, CLR01 inhibited tau seeding. In contrast to the mixed effect of CLR01 on the seeding of heparin-induced tau fibrils, the molecular tweezer inhibited the seeding of p-tau(S262A)^{RBE} seeds dose-dependently with $IC_{50} = 660 \pm 140$ nM (Figure 7A), further demonstrating the distinct behavior of the two types of tau aggregates and the problematic nature of using heparin-induced tau aggregates for testing aggregation and/or seeding modulators, such as CLR01.

A potential concern with interpretation of this experiment is that CLR01 could act directly on the seeds themselves and induce their dissociation once they get internalized by

the biosensor cells. However, the labile binding of CLR01 would not be expected to affect such fibrils, unless they are incubated with a large excess of the molecular tweezer for long periods, as has been shown previously for other amyloidogenic proteins.^{75,77,78} Indeed, morphological examination showed that when tau fibrils were incubated with CLR01 for 24 h in vitro, CLR01 did not cause appreciable changes in the fibril structure (Supplementary Figure S4)

NMR-based mapping of CLR01 binding to the different tau variants. To gain further insight into the way CLR01 interacts with tau, we used solution-state NMR. Full-length tau contains 44 Lys residues, each of which is a potential binding site for CLR01. However, the affinity of CLR01 for each Lys residue depends both on the flanking amino acid sequences and on the local and global conformation. Lys residues also are binding sites for heparin.^{42,79,80} Previous NMR mapping of tau-heparin interaction revealed that large heparins bind to most of the Lys residues in the MTBD and PRR, and to a low extent also to Lys residues in the extreme N-terminus.⁸¹ Using small heparins, specific hotspots have been identified allowing discrimination between sites with different binding affinities.⁴² Because binding of either CLR01 or heparin would be expected to cause perturbation of Lys residues, we did not attempt to use NMR for the characterization of the competition between the two molecules for binding to tau. Rather, we used NMR to determine whether phosphorylation alters the binding of CLR01 to tau.

Unlike heparin, CLR01 binds one Lys residue at a time by including the Lys side chain inside the cavity of the molecular tweezer, thus forming hydrophobic interactions with the tweezer's "arms" and electrostatic interactions

between the ϵ -NH₃⁺ group of Lys and the phosphate groups of the molecular tweezer. As CLR01 is a symmetric compound containing two phosphate groups, if another positively charged residue, such as Lys or Arg is nearby, it could form additional electrostatic interactions with the second phosphate group of CLR01 but those would be substantially weaker because only one Lys side chain can be included inside the cavity. Given this binding mode, we asked first if CLR01 had a preference for selective binding to certain Lys residues versus others. CLR01 also binds to Arg residues, but with 5–10-times lower affinity than to Lys residues.^{55,82,83} Therefore, we focused most of our analysis on Lys only using selective ¹⁵N-Lys labeling of tau, which helps reducing signal crowding in two-dimensional ¹H-¹⁵N HSQC spectra. The assignment of Lys resonances in the ¹H-¹⁵N HSQC spectrum of ¹⁵N-Lys labeled tau has been described previously.^{42,81}

We analyzed unphosphorylated tau first to establish a baseline. We titrated 200 μ M ¹⁵N _{α,ϵ} -Lys-labeled tau with increasing concentrations of CLR01 ranging from 20 to 400 μ M and recorded a ¹H-¹⁵N HSQC spectrum for each titration point. As the concentration of CLR01 increased,

we observed minimal chemical shift variations, but the HSQC signal intensity decreased gradually (Figure 8A; Supplementary Figure S5). Due to the dynamic nature of its on-off exchange, CLR01 binding induces a severe broadening of Lys resonances associated to signal intensity decrease, as has been reported previously in A β .⁷⁵ We observed a similar behavior in a short peptide derived from tau (data not shown). In ¹H spectra, we observed a substantial decrease in ¹H resonance intensities of aliphatic protons in Lys residues whereas other ¹H resonances decreased only by < 5% at the highest concentration of CLR01 (Figure 8B), confirming specific interactions of CLR01 with Lys residues. For example, ~50% of the Lys H _{ϵ} signal at 2.99 ppm disappeared at the highest tau:CLR01 ratio 1:2, respectively (Figure 8B). At this ratio, the CLR01:Lys ratio is 1:22. Thus, the strong decrease in the overall signal of Lys demonstrates the kinetically fast on-off rate of CLR01 binding to Lys. Interactions of CLR01 with more than one Lys residues at a time, leading to further signal broadening, also could account for the large signal decrease at low CLR01:Lys ratios.

Figure 5. Inhibition of tau aggregation by CLR01. A) Time-dependent ThT fluorescence in aggregation reactions containing 5 μ M tau, 1.25 μ M heparin, and increasing concentrations of CLR01 showing replicates and fitted association curves. B) Dose-response analysis of the fluorescence data in panel A were calculated by extracting the kinetic constant k from fitted curves and normalizing to k_0 , the kinetic constant in the absence of CLR01. C) Time-dependent ThT fluorescence in aggregation reactions containing 30 μ M p-tau(S262A)^{RBE} and increasing concentrations of CLR01 showing replicates and fitted association curves. “Tau-S262A” is a negative control containing this variant of tau in an unphosphorylated form. D) Dose-response analysis of the fluorescence data in panel C were calculated by extracting the kinetic constant k from fitted curves and normalizing to k_0 , the kinetic constant in the absence of CLR01.

Figure 6. Representative TEM images taken at the end of the aggregation reactions shown in Figure 5. A) Images of 5 μ M heparin-induced tau in the absence or presence of the indicated tau:CLR01 concentration ratios. B) Images of 30 μ M p-tau(S262A)^{RBE} in the absence or presence of the indicated p-tau(S262A)^{RBE}:CLR01 concentration ratios.

Mapping of the resonance intensity variations using the two-dimensional ¹H-¹⁵N HSQC spectra indicated that CLR01 bound preferentially to Lys residues in the MTBD and to a lower extent in the PRR. Most of the binding occurred in the region between K224 and K395 with the exception of K130 for which strong binding was found. Interestingly, little binding was detected to the Lys residues at the N- and C-termini (Figure 8C). A possible explanation for this preference is that the extreme N- and C-terminal regions are more acidic than the remaining of the protein (Supplementary Figure S6), which may lead to electrostatic repulsion of the negatively charged CLR01.

To gain additional insight into the binding of CLR01 within its main region of interaction in tau, we mapped its binding on the fragment F4, which encompasses residues 208–324 (Figure 8D and Supplementary Figure S5). F4 maintains many of the physiologic and pathologic functions of full-length tau, e.g., microtubule binding and the capacity to form PHF-like fibrils.^{84,85} F4 was produced with uniform ¹⁵N-labeling allowing mapping of intensity changes for all the residues of the fragment. The titration was performed

using the same increasing concentrations of CLR01 and under the same conditions used for full-length tau. F4 contains 17 of the 44 Lys residues of full-length tau, out of a total of 117 residues (i.e., Lys residues comprise 14.5% of the F4 sequence, compared to 10% of full-length tau) and 5 Arg residues (4.3% of the F4 sequence) all located in the PRR, at positions 209, 211, 221, 230 and 242. R209 and R211 were not detected in the HSQC spectrum and only low intensity variations, between 12% and 25% intensity loss, were observed for the other three Arg residues (Figure 8D) indicating weak binding of CLR01 to these Arg residues, in line with data described previously.^{55,82,83}

A gradual intensity loss was observed for some, but not all of the Lys residues (Figure 8D). Residues in the extreme N-terminus, e.g., S210, which was conjugated to a (His)₆-tag. His residues, e.g., H268 in the F4 sequence, showed substantial intensity changes, which likely reflected the high sensitivity of the His signal to small pH variations rather than CLR01 binding.

Figure 7. CLR01 inhibits tau seeding in biosensor cells.

Cells were incubated for 16 h in the absence or presence of increasing CLR01 concentrations, followed by addition of seeds. After additional 24 h of incubation, seeding was visualized by fluorescence microscopy and quantified using flow cytometry. A) Dose-response analysis of the seeding activity (Integrated FRET Density) of each tau form in the absence or presence of CLR01. B,C) High-magnification fluorescence microscopy images of cells incubated with heparin-induced tau fibrils after pre-treatment with media alone (B) or 50 μ M CLR01 (C). Bright YFP fluorescence indicates tau aggregates. Nuclei were stained with DAPI (blue).

Previously, the affinity of CLR01 for His was determined to be only ~10% of the affinity for Lys.⁵⁵ Except for these

residues, K224, K225, K240, K259, and K280 showed substantial intensity decrease, up to 50%, at equimolar concentration of CLR01 (Figure 8D). The most pronounced intensity decrease, 60 and 80%, respectively, was found for K290 and K321, both within KCGS motifs. However, it is difficult to determine whether this strong decrease represents the high sensitivity of Cys residues to changes in the environment or an actual higher affinity of CLR01 for the Lys residues in these positions. The data are in good agreement with a recent report, which found using top-down mass spectrometry, that a primary binding site for CLR01 in full-length tau was in a segment spanning residues K294–K331.⁸⁶ Importantly, Lys residues in F4 showed a comparable intensity decrease to their counterparts in

Figure 8. NMR mapping of CLR01 binding to tau and F4. A) NMR mapping of ¹⁵N-Lys-labeled tau in the absence (black spectrum) or presence (blue spectrum) of a two-fold excess CLR01. B) ¹H spectra of ¹⁵N-Lys-labeled tau with increasing concentration of CLR01. The H_ε resonance of Lys side chains at 2.99 ppm is depicted in the inset showing a selective decrease of ¹H signals of Lys residues. A similar intensity decrease was found for Lys H_δ resonances at 1.65 ppm. C,D) Mapping of resonance intensity of Lys residues along the ¹⁵N-Lys-tau (C) or ¹⁵N-F4 (D) sequences for different protein:CLR01 ratios. The region in grey in (C) corresponds to the F4 sequence.

full-length tau at the same protein:CLR01 concentration ratios (Supplementary Figure S5B), suggesting that the changes reflected the genuine affinity of CLR01 for these particular Lys residues.

After establishing the main binding region and specific primary binding sites of CLR01 on unphosphorylated tau, we asked how phosphorylation might affect CLR01 binding. We investigated the binding of CLR01 to p-tau^{ERK} and p-tau(S262A)^{RBE} by NMR in a similar manner to the unphosphorylated protein. First, we performed experiments using uniformly ¹⁵N,¹³C-labeled tau and selectively ¹⁵N_{αε}/¹³C-Lys,¹³C-labeled p-tau^{ERK}, which facilitates the mapping of resonance intensity upon CLR01 titration, to determine the phosphorylation pattern (Supplementary Figures S7). The findings are summarized in Table 1.

Spectral heterogeneity occurs upon incomplete phosphorylation of several Ser/Thr residues leading to a splitting of protein resonances for the phosphorylation sites and surrounding residues. Assignment of ¹H-¹⁵N resonances of ¹⁵N_{αε}/¹³C-Lys,¹³C-labeled p-tau^{ERK} was performed using a three-dimensional HN(CO)CACB experiment, which allows assigning Lys resonances that shift upon phosphorylation due to their proximity to phosphorylation sites (Supplementary Figure S7). Examination of the ¹H-¹⁵N HSQC NMR spectra of Lys-labeled p-tau^{ERK} revealed distinct behavior of certain Lys resonances. Some Lys resonances were split into two or more peaks due to phosphorylation heterogeneity, including K67, K190, K224, K225, and K234 and low-intensity resonances of the non-phosphorylated form were still detected. In contrast, for K44, K150, K174, K180, K240 and K395,

Figure 9. Binding site analysis of CLR01 on p-tau^{ERK}. A,B) Superimposition of ¹H-¹⁵N HSQC spectra of selectively ¹⁵N-Lys-labeled p-tau^{ERK} (A) and unphosphorylated tau (B) at 200 μ M in the absence (black) or presence of 200 μ M CLR01 (blue). Resonances of K240 corresponding to phosphorylated forms in panel A are annotated by K240-P1 and K240-P2. Note that the unphosphorylated form of K240 shown in panel B is absent in p-tau^{ERK}. The location of this missing resonance is indicated in panel A by "x". C) Variation of K240 resonance intensity upon increasing CLR01 concentrations for the unphosphorylated form in tau (open squares) and both phosphorylated forms in p-tau^{ERK} shown in panel A (filled diamonds and triangles). D) Changes in resonance intensity of Lys signals in p-tau^{ERK} upon CLR01 binding for protein:CLR01 ratios ranging from 1:0.1 to 1:2. For Lys residues close to phosphorylated residues, only intensity corresponding to the resonance in the phosphorylated form is indicated. Nt – N-terminus, Ct – C-terminus.

only peaks corresponding to the phosphorylated state were detected, as shown for K240; Figure 9A,B, indicating a high phosphorylation level for sites in the vicinity of these Lys residues (Supplementary Figure S7, Table 1).

The resonance intensity changes upon CLR01 binding to p-tau^{ERK} (Figure 9C) suggested that phosphorylation had

only a minor effect on the binding. The binding pattern of CLR01 to p-tau^{ERK} resembled the binding to unphosphorylated tau, targeting primarily the MTBD and to a lesser extent the flanking regions (Figure 9C and Supplementary Figure S9). Only binding to the PRR was slightly reduced (Supplementary Figures S9 and S10). This observation was consistent with the location of most of CLR01's binding sites in the MTBD, as opposed to most of

the phosphorylation sites in the PRR and the extreme N- and C-termini. The only exceptions were K150, K174 and K240, which are adjacent to the phosphorylation sites T153, T175 and T231/S235, respectively. Binding of CLR01 to these three Lys residues was significantly reduced upon phosphorylation (Figure 9C,D and Supplementary Figures S9 and S10).

The binding sites found for CLR01 on p-tau(S262A)^{RBE} were similar to those in unphosphorylated tau and p-tau^{ERK} (Supplementary Figures S9 and S10). Together, these data indicate that tau phosphorylation does not affect CLR01 binding substantially, allowing the molecular tweezer to remain an effective inhibitor of tau aggregation, oligomerization, and seeding.

Discussion

In Alzheimer's disease and other tauopathies, aberrant post-translational modification of tau leads to its self-assembly into neurotoxic oligomers and fibrils. Of the PTMs studied to date, hyperphosphorylation has been found most frequently and likely is the main PTM instigating aberrant tau self-assembly. The requirement of post-translational modifications for induction of aggregation makes studying this process in vitro or in cell culture difficult. To create experimental systems allowing such studies, researchers long have been using heparin and other polyanions to induce tau aggregation. These artificial models are particularly convenient for screening tau aggregation inhibitors and disaggregation inducers,⁸⁷ yet concerns about the artificial nature of such systems, and in particular about potential interference by the polyanions with testing potential modulators and inhibitors of aberrant tau self-assembly have been increasing. Recently, cryo-EM structures showed that heparin-induced 4R and 3R tau fibrils are structurally distinct from those in Alzheimer's and Pick's disease questioning the relevance of such *in-vitro* assays.⁸⁸ Furthermore, Fichou et al. have reported that heparin and RNA used to induce fibril formation co-assemble with tau in vitro.⁴⁷ Heparin-induced tau fibrils used in in-vitro seeding assays seeded freshly dissolved tau only in the presence of RNA whereas mouse-brain-derived tau fibrils could be used as seeds in the absence of co-factors in the first-generation seeding reaction and required RNA as a co-factor to increase the seeding activity of the second-generation reaction. Based on these findings, they suggested that in vivo, an unknown co-factor might be necessary for tau aggregation. In view of recent advances in generating in-vitro phosphorylated tau,^{29,30} here we set out to compare side-by-side this newer form of aggregation-prone tau with the common method of tau induced to aggregate by addition of heparin.

Induction of tau aggregation by addition of heparin allowed using a lower concentration of the protein and resulted in faster aggregation kinetics than those measured using p-tau(S262A)^{RBE} (Figure 1A). The morphology of the resulting fibrils was different – p-tau(S262A)^{RBE} were both thicker and longer than those of heparin-induced tau (*cf.* Figure 1B and D), demonstrating

that the two forms were structurally distinct. Our data suggest that such co-factors are not necessary for tau fibrillation and that hyperphosphorylation of tau is sufficient for induction of fibril formation.

Unlike p-tau(S262A)^{RBE}, p-tau^{ERK} did not form fibrils. NMR analysis showed that the two forms of in-vitro phosphorylated wild-type tau shared 15 common phosphorylated sites and differed in only three sites—S208, S262, and S356—which were present in RBE- but not in ERK-phosphorylated tau.²⁹ Comparison of these two forms suggested that the phosphorylation of S208 was the key site necessary for aggregation, as described by Despres et al.,³⁰ because phosphorylation of S262 and S356 had been reported previously to inhibit aggregation.²⁸ Therefore S262, a site that gets phosphorylated in vitro at a high level by RBE (in contrast to S356), was substituted by Ala in subsequent experiments to obtain an aggregation-prone isoform. Interestingly, comparison of the oligomerization behavior of these two forms showed that the aggregation resistant p-tau^{ERK} formed a variety of small (dimer through tetramer), medium (apparent mobility corresponding to 800–1,000 kDa), and large (apparent mobility corresponding to 1,100 kDa) oligomers, whereas the aggregation-prone p-tau(S262A)^{RBE} comprised a mixture in which individual oligomers were less discernible and high molecular weight species were prominent. Following incubation, most of the p-tau(S262A)^{RBE} preparation was no longer visible in native-PAGE/western blot analysis (Figure 1E), presumably due to formation of large aggregates that did not enter the gel.

Using the tau biosensor cell line developed by Diamond and co-workers, we found, as expected, that tau fibrils seeded intracellular tau aggregation, regardless of whether they were made of heparin-induced tau or p-tau(S262A)^{RBE} (Figure 2A–D). We were surprised, however, to discover that the morphology and cellular distribution of the two forms suggested that they belonged to distinct strains. Heparin-induced tau fibrils induced formation primarily of relatively large cytoplasmic/perinuclear aggregates (Figure 2E), whereas p-tau(S262A)^{RBE} seeded small, primarily nuclear aggregates (Figure 2F). These morphologies resembled the “ordered” and “speckles” strains described recently by Kaufman et al.⁵⁴ That study found that tau from different sources, including human AD brain, cell lines from patients with different tauopathies, brain extract from the P301S mouse model, and recombinant tau gave rise to various strains distinguished by the appearance of the aggregates in the biosensor cells. Our data suggest that heparin-induced tau fibrils prepared under the conditions described here and p-tau(S262A)^{RBE} may allow obtaining discrete strains reproducibly using recombinant tau for further structural and biological studies. In agreement with previous reports suggesting that tau seeding did not require fibril formation and could be mediated by oligomers, we found that both forms of freshly dissolved in-vitro phosphorylated tau, p-tau^{ERK} and p-tau(S262A)^{RBE}, though importantly not unphosphorylated tau, seeded tau aggregation in the biosensor cells.

Before the current study, only two pieces of information were available regarding the effect of CLR01 on tau. Using the ThT-fluorescence assay, equimolar concentration of CLR01 was shown to inhibit completely the aggregation of the shortest form of tau, 0N3R, which was induced to aggregate by arachidonic acid.⁷⁵ In the presence of arachidonic acid, tau aggregates into amorphous, rather than fibrillar aggregates, therefore the effect of CLR01 on fibril formation was not assessed in that study. The impact of CLR01 on tau aggregates *in vivo* was examined in the triple-transgenic AD mouse model expressing mutant forms of tau, amyloid β -protein precursor, and presenilin-1. These mice accumulate both amyloid plaques and NFTs in their brain with aging. Treatment of 14–15-month old mice, who already had both types of lesions, with a low dose of CLR01 led to a substantial reduction in both amyloid plaques and NFTs.⁹⁰ Because tau pathology is thought to be secondary to the neurotoxic effects of amyloid β -protein (A β), the study in the triple-transgenic mice could not reveal whether the decrease in neurofibrillary tangles was due to a direct effect of CLR01 on tau or an indirect effect due to inhibition of A β toxicity. Though the current study does not address this question directly, it reveals that CLR01 effectively inhibits tau aggregation and intracellular seeding. The data also demonstrate that the choice of the experimental system for studying the effect of the modulator is crucial.

When tau was induced to aggregate by heparin, CLR01 could interfere both with the induction itself and with the intermolecular interactions among tau monomers that mediate their further aggregation. Comparison of direct inhibition of tau-heparin binding using SPR (Figure 4) and inhibition of tau aggregation using ThT-fluorescence (Figure 5) suggested that the main inhibitory activity of CLR01 was effected on the self-assembly of tau polypeptides rather than on tau-heparin binding. Thus, the concentration of CLR01 needed for half-maximal inhibition of tau aggregation was only twice the concentration of tau, whereas 40-fold excess CLR01 was needed for half-maximal inhibition of tau binding to heparin. Consistent with the interpretation of efficient inhibition of tau self-assembly, CLR01 also efficiently inhibited the aggregation of p-tau(S262A)^{RBE}. Interestingly, however, the molecular tweezer had a more complex effect on the aggregation of the protein in this case. At low concentrations, an initial acceleration of ThT fluorescence increase was observed, whereas later in the aggregation reaction, the rate of increase and the final fluorescence values were lower than that of p-tau(S262A)^{RBE} alone. Examination of the morphology at the end of the reaction showed dose-dependent inhibition of fibril formation (Figure 6), suggesting that the initial increase in ThT fluorescence did not reflect formation of tau fibrils, but rather of a non-fibrillar, yet presumably β -sheet-rich structure. Similar non-fibrillar structures displaying increased ThT fluorescence have been observed before in the presence of CLR01 for other amyloidogenic proteins, including insulin, β_2 -microglobulin, transthyretin, and p53 mutants.^{75,76} The details of these structures are not known, but they seem to form when CLR01 interacts with structured, rather than unstructured amyloidogenic proteins. If this trend holds

true here too, it would suggest that phosphorylation of tau may induce formation of a more stable structure than that of unphosphorylated tau. Interestingly, formation of a partially structured conformation of tau recently has been suggested to transform tau from a benign to a seeding-competent form,⁶⁵ in agreement with this idea. NMR studies also have highlighted phosphorylation-induced conformational changes in the PRR around the binding phospho-epitopes of antibodies AT8 (pS202/pT205)⁹¹ and AT180 (pT231/pS235)^{92,93} in which phosphorylation stabilizes transient secondary structures. The double phosphorylation at the AT8 epitope was shown to induce a turn-like structure which protects tau from aggregation while a triple phosphorylation which includes pS208 destabilizes this conformation and stimulates aggregation of phosphorylated tau.³⁰

CLR01 binds to surface-exposed residues⁹⁴ and therefore in natively unstructured proteins would be expected to bind most or all of the Lys residues. However, previous studies with A β ⁷⁵ and α -synuclein⁹⁵ have shown that certain residues are preferred over others as binding sites. Here, we found that CLR01 binds preferentially to Lys residues in the MTBD (Figure 8, Supplementary Figure S5), possibly due to a high concentration of acidic residues at the N- and C-terminal regions that may repel the negatively charged molecular tweezer (Supplementary Figure S6). An alternative explanation is involvement of Lys residues in those regions in salt bridges or other structural elements that prevent them from interacting with CLR01. However, there is little evidence supporting the existence of such structural elements and the observation that the interaction of CLR01 with the Lys residues in tau was not affected substantially by phosphorylation (Figure 9, Supplementary Figures S9 and S10), which may increase structural order, suggest against this explanation.

Our study demonstrates that heparin-induced tau aggregation yields distinct aggregation kinetics, oligomerization patterns, and seeding behavior from those of *in-vitro* phosphorylated tau, which we hypothesize is a closer mimic of the hyperphosphorylated tau in toxic oligomers and NFTs in AD and other tauopathies. If our hypothesis is correct, interpretation of studies using tau induced to self-assemble by addition of heparin or other polyanions should be done with caution. Testing modulators and inhibitors of tau self-assembly and seeding in the presence of aggregation inducers is particularly problematic due to the competition between the inducer and the modulator for binding to tau. *In-vitro* phosphorylation of tau alleviates this problem and provides a better mimic of disease-associated hyperphosphorylated tau. Our data also suggest that CLR01 is an effective inhibitor of tau self-assembly and seeding. As CLR01 also has been shown to be a potent inhibitor of A β self-assembly and toxicity *in vitro* and *in vivo*,^{75,90,96,97} it may be a particularly attractive lead for developing disease-modifying therapy thanks to its ability to inhibit both of the offending proteins in AD simultaneously.

ASSOCIATED CONTENT

Supporting Information. Supplementary Figures S1–S10 are supplied as Supporting Information. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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ABBREVIATIONS

AD: Alzheimer's disease; DAPI: 4',6-diamidino-2-phenylindole; ERK2: Extracellular signal-regulated kinase 2; HSQC: heteronuclear single-quantum coherence; MEK3: Mitogen-activated protein-kinase kinase 3; MTBD: NFTs: neurofibrillary tangles; microtubule-binding domain; PHF: paired helical filaments; PTM: posttranslational modification; RBE: rat-brain extract; SPR: surface-plasmon resonance; TALS: Tau aggregation inhibitors; TEM: Transmission electron microscopy; ThT: Thioflavin T; TOCSY: total correlation spectroscopy; FRET: fluorescence-resonance energy transfer; MALDI-TOF MS: matrix-assisted laser desorption-ionization time-of-flight mass spectrometry.

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website.

Supplementary Figure 1: Morphological examination of p-tau(S262A)^{RBE} fibrils

Supplementary Figure 2: Comparison of fitting the plateau vs. the rate constant, k.

Supplementary Figure 3: Representative TEM images taken at the end of the aggregation reactions.

Supplementary Figure 4: Transmission electron microscopy of p-tau(S262A)^{RBE} fibrils in the absence or presence of CLR01.

Supplementary Figure 5: Titration of full-length tau and the F4 fragment with CLR01.

Supplementary Figure 6: Mean charge along the tau sequence.

Supplementary Figure 7: Phosphorylation of ¹⁵N-lysine, ¹³C-tau by ERK2.

Supplementary Figure 8: ¹H-¹⁵N HSQC spectra of p-tau(S262A)^{RBE} phosphorylated by a rat brain extract.

Supplementary Figure 9: Comparison of signal intensity variations in p-tau^{ERK}, p-tau(S262A)^{RBE}, and non-phosphorylated tau.

Supplementary Figure 10: Ratio of signal intensity of phosphorylated tau.

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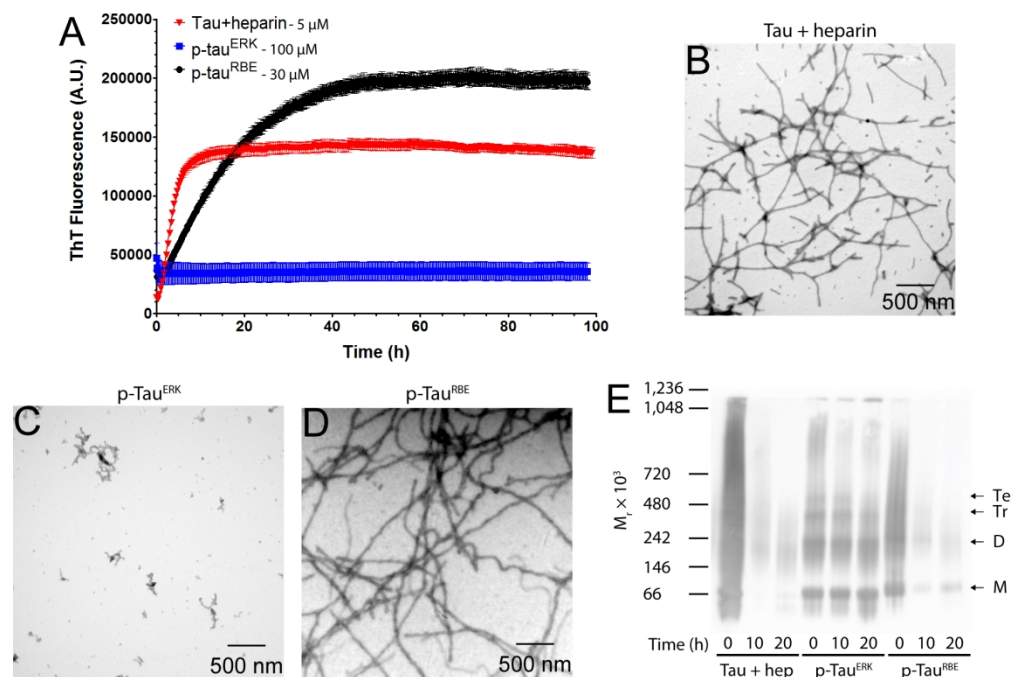


Figure 1. Self-assembly of tau induced by heparin, p-tauERK and p-tau(S262A)RBE. (A) Time-dependent ThT fluorescence of 5 μ M heparin-induced tau, 100 μ M p-tauERK, or 30 μ M p-tau(S262A)RBE. (B-D) Morphological examination by TEM at the end of the aggregation reaction (t = 100 h) of heparin-induced tau (B), p-tauERK (C) and p-tau(S262A)RBE (D). E) Native-PAGE analysis of 10 μ M each of heparin (hep)-induced tau, p-tauERK, and p-tau(S262A)RBE at t = 0, 10, or 20 h of incubation under the same conditions. For heparin-induced tau, t = 0 is just before addition of heparin. Molecular weight markers are shown on the left. The mobilities of putative monomer (M), dimer (D), trimer (Tr), and tetramer (Te) are indicated on the right.

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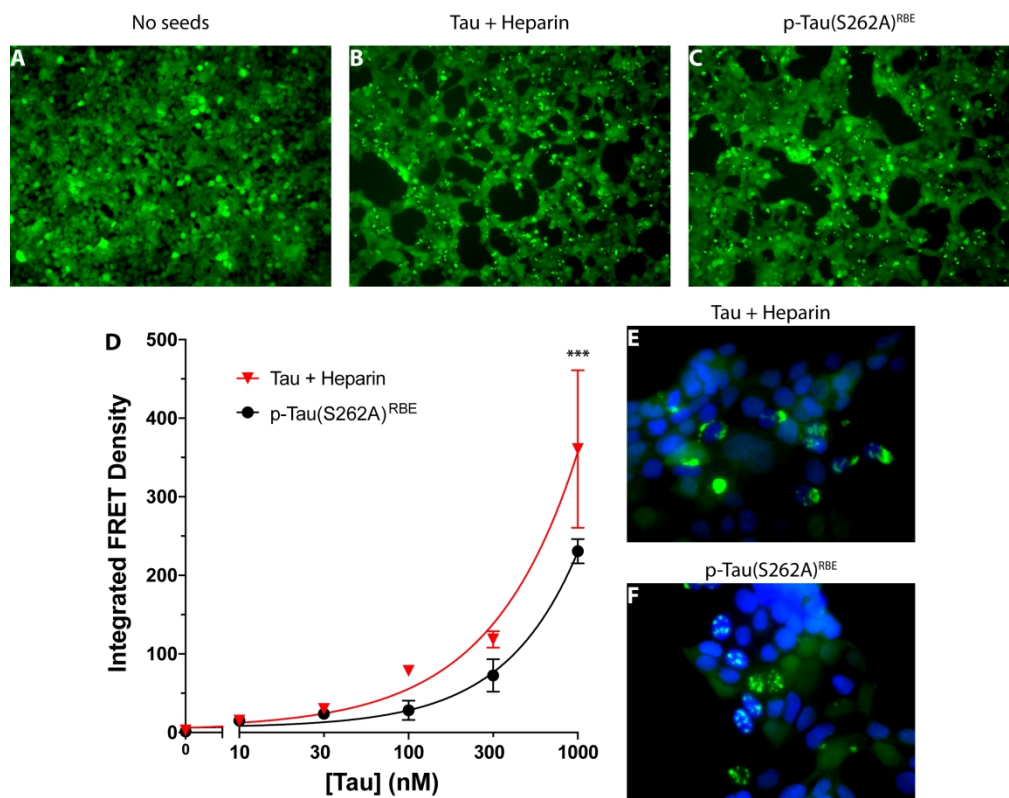


Figure 2. Induction of tau seeding by aggregated tau in biosensor cells. A–C) Fluorescence microscopy images of tau-biosensor cells. A) Negative control cells incubated with media alone. B) Cells incubated with heparin-induced tau fibrils. C) Cells incubated with p-tau(S262A)RBE fibrils. D) Dose-response analysis of the seeding activity (Integrated FRET Density) of each tau form. E,F) High-magnification fluorescence microscopy images of cells incubated with 1 μ M of either heparin-induced tau fibrils (E) or p-tau(S262A)RBE fibrils (F). Bright YFP fluorescence indicates tau aggregates. Nuclei were stained with 4',6-diamidino-2-phenylindole (DAPI, blue).

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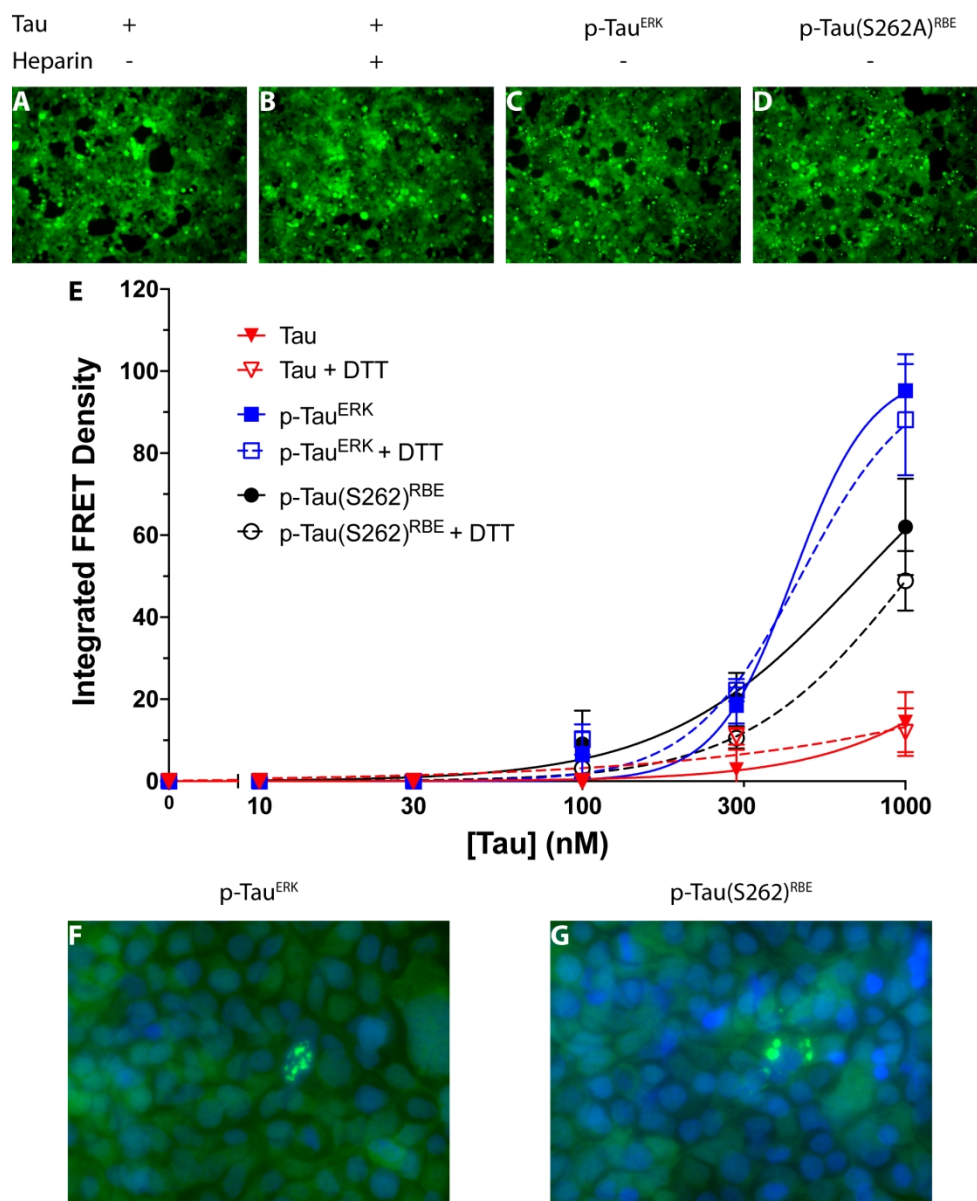


Figure 3. Induction of tau seeding by freshly dissolved tau in biosensor cells. A-D) Fluorescence microscopy images of tau-biosensor cells incubated with 1 μ M freshly dissolved tau (A), tau + heparin (B), p-tauERK (C), or p-tau(S262A)RBE (D). E) Dose-response analysis of the seeding activity (Integrated FRET Density) of each tau form in the absence or presence of DTT. F,G) High-magnification fluorescence microscopy images of cells incubated with 300 nM freshly dissolved p-tau ERK (F) or p-tau(S262A)RBE (G). Bright YFP fluorescence indicates tau aggregates. Nuclei were stained with DAPI (blue).

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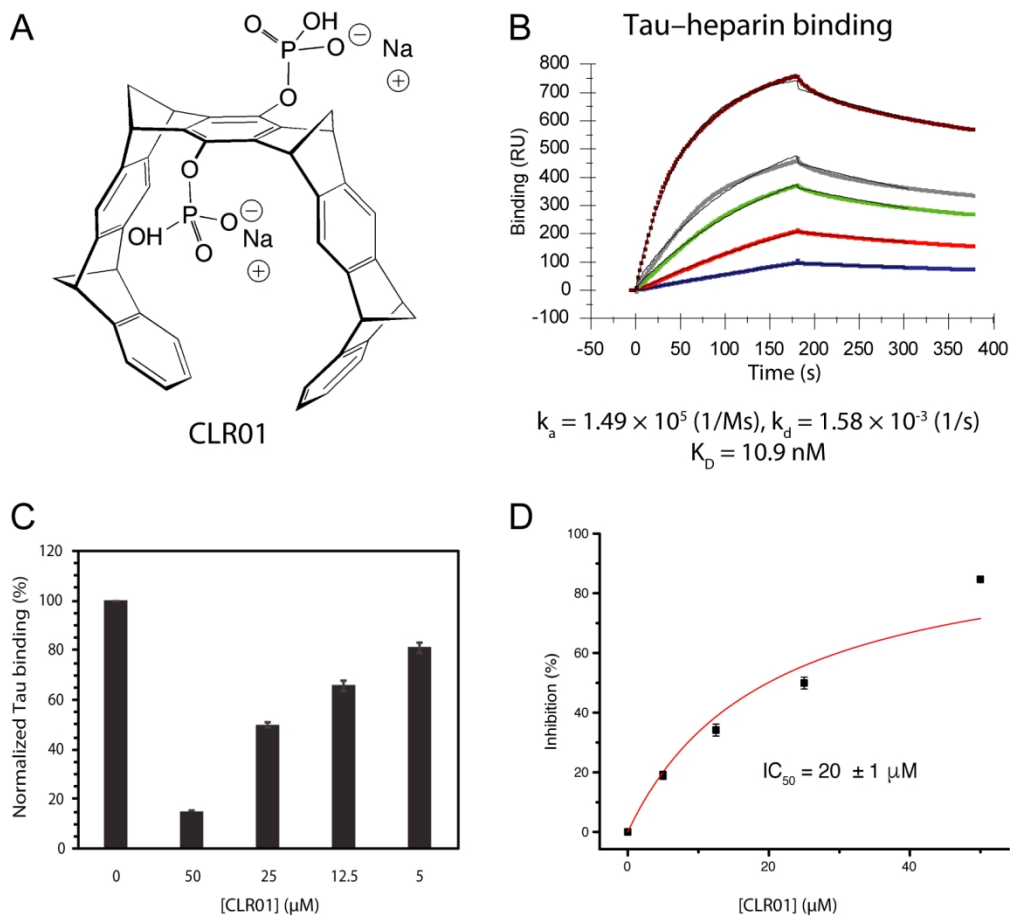


Figure 4. Inhibition of tau binding to heparin by CLR01. Binding of tau to immobilized heparin and its inhibition by CLR01 were measured using SPR. A) A schematic structure of CLR01. The compound is partially protonated at physiologic pH. B) SPR sensorgram of tau binding to heparin immobilized on a SPR chip. C) Normalized tau binding in the presence of increasing concentrations of CLR01. D) % Inhibition versus CLR01 concentration used for calculating the half-maximal inhibition (IC_{50}).

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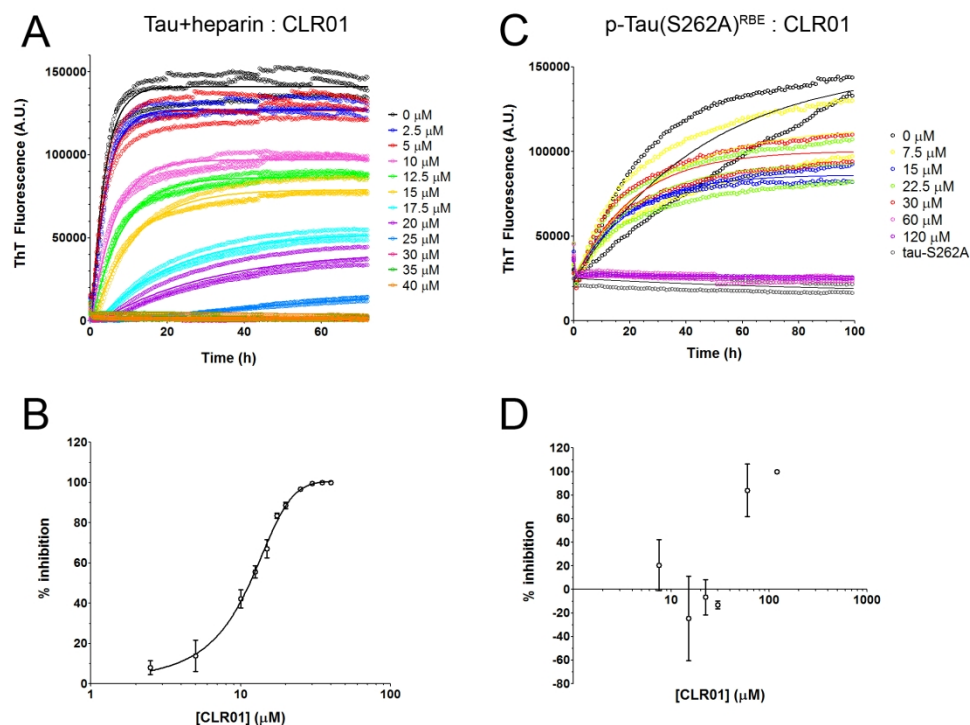


Figure 5. Inhibition of tau aggregation by CLR01. A) Time-dependent ThT fluorescence in aggregation reactions containing 5 μM tau, 1.25 μM heparin, and increasing concentrations of CLR01 showing replicates and fitted association curves. B) Dose-response analysis of the fluorescence data in panel A were calculated by extracting the kinetic constant k from fitted curves and normalizing to k_0 , the kinetic constant in the absence of CLR01. C) Time-dependent ThT fluorescence in aggregation reactions containing 30 μM p-tau(S262A)^{RBE} and increasing concentrations of CLR01 showing replicates and fitted association curves. "Tau-S262A" is a negative control containing this variant of tau in an unphosphorylated form. D) Dose-response analysis of the fluorescence data in panel C were calculated by extracting the kinetic constant k from fitted curves and normalizing to k_0 , the kinetic constant in the absence of CLR01.

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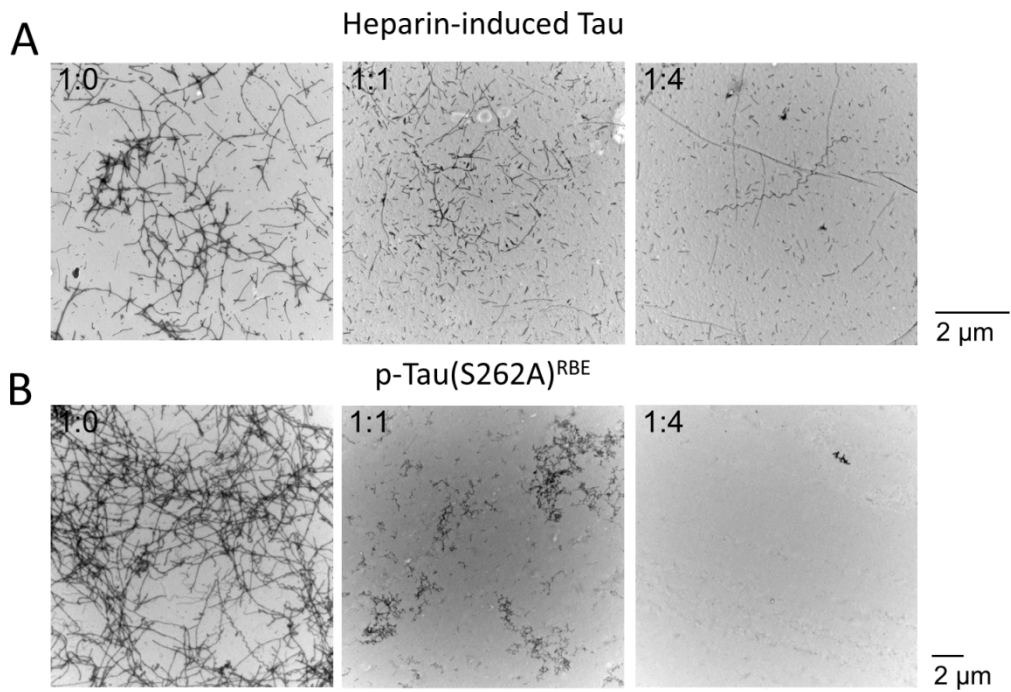


Figure 6. Representative TEM images taken at the end of the aggregation reactions shown in Figure 5. A) Images of 5 μM heparin-induced tau in the absence or presence of the indicated tau:CLR01 concentration ratios. B) Images of 30 μM p-tau(S262A)RBE in the absence or presence of the indicated p-tau(S262A)RBE:CLR01 concentration ratios.

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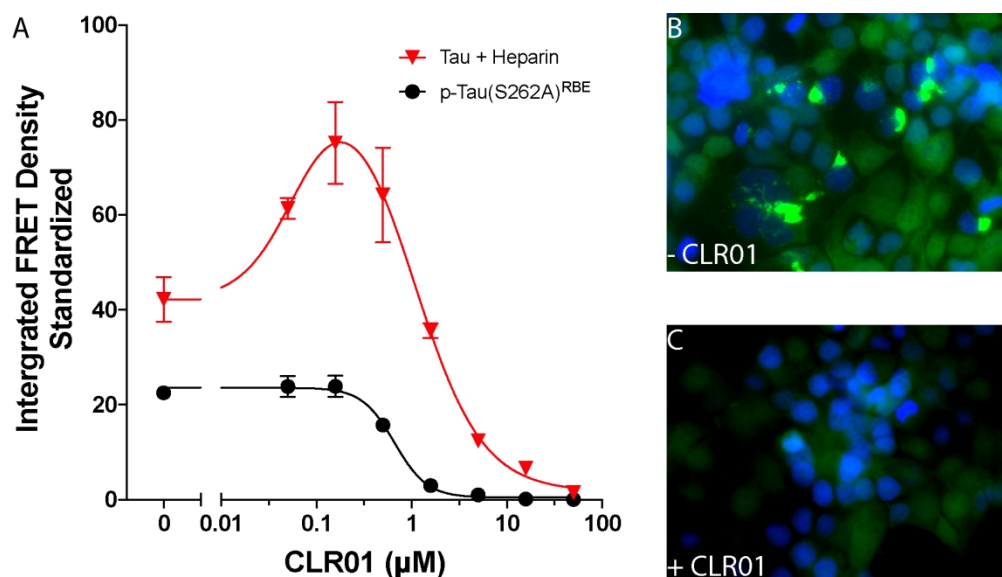


Figure 7. CLR01 inhibits tau seeding in biosensor cells. Cells were incubated for 16 h in the absence or presence of increasing CLR01 concentrations, followed by addition of seeds. After additional 24 h of incubation, seeding was visualized by fluorescence microscopy and quantified using flow cytometry. A) Dose-response analysis of the seeding activity (Integrated FRET Density) of each tau form in the absence or presence of CLR01. B,C) High-magnification fluorescence microscopy images of cells incubated with heparin-induced tau fibrils after pre-treatment with media alone (B) or 50 μM CLR01 (C). Bright YFP fluorescence indicates tau aggregates. Nuclei were stained with DAPI (blue).

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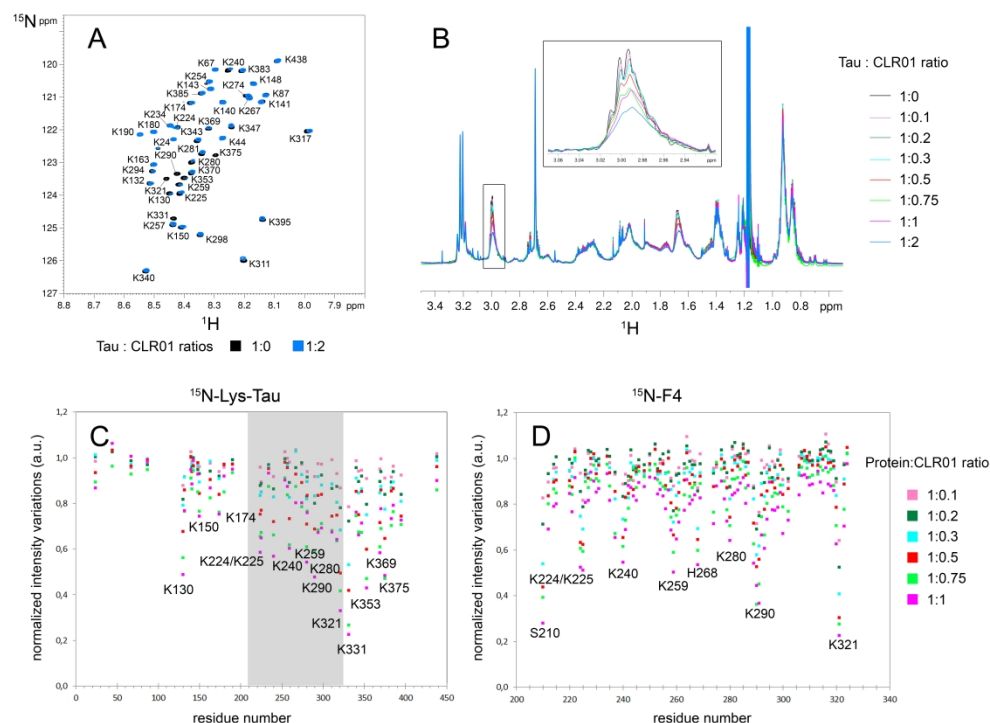


Figure 8. NMR mapping of CLR01 binding to tau and F4. A) NMR mapping of ^{15}N -Lys-labeled tau in the absence (black spectrum) or presence (blue spectrum) of a two-fold excess CLR01. B) ^1H spectra of ^{15}N -Lys-labeled tau with increasing concentration of CLR01. The $\text{H}\epsilon$ resonance of Lys side chains at 2.99 ppm is depicted in the inset showing a selective decrease of ^1H signals of Lys residues. A similar intensity decrease was found for Lys $\text{H}\delta$ resonances at 1.65 ppm. C,D) Mapping of resonance intensity of Lys residues along the 15N-Lys-tau (C) or 15N-F4 (D) sequences for different protein:CLR01 ratios. The region in grey in (C) corresponds to the F4 sequence.

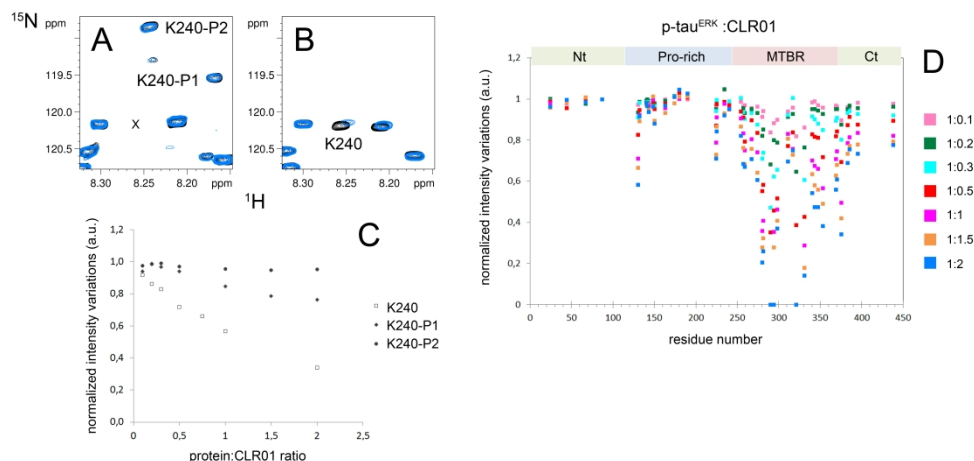


Figure 9. Binding site analysis of CLR01 on p-tauERK. A,B) Superimposition of ^1H - ^{15}N HSQC spectra of selectively ^{15}N -Lys-labeled p-tauERK (A) and unphosphorylated tau (B) at 200 μM in the absence (black) or presence of 200 μM CLR01 (blue). Resonances of K240 corresponding to phosphorylated forms in panel A are annotated by K240-P1 and K240-P2. Note that the unphosphorylated form of K240 shown in panel B is absent in p-tauERK. The location of this missing resonance is indicated in panel A by "X". C) Variation of K240 resonance intensity upon increasing CLR01 concentrations for the unphosphorylated form in tau (open squares) and both phosphorylated forms in p-tauERK shown in panel A (filled diamonds and triangles). D) Changes in resonance intensity of Lys signals in p-tauERK upon CLR01 binding for protein:CLR01 ratios ranging from 1:0.1 to 1:2. For Lys residues close to phosphorylated residues, only intensity corresponding to the resonance in the phosphorylated form is indicated. Nt – N-terminus, Ct – C-terminus.

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