Neurofibrillary tangle-bearing neurons are functionally integrated in cortical circuits in vivo

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Alzheimer’s disease (AD) is pathologically characterized by the deposition of extracellular amyloid-β plaques and intracellular aggregation of tau protein in neurofibrillary tangles (NFTs) (1, 2). Progression of NFT pathology is closely correlated with both increased neurodegeneration and cognitive decline in AD (3) and other tauopathies, such as frontotemporal dementia (4, 5). The assumption that mislocalization of tau into the somatodendritic compartment (6) and accumulation of fibrillar aggregates in NFTs mediates neurodegeneration underlies most current therapeutic strategies aimed at preventing NFT formation or disrupting existing NFTs (7, 8). Although several disease-associated mutations cause both aggregation of tau and neurodegeneration, whether NFTs per se contribute to neuronal and network dysfunction in vivo is unknown (9). Here we used awake in vivo two-photon calcium imaging to monitor neuronal function in adult rTg4510 mice that overexpress a human mutant form of tau (P301L) and develop cortical NFTs by the age of 7–8 mo (10). Unexpectedly, NFT-bearing neurons in the visual cortex appeared to be completely functionally intact, to be capable of integrating dendritic inputs and effectively encoding orientation and direction selectivity, and to have a stable baseline resting calcium level. These results suggest a reevaluation of the common assumption that insoluble tau aggregates are sufficient to disrupt neuronal function.

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crofibrillary tangles (NFTs) containing aggregated tau protein (1) have long been considered key players in the progressive neural dysfunction and neurodegeneration observed in Alzheimer’s disease (AD) (2, 3) and other tauopathies (4, 5). It is commonly assumed that NFT-bearing neurons exhibit deficits in synaptic integration and eventually lead to neurodegeneration (11, 12). However, the actual functional properties of NFT-bearing neurons in intact neural circuits have not been explored previously (13). We addressed this question directly using awake in vivo two-photon calcium imaging in a mouse model of NFT formation (rTg4510) by applying recently developed imaging approaches allowing for single-neuron-level and population-level assessment of neural activity in awake mice (14). Because two-photon calcium imaging allows for measurement of response properties in many neurons simultaneously, we were able to directly isolate the impact of NFT deposition in a neuronal microcircuit by evaluating population-level network dynamics and, more specifically, by differentiating the function of individual NFT-bearing and neighboring non–NFT-bearing neurons.

To assess the functional properties of neurons in the visual cortex, we used a genetically encoded ratiometric calcium indicator, yellow cameleon 3.6 (YC3.6), packaged in an adenovirus-associated viral vector (15, 16). To assess functional responses, we exploited the well-characterized functional architecture of visual cortex whereby neurons in mouse visual cortex modulate their activity during presentation of drifting gratings moving at specific orientations and directions (orientation and direction selectivity) (17, 18). We further used YC3.6 as a FRET-based ratiometric indicator to make quantitative measurements of resting calcium (15). Resting calcium is tightly regulated in the brain, and slight deviations can trigger chronic and severe degenerative pathways (15). Thus, measurement of resting calcium is an important and complementary functional assay for evaluating neuronal health. Importantly, performing experiments in awake, head-fixed animals eliminates the impact of anesthesia on response properties, resting calcium, and tau aggregation (Fig. 1 A and B and Movie S1) (19).

Results

We first confirmed that 8–10-mo-old control animals exhibited robust orientation and direction selectivity, similar to previous reports in young and aged mice (20) (Fig. 1 and Fig. 2 A–E, control traces). Average and single trial traces of responses to visual stimuli measured in a single neuron from a control animal are shown in Fig. 1C. In this example, the neuron (cell 1) preferentially responds to a visual stimulus orientation of 225° and has an orientation selectivity index (OSI) of 0.95 and a direction selectivity index (DSI) of 0.95 (Fig. 1 C, D, and F; 0 indicates no selectivity, and 1 indicates perfect selectivity).

Significance

Alzheimer’s disease is pathologically characterized by extracellular amyloid-β plaques and intracellular neurofibrillary tangles (NFTs). It has long been assumed that the accumulation of tau into NFTs causes neuronal dysfunction and death, and is a proximate cause of dementia in patients with Alzheimer’s disease. This assumption underlies the NFT-busting drugs currently in clinical trials and research efforts aimed at understanding tau aggregation. Our study tested the dogma that NFT-bearing neurons are indeed impaired in their ability to respond to complex sensory stimuli. Using two-photon imaging in awake mice with NFT pathology, we found that individual neurons with NFTs respond to visual stimuli and do not impair local circuits. These unexpected results suggest that the presence of an NFT does not inevitably lead to gross physiological alterations.


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We next explored the functional response profiles of neurons in the visual cortex of transgenic mice with a significant NFT load in the visual cortex at similar ages to control animals (Fig. 2A and Fig. S1) and compared these profiles with those of neurons from control brains (Fig. 2B–E). The rTg4510 mice exhibited numerous NFTs throughout the cortex, including the visual cortex (Fig. 2A), along with behavioral deficits typically seen at this age (10). We observed a marked uniformity in resting calcium levels across the population of neurons measured in control mice (Fig. 2B), consistent with previous reports of tightly regulated resting calcium concentration in the healthy cortex (15). Neurons in the rTg4510 mice and control mice exhibited nearly identical resting calcium profiles (control, 1.32; rTg4510, 1.31; cohort-level: P = 0.84, t test, n = 3 rTg4510 mice, n = 3 control mice; cell-specific: P = 0.22, Mann–Whitney U test, n = 898 rTg4510 neurons, n = 329 control neurons) (Fig. 2B).

We further explored whether other cortical brain regions exhibited disrupted calcium homeostasis by measuring resting calcium outside of the visual cortex (somatosensory areas), and found no significant difference (Fig. S2). Moreover, neurons in mice with a high NFT load exhibited the same robust visual responses (Fig. 2C–E) as were observed in controls including many neurons with sharp orientation and direction selectivity (OSI: P = 0.47, DSI: P = 0.82, Student t test; n = 6 Tg4510 mice, n = 6 control mice) (Fig. 2D and E and Fig. S3). Based on these data, we conclude that the neuronal network in the visual cortex of mice with a high NFT load appears to be functionally intact and comparable to that of control animals.

To examine whether disrupted calcium homeostasis or altered neuronal tuning in NFT-bearing neurons was masked by averaging the functional properties across all neurons and mice, we next compared resting calcium regulation and neuronal tuning in individual neurons with or without NFTs (Fig. 3). To identify NFT-bearing neurons in the visual cortex of rTg4510 mice, we aligned images obtained in vivo with images of the same neurons obtained by postmortem labeling for markers of NFTs (Fig. 3A–C) (21). The mice were perfused with paraformaldehyde (PFA) immediately after completion of in vivo imaging, followed by postmortem immunolabeling of YC3.6 (with anti-GFP antibody), hyperphosphorylated tau (with PHF1 antibody), and staining of mature NFTs using thioflavin-S (ThioS) in whole brains of rTg4510 mice (Fig. S4). All ThioS-positive NFT-bearing neurons (n = 32) showed PHF1 labeling of hyperphosphorylated tau, and very few neurons (n = 6) had hyperphosphorylated tau but no mature NFTs. For statistical analyses of NFT neuron responsiveness, ThioS stained cells were defined as NFT-bearing neurons. In a subset of animals, we also labeled NFTs in vivo using i.v. injections of the Congo red derivate methoxy-X04 in anesthetized animals (22, 23) (Fig. S5).

Remarkably, resting calcium (Fig. 3D) and neuronal tuning (Fig. 3G and H) were very similar in nearby (and often neighboring) NFT-bearing and non–NFT-bearing neurons from the same animal and at the same imaging volume. Average resting calcium levels were nearly identical, and resting calcium distributions of neighboring NFT-bearing neurons did not exhibit elevated calcium levels compared with nearby non–NFT-bearing neurons (n = 13–110 NFT-bearing neurons (per mouse), n = 22–1006 non-NFT-bearing neurons (per mouse), n = 3 mice; P = 0.99 for NFT vs. non-NFT within each animal, Mann–Whitney U test; P = 0.24 across cohort, paired t test). In addition, we found no difference between resting calcium levels in NFT-bearing and non–NFT-bearing neurons in other areas of the somatosensory cortex (Fig. S6). For neuronal responsiveness to visual stimuli, we detected no difference in the probability of response (n = 142 NFT-bearing neurons, n = 1,602 non-NFT-bearing neurons, n = 3 mice; P = 0.39, Student t test), orientation selectivity, or direction selectivity (n = 24 NFT-bearing neurons, n = 122 non–NFT-bearing neurons, n = 3 mice; OSI, P = 0.38; DSI, P = 0.36, two-way ANOVA) (Fig. 3F–H). For OSI and DSI, we performed statistical analyses across the cohort of three animals (Avg) and for each individual animal (Ms1–Ms3), to reduce any cohort-level averaging that might have obscured potential differences (Materials and Methods).
All experiments were performed using 8- to 10-mo-old transgenic (t) and control mice (n) with a high NFT load and control mice (n). Moreover, there is no significant difference when comparing at a cell-specific level (t) and rTg4510 mice (n). Dashed lines indicate the possibility exists that a more sensitive assay that challenges NFT-bearing neurons are capable of integrating sensory inputs, axonal projections. Moreover, although our findings suggest that NFT-bearing neurons are characterized by abnormal accumulation of fibrillar tau aggregates in the cell soma and proximal dendrites compared with the predominantly axonal localization of tau in healthy neurons (24). To isolate the consequences of tau aggregation and mislocalization to soma and dendrites (6), we focused on the primary visual cortex, a brain area in which the major driving input source, the lateral geniculate nucleus, is structurally intact (10, 25) (Fig. S7). We found that in the visual cortex, neurons containing conspicuous quantities of mislocalized and aggregated tau nonetheless appear to have a normal capacity to integrate dendritic inputs and respond robustly to visual stimuli and also maintain normal somatic baseline calcium levels. In particular, we show that individual NFT-bearing neurons can respond robustly after integrating sensory inputs and are functionally indistinguishable from neighboring non-NFT-bearing neurons. These results demonstrate that NFT-bearing neurons remain functionally integrated in cortical circuits.

The possibility remains that network elements downstream of NFT-bearing neurons may show deficits linked to axonal or presynaptic deficits arising from NFT-bearing neurons. This property may be particularly relevant in neurons with longer axonal projections. Moreover, although our findings suggest that NFT-bearing neurons are capable of integrating sensory inputs, the possibility exists that a more sensitive assay that challenges NFT-bearing neurons may reveal subtle alterations in neuronal function. It also is possible that NFT deposition exerts deleterious effects on neural system function at later stages in the disease; however, similar assays as those used in the present study have identified dysfunctional phenotypes in amyloid mouse models with clearly impaired neuronal tuning in animals with amyloid plaques (20) and profound alterations in baseline resting calcium levels (15).

In combination with a previous study showing that NFT-bearing neurons in the hippocampus of rTg4510 mice can induce expression of the immediate early gene Arc after behavioral stimulation (26) and another study showing relatively normal electrophysiological properties in an acute slice preparation (13), our results strengthen the hypothesis that NFT deposition resembles an “off-pathway” disease side effect that in itself does not cause significant disruption of network function (9, 10, 13, 27–29). Mouse models are inherently limited in their ability to capture all aspects of adult human AD because of differences in pathology structure, kinetics, and distribution. Future work will need to explore the longitudinal impact of tau deposition on single-neuron, local circuit, and neural system function: how neurons and networks change before, during, and after NFT deposition, and how prefibrillar and soluble oligomeric tau aggregates impact neuronal function. These types of studies will benefit greatly from new in vivo staining compounds that differentially bind soluble, prefibrillar, and aggregated tau. Even within these limitations, however, our present findings call into question therapeutic strategies aimed at preventing or disrupting fibrillar tau deposits (30), which may indeed sequester more toxic soluble tau species (31, 32).

**Materials and Methods**

Animals. All experiments were performed using 8- to 10-mo-old transgenic rTg4510 mice overexpressing full-length human four-repeat tau (ON4R) carrying the frontotemporal dementia-associated P301L mutation (10). By age 7–8 mo, rTg4510 mice had developed a large number of NFTs in the cortex. Control animals lacked the human tau transgene.
Fig. 3. NFT-bearing neurons exhibit normal visual response properties. (A) In vivo imaging sites were coregistered with postmortem images obtained from cleared brains in which neurons were labeled for YC3.6 (immunolabeled with anti-GFP) and NFTs (staining with ThioS; detailed workflow in Fig. S4). (B) In vivo image showing neighboring NFT-free (blue) and NFT-bearing (red) neurons, including two visually responsive neurons (white circles). (Scale bar: 50 µm.) (C) The presence of NFTs (white arrowheads) was identified based on postmortem NFT staining in clear brain. (Scale bar: 50 µm.) (D) Resting calcium distributions of non-NFT-bearing and NFT-bearing neurons in the visual cortex show no significant difference (cohort-level, paired t test; n = 3 mice; P = 0.24; intra-animal, Mann-Whitney U test right-sided, n = 13-110 NFT-bearing neurons (per mouse), n = 22-1006 non-NFT bearing neurons (per mouse), P = 0.99). (E) Average response traces of a broadly tuned non-NFT-bearing neuron and an NFT-bearing neuron (white circles in B and C labeled 1 and 2, respectively). (F) The average percentage of responding neurons (mean ± SEM) was not significantly different between non-NFT-bearing (blue, 1602 cells) and NFT-bearing (red, 142 cells) neurons (P = 0.4, Student t test; n = 3 mice), but was highly variable among mice (Ms1 = 0.99]. (G and H) Average (mean ± SEM) OSI (G) and DSI (H) of non-NFT-bearing (blue) and NFT-bearing (red) neurons, analyzed across all mice (Avg) and for intramouse comparisons (Ms1–Ms3) (two-way ANOVA with Tukey–Cramer post hoc test; n = 24 NFT-bearing neurons, n = 122 non-NFT-bearing neurons; 3 mice; OSI: P = 0.4 for NFT, P = 0.3 for interaction; DSI: P = 0.4 for NFT, P = 0.3 for interaction).

Surgical Procedures. A small craniotomy was performed over the visual cortex (15, 33). An adeno-associated virus (AAV) vector encoding the ratiometric calcium indicator YC3.6 (AAV8-CBA-YC3.6; Penn Vector Core) was stereotactically injected for expression of YC3.6 in layer 2/3 neurons in the right primary visual cortex V1 (from lambda: +0.5 mm anteroposterior, 2.7 mm lateral, 0.6 mm dorsoventral). Above the injection coordinates, a cranial window was implanted replacing a circular piece (3 mm diameter) of skull by a glass coverslip that was secured in place using a mix of dental cement and Krazy Glue. A custom-made stainless-steel headpost (Ponoko) was affixed to the skull using C&B Metabond dental cement (Parkell), and each animal was allowed to recover for at least 2-3 wk. Before imaging, animals were habituated to the custom-made head-fixation device (Thorlabs posts and Altos head clamps), during which they were allowed to run freely on a circular treadmill (Ponoko).

In Vivo Calcium Imaging. Two-photon fluorescence of YC3.6 was excited at 860 nm using a mode-locked Ti:sapphire laser (MaiTai; Spectra Physics) and detected in the blue channel (CFP emission) and the yellow channel (YFP emission). Imaging was performed with a multiphoton imaging system (Fluoview1000; Olympus) equipped with a water immersion objective (25x, NA = 1.05; Olympus), with the emission path shielded from external light contamination. To record time courses of visually stimulated neuronal activity-dependent calcium levels, awake animals were head-fixed under the microscope, and an LCD monitor displaying the visual stimuli was placed in front of the left eye (screen–eye distance, ∼20 cm; screen-midline angle, 60°). Visual stimuli were drifting (2 Hz) sine-wave gratings (80% or 100% contrast; black and white) presented for ∼7 s. Stimuli were presented at 60 Hz using a calibrated 19-inch LCD monitor (Viewsonic VP9308). Eight stimuli (at 45° orientation increments) were presented sequentially in counter-clockwise order with ∼7 s between stimuli. During visual stimulation, several ∼300 µm² areas in layer 2/3 of V1 containing multiple YC3.6-expressing neurons were selected and imaged (scan rate, ∼2.5 Hz, 0.429 uframe; laser power, ≤50 mW).

Image Processing and Data Analysis. Semiautomated data analysis was performed using custom-written software in ImageJ (National Institutes of Health) and MATLAB (MathWorks). Images were aligned for minor shifts in the x-y plane owing to movement, cellular regions of interest (ROI) were manually drawn, YFP:CFP ratios were created and spatially filtered, and raw time courses were extracted. Images were excluded if there was a significant
shift in the z-axis (rare; <5% of all time courses). For each cellular ROI, the change in directional response (ΔR; R – \( R_{\text{baseline}} \)) was compared across directions and against background. And, “responsive” cells were identified (\( P < 0.05 \), ANOVA with the Tukey-Cramer post hoc test and Bonferroni correction). Only responsive cells were used for further analysis of orientation and direction tuning.

The OSI and DSI were calculated as described previously (34). In short, OSI was calculated as the maximum directional response, \( \Delta R_{\text{max}} \), divided by the sum of responses in all other directions, \( \Sigma \Delta R_{\text{nonmax}} \leq OSI \leq 1 \); 1 perfectly orientation-tuned. And, the DSI was calculated at \( \Delta R_{\text{max}} \) divided by the sum of that and the antiparallel direction, \( \Delta R_{\text{max}} + \Delta R_{\text{antiparallel}} \) (0 ≤ DSI ≤ 1; perfectly direction-tuned).

Statistical Analysis. All statistical analyses were performed in MATLAB. All datasets were first tested for normality (Lilliefors test), after which appropriate statistical tests were applied (e.g., t-test/two-way ANOVA for normally distributed data, Mann–Whitney U test for nonparametric data). For comparisons of NFT versus non-NFT neuronal tuning in Fig. 3, two-way ANOVA was used to explore fixed effects due to interanatomical and intra-animal differences. Data were analyzed by cell, grouped by mouse (i.e., M1, M2, or M3) and the presence of NFTs (i.e., NFT-bearing or non–NFT-bearing). For cohort-level analyses, the P value for the NFT group was used. For intra-animal comparisons, the interaction term was used (mouse x NFT presence) and the Tukey–Cramer post hoc test was used to explore intra-animal comparisons of NFT vs. non-NFT. The average P value for intra-animal comparisons of OSI and DSI was highly insignificant, P = 0.47.

Postmortem Whole-Brain Staining, Brain Clearing, and In Vivo Correlations. In a subset of rTg4510 mice, postmortem identification of NFTs was achieved using brain immunostaining followed by brain “clearing” (21). Immediately after in vivo recording of calcium levels, mice were perfused with 4% PFA in PBS, and brains were postfixed in 4% PFA for 1 d and then stored in 0.25% PFA at 4 °C. Whole brains were permeabilized and blocked in PBS containing 2% Triton-X100, 1% BSA, and 5% normal goat serum (NGS) for 2 h at room temperature. To immobilize Y3.6-expressing cells and neurons with hyperphosphorylated tau, brains were incubated with primary antibodies (chicken anti-GFP (Invitrogen, 1:250) and mouse PHF1 (1:250)) in PBS containing 1% Triton, 1% BSA, and 2% NGs at 4 °C for 1 d. Secondary antibodies (goat anti-chicken Alexa Fluor 488 and goat anti-mouse Alexa Fluor 594 (Invitrogen, 1:250)) were applied for 7 d at 4 °C for clearing, brains were stepwise saturated with glyceraldehyde for 80:20, 60:40, 40:60, and 20:80 PBS %:glycerol % for 5 h each at room temperature. Final clearing was achieved by incubation in a clearing solution of 53% benzyl alcohol, 45% glyceraldehyde, and 2% DABCO (wt/vt) for 10 d at room temperature. Clear brain multiphoton imaging (excitation λ = 800 nm) of previously in vivo imaged cells allowed for single cell reconstruction and correlation of PHF1-positive cells with measured calcium levels. To label non-NFT-bearing brain clearings, mice were postfixed in 4% PFA in PBS containing 0.025% Thio for 2 d at room temperature, followed by reclearing in clearing solution for 3 d. Reimaging the same cortical area enabled reconstruction of NFT positions in vivo imaging and correlation with calcium responses.

Immunolabeling of Visual Cortex Sections. Eight- to 9-mo-old rTg4510 and control mice were perfused with 4% PFA in PBS. Brains were postfixed for 4% PFA, 1–2 d at 4 °C, incubated for 2 d in 30% sucrose in PBS for cryoprotection, and then cut into 50-μm-thick coronal sections. Sections containing visual cortex were permeabilized with Triton-X100 in PBS, blocked in 5% NGs in PBS, and incubated in primary antibody (mouse anti-human tau (TAU-13; Abcam, 1:1,000)) in 5% NGs in PBS overnight at 4 °C. Secondary antibody (goat anti-mouse Cy3 (Invitrogen, 1:1,000)) was applied in 5% NGs in PBS for 1 h at room temperature. After washing in PBS, brain slices were mounted on microscope slides, and NFTs were stained with 0.025% Thio for 5 h at 5% ethanol for 8 min. This was differentiated in 80% ethanol for 30s, sections were washed with water for 3 min, and coverslips were mounted using DAPI containing mounting medium ( Vectashield).

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