Aluminum and Tau in Neurofibrillary Tangles in Familial Alzheimer’s Disease

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Abstract.
Background: Familial Alzheimer’s disease (fAD) is driven by genetic predispositions affecting the expression and metabolism of the amyloid-β protein precursor. Aluminum is a non-essential yet biologically-reactive metal implicated in the etiology of AD. Recent research has identified aluminum intricately and unequivocally associated with amyloid-β in senile plaques and, more tentatively, co-deposited with neuropil-like threads in the brains of a Colombian cohort of donors with fAD.

Objective: Herein, we have assessed the co-localization of aluminum to immunolabelled phosphorylated tau to probe the potential preferential binding of aluminum to senile plaques or neurofibrillary tangles in the same Colombian kindred.

Methods: Herein, we have performed phosphorylated tau-specific immunolabelling followed by aluminum-specific fluorescence microscopy of the identical brain tissue sections via a sequential labelling method.

Results: Aluminum was co-localized with immunoreactive phosphorylated tau in the brains of donors with fAD. While aluminum was predominantly co-located to neurofibrillary tangles in the temporal cortex, aluminum was more frequently co-deposited with cortical senile plaques.

Conclusion: These data suggest that the co-deposition of aluminum with amyloid-β precedes that with neurofibrillary tangles. Extracellularly deposited amyloid-β may also be more immediately available to bind aluminum versus intracellular aggregates of tau. Therapeutic approaches to reduce tau have demonstrated the amelioration of its synergistic interactions with amyloid-β, ultimately reducing tau pathology and reducing neuronal loss. These data support the intricate associations of aluminum in the neuropathology of fAD, of which its subsequent reduction may further therapeutic benefits observed in ongoing clinical trials in vivo.

Keywords: Aluminum in human brain tissue, amyloid-β, familial Alzheimer’s disease, neurofibrillary tangles, senile plaques, tau

INTRODUCTION

The microtubule-associated protein tau in its normal physiological form supports the intracellular assembly of microtubules, stabilizing the neuronal cytoskeleton [1, 2]. Human tau is a natively unfolded protein with little secondary structure [3]. The six isoforms of human tau arise through alternative RNA splicing, which is defined by either three (3R) or four (4R) microtubule-binding domain repeats [3, 4]. Tau is predominantly found in neurons in the brain, though it is also expressed at lower glia levels [1, 2]. Tau binds to microtubules through its C-terminal domain with the N-terminus binding to cytosolic components, including the membrane-associated protein actin, the latter defining a role for tau in the organization of cytoskeletal networks [2, 3].
Post-translational modifications of tau and its phosphorylation by protein kinases, including glycogen synthase kinase 3β (GSK3β), have been shown to act as a critical mediator influencing both the function and stability of tau [1–3]. Under physiological conditions, the phosphorylation of tau modulates its binding to microtubules in competition with kinesin and dynein motors, thereby regulating the anterograde and retrograde transport of cellular cargoes, respectively [2, 5]. In healthy neuronal cells, tau is typically found along a concentration gradient, increasing toward the axonal terminals of the cells as maintained by the tau diffusion barrier of the axon initial segment (AIS) that acts to modulate against the retrograde transport of tau towards neuronal dendrites and cell soma [3, 6]. The low levels of tau found in axons are suggested to promote long-term potentiation via the strengthening of synaptic terminals [3, 7].

The excessive or hyperphosphorylation of tau under pathophysiological conditions have defined the tauopathies broadly encompassing neurodegenerative conditions, including Alzheimer’s disease (AD) [1]. The subsequent breakdown of the AIS that follows allows for the redistribution of tau from axons allowing its retrograde diffusion into the somatodendritic compartment of neurons [3, 6]. In combination with tau detachment from microtubules upon hyperphosphorylation, increasing intraneuronal concentrations of the protein propagate from the cell soma to dendrites causing the concomitant loss of dendritic spines and neurite outgrowth [1, 3]. The increased accumulation and hyperphosphorylation of tau lead to the formation of neurofibrillary tangles (NFTs) of the insoluble aggregated protein [1]. In concert with synaptic dysfunction and loss, the depolymerization of microtubules disrupts the cytoskeletal network, thus acting to drive the progressive and mechanistic loss of affected neurons [1, 5].

In addition to the large scale loss of neurons in the frontal cortex and astrogliosis, intracellular NFTs of tau protein and extracellular senile plaques of the amyloidogenic amyloid-β (Aβ) peptide constitute the principal neuropathological hallmarks of AD [8–10]. While sporadic forms of AD typically occur in patients aged 65 and over, familial Alzheimer’s disease (fAD) occurs at an earlier age of onset, driven by autosomal dominant mutations in the presenilin 1 (PSEN1) and 2 (PSEN2) proteins and the transmembrane amyloid-β protein precursor (AβPP) [11–13]. Aβ42 is the principal pathogenic peptide located in senile plaque cores [15, 16]. Mutations in PSENs are known to increase the production of the Aβ42 peptide via the enhanced and sequential proteolytic cleavage of AβPP via β-site APP cleaving enzyme 1 (BACE1) and γ-secretases [10, 12, 13].

Insoluble and aggregated forms of Aβ in senile plaques and hyperphosphorylated tau in NFTs dominate the pathological landscape of familial and sporadic AD brains. However, oligomeric forms of amyloidogenic proteins are thought to underlie neurotoxicity in vivo [17]. Therefore, such has continued to lend support to an updated and genetically driven view of the amyloid cascade hypothesis of AD, in which the deleterious effects observed in brain parenchyma have been cemented in the studies of fAD [11–13, 18]. In more recent studies, Aβ has been shown to modulate AIS function through its ability to activate its tethered GSK3β protein kinase [3, 6]. The downstream phosphorylation of non-tau targets that follows has been implicated in increased concentrations of endogenous and exogenous tau in the somatodendritic compartment of neurons due to impaired microtubule dynamics in the AIS [6, 19].

Furthermore, the ability of Aβ to synergistically activate cofilin, an enzyme that acts to depolymerize actin filaments (F-actin), results in the remodeling of cytoskeletal networks, causing the collapse of the AIS [6]. Such demonstrates the ability of Aβ and tau to synergistically drive the structural disintegration of neurons in the AD brain [1–3, 5, 6]. Interestingly, cross-seeding between Aβ and tau promotes aggregation of amyloidogenic β-pleated sheets and NFTs, forming mutated and stable complexes rich in aggregation-prone hydrophobic interactions [20, 21]. Those complexes formed are suggested to be aberrant and hence contributory to AD pathology [20].

Aluminum is non-essential yet in its free solvated and trivalent form, Al^{3+}_{aq}, is biologically reactive [22]. Our previous study of the brains of fAD donors carrying a PSEN1-E280A (Glu280Ala) mutation found aluminum co-localized with fibrillar Aβ in senile plaques [22, 23]. Donors in this kindred are unique in that this mutation is found in large numbers in the Antioquia region of Colombia [24]. Individuals with fAD carrying this mutation develop elevated cortical Aβ at an early age of onset of <50 years. The high brain aluminum content of fAD donors versus statistically lower amounts found in non-neurologically impaired individuals has highlighted the potential for aluminum to act as a nidus to overcome the precipitation of Aβ [23, 25].
In addition to a role for aluminum in senile plaque formation in fAD, we have investigated the potential for aluminum to bind to NFTs of hyperphosphorylated tau protein in the same kindred [26]. The large number of phosphate groups available on the surface of NFTs would be expected to bind intracellular aluminum with high avidity [27]. Herein, we have performed phosphorylated (Ser202 and Thr205) tau-specific immunolabelling to unequivocally identify NFTs in the brain tissue of Colombian fAD donors, carrying a PSEN1-E280A mutation [28]. The unequivocal identification of NFTs has been followed by aluminum-specific fluorescence microscopy to probe the potential preferential binding of aluminum to senile plaques or NFTs in fAD brain tissue.

**MATERIALS AND METHODS**

**Human brain tissue**

Human brain tissue consisting of formalin-fixed paraffin-embedded (FFPE) blocks were obtained following ethical approval from Universidad de Antioquia, Medellin, Colombia. Donor brain tissue was received from anonymized donors with fAD, all carrying the PSEN1-E280A mutation and research was performed following ethical approval obtained from Keele University, UK (ERP 2391). FFPE brain tissue blocks of a 45-year-old female, 57-year-old female, and 60-year-old male, all carrying a PSEN1 E280A mutation, were selected for imaging, based upon representative depictions of aluminum and NFT co-localization across frontal, parietal, temporal and occipital lobes.

**Microtomy**

Human brain tissue blocks were cooled on wet ice for 10 min. Ribbons of adjacent tissue sections were prepared at a thickness of 6 μm by use of a Leica RM2025 rotary microtome fitted with Surgipath® DB80 LX low-profile stainless-steel blades (Leica Microsystems, UK). Sections were floated onto ultrapure water (conductivity < 0.067 μS cm⁻¹) at 40°C and transferred onto Polysine adhesion slides (Thermo Scientific, UK). Sections were dried overnight at ambient temperature. Excess wax was removed from tissue sections by heating at 62°C for 20 min. Sections were allowed to cool fully before dewaxing and rehydration procedures were performed.

Human brain tissue sections were dewaxed with gentle agitation in Histo-Clear (National Diagnostics, US) for 3 min. Removal of any remaining wax was performed via a second wash in fresh Histo-Clear for 1 min. Sections were transferred into pure (HPLC grade) ethanol (Fisher Scientific, UK) for 2 min. Rehydration of tissue sections was achieved by transfer through an ethanol gradient (95, 70, 50, and 30% v/v) for 1 minute in each solvent. Sections were twice washed in ultrapure water for 30 s to remove the remaining ethanol before staining.

**Lumogallion staining**

Rehydrated tissue sections were placed in Coplin jars and fully immersed in 1 mM lumogallion (4-chloro-3-(2,4-dihydroxyphenylazo)-2-hydroxybenzene-1-sulphonic acid, TCI Europe N.V., Belgium) in 50 mM PIPES, pH 7.4. For the assessment of autofluorescence, the adjacent serial section was immersed in 50 mM PIPES buffer, pH 7.4 only. Sections were incubated for 6 h at ambient temperature away from light. Sections were subsequently washed six times in fresh PIPES buffer, rinsed for 30 s in ultrapure water and mounted under glass coverslips with the aqueous mounting medium, Fluormount™ (Sigma Aldrich, UK). Lumogallion staining, ThS counter-staining (ca 0.075% w/v ThS in 50% v/v ethanol) [23, 26], and AT8 immunolabelling were performed and analyzed in sequence, following each labelling method. Every such staining method implemented was performed on the identical tissue section.

**Immunohistochemistry**

All reagents for immunohistochemistry (IHC) were purchased from Abcam, UK, unless otherwise stated. Following analysis of lumogallion-stained tissue sections, coverslips were removed via immersion in ultrapure water with gentle agitation overnight. Sections were rinsed thoroughly with ultrapure water, outlined with a hydrophobic PAP pen and transferred into a humidity chamber for all subsequent staining procedures. Thereafter, IHC was performed according to the manufacturer’s instructions. Briefly, sections were immunolabelled with an AT8 recombinant primary anti-tau monoclonal antibody (rabbit) against human tau phosphorylated on
Ser202 and Thr205 (0.472 mg/mL: Abcam, UK: ab210703).

Antigen retrieval was performed by adding Proteinase K antigen retrieval solution to sections, followed by incubation at ambient temperature for 7 min. Sections were rinsed in 1 X Tris-buffered saline (TBS), followed by further rinses in TBS plus 0.025% v/v Triton X-100. Blocking was achieved by incubating sections in 10% w/v normal goat serum with 1% w/v bovine serum albumin (BSA) in TBS for 2 h at ambient temperature. Pre-optimized dilutions (1:1,000) of the unlabeled primary AT8 antibody in TBS, including 1% w/v BSA, were applied to sections and incubated overnight at 4°C to ensure complete saturation of antibody binding.

All staining procedures were performed at ambient temperature. Following AT8 labelling, sections were rinsed with TBS, including 0.025% v/v Triton X-100. Endogenous tissue peroxidase activity was blocked by incubation in 0.3% v/v H2O2 in TBS for 15 min. Sections were subsequently stained using a rabbit-specific horseradish peroxidase (HRP) 3,3′-diaminobenzidine (DAB) ABC IHC staining kit. A biotinylated goat anti-polyvalent secondary antibody was added to sections and incubated for 10 min. Sections were washed in TBS, and HRP-labelled streptavidin was applied for 10 min. After washing with TBS, DAB chromogen (30 µL DAB chromogen added to 1.5 mL DAB substrate) was applied to sections and developed by incubation for 10 min. Sections were washed in TBS and finally mounted under glass coverslips using the aqueous mounting medium, Faramount™ (Agilent Dako, UK).

**Microscopy**

Brightfield and fluorescence micrographs were acquired using an Olympus BX50 fluorescence microscope (mercury source) equipped with a vertical illuminator and BX-FLA reflected light attachment. A U-MNIB3 fluorescence filter cube (bandpassλ_{ex}: 470–495 nm, dichromatic mirror: 505 nm, longpassλ_{em}: 510 nm) was used to acquire lumogallion fluorescence (Olympus, UK). Images were captured using an Olympus DP74 (CMOS processor) color camera and the OLYMPUS cellSens (Standard 3.1) software suite. Optimal exposure times for collecting lumogallion fluorescence were set automatically by the superfluorescence (SFL) mode of the Olympus DP74 camera. The same optimized exposure settings were then subsequently fixed for each respective and corresponding autofluorescence micrograph.

Merging of fluorescence and brightfield channels was achieved using Photoshop (Adobe Systems Inc., US).

**RESULTS**

Aluminum and neurofibrillary tangles are found co-localized in familial Alzheimer’s disease

IHC labelling using an AT8 primary antibody demonstrated the presence of flame-like NFTs in predominantly cortical brain regions of Colombian fAD donors. Optimization of the IHC staining method using progressive serial dilutions of the AT8 primary antibody improved the signal to noise ratio sequentially. IHC of the parietal cortex of a 60-year-old male Colombian fAD donor using serial dilutions of the primary AT8 antibody of 1:40, 1:80, 1:200, and 1:1,000 identified NFTs via dark brown DAB staining at every antibody dilution tested. IHC performed in the absence of the AT8 primary antibody produced no detectable DAB staining (see Supplementary Figure 1).

In the temporal cortex of a 45-year-old female Colombian fAD donor, aluminum was found in neuronal cells close to extracellular deposits of aluminum as identified by an intense orange fluorescence emission (Fig. 1A) [26]. Punctate fluorescence was especially noted in the cell soma of neuronal cells and was readily differentiated from the yellow/green fluorescence of cytosolic lipofuscin deposits in neighboring cells (Fig. 1A) [26]. AT8 immunolabelling for phosphorylated tau revealed the dense and intracellular immunoprecipitation of DAB via dark brown staining (Fig. 1B).

Merging of the fluorescence (Fig. 1A) and brightfield (Fig. 1B) channels revealed the presence of intraneuronal aluminum and NFTs in a single neuron (Fig. 1C). Subsequent merging of the original ThS fluorescence channel [26] demonstrated the presence of a peripherally located intraneuronal green fluorescence emission, overlaying DAB immunostaining of phosphorylated tau. Higher magnifications demonstrated co-localization of intracellular particulate aluminum deposits and AT8-reactive phosphorylated tau (Fig. 1D). Lumogallion-reactive aluminum was also observed in neuronal cells within pia mater of the same donor that produced an orange fluorescence emission versus green autofluorescence of the non-stained section (see Supplementary Figure 2). Similarly, the merging of fluorescence and
brightfield channels identified aluminum co-localized with AT8-reactive DAB staining in the same neuron (see Supplementary Figure 2).

Co-localization of aluminum and phosphorylated tau predominates in the temporal cortex

Lumogallion staining of the temporal cortex of a 57-year-old female Colombian fAD donor revealed lumogallion-reactive aluminum in pyramidal neurons, as highlighted via an intracellular orange fluorescence emission (Fig. 2A). Intracellular lipofuscin was frequently observed as punctate yellow fluorescence in the cytoplasm of neuronal cells in both lumogallion-stained and non-stained tissue sections for autofluorescence. The yellow fluorescence emission of intracellular deposits of lipofuscin was readily distinguished from the uniform green autofluorescence of the adjacent serial section (Fig. 2C). Subsequent AT8 immunolabelling of the identical lumogallion-stained tissue section revealed dark brown DAB precipitation appearing as a classical flame-like NFT in a single pyramidal neuron (Fig. 2B). Merging of lumogallion fluorescence (Fig. 2A) and the brightfield channel (Fig. 2B) revealed that intracellular aluminum was occasionally found co-localized with AT8-reactive tau in the cell soma of pyramidal neurons (Fig. 2D).

Similarly, lumogallion-reactive aluminum was detected in neuronal cells in pia mater in the temporal cortex of the same donor, as highlighted via an intense orange fluorescence emission (Fig. 3A). AT8 counter-immunolabelling of the identical tissue section produced brown intraneuronal DAB
Fig. 2. Intracellular aluminum in pyramidal neurons in the temporal cortex of a 57-year-old female Colombian donor with familial Alzheimer’s disease. A) Intracellular aluminum (Al) in pyramidal neurons (orange). B) AT8 immunoreactive phosphorylated tau (pTau) in a single neuron (black arrow) located via DAB staining (brown) and the absence of staining in an adjacent neuron (white arrow). C). Autofluorescence (green) of the non-stained adjacent section highlighting deposits of intracellular lipofuscin (yellow). D) Merging of lumogallion and brightfield channels depicting aluminum and pTau in an identical neuron (black arrow) and the absence of DAB staining in an adjacent neuron (white arrow). Asterisks denote magnified inserts. Magnification: X 400, scale bars: 50 µm.

precipitation of an NFT-like morphology (Fig. 3B). The adjacent non-stained serial section yielded a green autofluorescence emission with yellow punctate lipofuscin deposits identified intracellularly (Fig. 3C). Merging of the lumogallion and brightfield channel demonstrated the co-localization of aluminum and AT8-reactive tau in the same neuron (Fig. 3D).

The co-localization of aluminum and immunoreactive tau was not restricted to the temporal lobe. Such deposition was also noted in the parietal (see Supplementary Figure 3) and frontal cortex (see Supplementary Figure 4) of the same 57-year-old female Colombian fAD donor. In both cases, aluminum was found intracellularly in neurons close to the pia mater. The orange fluorescence intensity produced was generally weaker in neurons in the frontal lobe, though markedly brighter than green autofluorescence typically noted for non-stained sections. AT8 immunolabelling produced dark brown staining, resembling intracellular NFTs. In the identical neurons, aluminum was found co-localized with AT8-labelled DAB-reactive tau deposits. In the occipital lobe, few neurons produced a positive lumogallion fluorescence signal for aluminum above background autofluorescence.

Additional brain tissue sections of the temporal lobe of fAD donors were investigated to support observations of aluminum-rich NFTs. In the temporal cortex of a 60-year-old male Colombian fAD donor, intracellular lumogallion-reactive aluminum produced an orange fluorescence emission that exhibited a flame-like morphology (Fig. 4A). IHC for phosphorylated tau revealed intracellular dark brown DAB staining in the same neuron, appearing as a characteristic flame-like NFT (Fig. 4B). Similar lipofuscin loading was noted in the same cell via punctate yellow autofluorescence of the non-stained adjacent tissue section (Fig. 4C). Merging of lumogallion fluorescence and the AT8 brightfield channel confirmed that both aluminum and immunolabelled tau were intraneuronal and co-localized, of which both adopted a flame-like structure, similar to that of NFTs (Fig. 4D). Such confirmed similar observations made
Fig. 3. Intraneuronal aluminum in the temporal cortex of a 57-year-old female Colombian donor with familial Alzheimer’s disease. A) Intracellular aluminum (Al) in a neuron (orange). B) AT8 immunoreactive phosphorylated tau (pTau) located via DAB staining (brown). C) Autofluorescence (green) of the non-stained adjacent section highlighting occasional deposits of intracellular lipofuscin (yellow). D) Merging of lumogallion and brightfield channels depicting aluminum and pTau in the identical neuron. Asterisks denote magnified inserts. Magnification: X 400, scale bars: 50 μm.

of aluminum co-localized with DAB-reactive NFTs, in different cortical regions of the same lobe (see Supplementary Figure 5).

Discussion

We have demonstrated the neuronal co-localization of aluminum and phosphorylated tau in NFTs in fAD donor brain tissue, carrying a PSEN1-E280A mutation. Aluminum-specific fluorescence microscopy utilizing the fluorophore lumogallion revealed the presence of aluminum in neurons...
Fig. 4. Intraneuronal aluminum in the temporal cortex of a 60-year-old male Colombian donor with familial Alzheimer’s disease. A) Intracellular aluminum (Al) in a neuron (orange). B) AT8 immunoreactive phosphorylated tau (pTau) located via DAB staining (brown) appearing as flame-like NFT. C) Autofluorescence (green) of the non-stained adjacent section highlighting occasional deposits of intracellular lipofuscin (yellow). D) Merging of lumogallion and brightfield channels depicting aluminum and pTau co-localized in the identical neuron. Asterisks denote magnified inserts. Magnification: X 400, scale bars: 50 µm.

Co-deposited with neuropil-like threads [26]. The latter was previously detected by the use of the benzothiazole-based dye, thioflavin S (ThS), which binds to β-pleated sheet-like moieties, including mature NFTs and Aβ fibrils in senile plaques [29]. While a key benefit of ThS labelling is its ability to produce fluorescence only with mature and pathologically-relevant misfolded amyloidogenic proteins, such is not specific to hyperphosphorylated tau in NFTs [29].

AT8 is an anti-tau primary antibody that recognizes the well-characterized Ser202 and Thr205 epitopes of phosphorylated tau [28]. Owing to the clear DAB labelling above background at 1:1,000 dilutions of the primary AT8 antibody, this concentration was used for all subsequent IHC experiments identifying NFTs. In cortical brain regions, AT8 revealed intracellular pathological NFT inclusions. As the most prominent of the AT-series monoclonal antibodies directed against phosphatase-sensitive epitopes, AT8 labels both intracellular and extracellular NFTs as distinguished by the presence of collapsed dendrites [30]. These late-stage or ghost-like tangles that form extracellularly following apoptosis of the affected neuronal cell show the distinctive absence of a nucleus with no discernable cytoplasm [29–31]. Late-stage ghost NFTs frequently possess a central void. Their characteristic presence adopts tangles of a paired helical filament morphology, located in the somatodendritic compartments of the ruptured cells [3, 30, 31].

Herein, we identified AT8-immunoreactive ghost NFTs co-localized with aluminum, confirming our previous observations of ThS-reactive tangles in identical neurons in the temporal cortex of a 45-year-old female fAD donor [26]. As expected, the use of IHC to detect phosphorylated tau was more sensitive than fluorescent staining using ThS and showed improvements in visualizing the various stages of tau tangle formation [29, 32]. Aluminum was observed co-located with NFTs across multiple brain regions, including the frontal, temporal, and parietal cortex, of which fewer positive staining examples were found in the occipital cortex.

Of all lobes investigated, aluminum was found most commonly co-located in NFTs in the temporal
Fig. 5. Aluminum in a senile plaque in the temporal cortex of a 60-year-old male Colombian donor with familial Alzheimer’s disease. A) Aluminum (Al) staining of a senile plaque (orange). B) AT8 immunoreactive phosphorylated tau (pTau) located via DAB staining (brown) in dystrophic neurites (arrows) at the periphery of the identical senile plaque. C) Autofluorescence (green) of the non-stained adjacent section highlighting occasional deposits of intracellular lipofuscin (yellow). D) Merging of lumogallion and brightfield channels depicting aluminum and pTau (arrows) in the identical senile plaque. Asterisks denote magnified inserts. Magnification: X 400, scale bars: 50 μm.

The temporal cortex is of the first brain regions in which tau is known to propagate as NFTs. The expansion of NFT pathology in the brain is graded via the Braak stages of severity, of which 0-IV ascribe the confinement of NFT pathology to the medial temporal lobe [33, 34]. At stage I, NFT pathology is confined to the transentorhinal region, which migrates to deeper entorhinal regions and into the hippocampus at stage II. Widespread hippocampal lesions are observed at stage III, spreading to the medial temporal gyrus at stage IV [34]. Later stages (V and VI) of disease progression are marked by NFT deposition in the striate and parastriate regions of the occipital cortex [34].

In living fAD donors carrying a PSEN1-E280A mutation, positron emission tomography has revealed that NFT formation ensues ten to fifteen years after marked increases of soluble tau are typically detected in the cerebrospinal fluid [35, 36]. Subsequent onset to severe dementia arises approximately seven years later, marking tau pathology as an essential biomarker of early disease progression. As a consequence of the late onset of tau pathology in the PSEN1-E280A brain, such may account for the early and widespread distribution of NFTs in the temporal cortex and perhaps their association with aluminum [35, 36].

Elevated soluble cortical Aβ in the same kindred is mirrored by widespread senile plaque deposition, preceding NFT formation in vivo [24, 35, 36]. Therefore, the abundance of extracellular Aβ depositing as β-pleated sheets in senile plaques would provide numerous binding sites for aluminum before potential competitive interactions with the intracellular phosphorylated binding sites available in NFTs [27]. Such a mechanism may account for the high levels of aluminum observed in the brain tissue of fAD donors carrying a PSEN1-E280A mutation versus donors with neither a clinical nor pathological diagnosis of neurodegenerative disease [23, 25, 37].

Interestingly, while numerous ghost NFTs were present in the cerebral cortex of fAD donors that produced positive DAB staining for phosphorylated tau, their co-localization with aluminum was detected less frequently. While immunoreactive phosphorylated tau was found co-located with aluminum in cortical neurons, they were fewer in incidence.
than co-deposits of aluminum in the core of extracellular senile plaques. Therefore, AT8 immunolabelling confirmed the preferential binding of aluminum exhibited in senile plaques versus NFTs [23, 26]. These data suggest that the co-deposition of aluminum with Aβ precedes that with NFTs. Extracellularly deposited Aβ may also be more immediately available to bind aluminum versus intracellular aggregates of tau. AT8 immunoreactive tau was found deposited at the periphery of senile plaques in both intracellular neuronal bodies and extracellular ghost NFTs. Lumogallion-reactive aluminum was frequently observed in the cores of neuritic plaques in which its deposition was less evident within surrounding tau-reactive neurons.

In addition to AT8-labelled dystrophic neurites, lipofuscin was also observed deposited at the periphery of senile plaques. As a marker of lysosomal degradation, autofluorescent lipid and misfolded protein-rich deposits are known to accumulate in the cytoplasm of neurons [38]. Aggregates of lipofuscin have previously been suggested to play an active role in neurodegeneration [38, 39]. As a marker of impeded lysosomal function and proteolytic activity, the release of lipofuscin into the extracellular milieu following neuronal death has been implicated in the formation of senile plaques [39].

Failure in the misfolded protein response promotes the formation of lipofuscin and its subsequent release from dying cells, and hence, the presence of lipofuscin in brain tissue sections increases upon ageing [38]. Such may explain the intimate associations observed between Aβ and coronal lipofuscin in senile plaques [37, 38, 40, 41]. In concert with Aβ, co-aggregated lipofuscin deposits bind aluminum that may act further to exacerbate the accumulation of aluminum in senile plaque cores, as has been previously reported in AD brain tissue [23, 37, 42]. Conversely, intracellular aluminum found in neurons heavily laden with lipofuscin was not always found co-located with NFTs. Therefore, intracellular lipofuscin presence may not necessarily dictate aluminum binding as was observed herein in pyramidal neurons in the temporal cortex of a 57-year-old female Colombian fAD donor.

AT8-immunoreactive phosphorylated tau protein and its silver-impregnated derivatives have previously been reported at the periphery of cored neuritic plaques in the hippocampus of AD donors and within dystrophic neurites surrounding senile plaques in the neighboring entorhinal cortex [43–45]. Early observations of Braak staging of NFT pathology in AD highlighted the presence of AT8-reactive NFTs in the amygdala of elderly (aged ≥ 70 years) AD donors [33]. The appearance of tau depositing as NFTs has been predominantly reported in pyramidal neurons upon labelling for phosphorylated tau. Therein, hyperphosphorylated tau filaments were observed near Aβ in approximately a third of AT8-immunolabelled senile plaques [43]. Furthermore, the co-localization of tau in senile plaques was not only restricted to AT8-reactive tau, as additional immunoreactivity to the phosphorylated-tau Ser396 isoform was always detected [43]. Therefore, our observations of coronal tau surrounding senile plaques would be expected to originate from phosphorylated forms of the protein in either dystrophic neurites or depositing as NFTs.

In summary, we have demonstrated the unequivocal association of aluminum and phosphorylated tau in NFTs in fAD. While aluminum binding to Aβ in senile plaques is expected in early disease progression concomitant with elevated amyloidogenic protein levels, an association with tau may occur at later disease stages [23, 24, 26, 35, 36]. Senile plaques containing dystrophic neurites are associated with neuronal death in AD, of which the ensuing neurotoxicity may be synergistically mediated by the co-existence of Aβ and aluminum in these neuropathological hallmarks [46–49]. Indeed, the latter potentiates the redox cycling of iron in the presence of Aβ2, generating reactive oxygen species, including the superoxide radical anion [50]. The subsequent extracellular release of NFTs from dying neurons may allow for a mechanism of aluminum binding in senile plaques to phosphorylated tau, exposed in filamentous regions of the misfolded protein [42]. Therapeutic approaches to reduce tau have been suggested to ameliorate its synergistic interactions with Aβ, reducing tau pathology and subsequent neuronal loss [51]. These data support the intricate associations of aluminum in the neuropathology of fAD, of which its subsequent reduction may further therapeutic benefits observed in ongoing clinical trials, in vivo [52].

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CONFLICT OF INTEREST

The authors have no conflict of interest to report.

SUPPLEMENTARY MATERIAL

The supplementary material is available in the electronic version of this article: https://dx.doi.org/10.3233/ADR-210011.

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