









# BRAIN COMMUNICATIONS

## REVIEW ARTICLE

# The role of neurofilament light in genetic frontotemporal lobar degeneration

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Genetic frontotemporal lobar degeneration caused by autosomal dominant gene mutations provides an opportunity for targeted drug development in a highly complex and clinically heterogeneous dementia. These neurodegenerative disorders can affect adults in their middle years, progress quickly relative to other dementias, are uniformly fatal and have no approved disease-modifying treatments. Frontotemporal dementia, caused by mutations in the *GRN* gene which encodes the protein progranulin, is an active area of interventional drug trials that are testing multiple strategies to restore progranulin protein deficiency. These and other trials are also examining neurofilament light as a potential biomarker of disease activity and disease progression and as a therapeutic endpoint based on the assumption that cerebrospinal fluid and blood neurofilament light levels are a surrogate for neuroaxonal damage. Reports from genetic frontotemporal dementia longitudinal studies indicate that elevated concentrations of blood neurofilament light reflect disease severity and are associated with faster brain atrophy. To better inform patient stratification and treatment response in current and upcoming clinical trials, a more nuanced interpretation of neurofilament light as a biomarker of neurodegeneration is now required, one that takes into account its relationship to other pathophysiological and topographic biomarkers of disease progression from early presymptomatic to later clinically symptomatic stages.

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Received June 16, 2022. Revised September 26, 2022. Accepted November 23, 2022. Advance access publication November 26, 2022

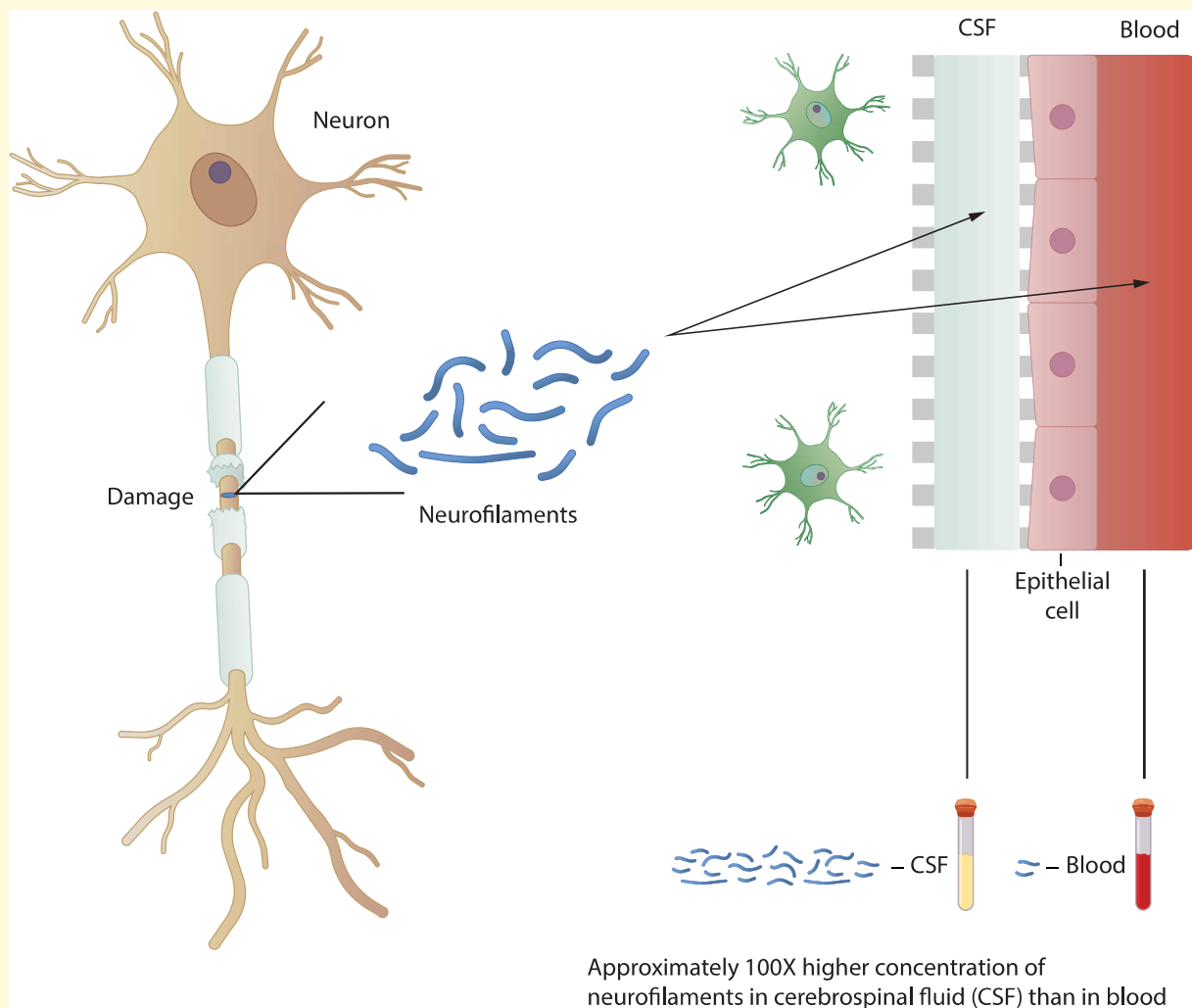
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**Keywords:** frontotemporal lobar degeneration; neurofilament light; trial stratification; disease progression; frontotemporal dementia

**Abbreviations:** ALLFTD=advancing research and treatment for frontotemporal lobar degeneration/longitudinal evaluation of familial frontotemporal dementia subjects; ALS=amyotrophic lateral sclerosis; ARC=annualized rate of change; bvFTD=behavioural variant frontotemporal dementia; *C9orf72*=chromosome 9 open reading frame 72 gene; FDA=US Food and Drug Administration; FTD=frontotemporal dementia; FTD-*C9orf72*=frontotemporal dementia caused by the *C9orf72* gene mutation; FTD-GRN=frontotemporal dementia caused by the progranulin gene mutation; FTLD=frontotemporal lobar degeneration; GENFI=genetic frontotemporal dementia initiative; GM=grey matter; GRN=granulin gene; m=messenger; MAPT=microtubule-associated protein tau gene; MD=mean diffusivity; NEFL=neurofilament light gene; NF=neurofilament; NFH=neurofilament heavy; NfL=neurofilament light; NFM=neurofilament medium; sAPP $\beta$ =soluble peptide amyloid precursor protein beta; TDP43=transactive response DNA-binding protein 43; WM=white matter

### Graphical Abstract



Neurofilament light (NfL) is a surrogate marker of neuroaxonal degeneration that is increased following acute injury, with age, and in neurodegenerative disease

## Introduction

This review is the synthesis of a meeting convened by Alector, Inc., and the Bluefield Project to Cure Frontotemporal

Dementia in December 2020. Key opinion leaders from academia with expertise in genetic frontotemporal dementia (FTD), multiple sclerosis, biomarkers and drug development, and the use of neurofilament light (NfL) as a biomarker in

neurodegenerative diseases were invited speakers. Meeting attendees (list provided in manuscript addendum) represented various companies in addition to clinical and basic scientists with interest in biomarkers and drug development for neurological diseases.

Frontotemporal lobar degeneration (FTLD) underlies a group of clinically, genetically and pathologically diverse clinical syndromes involving proteinopathies that are associated with dysfunction of the frontotemporal networks. FTD refers to canonical presentations of FTLD at the symptomatic (dementia) stage. The clinical syndromes are often, but not always, associated with a progressive neurodegenerative pathology; however, prediction of the underlying pathology based on clinical presentation is problematic. These disorders are challenging to diagnose, particularly in the preclinical stage through advancement to the prodromal stage. Optimally, disease-modifying treatments should be tested at these earlier stages, while judgment is preserved, and individuals still retain their independence. Ideally, fluid and imaging biomarkers could provide information on the start of pathological changes, but in sporadic FTLD, biomarkers are currently used to support a diagnosis based on established clinical consensus criteria. In cases where there is a suggestive family history, an estimate of years to disease onset may be feasible, and genetic testing can be done to reveal a causal mutation to support the diagnosis. Fluid and imaging markers could then be employed to determine the sequence of events contributing to disease pathobiology and their correlation with clinical progression.<sup>1</sup>

Autosomal dominant mutations in the progranulin (*GRN*), chromosome 9 open reading frame 72 (*C9orf72*) and microtubule associated protein tau (*MAPT*) genes are recognized as the major causes of the highly heritable forms of genetic FTLD. It is estimated that these three genes are responsible for approximately half of familial FTD cases and approximately 10–30% of all FTD cases, whereas the remainder is considered sporadic.<sup>2</sup> FTLD is pathologically associated primarily with either misfolded transactive response (TAR) DNA-binding protein 43 (TDP43) or misfolded tau protein aggregates in neurons. Longitudinal data from the Genetic FTD Initiative (GENFI; [www.genfi.org](http://www.genfi.org)) and Advancing Research and Treatment for Frontotemporal Lobar Degeneration/Longitudinal Evaluation of Familial FrontoTemporal Dementia Subjects (ALLFTD; [www.allftd.org](http://www.allftd.org)) networks,<sup>3,4</sup> as well as a network of French amyotrophic lateral sclerosis (ALS) and FTD clinics,<sup>5</sup> suggest that NfL may be a valuable blood-based biomarker, reporting that NfL baseline levels and rate of change with disease progression differ among the FTLD gene mutation subtypes as compared with healthy controls.

Several clinical trials testing experimental, potentially disease-modifying treatments are underway for FTD caused by the progranulin gene mutation (FTD-GRN), employing unique strategies to restore progranulin levels, including a monoclonal antibody trial to block the progranulin/sortilin pathway (NCT04374136), two separate gene therapy trials using adeno-associated viral vectors (NCT04408625;

NCT04747431), and a protein replacement therapy trial (NCT05262023). Interventional trials for FTD caused by the *C9orf72* gene expansion (FTD-*C9orf72*) have also begun and include a novel antisense oligonucleotide trial (NCT04931862) and a monoclonal antibody trial (NCT03987295). Both FTD-GRN and FTD-*C9orf72* are characterized by a TDP43 proteinopathy. TDP43 is an RNA/DNA-binding protein involved in RNA metabolism that can aggregate in neuronal and glial cell cytoplasm.<sup>6</sup> It is assumed that increases in blood NfL levels may signal an undesirable change based on its steady increase with aging, as well as its elevation in acute injury and chronic neurological diseases when compared with healthy control populations.<sup>7,8</sup> Interpretation of short-term changes in absolute levels of blood-based NfL in healthy populations and clinical trials of disease-modifying therapies for FTLD disorders is an emerging need. Longitudinal NfL dynamics relative to the time course of structural changes in FTLD to improve our comprehension of advancing neurodegeneration through the presymptomatic, proximity to conversion (prodromal), and clinically symptomatic stages of genetic FTLD are also needed, in addition to other pathophysiological markers. Additionally, a better understanding is required of NfL dynamics and turnover (e.g. the potential role of microglia in NfL clearance) and the relationship between NfL and comorbidities such as subclinical brain injuries, cardiovascular events, diabetes, and prescribed medication use, which can all increase blood-based NfL levels.<sup>9,10</sup>

## Axonal damage after acute injury and in chronic neurodegenerative disease

Neurofilaments (NFs) are class IV intermediate filaments composed of three subunits of different molecular sizes: 68 kDa NfL, 160 kDa neurofilament medium (NFM), and 200 kDa neurofilament heavy (NFH), combined with  $\alpha$ -internexin as the fourth subunit in CNS axons, and function as one of the cytoskeletal elements of neurons.<sup>11</sup> NFs are released into the extracellular space following axonal injury and have been hypothesized to act as surrogate markers of neurodegeneration. Release of NFs, particularly NfL and NFH, into the CSF, and eventually the bloodstream is proposed to occur after axotomy and secondary axonal injury.<sup>11,12</sup> Some adult animal models show NfL accumulation before NFM and NFH, indicating an organized, temporal sequence of events postinjury.<sup>13</sup> Case studies in athletes have demonstrated that elevated NfL levels can persist at least 30 weeks in the CSF following some acute injuries,<sup>14,15</sup> arguing that absolute NfL levels, as well as duration of elevation, may reflect continuous axonal injury. An examination of acute, mild traumatic brain injury cases suggests that the delayed peak in plasma NfL levels after injury may reflect ongoing secondary injury or repair mechanisms.<sup>16</sup>

Post-mortem studies in ALS revealed that degenerating neurons had a 70% decrease in NfL mRNA, as well as decreases in  $\alpha$ -internexin and peripherin.<sup>17</sup> TDP43, the pathogenic intracellular protein aggregate common to ALS (*C9orf72*) and FTD (*C9orf72*, *GRN*), is able to bind and destabilize, or sequester, neurofilament light gene (*NEFL*) mRNA by direct interaction with the three prime untranslated region.<sup>18,19</sup> This has implications for altered RNA processing in ALS, which can ultimately impact NfL protein synthesis, stoichiometry, and aggregation.<sup>20</sup> Pathology studies have elucidated the trajectory of nerve cell degeneration in chronic neurodegenerative diseases, whereby misfolded proteins like TDP43 spread through a disease-specific neural connectome and aggregate in distinct neuronal populations, ultimately resulting in functional deficits and death.<sup>21</sup> It is not known whether specific messenger RNA-binding proteins that normally regulate *NEFL* RNA processing, trafficking, and local protein synthesis within axons and dendrites play an active role in chronic neurodegenerative diseases or secondary axonal injury.<sup>22</sup> Animal models have demonstrated that the specialized transcriptome in axons allows them to quickly alter the axonal proteome via local translation and protein degradation.<sup>23–25</sup> This capacity for local translation in axons can provide cells with the ability to respond to external cues that impact axon elongation, synaptic function, and maintenance during development and adulthood.<sup>26,27</sup>

## NfL as a biomarker of disease progression and drug efficacy in multiple sclerosis

Our most comprehensive understanding to date of how NfL may be implemented as a disease activity/progression marker and disease treatment efficacy marker comes from multiple sclerosis studies. Multiple sclerosis has a strong neuroaxonal injury component, which is the likely substrate of the observed clinical disability. The neuropathology includes axonal bulbs, transection, and demyelination of the axon,<sup>28</sup> with acute axonal degeneration appearing early in the disease course and most prominently within the first year after disease onset.<sup>29</sup> Chronic spinal plaques in multiple sclerosis reflect 58–68% axonal loss.<sup>30</sup> Although several anti-inflammatory treatments are available for relapsing multiple sclerosis, there is a struggle to understand if treatments suppress subclinical, ‘benign’ multiple sclerosis.<sup>31</sup> A precision medicine approach would be preferred to treat individuals, incorporating biomarkers as well as companion treatments that would lead to a demonstrable halt or slowed rate of neurodegeneration. Patients with progressive or relapsing multiple sclerosis show higher serum NfL levels than healthy controls, and the correlation is strong between CSF and serum NfL, with disease-modifying treatments demonstrating similar effect on NfL in both fluids.<sup>32</sup> Recent trials in progressive multiple sclerosis also demonstrate that increased

NfL levels are associated with ongoing disease activity, as well as long-term outcome.<sup>33</sup> A multisite, longitudinal study involving >10 000 serum samples from >5000 participants without evidence of CNS disease spanning over six decades of life is underway in an effort to generate age-adjusted reference values for NfL.<sup>34</sup> When compared with reliable, age-adjusted reference values, blood-based NfL may be considered a first-in-class biomarker for multiple sclerosis disease activity with prognostic value,<sup>35</sup> and some hope for a potential clinical application. Short-term neuroprotective studies using area under the curve analysis of serum NfL levels as an outcome for screening effective neuroprotective drugs for late phase 3 clinical development may prove a valuable concept. A Letter of Support was granted to the International Progressive multiple sclerosis Alliance (<https://www.progressivemsa.org/>) by the US Food and Drug Administration (FDA; <https://www.fda.gov/media/149608/download>) in June 2021 to encourage further study and use of NfL in plasma or serum as a pharmacodynamic/response biomarker for early clinical trials in progressive multiple sclerosis.

## Multidecade and individual variability of NfL over time in aging and dementia cohorts

Data collected every five years in the 30-year Betula study in Sweden reported that age-related changes in plasma NfL levels (measured by Simoa) are linked to brain white matter (WM) alterations using T1- and T2-weighted images.<sup>36</sup> Alzheimer’s disease cases showed elevated levels compared with controls, but no significant differences from the preclinical phase. NfL did not predict age-related cognitive impairment or impending Alzheimer’s disease. By contrast, the Dominantly Inherited Alzheimer Network modelled that serum NfL was elevated in the presymptomatic stages of familial Alzheimer’s disease; specifically, longitudinal, within-person analysis showed that the rate of change of NfL discriminated mutation carriers from non-mutation carriers almost 10 years earlier than cross-sectional, absolute NfL levels.<sup>37</sup>

The single-site Austrian Stroke Prevention Study monitored a cohort of 335 adults with no obvious neurological disorder ranging in age from approximately 38 to 85 years over an approximately six-year period for serum NfL changes using a single-molecule array (Simoa assay) with UmanDiagnostics anti-core domain antibodies.<sup>38</sup> Although median NfL levels were relatively similar among individuals in the fourth and fifth decades of this normal aging cohort, NfL rose non-linearly and demonstrated more variability in participants aged 60 and older. These data support the hypothesis that NfL increases with age, and those age-related degenerative processes may also contribute to NfL increase in the absence of overt clinical symptoms. Subclinical brain pathology with volume loss was supported by longitudinal

and cross-sectional T1-weighted MRI of brain volume in participants and showed a close correlation with serum NfL levels.

A survey of the Alzheimer Center Amsterdam clinics revealed that clinicians perceive serum NfL to be more useful in patients younger than 60 years to confirm or exclude neurodegenerative disease versus a psychiatric diagnosis.<sup>39</sup> As a result, NfL measures are now being requested to support an FTD diagnosis in this age group. A Shiny app has been developed to provide a quick comparison between disease readout and age-dependent reference values (<https://networkinstitute.org/2020/11/23/neurofilament-light-application-online>).

Across sporadic and genetic FTD, CSF NfL levels are higher compared to patients with Alzheimer's disease or controls.<sup>7,8,40</sup> Serum NfL levels follow a similar pattern to CSF, with FTD-GRN reported as higher on average (61.5 pg/ml) compared with FTD-C9orf72 (33.9 pg/ml). However, variation can be considerable in absolute blood NfL levels within all genetic FTD groups, as well as in control subjects. On occasion, non-carriers may have a higher baseline serum NfL than a presymptomatic or symptomatic carrier. As higher NfL levels decrease to normal in these few cases, repeated measurements over time may help to interpret these outliers.<sup>41</sup> In the short term, examining cohorts for variations between consecutive measurements of blood NfL levels will be valuable to understand what constitutes a clinically meaningful variation in untreated patients, patients undergoing treatment as part of their clinical care and management, and participants in clinical trials. One such study analyzed serum NfL at baseline and at six-month follow-up using the Simoa NF-Light kit in neurologically healthy controls, untreated relapsing multiple sclerosis, and secondary and primary progressive multiple sclerosis.<sup>42</sup> Elevation of NfL in healthy controls was positively associated with increased age, and at follow-up, NfL was elevated compared with baseline. Baseline NfL was elevated in multiple sclerosis compared with healthy controls but was lower at follow-up in both treated and untreated relapsing multiple sclerosis. The authors concluded that variations between two consecutive measurements within individuals occur to a similar extent in healthy controls and subjects with multiple sclerosis.

## Brain structural changes, NfL, and cognition in Alzheimer's disease dementia

There is a need to better understand NfL regulation in health and disease, particularly the relationship between NfL and cognitive decline, to support recommendations for the use of NfL as a surrogate biomarker in FTLD clinical trials. A scoping review of neurological disease literature revealed

that higher levels of NfL (CSF, plasma, or serum) were generally associated with decreased cognition.<sup>43</sup> However, inconsistencies in the relationship between elevated NfL and declining Mini-Mental State Examination performance were found within and across Alzheimer's disease and other dementias. There may be dynamic relationships among cognitive performance, structural changes in the cortex and NfL levels obtained from blood and CSF, which may vary over the course of different dementias. There is evidence to suggest that mean diffusivity (MD) can evaluate changes in grey matter (GM) at a microscopic level, reflecting neuronal loss and the breakdown of myelin, cell membranes, and organelles, which results in a measurable difference in the diffusion of water molecules. MD, a metric of diffusion weighted imaging, assesses diffusion in all directions and is therefore uniquely suited to measuring microstructural change in GM.<sup>44</sup> Two studies that examined cortical thickness and cortical MD in autosomal dominant Alzheimer's disease reported a biphasic trajectory of macro- and microstructural changes.<sup>45,46</sup> A close association was found between increasing MD and decreasing cortical thickness, suggesting that MD as a marker of microlevel cortical integrity may be part of the same disease pathology continuum as macrolevel neurodegeneration.<sup>45</sup> Importantly, increased cortical thickness and decreased cortical MD were observed about 15–20 years prior to estimated symptom onset, followed by cortical thinning and increased cortical MD in later prodromal and symptomatic stages of autosomal dominant Alzheimer's disease.<sup>46</sup> In a separate study, NfL in the CSF was measured in 221 participants of the Australian Imaging, Biomarkers & Lifestyle Flagship Study of Ageing and was found to be significantly elevated in Alzheimer's disease compared to healthy controls. Further, NfL levels predicted baseline cognition and cortical GM volume and were significantly correlated with tau pathology assessed by CSF total-tau and phosphorylated-tau.<sup>47</sup>

## Neurodegeneration associated with FTD-GRN and FTD-C9orf72

Mutations in the GRN gene produce haploinsufficiency, resulting in an approximately 50% decrease in progranulin protein levels and, ultimately, aggregation of misfolded TDP43 protein in vulnerable neurons. Blood progranulin levels are significantly reduced in pathogenic GRN mutation carriers compared with non-mutation carrier family members from childhood through adulthood in presymptomatic FTD-GRN. Thus, blood progranulin level provides a sensitive diagnostic biomarker, but does not appear to have a direct correlation with progressive brain atrophy.<sup>48</sup>

Age of onset is highly variable in FTD-GRN, ranging from 25 to 90 years, with a mean age of symptom onset of  $61 \pm 8.8$  years and a mean disease duration of 7.1 years ( $n = 548$ ).<sup>49</sup> GRN mutations appear to be nearly 100% penetrant by

80 years of age.<sup>50</sup> WM hyperintensities have been associated with FTD-GRN<sup>51,52</sup> but not with FTD-C9orf72. Neuropathological findings suggest that these WM hyperintensities are associated with microglial activation and microglial dystrophy.<sup>53</sup> GM atrophy and hypometabolism have been reported about 10–15 years prior to clinical symptom onset.<sup>54–57</sup>

FTD-C9orf72 is the most common genetic form of FTD in North America and Europe, responsible for approximately 13% of all FTD cases and approximately 11% of all ALS (ALS-C9orf72) cases; it can also cause a combination of ALS and FTD.<sup>58</sup> FTD-C9orf72 results from a hexanucleotide repeat expansion within the C9orf72 gene, and there is compelling evidence to support several different mechanisms that cause this repeat to result in TDP43 neuropathology and degeneration. Symptom onset can range from 20 to 90 years of age and the mean age of onset of FTD-C9orf72 is  $58.2 \pm 9.8$  years, with a mean disease duration of 6.4 years ( $n = 618$ ).<sup>49</sup> The pathogenic expansion is almost fully penetrant by 80 years of age.<sup>59</sup> Brain atrophy in FTD-C9orf72 carriers occurs earlier than in other mutation groups, appearing before age 40 and potentially 25 years before clinical symptoms.<sup>60</sup>

Structural imaging studies have demonstrated cortical thinning in presymptomatic FTD-GRN carriers compared with non-carriers.<sup>61,62</sup> A recent study comparing GRN carriers and non-carriers reported age-related cortical thinning in both groups but with significantly greater thinning in the presymptomatic GRN carriers.<sup>63</sup> Presymptomatic C9orf72 carriers also demonstrated greater cortical thinning compared with controls.<sup>64</sup> NfL has been considered to reflect WM changes, as NFs are primarily found in axons.<sup>65,66</sup> Longitudinal GM and WM changes were analyzed in presymptomatic GRN and C9orf72 carriers in GENFI cohorts over 2 years, and although C9orf72 carriers showed lower GM volume in the cerebellum and insula and WM differences in the anterior thalamic radiation compared with non-carriers, presymptomatic GRN carriers did not show presymptomatic GM or WM degeneration.<sup>67</sup> Of interest is a study that used diffusion tensor imaging to provide evidence of a correlation between plasma NfL in behavioural variant FTD (bvFTD) and WM changes.<sup>68</sup> Although it was a small study of 20 subjects, a reduction in fractional anisotropy was associated with increased plasma NfL concentration, an elevated CDR® plus NACC FTL D score, and a global level of degeneration in WM tracts that included the superior longitudinal fasciculus, the fronto-occipital fasciculus, the anterior thalamic radiation, and the dorsal cingulum bundle.

Cortical macrostructural changes, as measured by voxel-based morphometry or cortical thickness with surface-based analysis, have been the standard for neuroimaging in FTD. However, a recent multicentre study examined cortical MD and cortical thickness in a bvFTD cohort that included possible and probable bvFTD subgroups. In the whole bvFTD group, cortical MD and cortical thickness correlated with the CDR® plus NACC FTL D; but in the possible

bvFTD group, only cortical MD correlated with the CDR rating scale, suggesting that cortical MD may be a more sensitive biomarker for microstructural neurodegenerative changes that can occur earlier than macrostructural changes in disease trajectory. Only minimal cortical thinning was found in the possible bvFTD subgroup compared to controls. In the whole bvFTD group, cortical thickness and cortical MD both correlated with measures of disease severity and CSF biomarkers, but the areas of correlation with cortical MD were more extensive. CSF soluble peptide amyloid precursor protein beta (sAPP $\beta$ ), CSF NfL, and CDR measures had a more widespread correlation with cortical MD than cortical thickness.<sup>69</sup> Cortical MD was also found to be increased in mild primary progressive aphasia with GRN mutation, while only minimal cortical thinning was observed. Together, these data suggest that cortical MD may be more sensitive than cortical thickness in the detection of the earliest neurodegenerative changes in FTL D and that a biphasic trajectory of neurodegeneration should be considered.<sup>70</sup>

Data from the GENFI network revealed a relationship between NfL and executive function, rather than cognitive performance, in FTD.<sup>71</sup> And a recent case-control study supports a relationship between elevated CSF NfL in FTL D with known TDP43 pathology and a longitudinal decline in specific executive and language measures compared with controls. Further, it was shown that patients with FTL D-TDP pathology had significantly greater CSF NfL than those with FTL D-tau pathology.<sup>72</sup> Longitudinal studies of genetic FTL D that include pathophysiological markers (such as NfL in CSF and blood) combined with topographical markers of neurodegeneration are needed to better understand the relationship between NfL and FTL D. In particular, examination of presymptomatic, prodromal, and symptomatic gene mutation carriers could indicate the presence of a biphasic trajectory of neurodegenerative change in FTL D, inform on absolute cutoffs between stages, and reveal potential differences in disease trajectory across the different autosomal dominant gene mutations. Longitudinal studies could also provide insight into the dynamic relationship between NfL and other biomarkers, their correlation with clinical status, and what combinations of biomarkers might improve the prognostic value of NfL over the course of these complex, highly heterogeneous FTL D disorders.

## NfL rate of change in FTD-GRN versus FTD-C9orf72

In the GENFI and ALLFTD cohorts, the conversion (prodromal) period from clinically presymptomatic to symptomatic status is accompanied by the highest rate of change in serum NfL levels at any time during the disease, and GRN mutation carriers demonstrate a higher rate of NfL change than FTD-C9orf72 mutation carriers. Higher concentrations

of serum NfL reflect disease severity, as they are associated with faster rates of brain atrophy, and baseline NfL was significantly elevated in subjects who converted from presymptomatic to symptomatic or showed disease progression versus non-progressors. Overall, baseline plasma NfL was highly predictive of change in clinical status over the following 2 years. Furthermore, higher baseline NfL correlated with worse longitudinal clinical measures and brain atrophy in all FTD genetic mutation types, arguing that blood-based NfL is predictive of short-term risk for FTD disease progression.<sup>3,4,73,74</sup>

Longitudinal changes in plasma NfL in presymptomatic and symptomatic *GRN* and *C9orf72* mutation carriers and controls (aged 21–83 years) were measured in cohorts from the French Research Network on FTD/FTD-ALS (Inserm RBM02–59), Predict to Prevent Frontotemporal Lobar Degeneration and ALS (PREV-DEMALS), and Predict-PGRN.<sup>5</sup> Subjects were followed for more than 2 years, and the mean annualized rate of change (ARC) of plasma NfL was calculated along with absolute levels of plasma NfL at baseline and follow-up. *GRN* carriers' baseline NfL was 86.21 pg/ml versus 39.49 pg/ml for *C9orf72* ( $P = 0.014$ ) and demonstrated greater progression on ARC (+29.3%) compared with *C9orf72* carriers (+24.7%;  $P = 0.016$ ). Of interest, ARC differed with phenotype in *C9orf72* carriers: ALS had a mean ARC of +37%, FTD was +21.7%, and psychiatric presentations were +8.3%. However, slow-progressing *C9orf72* carriers had an ARC of +2.5% compared to *C9orf72* patients with standard progression. *GRN* carriers did not display such differences, possibly because of less variability in *GRN* disease progression. Separate cutoff thresholds of 19.00 pg/ml and 27.48 pg/ml for *C9orf72* and *GRN* carriers, respectively, differentiated them from controls. In the ALLFTD cohorts, plasma NfL discriminated the presymptomatic from symptomatic mutation carriers, with a cutoff of 13.6 pg/ml in the original cohort and 19.8 pg/ml in the validation group.<sup>3</sup> Data from these international cohorts demonstrate that neurodegeneration in genetic FTLD, as measured by blood NfL, has the potential to reflect disease progression and that the specific FTD gene mutation may dictate recommended cutoff thresholds.

## NfL as a pharmacodynamic marker in genetic FTLD

NfL as a pharmacodynamic marker may provide a means to distinguish treatment responders in clinical trials and, potentially, a way to gauge fast versus slow disease progressors.<sup>5,75–77</sup> An individual's disease status, as determined via blood NfL and NfH,<sup>4</sup> may provide added value when combined with macro- and microstructural imaging of regional neurodegenerative changes in the brain, potentially offering greater sensitivity to earlier neurodegenerative changes in presymptomatic and prodromal stage carriers. However, longitudinal studies are required to reveal absolute

levels or rate of change in NfL in relation to the trajectory of neurodegeneration measured by MD and cortical thickness neuroimaging methods. To use NfL as a pharmacodynamic marker, detailed knowledge of how the drug treatment itself influences NfL turnover is needed. Preliminary, unpublished observations suggest that NfL turnover in extracellular fluids may be influenced by microglial activity; this potential confounder would be particularly important to examine for drugs affecting microglial activation.

When compared with the apparently more gradual changes in brain GM or WM as measured with current neuroimaging methods, a fluid biomarker can provide an earlier and more accessible dynamic readout in trials that may suggest movement towards clinical improvement, to be supported later by clinical measures at 6–12 months that would reveal whether the change is statistically significant. Currently, blood-based NfL levels are supporting decisions about proceeding from investigational phase 2 studies to phase 3 trials. NfL in genetic FTLD could potentially be a more sensitive biomarker of change, but more data are needed to determine whether blood-based NfL, in combination with one or more markers, could provide greater confidence in moving forward to a pivotal trial. Longitudinal fluid biomarkers reflecting changes in the autophagy-lysosome pathway,<sup>78</sup> microglial<sup>79</sup> and astroglial activation markers,<sup>80–83</sup> and potentially TDP43 burden,<sup>84</sup> could provide additional insights into the pathobiology of FTLD and status of disease progression. Identifying the right combination of biomarkers that elucidates and refines our understanding of NfL is a work in progress, as different exploratory fluid markers are being tested in interventional trials enrolling presymptomatic and symptomatic *GRN* and *C9orf72* mutation carriers.<sup>85</sup> However, such exploratory markers must also be tested in longitudinal studies in larger multicentre research cohorts and compared against healthy controls to establish their progression. For phase 3 studies, surrogacy of NfL is not yet defined, and the dynamic range of blood NfL across FTD genotypes will require absolute cutoff levels and/or rates of change defined for each FTD gene mutation population. It remains to be determined whether normalization to prodromal blood NfL ranges in genetic FTD, or prevention of further increases in absolute levels of blood NfL (or rate of change), are possible. Hopefully, the time course of NfL changes after treatment initiation compared with controls will be revealed by ongoing interventional drug trials.<sup>3,77</sup>

In testing NfL as a pharmacodynamic marker in genetic FTD treatment trials, the percentage change indicating clinically meaningful NfL remains unknown. The current approach is to determine if the observed changes are statistically significant in the cohort studied, but there is a need to identify and establish disease-specific cutoff points for NfL levels, or even gene mutation-specific cutoff points. In other diseases, it is known that normalization or close-to-baseline decrease in NfL is possible with treatment, but the time lag between treatment initiation and time to stable decline of NfL appears

highly variable among diseases and treatment modalities when considering antisense oligonucleotide, nusinersen-treated paediatric spinal muscular atrophy versus treated adolescents or adults,<sup>86–88</sup> hematopoietic stem cell–transplanted X-linked adrenoleukodystrophy,<sup>89</sup> antiviral-treated adult HIV dementia,<sup>90–92</sup> or monoclonal antibody-treated adult multiple sclerosis.<sup>32,93</sup>

FTD-GRN, the most rapidly progressing of the FTD autosomal dominant gene mutations, show the highest rate of change in blood NfL moving from presymptomatic to symptomatic clinical status, as well as the highest median baseline. NfL may help define a window of opportunity for enrolling participants into investigational disease-modifying drug trials by employing an absolute, elevated NfL concentration as a cutoff that reflects transition from presymptomatic to prodromal stage. Longitudinal data suggest that the rate of change in blood NfL from conversion stage to symptomatic stage may be of prognostic value in gauging the rate of disease progression or drug efficacy.<sup>3,4</sup>

NfL levels observed over the first 12 months of an investigational treatment trial for FTD-GRN reported that both CSF and plasma NfL levels remained stable over this time period, reflecting a consistent trend in response to experimental drug treatments.<sup>94</sup> Decreased levels of blood or CSF NfL in other treatment trials have shown that a downward trend toward baseline values can take 12–24 months.<sup>35</sup> Many important questions remain regarding the use of NfL as a biomarker in genetic FTD and ongoing international longitudinal studies, as well as new investigational treatment trials, are anticipated to provide new insights. At present, we may find some answers regarding trends in NfL from observational studies in neurologically healthy aging populations and treatment trials in multiple sclerosis. The Foundation for the National Institutes of Health Biomarkers Consortium Neurofilament Project is spearheading a collaborative public-private partnership effort to examine and compare available NfL assays and platforms. A goal of this partnership is to work toward a global calibration of assay readouts and to prepare an NfL biomarker qualification letter of intent to be submitted to the FDA and European Medicines Agency (<https://fnih.org/what-we-do/biomarkers-consortium>).

## Summary and recommendations on use of NfL as a biomarker in genetic FTL D

Present data suggest that NfL can serve as a biomarker for onset and intensity of neurodegeneration in genetic FTL D. Given that it is a general biomarker for neurodegeneration, its diagnostic capacity is limited; however, this limitation does not apply to genetic FTL D, as the genotype gives the diagnosis. In clinical trials of disease-modifying treatments, the direction of change in fluid biomarker levels remains to

be discovered. Lowering and/or stabilization of NfL in genetic FTD would be an anticipated outcome if the treatment were to slow or stop neurodegeneration in the brain. Longitudinal FTL D studies that combine pathophysiological markers with sensitive topographic biomarkers correlated with established clinical instruments to reflect disease progression are required. The timing and effect size of NfL change is difficult to address. Here, stabilization and/or lowering of NfL to levels seen in same-age healthy controls, such as those observed in patients with multiple sclerosis under highly effective treatments, may predict clinical improvement. Considering the heterogeneity of genetic and sporadic FTL D, it would be advisable to combine NfL with other pathophysiological markers to provide a more robust readout of disease progression. Lysosomal/endosomal processing, microglial and astroglial activation, and potentially changes in total-tau and phosphorylated-tau protein may help identify possible inflection points moving from presymptomatic to prodromal to symptomatic stages.<sup>95</sup> In clinical trials, NfL can be used as both an inclusion criterion to ascertain that the patient has entered the neurodegenerative stage of the disease, and a prognostic biomarker. NfL in combination with other markers could potentially offer more precision on disease trajectory and interpretation of disease-modifying treatments in clinical trials. Cutoff criteria for elevated NfL levels in FTD-GRN and FTD-C9orf72 should now be established. Most data suggest that CSF and plasma/serum NfL levels and rates of change are higher in GRN carriers but may otherwise give similar information in trials across the genetic FTL D spectrum. Plasma NfL levels may provide additional value as these levels relate to patient stratification in trials. Disclosing NfL status to patients with FTD can be complicated, since trial inclusion of participants with elevated NfL implies that their disease status has progressed (although additional markers are likely needed to establish an individual's disease progression). Conversely, exclusion of patients from a trial because of 'low' NfL levels may result in a difficult conversation with the prospective participants because they have a disease that progresses quickly and will ultimately take away their independence.

## Acknowledgements

The Neurofilament Light Task Force, an international group of key opinion leaders, was convened by Alector, Inc., and The Bluefield Project to Cure Frontotemporal Dementia in late 2020 to synthesize recent knowledge about FTD and other neurological diseases to help develop recommendations on the use of NfL as a biomarker in genetic FTD.

## Funding

This work was supported by Alector, Inc.



## Competing interests

A.B. received grant support from NIH U19AG063911, U54NS092089, R01AG038791, U01AG045390, Tau Research Consortium, Bluefield Project to Cure FTD, University of California Cures Alzheimer's disease Program, Association for Frontotemporal Degeneration, CBD Solutions, Alzheimer's Drug Discovery Foundation, Alzheimer's Association, and UCSF Parkinson's Spectrum Disorders Center. A.B. received industry research support (funding to UCSF) from Biogen, Eisai, and Regeneron; A.B. is an industry consultant for AGTC, Alector, Arkuda, Arvinas, AZTherapeutics, GSK, Lundbeck, Oligomerix, Ono, Roche, Samumed, Stealth, Third Rock, Transposon, and Wave.

Jens Kuhle received speaker fees, research support, travel support, and/or served on advisory boards by Swiss MS Society, Swiss National Research Foundation (320030\_189140/1), University of Basel, Progressive MS Alliance, Bayer, Biogen, Celgene, Merck, Novartis, Octave Bioscience, Roche, and Sanofi. S.M.M. is an employee of Alector, LLC and may have an equity interest in Alector, Inc. J.D.R. has been a member of a medical advisory board for Alector, Prevail Therapeutics, and Wave Life Sciences and has provided consultancy for Arkuda Therapeutics, Astex, UCB, Takeda, and Biogen. N.T. is an employee of Alector, LLC and may have an equity interest in Alector, Inc. C.T.'s research of CET is supported by the European Commission [Marie Curie International Training Network, grant agreement No. 860197 (MIRIADE), and JPND], Health Holland, the Dutch Research Council (ZonMW), Alzheimer Drug Discovery Foundation (ADDF), The Selfridges Group Foundation, Alzheimer Netherlands, Alzheimer Association. C.T. is a recipient of ABOARD, which is a public-private partnership receiving funding from ZonMW (#73305095007) and Health~Holland, Topsector Life Sciences & Health (PPP-allowance; #LSHM20106). More than 30 partners participate in ABOARD. ABOARD also receives funding from Edwin Bouw Fonds and Gieskes-Strijbisfonds. IV is appointed on a research grant by Alzheimer Nederland (NL-17004). C.T. has a collaboration contract with ADx Neurosciences and Quanterix, and has performed contract research or received grants from AC-Immune, Axon Neurosciences, Biogen, Brainstorm Therapeutics, Celgene, EIP Pharma, Eisai, Olink, PeopleBio, Roche, Toyama, and Vivoryon. H.Z. has served on scientific advisory boards and/or as a consultant for Abbvie, Acumen, Alector, ALZPath, Annexon, Apellis, Artery Therapeutics, AZTherapies, CogRx, Denali, Eisai, Nervgen, Novo Nordisk, Passage Bio, Pinteon Therapeutics, Red Abbey Labs, reMYND, Roche, Samumed, Siemens Healthineers, Triplet Therapeutics, and Wave, has given lectures in symposia sponsored by Cellectricon, Fujirebio, Alzecure, Biogen, and Roche and is a co-founder of Brain Biomarker Solutions in Gothenburg AB (BBS), which is a part of the GU Ventures Incubator Program. H.Z. is a Wallenberg Scholar supported by grants from the Swedish

Research Council (#2018-02532), the European Research Council (#681712 and #101053962), Swedish State Support for Clinical Research (#ALFGBG-71320), the ADDF, USA (#201809-2016862), the Alzheimer's disease Strategic Fund and the Alzheimer's Association (#ADSF-21-831376-C, #ADSF-21-831381-C, and #ADSF-21-831377-C), the Bluefield Project, the Olav Thon Foundation, the Erling-Persson Family Foundation, Stiftelsen för Gamla Tjänarinnor, Hjärnfonden, Sweden (#FO2022-0270), the European Union's Horizon 2020 research and innovation program under the Marie Skłodowska-Curie grant agreement No. 860197 (MIRIADE), the European Union Joint Programme—Neurodegenerative Disease Research (JPND2021-00694), and the UK Dementia Research Institute at UCL (UKDRI-1003).

## Supplementary material

Supplementary material is available at *Brain Communications* online.

## Data availability

The names and affiliations of the Neurofilament task force participants is provided in [Supplementary\\_Data-docx](#).

## Appendix I

### NfL Task Force Participants

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## References

- Boeve BF, Boxer AL, Kumfor F, Pijnenburg Y, Rohrer JD. Advances and controversies in frontotemporal dementia: Diagnosis, biomarkers, and therapeutic considerations. *Lancet Neurol.* 2022;21(3): 258-272.
- Greaves CV, Rohrer JD. An update on genetic frontotemporal dementia. *J Neurol.* 2019;266(8):2075-2086.
- Rojas JC, Wang P, Staffaroni AM, et al. Plasma neurofilament light for prediction of disease progression in familial frontotemporal lobar degeneration. *Neurology.* 2021;96(18):e2296-e2312.
- Wilke C, Reich S, van Swieten JC, et al. Stratifying the presymptomatic phase of genetic frontotemporal dementia by serum NfL and pNfH: A longitudinal multicentre study. *Ann Neurol.* 2022; 91(1):33-47.

5. Saracino D, Dorgham K, Camuzat A, et al. Plasma NfL levels and longitudinal change rates in C9orf72 and GRN-associated diseases: From tailored references to clinical applications. *J Neurol Neurosurg Psychiatry*. 2021;92(12):1278–1288.
6. Jo M, Lee S, Jeon YM, Kim S, Kwon Y, Kim HJ. The role of TDP-43 propagation in neurodegenerative diseases: Integrating insights from clinical and experimental studies. *Exp Mol Med*. 2020;52(10):1652–1662.
7. Bridel C, van Wieringen WN, Zetterberg H, et al. Diagnostic value of cerebrospinal fluid neurofilament light protein in neurology: A systematic review and meta-analysis. *JAMA Neurol*. 2019;76(9):1035–1048.
8. Forgrave LM, Ma M, Best JR, Demarco ML. The diagnostic performance of neurofilament light chain in CSF and blood for Alzheimer's disease, frontotemporal dementia, and amyotrophic lateral sclerosis: A systematic review and meta-analysis. *Alzheimers Dement*. 2019;11:730–743.
9. Moseby-Knappe M, Mattsson N, Nielsen N, et al. Serum neurofilament light chain for prognosis of outcome after cardiac arrest. *JAMA Neurol*. 2019;76(1):64–71.
10. Polymeris AA, Coslovksy M, Aeschbacher S, et al. Serum neurofilament light in atrial fibrillation: Clinical, neuroimaging and cognitive correlates. *Brain Commun*. 2020;2(2):fcaa166.
11. Petzold A. Neurofilament phosphoforms: Surrogate markers for axonal injury, degeneration and loss. *J Neurol Sci*. 2005;233(1-2):183–198.
12. Steinacker P, Feneberg E, Weishaupt J, et al. Neurofilaments in the diagnosis of motoneuron diseases: A prospective study on 455 patients. *J Neurol Neurosurg Psychiatry*. 2016;87(1):12–20.
13. Chen XH, Meaney DF, Xu BN, et al. Evolution of neurofilament subtype accumulation in axons following diffuse brain injury in the pig. *J Neuropathol Exp Neurol*. 1999;58(6):588–596.
14. Neselius S, Brisby H, Marcusson J, Zetterberg H, Blennow K, Karlsson T. Neurological assessment and its relationship to CSF biomarkers in amateur boxers. *PLoS One*. 2014;9(6):e99870.
15. Taghdiri F, Multani N, Ozzoude M, et al. Neurofilament-light in former athletes: A potential biomarker of neurodegeneration and progression. *Eur J Neurol*. 2020;27(7):1170–1177.
16. Clarke GJB, Skandsen T, Zetterberg H, et al. One-year prospective study of plasma biomarkers from CNS in patients with mild traumatic brain injury. *Front Neurol*. 2021;12:643743.
17. Wong NKY, He BP, Strong MJ. Characterization of neuronal intermediate filament protein expression in cervical spinal motor neurons in sporadic amyotrophic lateral sclerosis (ALS). *J Neuropathol Exp Neurol*. 2000;59(11):972–982.
18. Strong MJ, Volkening K, Hammond R, et al. TDP43 is a human low molecular weight neurofilament (hNFL) mRNA-binding protein. *Mol Cell Neurosci*. 2007;35(2):320–327.
19. Volkening K, Leystra-Lantz C, Yang W, Jaffee H, Strong MJ. Tar DNA binding protein of 43 kDa (TDP-43), 14-3-3 proteins and copper/zinc superoxide dismutase (SOD1) interact to modulate NFL mRNA stability. Implications for altered RNA processing in amyotrophic lateral sclerosis (ALS). *Brain Res*. 2009;1305:168–182.
20. Rosengren LE, Karlsson JE, Karlsson JO, Persson LI, Wikkelso C. Patients with amyotrophic lateral sclerosis and other neurodegenerative diseases have increased levels of neurofilament protein in CSF. *J Neurochem*. 1996;67(5):2013–2018.
21. Sealey WW. Selective functional, regional, and neuronal vulnerability in frontotemporal dementia. *Curr Opin Neurol*. 2008;21(6):701–707.
22. Khalil B, Morderer D, Price PL, Liu F, Rossoll W. mRNP assembly, axonal transport, and local translation in neurodegenerative diseases. *Brain Res*. 2018;1693(Pt A):75–91.
23. Briese M, Saal L, Appenzeller S, Moradi M, Baluapuri A, Sendtner M. Whole transcriptome profiling reveals the RNA content of motor axons. *Nucleic Acids Res*. 2016;44(4):e33.
24. Cajigas JJ, Tushev G, Will TJ, tom Dieck S, Fuerst N, Schuman EM. The local transcriptome in the synaptic neuropil revealed by deep sequencing and high-resolution imaging. *Neuron*. 2012;74(3):453–466.
25. Shigeoka T, Jung H, Jung J, et al. Dynamic axonal translation in developing and mature visual circuits. *Cell*. 2016;166(1):181–192.
26. Aerts J, Nys J, Moons L, Hu TT, Arckens L. Altered neuronal architecture and plasticity in the visual cortex of adult MMP-3-deficient mice. *Brain Struct Funct*. 2015;220(5):2675–2689.
27. Twiss JL, Kalinski AL, Sachdeva R, Houle JD. Intra-axonal protein synthesis—A new target for neural repair? *Neural Regen Res*. 2016;11(9):1365–1367.
28. Trapp BD, Peterson J, Ransohoff RM, Rudick R, Mörk S, Bö L. Axonal transection in the lesions of multiple sclerosis. *N Engl J Med*. 1998;338(5):278–285.
29. Kuhlmann T. Acute axonal damage in multiple sclerosis is most extensive in early disease stages and decreases over time. *Brain*. 2002;125(10):2202–2212.
30. Bjartmar C, Kidd G, Mork S, Rudick R, Trapp BD. Neurological disability correlates with spinal cord axonal loss and reduced N-acetyl aspartate in chronic multiple sclerosis patients. *Ann Neurol*. 2000;48(6):893–901.
31. Strasser-Fuchs S, Enzinger C, Ropele S, Wallner M, Fazekas F. Clinically benign multiple sclerosis despite large T2 lesion load: Can we explain this paradox? *Mult Scler*. 2008;14(2):205–211.
32. Novakova L, Zetterberg H, Sundström P, et al. Monitoring disease activity in multiple sclerosis using serum neurofilament light protein. *Neurology*. 2017;89(22):2230–2237.
33. Kuhle J, Plavina T, Barro C, et al. Neurofilament light levels are associated with long-term outcomes in multiple sclerosis. *Mult Scler*. 2020;26(13):1691–1699.
34. Delcoigne B, Manouchehrinia A, Barro C, et al. Blood neurofilament light levels segregate treatment effects in multiple sclerosis. *Neurology*. 2020;94(11):e1201–e1212.
35. Kapoor R, Smith KE, Allegratta M, et al. Serum neurofilament light as a biomarker in progressive multiple sclerosis. *Neurology*. 2020;95(10):436–444.
36. Nyberg L, Lundquist A, Nordin Adolfsson A, et al. Elevated plasma neurofilament light in aging reflects brain white-matter alterations but does not predict cognitive decline or Alzheimer's disease. *Alzheimers Dement*. 2020;12(1):e12050.
37. Preische O, Schultz SA, Apel A, et al. Serum neurofilament dynamics predicts neurodegeneration and clinical progression in presymptomatic Alzheimer's disease. *Nat Med*. 2019;25(2):277–283.
38. Khalil M, Pirpamer L, Hofer E, et al. Serum neurofilament light levels in Normal aging and their association with morphologic brain changes. *Nat Commun*. 2020;11(1):812.
39. Willems EAJ, Scheltens P, Teunissen CE, Vijverberg EGB. A neurologist's perspective on serum neurofilament light in the memory clinic: A prospective implementation study. *Alzheimers Res Ther*. 2021;13(1):101.
40. Scherling CS, Hall T, Berisha F, et al. Cerebrospinal fluid neurofilament concentration reflects disease severity in frontotemporal degeneration. *Ann Neurol*. 2014;75(1):116–126.
41. van der Ende EL, Meeter LH, Poos JM, et al. Serum neurofilament light chain in genetic frontotemporal dementia: A longitudinal, multicentre cohort study. *Lancet Neurol*. 2019;18(12):1103–1111.
42. Bridel C, Verberk IMW, Heijst JJA, Killestein J, Teunissen CE. Variations in consecutive serum neurofilament light levels in healthy controls and multiple sclerosis patients. *Mult Scler Relat Disord*. 2021;47:102666.
43. Ramani S, Berard JA, Walker LAS. The relationship between neurofilament light chain and cognition in neurological disorders: A scoping review. *J Neurol Sci*. 2021;420:117229.
44. Le Bihan D. Looking into the functional architecture of the brain with diffusion MRI. *Nat Rev Neurosci*. 2003;4(6):469–480.
45. Weston PSJ, Poole T, Nicholas JM, et al. Measuring cortical mean diffusivity to assess early microstructural cortical change in presymptomatic familial Alzheimer's disease. *Alzheimers Res Ther*. 2020;12(1):112.

46. Montal V, Vilaplana E, Pegueroles J, *et al.* Biphasic cortical macro- and microstructural changes in autosomal dominant Alzheimer's disease. *Alzheimers Dement.* 2021;17(4):618-628.
47. Dhiman K, Gupta VB, Villemagne VL, *et al.* Cerebrospinal fluid neurofilament light concentration predicts brain atrophy and cognition in Alzheimer's disease. *Alzheimers Dement (Amst).* 2020;12(1): e12005.
48. Galimberti D, Fumagalli GG, Fenoglio C, *et al.* Progranulin plasma levels predict the presence of GRN mutations in asymptomatic subjects and do not correlate with brain atrophy: Results from the GENFI study. *Neurobiol Aging.* 2018;62:245.e249-245.e212.
49. Moore KM, Nicholas J, Grossman M, *et al.* Age at symptom onset and death and disease duration in genetic frontotemporal dementia: An international retrospective cohort study. *Lancet Neurol.* 2020; 19(2):145-156.
50. Goldman JS, Van Deerlin VM. Alzheimer's disease and frontotemporal dementia: The current state of genetics and genetic testing since the advent of next-generation sequencing. *Mol Diagn Ther.* 2018;22(5):505-513.
51. Caroppo P, Le Ber I, Camuzat A, *et al.* Extensive white matter involvement in patients with frontotemporal lobar degeneration. *JAMA Neurol.* 2014;71(12):1562.
52. Sudre CH, Bocchetta M, Cash D, *et al.* White matter hyperintensities are seen only in GRN mutation carriers in the GENFI cohort. *Neuroimage Clin.* 2017;15:171-180.
53. Woollacott IOC, Bocchetta M, Sudre CH, *et al.* Pathological correlates of white matter hyperintensities in a case of progranulin mutation associated frontotemporal dementia. *Neurocase.* 2018;24(3): 166-174.
54. Rohrer JD, Warren JD, Fox NC, Rossor MN. Presymptomatic studies in genetic frontotemporal dementia. *Rev Neurol (Paris).* 2013; 169(10):820-824.
55. Cash DM, Bocchetta M, Thomas DL, *et al.* Patterns of gray matter atrophy in genetic frontotemporal dementia: Results from the GENFI study. *Neurobiol Aging.* 2018;62:191-196.
56. Chen Q, Kantarci K. Imaging biomarkers for neurodegeneration in presymptomatic familial frontotemporal lobar degeneration. *Front Neurol.* 2020;11:80.
57. Jacova C, Hsiung GY, Tawankanjanachot I, *et al.* Anterior brain glucose hypometabolism predates dementia in progranulin mutation carriers. *Neurology.* 2013;81(15):1322-1331.
58. Abramzon YA, Fratta P, Traynor BJ, Chia R. The overlapping genetics of amyotrophic lateral sclerosis and frontotemporal dementia. *Front Neurosci.* 2020;14:42.
59. Majounie E, Renton AE, Mok K, *et al.* Frequency of the C9orf72 hexanucleotide repeat expansion in patients with amyotrophic lateral sclerosis and frontotemporal dementia: A cross-sectional study. *Lancet Neurol.* 2012;11(4):323-330.
60. Rohrer JD, Nicholas JM, Cash DM, *et al.* Presymptomatic cognitive and neuroanatomical changes in genetic frontotemporal dementia in the genetic frontotemporal dementia initiative (GENFI) study: A cross-sectional analysis. *Lancet Neurol.* 2015;14(3):253-262.
61. Moreno F, Sala-Llonch R, Barandiaran M, *et al.* Distinctive age-related temporal cortical thinning in asymptomatic granulin gene mutation carriers. *Neurobiol Aging.* 2013;34(5):1462-1468.
62. Verde F, Otto M, Silani V. Neurofilament light chain as biomarker for amyotrophic lateral sclerosis and frontotemporal dementia. *Front Neurosci.* 2021;15:679199.
63. Borrego-Ecija S, Sala-Llonch R, van Swieten J, *et al.* Disease-related cortical thinning in presymptomatic granulin mutation carriers. *Neuroimage Clin.* 2021;29:102540.
64. Le Blanc G, Jette Pomerleau V, McCarthy J, *et al.* Faster cortical thinning and surface area loss in presymptomatic and symptomatic C9orf72 repeat expansion adult carriers. *Ann Neurol.* 2020;88(1): 113-122.
65. Mahoney CJ, Ridgway GR, Malone IB, *et al.* Profiles of white matter tract pathology in frontotemporal dementia. *Hum Brain Mapp.* 2014;35(8):4163-4179.
66. Sjogren M, Blomberg M, Jonsson M, *et al.* Neurofilament protein in cerebrospinal fluid: A marker of white matter changes. *J Neurosci Res.* 2001;66(3):510-516.
67. Panman JL, Jiskoot LC, Bouts M, *et al.* Gray and white matter changes in presymptomatic genetic frontotemporal dementia: A longitudinal MRI study. *Neurobiol Aging.* 2019;76:115-124.
68. Spotorno N, Lindberg O, Nilsson C, *et al.* Plasma neurofilament light protein correlates with diffusion tensor imaging metrics in frontotemporal dementia. *PLoS One.* 2020;15(10):e0236384.
69. Illan-Gala I, Montal V, Borrego-Ecija S, *et al.* Cortical microstructure in the behavioural variant of frontotemporal dementia: Looking beyond atrophy. *Brain.* 2019;142(4): 1121-1133.
70. Illan-Gala I, Montal V, Borrego-Ecija S, *et al.* Cortical microstructure in primary progressive aphasia: A multicenter study. *Alzheimers Res Ther.* 2022;14(1):27.
71. Rohrer JD, Woollacott IO, Dick KM, *et al.* Serum neurofilament light chain protein is a measure of disease intensity in frontotemporal dementia. *Neurology.* 2016;87(13):1329-1336.
72. Zhang JV, Irwin DJ, Blennow K, *et al.* Neurofilament light chain related to longitudinal decline in frontotemporal lobar degeneration. *Neurol Clin Pract.* 2021;11(2):105-116.
73. Meeter LH, Dopfer EG, Jiskoot LC, *et al.* Neurofilament light chain: A biomarker for genetic frontotemporal dementia. *Ann Clin Transl Neurol.* 2016;3(8):623-636.
74. Illan-Gala I, Lleo A, Karydas A, *et al.* Plasma tau and neurofilament light in frontotemporal lobar degeneration and Alzheimer disease. *Neurology.* 2021;96(5):e671-e683.
75. Trageser KJ, Smith C, Herman FJ, Ono K, Pasinetti GM. Mechanisms of immune activation by c9orf72-expansions in amyotrophic lateral sclerosis and frontotemporal dementia. *Front Neurosci.* 2019;13:1298.
76. Huang J, Khademi M, Fugger L, *et al.* Inflammation-related plasma and CSF biomarkers for multiple sclerosis. *Proc Natl Acad Sci USA.* 2020;117(23):12952-12960.
77. Panman JL, Venkatraghavan V, van der Ende EL, *et al.* Modelling the cascade of biomarker changes in GRN-related frontotemporal dementia. *J Neurol Neurosurg Psychiatry.* 2021;92(5):494-501.
78. Beckers J, Tharkeshwar AK, Van Damme P. C9orf72 ALS-FTD: Recent evidence for dysregulation of the autophagy-lysosome pathway at multiple levels. *Autophagy.* 2021:1-17.
79. Parbo P, Madsen LS, Ismail R, *et al.* Low plasma neurofilament light levels associated with raised cortical microglial activation suggest inflammation acts to protect prodromal Alzheimer's disease. *Alzheimers Res Ther.* 2020;12(1):3.
80. Oeckl P, Weydt P, Steinacker P, *et al.* Different neuroinflammatory profile in amyotrophic lateral sclerosis and frontotemporal dementia is linked to the clinical phase. *J Neurol Neurosurg Psychiatry.* 2019;90(1):4-10.
81. Nowacki P, Koziarska D, Masztalewicz M. Microglia and astroglia proliferation within the normal appearing white matter in histologically active and inactive multiple sclerosis. *Folia Neuropathologica.* 2019;57(3):249-257.
82. Katisko K, Cajanus A, Huber N, *et al.* GFAP As a biomarker in frontotemporal dementia and primary psychiatric disorders: Diagnostic and prognostic performance. *J Neurol Neurosurg Psychiatry.* 2021; 92(12):1305-1312.
83. Heller C, Foiani MS, Moore K, *et al.* Plasma glial fibrillary acidic protein is raised in progranulin-associated frontotemporal dementia. *J Neurol Neurosurg Psychiatry.* 2020;91(3): 263-270.
84. Borrego-Écija S, Turon-Sans J, Ximelis T, *et al.* Cognitive decline in amyotrophic lateral sclerosis: Neuropathological substrate and genetic determinants. *Brain Pathol.* 2021;31(3):e12942.
85. Boxer AL, Gold M, Feldman H, *et al.* New directions in clinical trials for frontotemporal lobar degeneration: Methods and outcome measures. *Alzheimers Dement.* 2020;16:131-143.

86. Gidaro T, Servais L. Nusinersen treatment of spinal muscular atrophy: Current knowledge and existing gaps. *Dev MedChild Neurol.* 2019;61(1):19-24.
87. Faravelli I, Meneri M, Saccomanno D, *et al.* Nusinersen treatment and cerebrospinal fluid neurofilaments: An explorative study on spinal muscular atrophy type 3 patients. *J Cell Mol Med.* 2020;24(5):3034-3039.
88. Olsson B, Alberg L, Cullen NC, *et al.* NFL is a marker of treatment response in children with SMA treated with nusinersen. *J Neurol.* 2019;266(9):2129-2136.
89. Weinhofer I, Rommer P, Zierfuss B, *et al.* Neurofilament light chain as a potential biomarker for monitoring neurodegeneration in X-linked adrenoleukodystrophy. *Nat Commun.* 2021;12(1):1816.
90. Guha D, Mukerji SS, Chettimada S, *et al.* Cerebrospinal fluid extracellular vesicles and neurofilament light protein as biomarkers of central nervous system injury in HIV-infected patients on antiretroviral therapy. *Aids.* 2019;33(4):615-625.
91. Gisslén M, Price RW, Andreasson U, *et al.* Plasma concentration of the neurofilament light protein (NFL) is a biomarker of CNS injury in HIV infection: A cross-sectional study. *EBioMedicine.* 2016;3:135-140.
92. Anderson AM, Easley KA, Kasher N, *et al.* Neurofilament light chain in blood is negatively associated with neuropsychological performance in HIV-infected adults and declines with initiation of antiretroviral therapy. *J Neurovirol.* 2018;24(6):695-701.
93. Disanto G, Barro C, Benkert P, *et al.* Serum neurofilament light: A biomarker of neuronal damage in multiple sclerosis. *Ann Neurol.* 2017;81(6):857-870.
94. Paul R, Yeh F, Ward M, *et al.* Update on the phase 2 study of AL001 in frontotemporal dementia patients carrying a granulin mutation. 14th Clinical Trials on Alzheimer's Disease (CTAD), November 9–12, 2021.
95. Agnello L, Colletti T, Lo Sasso B, *et al.* Tau protein as a diagnostic and prognostic biomarker in amyotrophic lateral sclerosis. *Eur J Neurol.* 2021;28(6):1868-1875.