Mitotic defects lead to neuronal aneuploidy and apoptosis in frontotemporal lobar degeneration caused by MAPT mutations

Julbert Caneus,a,b,c,† Antoneta Granic,a,b,d,e,f,† Rosa Rademakers,g Dennis W. Dickson,g Christina M. Coughlan,a,b, Heidi J. Chia,b,b, and Huntington Potter,a,b,c,*
-Department of Neurology, Rocky Mountain Alzheimer’s Disease Center, and †Linda Crnic Institute for Down Syndrome, University of Colorado School of Medicine, Aurora, CO 80045; -Neuroscience Program, University of Colorado, Anschutz Medical Campus, Aurora, CO 80045; ‡AGE Research Group, Institute of Neuroscience, Campus for Ageing and Vitality, Biomedical Research Building, Newcastle University, Newcastle upon Tyne NE4 5PL, United Kingdom; †NIHR Newcastle Biomedical Research Centre, Newcastle upon Tyne Hospitals NHS Foundation Trust and Newcastle University, Newcastle upon Tyne NE4 5PL, United Kingdom; ‡Newcastle University Institute for Ageing, Newcastle upon Tyne NE4 5PL, United Kingdom; *Department of Neuroscience, Mayo Clinic, Jacksonville, FL 32224

ABSTRACT Mutant Tau (MAPT) can lead to frontotemporal lobar degeneration (FTLD). Previous studies associated MAPT mutations and altered function with aneuploidy and chromosome instability in human lymphocytes and in Drosophila development. Here we examine whether FTLD-causing mutations in human MAPT induce aneuploidy and apoptosis in the mammalian brain. First, aneuploidy was found in brain cells from MAPT mutant transgenic mice expressing FTLD mutant human MAPT. Then brain neurons from mice homozygous or heterozygous for the Tau (Mapt) null allele were found to exhibit increasing levels of aneuploidy with decreasing Tau gene dosage. To determine whether aneuploidy leads to neurodegeneration in FTLD, we measured aneuploidy and apoptosis in brain cells from patients with MAPT mutations and identified both increased aneuploidy and apoptosis in the same brain neurons and glia. To determine whether there is a direct relationship between MAPT-induced aneuploidy and apoptosis, we expressed FTLD-causing mutant forms of MAPT in karyotypically normal human cells and found that they cause aneuploidy and mitotic spindle defects that then result in apoptosis. Collectively, our findings reveal a neurodegenerative pathway in FTLD-MAPT in which neurons and glia exhibit mitotic spindle abnormalities, chromosome mis-segregation, and aneuploidy, which then lead to apoptosis.

INTRODUCTION Frontotemporal lobar degeneration (FTLD), also termed frontotemporal dementia (FTD), is most often an early-onset neurodegenerative disease in which a subset of cases has tau-positive neuronal and glial inclusions in the absence of Alzheimer’s disease (AD)-like amyloid deposits, whereas other cases have TDP-43 neuronal and glial inclusions (Rademakers et al., 2012). Familial forms of FTLD are associated with mutations in several genes, and 50–80% of FTLD cases are sporadic. Some autosomal-dominant familial forms of FTLD and Parkinsonism are linked to chromosome 17 (FTDP-17) and are caused by mutations in one of two nearby genes, the tau gene (MAPT) or the progranulin gene (PGRN). A large proportion of familial FTLD and of amyotrophic lateral sclerosis (ALS) is also caused by abnormal expansions of a hexanucleotide repeat in the C9orf72 gene (Rademakers et al., 2012).

A potential mechanism by which FTLD arises in patients with MAPT mutations is suggested by work on AD. We previously proposed that trisomy 21 mosaicism may underlie the development of AD (Potter, 1991), and subsequent studies demonstrated that people with both sporadic and familial forms of AD exhibit trisomy 21 mosaicism and other aneuploidy in all tissues examined, including...
in brain neurons and glia (Potter et al., 1995; Migliore et al., 1997; Geller and Potter, 1999; Kingsbury et al., 2006; Mosch et al., 2007; Thomas and Fenech, 2008; Iourov et al., 2009; Arendt et al., 2010, 2015; Arendt, 2012). In addition, AD-causing mutations in the amyloid precursor protein (APP) and presenilin (PS) genes as well as exposure to amyloid-beta (Aβ) lead to mitotic spindle abnormalities and chromosome mis-segregation that depend on MAPT/Tau (Boeras et al., 2008; Granic et al., 2010). Furthermore, we discovered that a likely mechanism by which Aβ causes spindle abnormalities and aneuploidy is through its competitive inhibition of the microtubule-dependent motor Kinesin-5/Eg5, which can be modulated by addition of MAPT/Tau (Borysov et al., 2011). These findings suggested that an FTLD-associated alteration in MAPT function may include chromosome mis-segregation as a contributing or even a necessary step in the neurodegenerative pathway. Indeed, lymphocytes from FTLD patients or from transgenic mice expressing human MAPT harboring FTLD-causing mutations exhibit chromosome instability, including aneuploidy (usually chromosome loss) (Rossi et al., 2008). However, whether and how FTLD-causing mutations in MAPT affect mitosis, whether they affect chromosome segregation in the brain, and whether such cell-cycle defects contribute to neurodegeneration in FTLD are unknown.

Herein we examined the effects of FTLD-causing MAPT mutations and loss of MAPT function in brain cell populations and/or in transfected cells and determined that defects in MAPT lead to aberrant mitotic spindle function, abnormal chromosome segregation, and apoptosis. Together, the data indicate that, as in AD, aneuploid neurons arise in the FTLD-MAPT brain, are prone to apoptosis, and thus can contribute to the development of neurodegeneration and dementia.

RESULTS

Aneuploidy induced by expression of human MAPT harboring FTLD-causing mutations in mice

Although overexpression of the human MAPT gene harboring familial FTLD-causing mutations (P301L or P301S) in mice has been shown to result in increased aneuploidy in splenic lymphocytes (Rossi et al., 2014), a key unanswered question with high disease relevance is whether FTLD-causing MAPT mutations also lead to aneuploidy in brain cells. We generated brain cell suspensions from young and older transgenic mice expressing human MAPT P301S or P301L and from age-matched control mice and determined aneuploidy levels by fluorescence in situ hybridization (FISH) using a bacterial artificial chromosome (BAC) probe for mouse chromosome 16 (Kulnane et al., 2002), which is partially syntenic with human chromosome 21. The data showed that expression of human mutant MAPT P301L induces a small but significant increase in chromosome 16 trisomy and total aneuploidy in the brain as early as 8 weeks of age (Figure 1) with nearly undetectable levels of TUNEL-positive (apoptotic) cells (data not shown). A similar small but significant increase in aneuploidy was also detected in both brain cells (Figure 2A) and splenocytes (Figure 2B) from 8-mo-old transgenic mice expressing human MAPT P301S relative to nontransgenic control mice.

Total or partial loss of tau function induces aneuploidy in mouse brain neurons

Our finding that expression in mice of human MAPT harboring FTLD-causing mutations resulted in increased levels of chromosome 16 aneuploidy in brain cells provided the first evidence that this mutation is not only associated with aneuploidy in peripheral cells (Rossi et al., 2008) but also with aneuploidy in the brain. Previously, we showed that splenocytes from tau (mapt) heterozygous (tau+/−) or homozygous (tau−/−) knockout mice exhibited 2.5% or 4.9% trisomy 16, respectively, compared with 1.1% stochastic trisomy in splenocytes from age-matched tau+/+ control mice (Granic et al., 2010). To further investigate the role of wild-type mouse Tau in chromosome segregation in the brain during neurogenesis and during

FIGURE 1: Increased percentage of cells with abnormal chromosome numbers in brain tissues from young transgenic mice expressing a human mutant FTLD-MAPT gene. FISH analysis using a mouse chromosome 16 probe was carried out using single-nuclei suspensions prepared with brain tissues from 8-wk-old transgenic mice expressing a human mutant MAPT transgene (MAPT-P301L, n = 4). Brain tissues from the transgenic mice exhibited elevated levels of trisomy 16 (A) and total chromosome 16 aneuploidy (including monosomy plus trisomy) (B) in comparison to control nontransgenic mice (control, n = 4). Both neurons [NeuN(+), green] and nonneuronal cells [NeuN(−)] showed abnormal chromosome copy numbers in the FTLD-MAPT mice (C). For statistical analyses, more than 300 cells per slide and three slides per brain sample were counted. Statistical analyses were conducted using a Student’s t test. Error bars indicate SEM, and * indicates p < 0.05.
MAPT lead to aneuploidy and subsequent apoptosis in the human brain, where they could affect neurodegeneration, is unknown. To address this question, we prepared single-brain-cell suspensions using frozen samples of cortex from familial FTLD patients with one of three different MAPT mutations, specifically MAPT P301L, MAPT N279K, or MAPT E372G (Table 1), or from normal controls and determined aneuploidy levels using a dual-color DNA probe set to visualize chromosomes 12 and 21 via FISH. We found that cortical brain cells from FTLD patients with MAPT mutations exhibit significantly higher levels of trisomy 12 and trisomy 21 relative to cortical brain cells from normal controls (Figure 4A). Furthermore, we observed significantly higher levels of mosaic aneuploidy (i.e., either monosomy or trisomy) for chromosomes 12 and 21 in both neuronal [NeuN(+)] and nonneuronal [NeuN(–)] cells from the cortex relative to normal controls (Figure 4B). These results show that familial FTLD caused by MAPT mutations is characterized by mosaic aneuploidy, including trisomy 21, in both neurons and glia in the brain.

Increased apoptosis in brain neurons and glia from FTLD patients with MAPT mutations correlates with induced aneuploidy

To ask directly whether aneuploidy and neurodegeneration are linked in FTLD, we measured apoptosis in cortical brain cells from the patients with MAPT mutations. Simultaneous terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL) to detect apoptotic cells, NeuN staining to identify neurons, and chromosome

Elevated levels of mosaic aneuploidy in brain neurons and glia from FTLD patients with MAPT mutations

Although we observed aneuploidy in mouse brain cells that was induced by expression of the P301L or P301S FTLD-causing form of human MAPT and in mouse brain neurons with complete or partial loss of Tau (Mapt) function, whether FLTD-associated mutations in

MAPT lead to aneuploidy and subsequent apoptosis in the human brain, where they could affect neurodegeneration, is unknown. To address this question, we prepared single-brain-cell suspensions using frozen samples of cortex from familial FTLD patients with one of three different MAPT mutations, specifically MAPT P301L, MAPT N279K, or MAPT E372G (Table 1), or from normal controls and determined aneuploidy levels using a dual-color DNA probe set to visualize chromosomes 12 and 21 via FISH. We found that cortical brain cells from FTLD patients with MAPT mutations exhibit significantly higher levels of trisomy 12 and trisomy 21 relative to cortical brain cells from normal controls (Figure 4A). Furthermore, we observed significantly higher levels of mosaic aneuploidy (i.e., either monosomy or trisomy) for chromosomes 12 and 21 in both neuronal [NeuN(+)] and nonneuronal [NeuN(–)] cells from the cortex relative to normal controls (Figure 4B). These results show that familial FTLD caused by MAPT mutations is characterized by mosaic aneuploidy, including trisomy 21, in both neurons and glia in the brain.

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TABLE 1: Demographics of familial MAPT FTLD cases and age-matched control individuals without neurologic disease.

| Diagnosis | MAPT mutation | Age (yr) | Sex |
|-----------|---------------|----------|-----|
| FTLD      | P301L         | 52       | M   |
|           | P301L         | 62       | M   |
|           | N279K         | 51       | F   |
|           | E372G         | 58       | M   |
| Controls  | WT            | 59       | F   |
|           | WT            | 61       | M   |
|           | WT            | 63       | M   |
|           | WT            | 63       | M   |
|           | WT            | 53       | M   |
|           | WT            | 79       | M   |
|           | WT            | 74       | M   |

TABLE 4: Increased levels of mosaic aneuploidy in brain neurons and glia from FTLD patients with MAPT mutations. Quantitative FISH analysis of single-cell suspensions prepared from frozen cortical brain samples of individuals genetically identified as having familial FTLD with a mutation in MAPT (MAPT-FTLD) (n = 4) compared with normal controls (n = 7) revealed significantly higher percentages of brain cells with trisomy 12 or 21 in FTLD-MAPT patients (A). Separated analysis of neurons (NeuN(+)) from other brain cells (NeuN(−)) showed increased total aneuploidy (i.e., monosomy or trisomy for chromosome 12 or 21) in both cortical neurons and glia from familial FTLD patients relative to normal controls (B). The graphs show mean difference and significance (*) using one-way ANOVA with Tukey post hoc test. Error bars indicate SEM. *p < 0.05, **p < 0.005, and ***p < 0.0005. A minimum of ~1000 nuclei/slide and three slides/sample were counted and recorded for statistical analyses.

Expression of human MAPT genes harboring FTLD-causing mutations induces aneuploidy and subsequent apoptosis in karyotypically normal human cells

To rule out the possibility that pathological processes that arise in human FTLD-MAPT and in mutant MAPT mouse models are indirectly responsible for generating aneuploid neurons and glia, we transiently expressed human MAPT genes harboring one of two different FTLD-causing mutations in the karyotypically normal, nonneuronal hTERT-HME1 human cell line and assessed the levels of induced aneuploidy and apoptosis. As shown in Figure 6A and Supplemental Figure 1, A–C, and illustrated in Figure 6B and Supplemental Figure 1B, expression of FTLD-causing mutant forms of MAPT (P301L or V337M) induced a significant increase in aneuploidy within 48 h posttransfection, compared with control cells transfected with the vector alone. Levels of aneuploidy in cells transfected with wild-type MAPT were intermediate between those of cells transfected with the vector alone and cells transfected with the FTLD-causing MAPT mutants. At this 48-h time point, levels of apoptosis were essentially undetectable (data not shown).

TUNEL staining at a later time point of 72 h posttransfection revealed significantly higher levels of apoptosis in cells expressing either MAPT P301L or MAPT V337M, relative to control cells transfected with the vector alone (Figure 6C). Quantitative analysis of cells costained for TUNEL and FISH for chromosomes 12 and 21 showed that the majority of TUNEL(+) cells at 72 h were aneuploid, regardless of whether they were negative controls or were expressing wild-type or mutant MAPT (Figure 6, D and E). Analysis of the frequency of apoptosis among aneuploid cells and of aneuploidy among apoptotic cells at 72 h indicated that ~80%...
of apoptotic cells were aneuploid, but less than 20% of aneuploid cells were apoptotic (Figure 6F). This relationship between aneuploidy and apoptosis was true both in negative control cells, which had low levels of aneuploidy and apoptosis, and in cells expressing the FLTD-causing mutant MAPT, which had significantly higher levels of aneuploidy and apoptosis. Notably, although the cells expressing the FLTD-causing mutant MAPT genes showed increased levels of aneuploidy at 48 h posttransfection, they did not exhibit increased levels of apoptosis until 72 h posttransfection. The cells transfected with either of the MAPT mutants exhibited increased levels of various types of mitotic spindle abnormalities, including lagging chromosomes, multiple centrosomes, and microtubule aggregates at 48 h posttransfection (Figure 7, D–J), when aneuploidy is observed but before apoptosis is observed. These data provide strong support for the conclusion that microtubule instability and consequent mitotic spindle dysfunction are a mechanism by which mutant MAPT induces chromosome mis-segregation, aneuploidy, and, ultimately, apoptosis.

**Apoptosis induced by expression of MAPT harboring FTLD-causing mutations is not caused by oxidative stress**

Our data thus far show that FTLD-causing forms of human MAPT induce aneuploidy and apoptosis in human brain cells and when expressed in a karyotypically normal human cell line. The temporal order of aneuploidy followed by apoptosis in our cell culture experiments suggests a cause–effect relationship. However, it is theoretically possible that FTLD-associated MAPT mutations cause both aneuploidy and apoptosis through some common intermediate step. Indeed, oxidative stress has been described in transgenic mice expressing MAPT harboring the P301S FTLD mutation (Lopez-Gonzalez et al., 2015). To test the possibility that FTLD-causing forms of MAPT induce oxidative stress, which in turn leads to both aneuploidy and apoptosis, we measured oxidative stress levels, including reactive oxygen species (ROS) and superoxide, in cells expressing the FTLD mutant forms of MAPT. In these experiments, we found no evidence of increased stress relative to controls (Supplemental Figure 2). These findings show that the effects of FTLD mutant MAPT on aneuploidy and apoptosis are not likely to be caused by oxidative stress.

**Expression of MAPT harboring FTLD-causing mutations induces mitotic spindle defects in karyotypically normal human cells**

To investigate the mechanism by which MAPT dysfunction leads to chromosome mis-segregation, aneuploidy, and subsequent apoptosis, we analyzed mitotic spindle structures in hTERT-HME1 cells transfected with a vector for expressing wild-type MAPT, MAPT P301L, or MAPT V337M or with the vector alone as a negative control (Figure 7).
Mutant MAPT-induced aneuploidy and apoptosis both depend on cell division

The data presented above are most consistent with the defects in mitosis and consequent chromosome mis-segregation and aneuploidy preceding and thus likely leading to apoptosis (Figure 8). As a more direct test of whether the aneuploidy caused by mutant MAPT expression induces apoptosis or whether the apoptosis caused by mutant MAPT expression occurs independently of the aneuploidy, hTERT-HME1 cells transfected with the pcdNA vector for expressing MAPT P301L or MAPT V337M, or with the pcdNA vector alone as a negative control, were treated for 48 h with Nutlin-3 at 24 h posttransfection to halt cell division and inhibit the generation of aneuploid cells. As expected, cells expressing mutant MAPT (P301L or V337M) exhibited significantly higher levels of aneuploidy (Supplemental Figure 3A) and apoptosis (Supplemental Figure 3B) relative to mock transfected cells, cells transfected with the pcdNA vector, or cells expressing wild-type MAPT; however, Nutlin-3 treatment of cells expressing mutant MAPT (P301L or V337M) prevented them from becoming aneuploid, as measured using a FISH probe for chromosome 21 (Supplemental Figure 3C), and it also prevented them from undergoing apoptosis, as measured by TUNEL staining (Supplemental Figure 3D). If the expression of mutant MAPT induces apoptosis independently of the aneuploidy, then we would have expected to observe increased apoptosis but not increased aneuploidy in the cells expressing mutant MAPT, since the Nutlin-3 treatment would prevent them from dividing and becoming aneuploid. However, the data clearly show that both mutant MAPT-induced aneuploidy and apoptosis were suppressed by Nutlin-3 treatment, indicating that the apoptosis is dependent on the aneuploidy caused by abnormal cell division (Supplemental Figure 3).

DISCUSSION
Aneuploidy and apoptosis in various neurodegenerative diseases

Previous studies have shown that aneuploidy occurs in neurons, glia, and other cells in sporadic and all familial forms of AD (Potter, 1991; Potter et al., 1995; Migliore et al., 1997; Geller and Potter, 1999; Kingsbury et al., 1991; Potter et al., 2006; Mosch et al., 2007; Thomas and Fenech, 2008; Iourov et al., 2009) and that the specific degeneration of aneuploid neurons that occurs between midstage and advanced AD accounts for 90% of total neuronal cell loss observed at autopsy (Arendt et al., 2010). Although chromosome instability was described previously in peripheral cells from FTLD patients with MAPT mutations (Rossi et al., 2008), aneuploidy levels in the brain were not determined.

Here we show that the brains of familial FTLD patients with MAPT mutations exhibit significantly increased levels of mosaic aneuploidy for chromosomes 12 and 21, together with a significant
FIGURE 8: Proposed theoretical mechanistic pathway of aneuploidy and apoptosis in FTLD-MAPT and other tauopathies. On the basis of the evidence presented here, which shows that mutant MAPT-induced aneuploidy and apoptosis require cell division, we hypothesize that the expression of mutant forms of MAPT in the cells of FTLD-MAPT patients (or in patients with other tauopathies) would induce mitotic spindle defects, which would in turn lead to the accumulation of aneuploid cells over time as a result of aberrant cell division. An increase in the aneuploid cell population would not only place a heavy burden on the aneuploid cells to overcome the detrimental effects of altered gene dosage but also render them more vulnerable and prone to undergoing apoptosis.

FIGURE 7: Mitotic spindle defects induced by expression of MAPT harboring FTLD-causing mutations in a karyotypically normal human cell line. The hTERT-HME1 cell line was transfected in the absence of vector (Reagent), with the pcDNA3.1 vector alone (pcDNA), or with the pcDNA3.1 vector harboring wild-type MAPT (WT-MAPT), MAPT with the P301L mutation (P301L), or MAPT with the V337M mutation (V337M), and microscopy images were taken during the metaphase and anaphase stages of mitosis at 48 h posttransfection; microtubules are shown in red, and DNA is shown in blue. Normal spindles: the chromosomes are aligned properly in metaphase and are also undergoing proper segregation in anaphase (A–C). Abnormal spindles: lagging chromosomes, the majority of the chromosomes are aligned properly but some remain unattached and unaligned (D, E); multiple centrosomes: more than two microtubule arrays are present during mitosis (F, G); and microtubule aggregate: the tubulin subunits do not polymerize properly but instead form a cluster inside the nucleus (H, I). Cells expressing the mutant MAPT genes (P301L or V337M) show a significant increase in MT abnormalities in comparison to samples transfected with the pcDNA vector alone or in the absence of vector (J). The graphs show mean difference and significance (*) of the cells transfected with the mutant MAPT genes compared with the pcDNA3.1 vector alone (sham-control) using ANOVA with a Tukey post hoc test. Error bars indicate SEM. **p < 0.005.

Our finding that apoptosis occurs preferentially in aneuploid cells from familial FTLD patients with MAPT mutations, including neurons, combined with our observation that a karyotypically normal human cell line expressing MAPT with an FTLD-causing mutation first exhibits aneuploidy without apoptosis at 48 h and then undergoes apoptosis at 72 h suggest that apoptosis arises from aneuploidy and not vice versa. This conclusion was tested directly by experiments with Nutlin-3 showing that the aneuploidy and the increase in apoptosis in both neuronal and nonneuronal cells. Furthermore, ~80% of the neuronal apoptosis occurs in the aneuploid neurons, which parallels the finding described above that 90% of neuronal cell death in advanced AD can be attributed to the specific loss of aneuploid neurons (Arendt et al., 2010).

Our finding that aneuploidy precedes and is causally implicated in the development of FTLD suggests a common mechanism for these conditions that may explain the genetic and environmental factors that contribute to their pathogenesis.
apoptosis induced by mutant MAPT expression in cultured cells are not independent of each other, because both require cell division. Our additional finding of mosaic aneuploidy in nonneuronal brain cells from FTLD patients with MAPT mutations reinforces the previous finding of chromosome instability in peripheral cells of FTLD patients with MAPT mutations (Rossi et al., 2008) and indicates that chromosome mis-segregation is likely a general feature of all cells in patients with FTLD-MAPT. The fact that mitotic spindle abnormalities and aneuploidy arise within two mitotic cell cycles (at 48 h post-transfection) in karyotypically normal human cells expressing FTLD-causing mutant forms of MAPT and later lead to apoptosis provides a mechanistic link between microtubule dysfunction and neurodegeneration in FTLD-MAPT. Over time, an individual with FTLD-MAPT who has carried one of these MAPT mutations in all of their cells for their entire life would be expected to also show significant levels of mosaic aneuploidy in cells throughout their body, and their aneuploid neurons would be at significant risk for apoptosis.

The general cell-cycle dysfunction shown in our FTLD-MAPT experiments and the similar findings in AD indicate that multiple chromosomes undergo mis-segregation in these and potentially other neurodegenerative diseases (Potter et al., 1995; Migliore et al., 1997; Geller and Potter, 1999; Kingsbury et al., 2006; Mosch et al., 2007; Boeras et al., 2008; Thomas and Fenech, 2008; Iourov et al., 2009; Arendt et al., 2010, 2015; Arendt, 2012; Granic et al., 2010; Borysov et al., 2011; Yurov et al., 2014). For example, we have found that Niemann-Pick Type C1, another tauopathy, caused by defects in cholesterol metabolism, is characterized by aneuploid neurons and other cells (Granic and Potter, 2013). In addition, a recent study of Lewy body dementia showed an increase in neuronal DNA content (hyperdiploidy) that correlated with neuronal loss in all brain regions examined (Yang et al., 2016). These results together with our studies of AD and FTLD-MAPT support the general conclusion that aneuploid neurons arise by different mechanisms in multiple forms of neurodegenerative disease and that the increasing susceptibility of aneuploid neurons to apoptosis may underlie much of the neuronal loss and cognitive decline that characterize these disorders.

Potential means by which aneuploid neurons arise in MAPT-FTLD

The generation and accumulation of aneuploidy in dividing cell populations might, in principle, arise by both genetic and environmental stressors at any time in life (discussed in Potter, 1991, and Oromendia and Amon, 2014). However, it has long been believed that neurons are postmitotic cells and furthermore that vertebrates are born with a finite number of neurons with little or no neurogenesis occurring in the adult brain (Bhardwaj et al., 2006). These long-held beliefs raise the question of when and how aneuploid neurons arise during age-associated neurodegenerative disease.

Aneuploidy generation during embryogenesis. One explanation for neuronal aneuploidy in neurodegenerative diseases is that it arises in neuronal precursor cells during embryogenesis and then later leads to neurodegeneration. Certainly, our data show that having an FTLD-causing mutation in MAPT is sufficient to induce abnormal mitotic spindles and subsequent aneuploidy. However, the mechanism by which age may contribute to the process remains unclear. It is possible that aging may promote the development of additional processes that promote the degeneration of aneuploid neurons in these disorders (Potter, 1991; Potter et al., 2016).

Aneuploidy generated during adult neurogenesis and aging. Although neurogenesis appears to be rare in normal adult brain, three processes may generate the neuronal aneuploidy observed at autopsy in patients with neurodegenerative diseases and may arise in the adult and be promoted by aging. First, data from many studies over the past couple of decades have provided evidence that neurogenesis can be induced in many brain regions in adult mice and rats in response to brain damage (Zhao et al., 2008; Spalding et al., 2013; Zheng et al., 2013; Ibrahim et al., 2016) and attempted self-repair by the brain or as part of an ongoing process in the subventricular/granular zone of the brain (Eriksson et al., 1998; Hallbergson et al., 2003; Sakamoto et al., 2014). Therefore, neuronal damage and the mitotic defect in FTLD-MAPT could produce aneuploid replacement neurons, which would not be fully functional and would be particularly prone to apoptosis and degeneration, as our experiments have shown. Aneuploidy has been shown to promote degeneration in many other experimental systems, possibly by inducing proteotoxic stress (Rajendran et al., 2008; Kai et al., 2009; Arendt et al., 2010; Oromendia and Amon, 2014).

The second potential mechanism for aneuploidy generation during aging and neurodegeneration is cell-cycle reentry. Neurons in the AD brain express phosphoproteins usually detected only during mitosis (McShea et al., 1997; Vincent et al., 1997; Arendt, 2012), and in AD mice, the loss of preexisting neurons induces neuronal reentry into the cell cycle (Lopes et al., 2009; Seward et al., 2013). Indeed, Aβ induces the expression of mitotic proteins and cell-cycle reentry in mature neurons in culture (Majid et al., 2008; Seward et al., 2013). A similar process may be occurring in familial FTLD associated with MAPT mutations.

Finally, the recent discovery that striatal astrocytes can transdifferentiate into new neurons capable of forming functional neuronal circuits with preexisting neurons following ischemic brain injury (Magnusson et al., 2014; Duan et al., 2015) suggests that the aneuploid glia we observed in FTLD brains may have the potential to transdifferentiate into the aneuploid neurons that we also observed. It seems very likely that the aging process may exacerbate any or all of these processes that have the potential to generate aneuploid neurons and other cells. Indeed, aneuploidy has been proposed and found to increase with age (Arendt et al., 2009; Yurov et al., 2009, 2010; Fischer et al., 2012).

Evidence that MAPT plays a role in chromosome segregation in both FTLD and AD

Our studies showing that MAPT plays a role in chromosome segregation in some forms of FTLD is consistent with previous findings in AD, which showed that mutant forms of APP or PSEN1 or Aβ itself also induce chromosome mis-segregation in a MAPT-dependent manner (Boeras et al., 2008; Granic et al., 2010; Borysov et al., 2011). Additionally, the fact that MAPT and APP phosphorylation and processing are cell-cycle-dependent modifications that increase during mitosis, and that MAPT, APP, and presenilin are localized to mitotic structures, such as centrosomes (Li et al., 1997; Sjöberg et al., 2006; Rossi et al., 2008; Judge et al., 2011), further implicates both MAPT and Aβ function in the process of chromosome segregation.

Other studies linking aneuploidy, neurodegeneration, and MAPT

Evidence linking neuronal aneuploidy, neurodegeneration, and MAPT was reported recently by two other groups in Drosophila. Specifically, a study by Bougé and Parmentier (2016) showed that excess Tau causes mitotic spindle defects, aneuploidy, and apoptosis through inhibition of the microtubule-dependent motor protein Kinesin-5/Eg5. Similar results have been observed by Malmanche et al. (2017), who examined photoreceptors and brain
neurons in Drosophila and found that adult-onset neurodegeneration mediated by MAPT overexpression included the generation of aneuploid cells (Malmanche et al., 2017). The former result is of particular interest in light of our prior finding that Aβ induces chromosome mis-segregation and aneuploidy though competitive inhibition of Kinesin-5/Eg5 (Borysov et al., 2011). Thus, causal mutations leading to AD and FTLD-MAPT appear to lead to chromosome mis-segregation, aneuploidy, and apoptosis through inhibition of the same target enzyme: Kinesin-5/Eg5.

Possible explanations for how a general cell-cycle defect can lead to regional neurodegeneration

Both AD and FTLD and indeed all neurodegenerative disease show pathology and cell loss in specific regions of the brain, while sparing other regions, and this regional specificity correlates roughly with the clinical symptoms (e.g., tau pathology in AD). Because APP and MAPT are expressed widely, other factors must control essential steps in the pathogenic pathway that results in regional specificity. In AD, the specificity arises from differences in the expression of modulator proteins, such as the amyloid catalysts alpha-1-antichymotrypsin (ACT) and ApoE, which are essential for the conversion of Aβ to oligomers and filaments. Indeed, only Aβ oligomers cause chromosome mis-segregation and aneuploidy (Granic et al., 2010; Borysov et al., 2011). Both ACT and ApoE are inflammation-associated proteins in the brain that are not expressed highly in the cerebellum in AD, for example, where amyloid pathology also does not arise. Prion disease research has also found that species and regional specificity must involve a specific component, termed “protein X,” that is needed for pathology (Yehiel et al., 1997). Protein X has yet to be definitively identified, but one study showed that it is not ApoE (Tatzelt et al., 1996).

The finding that APP and Presenilin mutations and their product, oligomeric Aβ peptide, cause chromosome mis-segregation in cells, transgenic mice, and humans through disruption of the mitotic spindle and that the consequent aneuploidy may underlie much of the neurodegeneration, as discussed above, also begs the same question of how an effect on a common physiological process such as mitosis can lead to region specific neurodegeneration. As anticipated above, the explanation in AD may lie in the regional expression of ApoE.

Similarly, we suggest that one potential explanation for the regional specificity of FTLD is that another protein that is expressed in the cortex interacts with mutant MAPT to promote its pathological effect on microtubules. It is also possible that such a protein or even the direct effect of mutant MAPT only occurs in dividing cells, as our experiment with Nutlin-3 indicates. In that case, only neurons undergoing cell-cycle reentry or being replaced by neurogenesis would be sensitive to mutant MAPT. Alternatively, other microtubule-binding proteins that can substitute for damaged tau may be normally preferentially expressed in certain brain regions (i.e., those not affected in FTLD), thereby underlying region-specific protection.

Targeting microtubule and mitotic spindle dysfunction to prevent neurodegeneration

Taken together with the results of our previous work and recent reports, the data presented herein provide a novel mechanism for neurodegeneration and neuronal dysfunction in FTLD based on microtubule and mitotic spindle dysfunction (Figure 8). Further, the findings may lead to new general treatments or preventative strategies for a wide range of neurodegenerative disorders, including FTLD, AD, Niemann Pick Type C1, ALS, Lewy body dementia, and Down syndrome. For example, microtubule stabilization, by drug therapy, by inhibition of MAPT phosphorylation, or by rescue of inhibited Kinesin-5/Eg5 activity, may improve chromosome segregation and lead to reduced levels of aneuploidy and subsequent neurodegeneration.

MATERIALS AND METHODS

Cell line

The hTERT-HME1 (Clontech) is a telomerase-immortalized normal human mammary epithelial cell line that divides indefinitely and retains normal function, phenotype, and karyotype (Jiang et al., 1999). The cell line was cultured in supplemented mammary epithelium basal medium (MEBM; Lonza), as described previously (Boerars et al., 2008; Granic et al., 2010). Early cell passages were used for the experiments.

Primary mouse cells

Mouse spleens and brains were harvested and processed for culture or immediately fixed as single-cell suspensions for FISH and FISH/immunocytochemistry, as described previously (Boerars et al., 2008; Granic et al., 2010; Granic and Potter, 2013; Caneus et al., 2017).

Human brain cells

Approximately 1–2 g of frozen human brain cortices from patients diagnosed with familial FTLD associated with MAPT mutations (aged 56 ± 5 yr) and controls (aged 65 ± 9 yr; p = 0.054) was obtained from the Mayo Clinic Brain Bank and the Banner Sun Health Research Institute and Brain and Body Donation Program, respectively. The tissues were dissociated into single-nuclei suspensions, fixed, and processed for FISH in combination with NeuN and/or TUNEL immunocytochemistry, as described previously (Granic et al., 2010; Granic and Potter, 2013).

Mice

Tau−/- knockout mice (obtained from the Jackson Laboratory, B6.129 × 1-MAPT<sup>tm</sup>V337M/J) [Dawson et al., 2001] and their heterozygous (tau+/−) and wild-type (tau+/+) littermates were 6–7 mo old (background strain: C57BL/6). Transgenic mice expressing a human FTLD-causing mutation in the MAPT gene (P301S) from a prion promoter and their nontransgenic control (NON) were 8–9 mo old and were obtained from the Jackson Laboratory (B6.C3-Tg[Prnp-MAPT*P301S]PS19Vle/J). In addition, brain tissue samples from an FTLD mouse model expressing the human mutant MAPT (P301L) transgene under the control of the prion promoter and control nontransgenic mice (−8 wk old) were purchased from the Jackson Laboratory (FVB-Tg[tetO-MAPT*P301L]#Kha/J#Jwsl and FVB/NJ, respectively).

Plasmids

Plasmids constructed by inserting the wild-type human MAPT cDNA or the human MAPT cDNA harboring an FTLD-causing mutation (P301L or V337M) into the pcDNA3.1 expression vector were gifts from Chad Dickey (University of South Florida, Tampa). The Nucleo-Bond Plasmid Purification Kit (BD Bioscience) was used for nucleic acid extraction from NovaBlue Singles competent cells (Novagen).

Antibodies, kits, and DNA probes

Mouse/rabbit anti-neuronal nuclei (NeuN) AlexaFluor antibodies (Millipore, MAB377X; Abcam, ab190195) and DAPI (4',6-diamidino-2-phenylindole) II (Abbott, Vyysis) or VectaShield with DAPI (Vector Laboratories) were used to label neurons and nuclei, respectively. A BAC probe to mouse chromosome 16 was either purchased (Mouse iDetect Chromosome 16 D16Mit88 Point Probe, RP24–288N19, Empire Genomics) or prepared directly using a Nick Translation Kit
cells (Roche) with Spectrum Green or Red dUTP (Vysis) using a chromoso-

some 16 BAC plasmid (a gift from Bruce Lamb, The Cleveland

Clinic), as described previously (Boeras et al., 2008; Granic et al.,

2010; Granic and Potter, 2013). A mixture of fluorescently labeled

DNA probes for the detection of human chromosomes 12 and 21

(LSI Tel/AML1 ES Dual Color Translocation Probe) or individual

probes for the detection of human chromosome 21 (Vysis Tel/Vysis

21q SpectrumOrange Probe, Vysis LSI ERG [Tel] SpectrumGreen

Probe, Vysis LSI ERG [Cen] SpectrumRed Probe) were obtained from

Abbott, Vysis. TUNEL assay was carried out using the Click-iT Plus

TUNEL Assay for in situ apoptosis detection with Alexa Fluor dyes

(molecular probes) on brain samples and cultured cells in with or

without Nutlin-3 (Selleckchem; Cat. No. S1061). A cellular ROS/

Superoxide Detection Assay Kit (Abcam; ab139476) was used to

measure oxidative stress levels in cultured cells.

FISH analysis

Single nuclei suspensions were dropped on glass slides (Fisher-

Brand) and air-dried/aged at room temperature overnight. The

slides/nuclei were then subjected to hybridization procedures ac-

curring to the manufacturer (Abbott, Vysis) using established proto-

cols with the mouse and human probes described above and previ-

ously, including hybridization in a HyBrite chamber (Vysis). In

subsequent steps, the slides were either counterstained with Vecta-

shield with DAPI immediately (cultured cells) or processed for NeuN

immunostaining followed by DAPI counterstain (brain samples)

(Boeras et al., 2008; Granic et al., 2010; Granic and Potter, 2013;

Caneus et al., 2017).

TUNEL staining combined with FISH and

immunocytochemistry for brain cells

Single-nuclei suspensions prepared from either frozen human brain tis-

sues or from hTERT cells transfected for 72 h with mutant MAPT genes

were fixed, dropped, and air-dried overnight on glass slides and sub-

jected to TUNEL staining, which involved incubation with a recombi-

nant terminal deoxynucleotidyl transferase (TdT) and dUTPs with Alexa

Fluor dyes for a combined 1 h, 40 min at 37°C, according to the manu-

ufacturer’s recommended procedure (Click-iT Plus TUNEL Assay; Life

Technologies). Following TUNEL staining, the cells were processed for

FISH with DNA probes for chromosomes 12 and 21 and immunocyto-

chemistry with anti-NeuN antibodies (Millipore and Abcam), as de-

scribed previously. On average, more than 500 cells/slide and three

slides/sample were counted for the transfected hTERT cells, and ~1000

cells/slide and two slides were counted for each human subject.

Transient transfection

The hTERT-HME1 cells (~1.8 x 10⁶ cells/2 ml) were plated into a six-

well plate and transfected with an empty pcDNA3.1 vector, with the

pcDNA3.1 vector harboring a wild-type or mutant (P301L or V337M)

human MAPT cDNA, or with a mixture of reduced Opti-MEM and

FuGENE 6 transfection reagent (Promega), according to the manu-

ufacturer’s protocol, as described previously (Boeras et al., 2008;

Granic et al., 2010). Cells from three parallel transfections were

treated with Nutlin-3 at 24 h posttransfection and incubated for an

additional 48 h. The samples were then harvested, fixed, and scored for

aneuploidy using chromosome 21 FISH and apoptosis using

TUNEL at 72 h posttransfection.

Immunocytochemistry of hTERT-HME1 cells for

microtubule analyses

Confluent hTERT cells were harvested, plated at a density of

~1.5 x 10⁵ cells/2 ml, and treated with the FuGENE 6 reagent

alone or transfected with a FuGENE 6-DNA complex of either the

plasmid vector (pcDNA3.1) or the pcDNA3.1 plasmid vector har-

boring wild-type MAPT, MAPT V337M, or MAPT P301L and cul-

tured on glass chamber slides. The cells were allowed to pro-

gress through the cell cycle and harvested at the metaphase/ anaphase

stages of mitosis, fixed in methanol for 5 min at ~20°C, subjected to

indirect immunofluorescence staining with a mouse anti-alpha-tubulin

antibody (Abcam; Cat. Ab28439) and Alexa Fluor-conjugated goat anti-
mouse antibody, and mounted in Vectashield with DAPI.

Image acquisition and analysis

FISH hybridization signals were analyzed following Abbott/Vysis

guidelines as described (Boeras et al., 2008; Granic et al., 2010).

Between 800 and 1000 interphases per sample per treatment were

scored for aneuploidy, and at least 400 NeuN(+) and NeuN(-)

mouse brain cells were analyzed for BAC DNA signals. In addition, the

percentage of cells that were positive for TUNEL labeling was

recorded and scored for apoptosis analysis. Mitotic spindles and

data were visualized and scored for quantitative analysis using ZEN

2011 SP1 imaging software. Each of the in vitro experiments was

done in triplicate.

Statistics

Normality of data was determined by Shapiro-Wilk test. Independent

t tests were used to compare aneuploidy in FTLD and control brain

and spleen cells in the experiments using transgenic mice expressing

human MAPT P301S. One-way analysis of variance (ANOVA) with

Tukey HSD and Dunnett test was used to establish significance in all

other studies, as indicated by (*) or presented in the figures. The ef-

fect size (r² for ANOVA and Cohen’s d for t test) was calculated for

the main outcomes. The analyses were performed in Microsoft Excel

2013 and IBM SPSS Statistics version 22.

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