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# Truncated stathmin-2 is a marker of TDP-43 pathology in frontotemporal dementia

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**No treatment for frontotemporal dementia (FTD), the second most common type of early-onset dementia, is available, but therapeutics are being investigated to target the 2 main proteins associated with FTD pathological subtypes: TDP-43 (FTLD-TDP) and tau (FTLD-tau). Testing potential therapies in clinical trials is hampered by our inability to distinguish between patients with FTLD-TDP and FTLD-tau. Therefore, we evaluated truncated stathmin-2 (*STMN2*) as a proxy of TDP-43 pathology, given the reports that TDP-43 dysfunction causes truncated *STMN2* accumulation. Truncated *STMN2* accumulated in human induced pluripotent stem cell–derived neurons depleted of TDP-43, but not in those with pathogenic *TARDBP* mutations in the absence of TDP-43 aggregation or loss of nuclear protein. In RNA-Seq analyses of human brain samples from the NYGC ALS cohort, truncated *STMN2* RNA was confined to tissues and disease subtypes marked by TDP-43 inclusions. Last, we validated that truncated *STMN2* RNA was elevated in the frontal cortex of a cohort of patients with FTLD-TDP but not in controls or patients with progressive supranuclear palsy, a type of FTLD-tau. Further, in patients with FTLD-TDP, we observed significant associations of truncated *STMN2* RNA with phosphorylated TDP-43 levels and an earlier age of disease onset. Overall, our data uncovered truncated *STMN2* as a marker for TDP-43 dysfunction in FTD.**

## Introduction

Tar DNA-binding protein 43 (TDP-43) regulates multiple aspects of RNA metabolism, including RNA stabilization, transcription, splicing, and transport (1). TDP-43 binds thousands of RNAs (2,

3) and, for particular genes, represses nonconserved cryptic exons in a species- (4) and cell-type-specific manner (5, 6). Recently, 2 groups independently found that TDP-43 is required for the correct splicing and expression of stathmin-2 (*STMN2*) (7, 8), a microtubule-associated protein involved in axonal regeneration (9). As shown schematically in Figure 1, when TDP-43 is present and functional, it binds to GU-rich sequences in the first intron of the *STMN2* transcript, allowing normal splicing of intron 1 and repressing the usage of an alternative or cryptic polyadenylation site. Reduced TDP-43 binding to *STMN2* pre-mRNA, which could occur as a result of TDP-43 nuclear depletion, leads to the inclusion of a region within intron 1 (exon 2a) and introduces a premature stop codon and polyadenylation. This causes the accumulation of a truncated variant of *STMN2* that lacks exons 2 through 5.

### ► Related Commentary: p. 5677

**Authorship note:** MP, JH, and SP are co-first authors and contributed equally to this work.

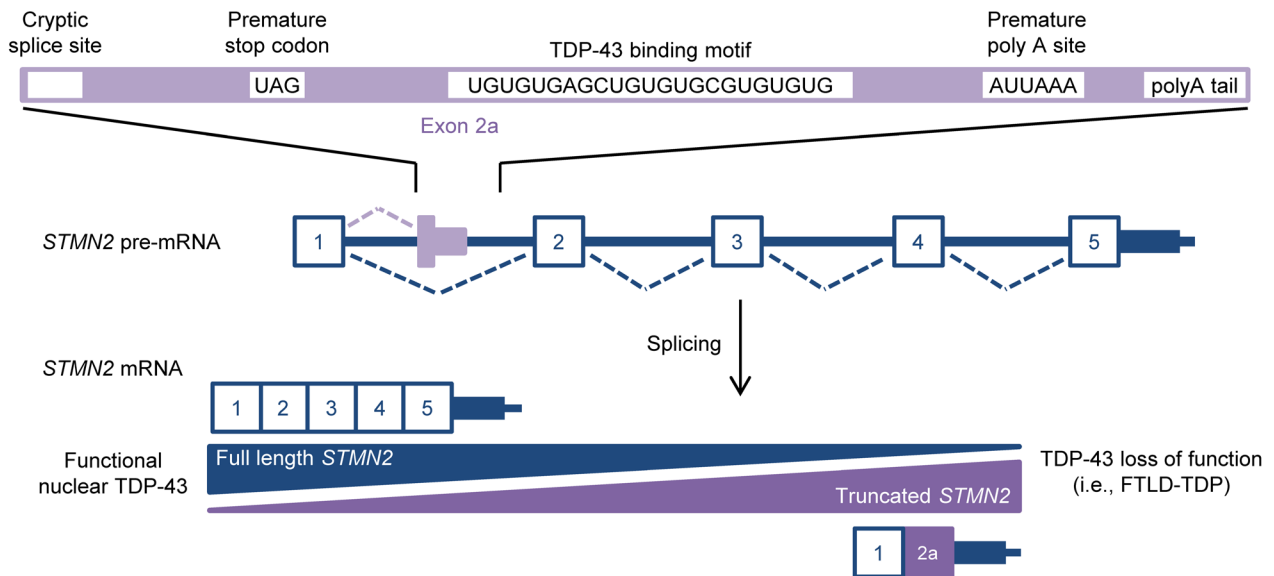
**Conflict of interest:** RCP is a consultant for Roche Inc., Merck Inc., Biogen Inc., and Eisai Inc.; is a member of the data and safety monitoring board (DSMB) of Genentech; and has provided educational talks for GE Healthcare.

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**Figure 1. Truncated *STMN2* RNA is generated with loss of nuclear TDP-43.** Schematic of TDP-43-regulated *STMN2* splicing. When TDP-43 is present and functional, it binds to GU-rich sequences in the first intron of the *STMN2* transcript and allows normal splicing of intron 1. Reduced TDP-43 binding to *STMN2* RNA, from TDP-43 aggregation or depletion from the nucleus, leads to mis-splicing of *STMN2* RNA, resulting in the inclusion of a novel exon encoded within intron 1 (termed exon 2a) and containing an alternative or cryptic polyadenylation site. The production of this alternative variant of *STMN2*, which lacks exons 2 through 5 (referred to as truncated *STMN2*), is at the expense of full-length *STMN2*, which is reduced upon TDP-43 downregulation.

Neuronal and glial inclusions of TDP-43 are a pathological hallmark of the majority (~97%) of patients with ALS and of approximately 50% of cases with the pathological subtype of frontotemporal dementia (FTD), known as frontotemporal lobar degeneration-TDP (FTLD-TDP) (1). ALS is a motor neuron disease (MND), whereas patients with FTD present clinically with 1 or more of the FTD syndromes characterized by behavior, executive function, language, and motor impairments (10). TDP-43 abnormalities are proposed to play a direct role in these diseases, since mutations in *TARDBP*, the gene that encodes TDP-43, were discovered in sporadic and familial ALS and FTD (11, 12). It is also notable that TDP-43 morphology and distribution are associated with clinical phenotypes and neurodegeneration in key anatomical regions, such as frontal and temporal cortices in FTLD-TDP (13). Cells with TDP-43 pathology feature a redistribution of TDP-43 from the nucleus to the cytoplasm, where TDP-43 is ubiquitinated, hyperphosphorylated, and cleaved into C-terminal fragments (14). Patients with FTLD-TDP can be further classified into different pathological subtypes on the basis of the morphology and distribution of the TDP-43 inclusions observed (15, 16). These can take the form of neuronal cytoplasmic inclusions, dystrophic neurites, neuronal intranuclear inclusions, glial inclusions, perivascular inclusions, and fine neurites in the hippocampus (14, 17–20). At present, 5 subtypes of FTLD-TDP (A–E) have been identified (15, 20–23), with types A–C accounting for more than 95% of all FTLD-TDP cases (24). FTLD-TDP type A, the most common subtype, is characterized by the presence of all the inclusion types mentioned above, whereas types B–E are characterized by the presence of a predominant inclusion type (19, 22). Furthermore, some FTLD-TDP subtypes are associated with particular mutations (20, 25, 26).

While approximately 50% of patients with an FTD syndrome have underlying TDP-43 pathology, such pathology can, at present, only be definitively confirmed at autopsy. Given the efforts to develop therapies specific to FTLD pathological subtypes (i.e., FTD with TDP-43 vs. tau pathology), there is a clear need for biomarkers that discriminate between them. Further, biomarkers for diagnosis or prognosis or that can monitor TDP-43 activity in response to TDP-43-tailored therapies are also sorely lacking. Although TDP-43 represents an attractive biomarker candidate to fulfill these needs, efforts to develop sufficiently sensitive immunoassays for the detection of TDP-43 or its pathological forms in biofluids and that reliably differentiate between individuals with and without TDP-43 pathology have proven disappointing (27). However, nuclear clearance of TDP-43, in the absence of TDP-43 inclusions, has been associated with neuronal atrophy in patients with FTLD-TDP or ALS-TDP (28), suggesting that early mis-splicing events due to TDP-43 loss of function may represent a robust and early disease biomarker.

We thus propose that the mis-splicing of *STMN2* may provide a novel strategy for the indirect assessment of TDP-43 dysfunction. To test this hypothesis, we first validated the *STMN2* splicing event in induced pluripotent stem cell-derived (iPSC-derived) cortical and motor neurons that were TDP-43 depleted. Then, we evaluated whether the truncated *STMN2* isoform accumulates in diseases and tissues characterized by TDP-43 pathology using data sets from RNA-Seq studies of brain tissues from a large cohort of control subjects and of patients with FTD or ALS with or without TDP-43 pathology. Furthermore, we measured truncated and full-length *STMN2* RNA levels in postmortem medial frontal cortex tissue from FTLD-TDP, progressive nuclear palsy (PSP), and pathologically normal control brains. PSP was selected as the disease control, given that TDP-43 pathology is not found in the frontal cortex of these patients with FTLD-tau (29). We addition-

ally evaluated whether truncated *STMN2* levels in the medial frontal cortex discriminated between patients with FTLT-DTP and control subjects and assessed associations of truncated *STMN2* with phosphorylated TDP-43 (p-TDP-43) and clinical features.

## Results

*TDP-43 depletion in human iPSC-derived neurons leads to aberrant STMN2 splicing.* To confirm recent reports that correct *STMN2* splicing requires TDP-43 (7, 8), we used clustered regularly interspaced short palindromic repeat inactivation (CRISPRi) to knock down TDP-43 expression in human iPSC-derived cortical neurons that constitutively express nuclease deactivated Cas9 (dCas9) fused to blue fluorescent protein (BFP) and the transcriptional repressor Krüppel-associated box (KRAB) (dCas9-BFP-KRAB) (30). To achieve this, cells were transduced with a lentivirus expressing either an sgRNA against the transcriptional start site of *TARDBP* (TDP-43 KD), or a control sgRNA predicted not to bind anywhere in the human genome (controls). We confirmed silencing of *TARDBP* RNA by quantitative reverse transcription PCR (qRT-PCR) and RNA-Seq (data set labeled “a”), which showed TDP-43-KD efficiencies of approximately 50% (47.4% for qRT-PCR, 63.9% for RNA-Seq) (Figure 2A). Reduced *TARDBP* levels resulted in significant reductions of full-length *STMN2* RNA (56.7% for qRT-PCR, 65.5% for RNA-Seq) (Figure 2B) and a significant increase in truncated *STMN2* RNA (qRT-PCR, 32.3-fold increase) (Figure 2C) and RNA-Seq (Figure 2D, data set labeled “a”) compared with levels in neurons expressing control sgRNA. In parallel, we analyzed published RNA-Seq data from Klim et al. on iPSC-derived motor neurons in which *TARDBP* was knocked down (7). Knockdown of *TARDBP* (Supplemental Figure 1A; supplemental material available online with this article; <https://doi.org/10.1172/JCI139741DS1>) was confirmed in the Klim data set (data set labeled “b”) and, as expected, was accompanied by a reduction in full-length *STMN2* RNA levels (Supplemental Figure 1B) and an increase in truncated *STMN2* RNA levels (Figure 2D, data set labeled “b”).

We also investigated whether disease-causing mutations of *TARDBP*, in the absence of TDP-43 protein mislocalization or a reduction of nuclear levels, are sufficient to impact *STMN2* splicing. We evaluated truncated *STMN2* expression in RNA-Seq data from 2 independent studies that used iPSC-derived motor neurons from patients carrying ALS pathogenic *TARDBP* mutations and from control subjects (Supplemental Table 1). In contrast to neurons in which *TARDBP* was downregulated, we observed no increase in truncated *STMN2* in motor neurons with *TARDBP* mutations (Figure 2D, data sets labeled “c” and “d”). This observation was consistent with the lack of change in TDP-43 localization (Figure 2E), *TARDBP* RNA levels (Supplemental Figure 2), and TDP-43 protein levels (Supplemental Figure 3) in *TARDBP*-mutant motor neurons. Overall, these data suggest that truncated *STMN2* accumulation in iPSC-derived neurons results from a loss of TDP-43 function due to TDP-43 depletion and not from splicing function changes arising from mutations in *TARDBP* (31). The redistribution of TDP-43 from the nucleus and its sequestration into cytoplasmic inclusions, a pathological hallmark of FTD and ALS, with and without a genetic link to *TARDBP*, are consistent with loss-of-function mechanisms that would lead to *STMN2* mis-splicing and an upregulation of truncated *STMN2* expression and prompted us to further examine this in human tissue from FTD and ALS patients.

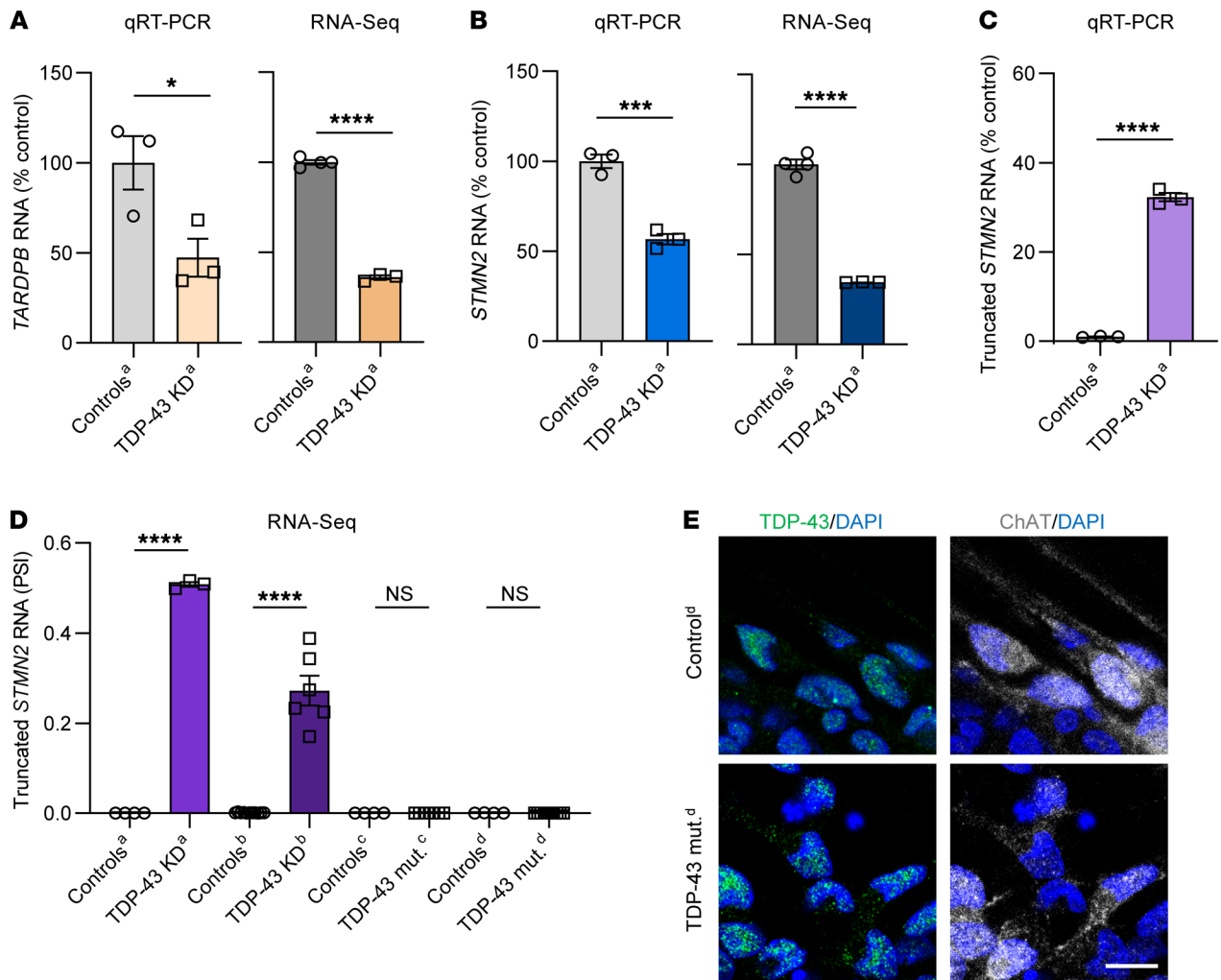
*Analyses of bulk brain tissue RNA-Seq data uncover tissue-specific truncated STMN2 RNA accumulation in diseases characterized by TDP-43 pathology.* We quantified the expression of full-length and truncated *STMN2* in the NYGC ALS Consortium RNA-Seq data set across multiple neuroanatomical regions and diseases. As of March 2020, we were able to examine 11 tissues, with 1659 RNA-Seq samples from 439 donors matching established criteria for inclusion (see Methods). Donors were diagnosed as non-neurological disease controls or as having FTD, ALS, FTD with ALS (ALS-FTLD), or ALS with suspected Alzheimer’s disease (ALS-AD). Patients with FTD were classified on the basis of their neuropathological diagnosis: FTLT-DTP, FTLT-tau, or FTLT with mutations in *FUS* (FTLT-FUS). ALS samples were divided into those with a *SOD1* mutation (ALS-SOD1) versus those without a *SOD1* mutation and thus assumed to have TDP-43 pathology (ALS-TDP). However, pathology confirming the presence of TDP-43 mislocalization in patients with ALS-TDP was not systematically reported (see Methods).

For each sample, we estimated the expression of full-length *STMN2* using all RNA-Seq reads covering known *STMN2* exons, whereas the truncated *STMN2* isoform was quantified using high-confidence spliced RNA-Seq reads that unambiguously connected *STMN2* exon 1 with the cryptic exon 2a (exon 1–exon 2a) (Figure 3A). We found that full-length *STMN2* was highly expressed in both cortical and cerebellar tissues (median transcripts per million [TPM] in frontal cortex = 58.0 and in cerebellum = 73.1) compared with expression levels in spinal cord (median TPM in lumbar spinal cord = 6.7), presumably because its expression is restricted to neurons and because of the higher white matter composition in the spinal cord (Supplemental Figure 4).

We then estimated the expression of truncated *STMN2* in the NYGC ALS cohort in disease groups across tissues. At least 2 reads spanning the exon 1–exon 2a junction could be detected in 398 of the 1659 samples in the data set (24.0%) (Supplemental Figure 5). Samples that expressed the splice junction did so at a median read count of 6 (0.13 reads per million) and a maximum read count of 120 (2.5 reads per million) (Supplemental Figure 6). We therefore chose to model truncated *STMN2* expression as a binary outcome, with a detection threshold of 2 junction reads uniquely mapped across the intron boundary.

Truncated *STMN2* was present in disease subtypes known to have TDP-43 pathology, as it was detected in the frontal cortex of 82.4% of patients with FTLT-DTP but was absent in those with FTLT-FUS or FTLT-tau and in controls (Figure 3B). In ALS lumbar spinal cord, we detected truncated *STMN2* in 62.2% of patients with ALS with presumed TDP-43 pathology (ALS-TDP) but not in patients with ALS-SOD1 or in control individuals (Figure 3C).

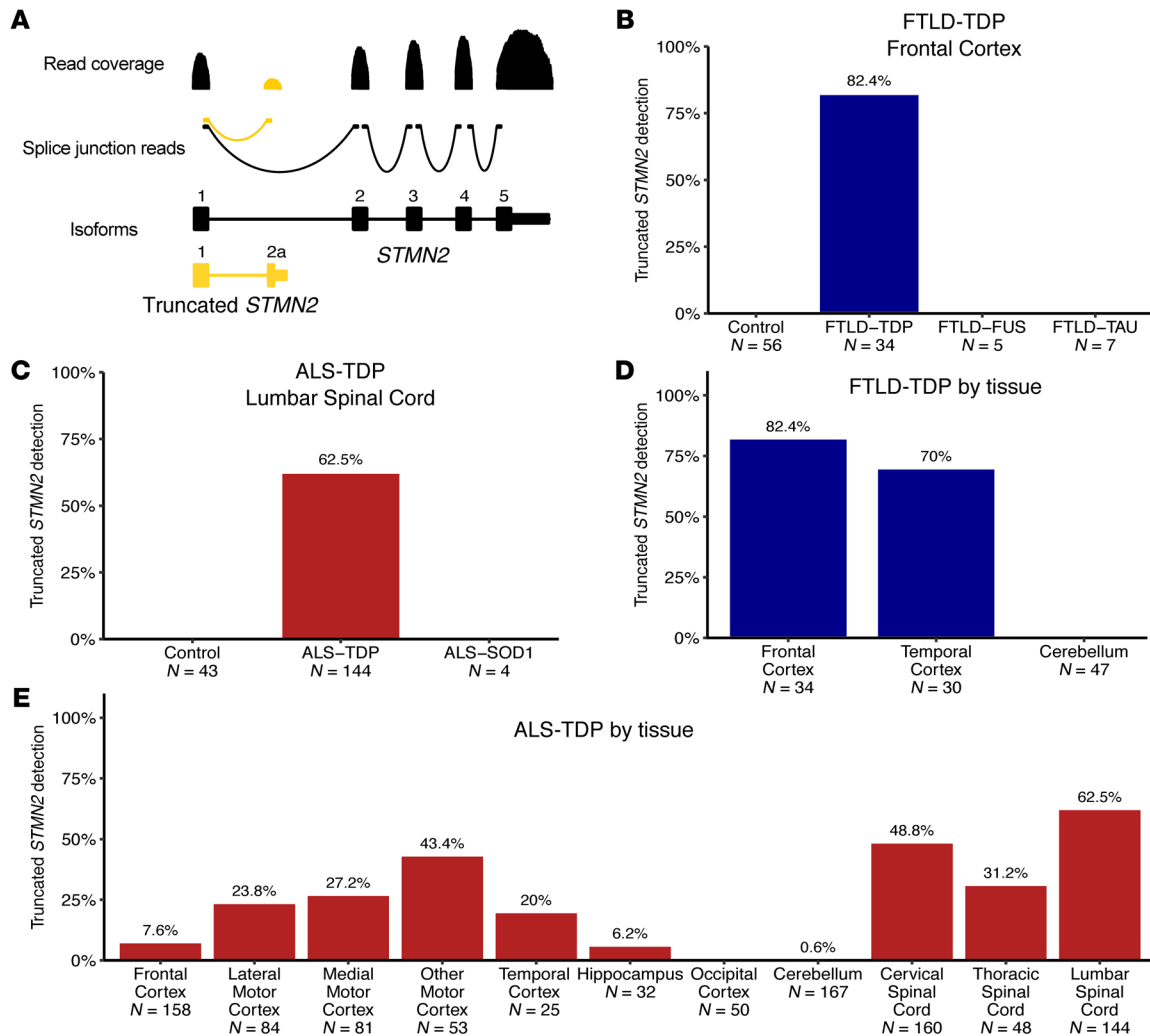
We observed an exquisite specificity of truncated *STMN2* expression in tissues known to have TDP-43 pathology. In patients with FTLT-DTP, for whom frontal cortex, temporal cortex, and cerebellum data sets were available, we detected truncated *STMN2* in frontal and temporal cortices but not in cerebellum (Figure 3D). In ALS-TDP tissues, which also included hippocampus, temporal cortex, motor cortex, occipital cortex, and spinal cord samples, truncated *STMN2* expression was most prevalent in the spinal cord, followed by the motor cortex and frontal cortex, with little to no expression detected in the occipital cortex or cerebellum,



**Figure 2. Truncated *STMN2* RNA is elevated in iPSC-derived neurons with reduced levels of TDP-43.** (A–C) iPSC-derived cortical neurons constitutively expressing CRISPR inactivation machinery (data set labeled “a”) were transduced with a lentivirus expressing a sgRNA against *TARDBP* (TDP-43 KD) or a control sgRNA (Controls) and subjected to analysis by qRT-PCR ( $n = 3$  TDP-43 KD,  $n = 3$  controls) and/or RNA-Seq ( $n = 3$  TDP-43 KD,  $n = 4$  controls). The graphs show a decrease in *TARDBP* transcripts (A), a reduction of the full-length *STMN2* transcript (B), and an increase in the truncated *STMN2* transcript (C) upon TDP-43 depletion. (D) Detection of *STMN2* cryptic exon 2a inclusion expressed as PSI by RNA-Seq in various iPSC-derived neurons: iPSC-derived cortical neurons (data set labeled “a”) described above and performed in this study; iPSC-derived motor neurons treated with *TARDBP* siRNA (TDP-43 KD) or control siRNA (controls) from Klim et al. (data set labeled “b”) ( $n = 6$  TDP-43 KD,  $n = 11$  controls) (7); iPSC-derived motor neurons from patients with *TARDBP* mutations (TDP-43 mut.) or controls from 2 independent groups in Edinburgh (data set labeled “c”) ( $n = 7$  TDP-43 KD,  $n = 4$  controls) and Oxford (data set labeled “d”) ( $n = 10$  TDP-43 KD,  $n = 4$  controls) (72). Data are presented as the mean  $\pm$  SEM and were normalized to the control groups in A and B (set to 100%).  $*P < 0.05$ ,  $***P < 0.005$ , and  $****P < 0.001$ , by Student’s *t* test (A–C) or 1-way ANOVA (D). (E) Representative images of iPSC-derived motor neurons from patients with *TARDBP* mutations and control subjects from a group in Oxford (data set labeled “d”) in Figure 1D) showed similar nuclear-cytoplasmic distribution of TDP-43 (green) on day 30 of differentiation (DAPI in blue, choline acetyltransferase [ChAT] in white). Scale bar: 10  $\mu$ m. All panels of Figure 2E are reshown in Supplemental Figure 3A. Additional data associated with this figure can also be found in Supplemental Figure 3. See also the complete unedited blots in the supplemental material.

respectively (Figure 3E and Supplemental Figure 5). As anticipated, patients with concurrent FTD and ALS (ALS-FTLD) had truncated *STMN2* expression in frontal and temporal cortices, as well as in the motor cortex and spinal cord (Supplemental Figure 7A). In contrast, patients with ALS-AD strongly resembled patients with ALS-TDP in their truncated *STMN2* detection patterns, with most truncated *STMN2* expression found in spinal cord and reduced or no expression in cortical brain regions (Supplemental Figure 7B). Together, these findings demonstrate that truncated *STMN2* RNA is detected in diseases and tissues characterized by TDP-43 pathology, supporting its potential as a marker for TDP-43 dysfunction.

*Truncated STMN2 RNA expression is elevated in FTLD-TDP frontal cortex.* Given that we detected truncated *STMN2* RNA in patients with FTLD-TDP but not in those with FTLD-FUS or FTLD-tau (Figure 3B), this aberrant splicing event may provide a means to identify patients with FTD with marked TDP-43 pathology. To further examine the relationship between TDP-43 and *STMN2* in patients with FTD, we used the NanoString PlexSet platform to systematically quantify truncated and full-length *STMN2* RNA levels in frontal cortex tissue from a cohort of 238 patients with FTD with immunohistologically confirmed TDP-43-immunoreactive inclusions, a cohort of 33 cognitively normal controls who were TDP-43 negative



**Figure 3. Truncated *STMN2* RNA is detected in bulk RNA-Seq from FTLD/ALS tissues with TDP-43 pathology.** (A) The NYGC ALS Consortium RNA-Seq data set was analyzed for the presence of truncated *STMN2* transcripts, which were identified by RNA-Seq reads spanning the exon 1–exon 2a splice junction. (B–E) Bar graphs represent the proportion of individuals (percentage) with at least 2 reads spanning the exon 1–exon 2a junction. The number of individuals for the samples of the indicated tissues/diseases is indicated. (B) Truncated *STMN2* was detected in the frontal cortex of patients with FTLD-TDP but not in control subjects or patients with FTLD-FUS or FTLD-tau. (C) In the lumbar spinal cord of patients with ALS, truncated *STMN2* RNA was only detected in ALS-TDP samples but not in ALS-SOD1 or control samples. (D) Among FTLD-TDP tissues, truncated *STMN2* was only detected in those known to have TDP-43 pathology (frontal and temporal cortex), whereas (E) within ALS-TDP tissues, truncated *STMN2* RNA was seen in motor cortex and spinal cord tissues but not in regions without TDP-43 pathology.

(henceforth referred to as controls), and a cohort of 41 patients with PSP (Table 1). The latter cohort of patients were characterized by tau pathology and neurodegeneration in the frontal cortex.

As mentioned, FTLD-TDP can be further classified on the basis of morphology and distribution of TDP-43 inclusions. As such, our cohort of patients with FTLD-TDP were classified into different TDP-43 pathological subtypes: 117 patients were type A (27 and 34 patients carried a mutation in *C9orf72* or *GRN*, respectively), 66 were type B (21 and 1 carried a mutation in *C9orf72* or *GRN*, respectively), 43 were type C (2 and 1 carried a mutation in *C9orf72* or *GRN*, respectively), and 2 were type D (both patients had *VCP* mutants) (Table 1). In addition to conducting our analyses of all FTLD-TDP cases combined, we also evaluated whether truncated *STMN2* accumulation was preferentially associated with a particular TDP-43 subtype. In the frontal cortex, we found

that truncated *STMN2* RNA levels were significantly elevated in all patients with FTLD-TDP compared with levels in the control subjects without disease or those with PSP disease in both unadjusted analysis ( $P < 0.001$ ) (Figure 4A, Table 2, and Supplemental Table 2) and analysis adjusted for age at death, sex, and RNA integrity number (RIN) ( $P = 0.009$  vs. controls,  $P < 0.001$  vs. patients with PSP) (Figure 4A, Table 2, and Supplemental Table 2). We also noted that the levels of truncated *STMN2* in individuals with PSP were significantly lower than those in control subjects ( $P = 0.028$ , adjusted analyses for age, sex, and RIN), possibly because of the neuronal loss in the patients with PSP.

As in all patients with FTLD-TDP combined, truncated *STMN2* RNA levels in the frontal cortex were increased in FTLD-TDP for each TDP-43 type (A–D) compared with levels in patients with PSP in both unadjusted and adjusted analyses (Table 2 and Sup-

**Table 1. Patient characteristics in the postmortem cohort**

Variable	Controls (n = 33)	PSP (n = 41)	All FTLT-DTP (n = 238)	FTLD-TDP type A (n = 117)	FTLD-TDP type B (n = 66)	FTLD-TDP type C (n = 43)	FTLD-TDP type D (n = 2)
Age at death (yr)	82 (55, 99)	69 (52, 86)	72 (49, 100)	76 (50, 100)	67 (49, 98)	74 (60, 84)	63 (61, 64)
Sex							
Male	20 (60.6%)	13 (51.2%)	130 (54.6%)	63 (53.8%)	37 (56.1%)	23 (53.5%)	1 (50.0%)
Female	13 (39.4%)	20 (48.8%)	108 (45.4%)	54 (46.2%)	29 (43.9%)	20 (46.5%)	1 (50.0%)
Survival after onset (yr)	NA	7.0 (4.0, 14.0)	7.0 (0.6, 25.0)	7.0 (1.0, 25.0)	4.0 (0.6, 20.0)	10.6 (4.0, 25.0)	8.5 (7.0, 10.0)
Age at onset (yr)	NA	61.5 (46.0, 79.5)	64.5 (40.0, 85.8)	67.8 (40.0, 85.8)	61.4 (44.0, 85.7)	62.7 (46.0, 76.0)	54.1 (54.0, 54.2)
RIN	9.6 (7.1, 10.0)	9.5 (7.1, 10.0)	9.5 (7.0, 10.0)	9.4 (7.0, 10.0)	9.6 (7.0, 10.0)	9.5 (7.8, 10.0)	9.9 (9.8, 10.0)

The sample median (minimum, maximum) is given for continuous variables. Information was unavailable regarding TDP-43 subtype (all FTLT-DTP:  $n = 10$ ), age at death (all FTLT-DTP:  $n = 1$ ), p-TDP-43 burden in the frontal cortex (all FTLT-DTP:  $n = 1$ ), survival after onset (PSP:  $n = 4$ ; all FTLT-DTP:  $n = 20$ ), and age at onset (PSP:  $n = 4$ ; all FTLT-DTP:  $n = 20$ ). Among the patients with FTLT-DTP, 50 had *C9orf72* mutations ( $n = 27$  TDP-43 type A,  $n = 21$  TDP-43 type B,  $n = 2$  TDP-43 type C), 36 had *GRN* mutations ( $n = 34$  TDP-43 type A,  $n = 1$  TDP-43 type B,  $n = 1$  TDP-43 type C), and 2 had *VCP* mutations (TDP-43 type D).

plemental Table 2). Compared with controls, truncated *STMN2* levels were significantly increased in patients with FTLT-DTP types A, B, and C, but not in those with type D, in unadjusted analyses (Table 2 and Supplemental Table 2), although the levels only in those with type A FTLT-DTP remained significant in analyses adjusted for age, sex, and RIN (Table 2 and Supplemental Table 2).

We observed that truncated *STMN2* levels in the frontal cortex discriminated between the patients with FTLT-DTP and the PSP control subjects, with an area under the receiver operating characteristic curve (AUC) of 0.76 (95% CI = 0.69–0.84) (Table 2). Frontal cortex truncated *STMN2* also discriminated between FTLT-DTP patients and cognitively normal controls with a more modest AUC of 0.67 (95% CI = 0.57–0.78) (Supplemental Table 2). Furthermore, truncated *STMN2* levels similarly discriminated between controls (cognitively normal or PSP disease controls) and patients with FTLT-DTP of all TDP-43 types, most prominently for types A, C, and D (Table 2 and Supplemental Table 2). In contrast, truncated *STMN2* levels did not effectively distinguish patients with PSP from control subjects (AUC: 0.60, 95% CI = 0.47–0.73).

The increase in truncated *STMN2* levels in FTLT-DTP frontal cortex was paired with a decrease in full-length *STMN2* levels ( $P < 0.001$ ) (Figure 4B and Supplemental Table 3). Of interest, full-length *STMN2* expression was also decreased in patients with PSP compared with expression levels in controls ( $P < 0.001$ ) (Figure 4B and Supplemental Table 3), probably because of neurodegeneration in this region. Full-length *STMN2* levels also distinguished control subjects from patients with FTLT-DTP (AUC: 0.74) and PSP (AUC: 0.83), but were not as effective in distinguishing patients with FTLT-DTP from those with PSP (AUC: 0.60, 95% CI = 0.51–0.68; Supplemental Table 3).

Taken together, our data show that truncated *STMN2* RNA levels were significantly elevated in patients with FTLT-DTP compared with levels in controls and may be used to distinguish FTD patients with TDP-43 pathology from both cognitively normal individuals and disease controls.

*Truncated STMN2 RNA associates with p-TDP-43 burden in the frontal cortex of patients with FTLT-DTP.* That truncated *STMN2* levels were elevated in the frontal cortex of patients with FTLT-DTP but not in those with PSP (Figure 4A) or other forms of FTLT

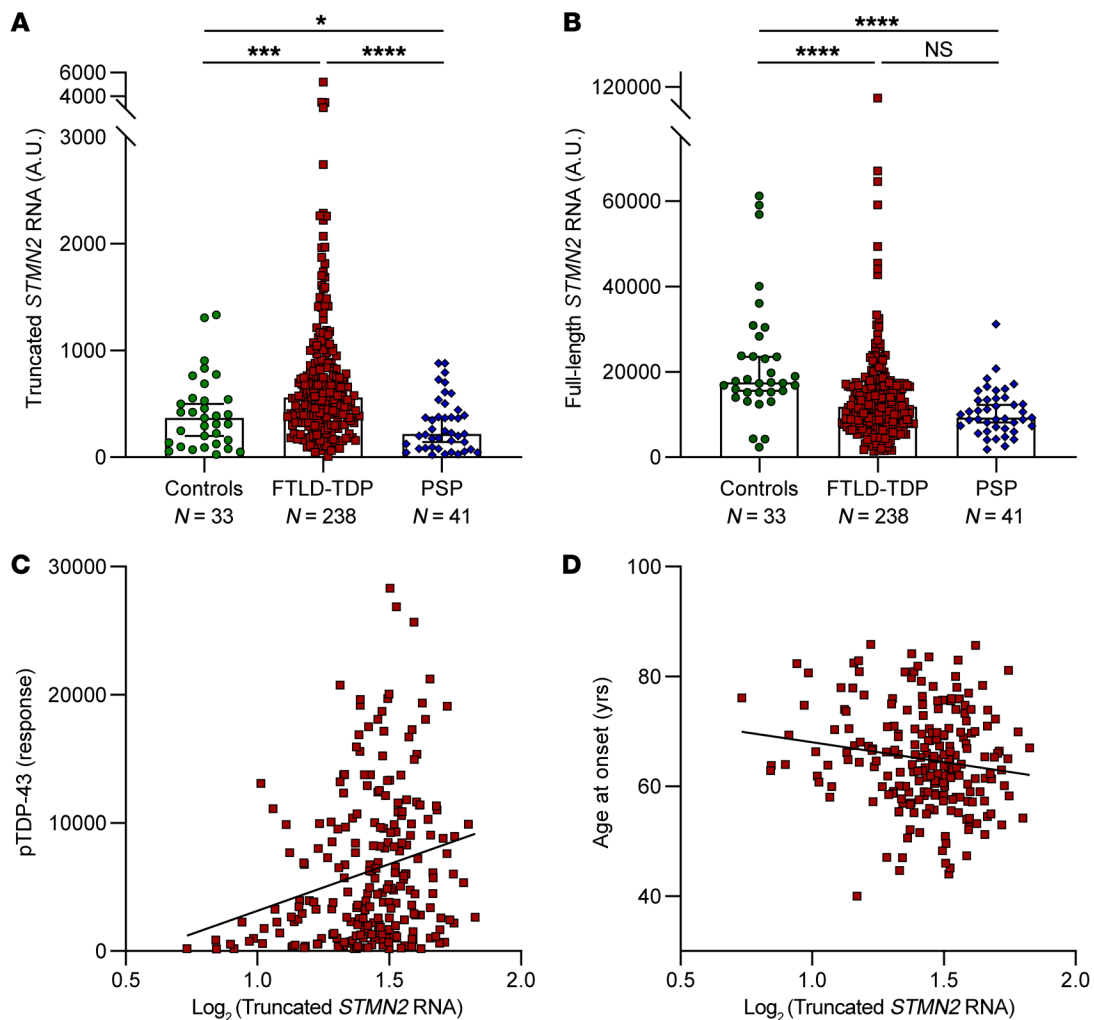
(FTLT-tau, FTLT-FUS) or ALS (ALS-SOD1) (Figure 3) suggests that truncated *STMN2* production is caused by a loss of TDP-43 nuclear function resulting from its mislocalization to the cytoplasm and/or its sequestration into inclusions — hallmark features of FTLT-DTP. To evaluate this possibility further, we examined whether full-length or truncated *STMN2* RNA levels in the frontal cortex correlated with p-TDP-43 accumulation. To this end, we prepared urea-soluble protein lysates from the same tissues used to measure *STMN2* isoforms and quantified p-TDP-43 in these lysates using a previously described immunoassay (32–34). Although the levels of full-length *STMN2* did not associate with p-TDP-43 levels ( $\beta$ : 0.02,  $P = 0.53$ ) (Supplemental Table 4), higher p-TDP-43 levels were associated with higher truncated *STMN2* RNA levels (Figure 4C) in both unadjusted analyses ( $\beta$ : 0.20,  $P < 0.001$ ) (Table 3) and in a multivariable analysis adjusted for age, sex, and TDP-43 subtype ( $\beta$ : 0.19,  $P < 0.001$ ) (Table 3). Taken together, our data suggest that truncated *STMN2* RNA, but not full-length *STMN2*, accumulates in brain regions marked by TDP-43 pathology and associates with p-TDP-43 burden.

*Truncated STMN2 RNA levels associate with an earlier age of FTLT-DTP onset.* We additionally examined whether truncated or full-length *STMN2* levels in the frontal cortex associate with clinical characteristics of individuals with FTLT-DTP. We found a significant inverse correlation between truncated *STMN2* RNA levels and age at disease onset (Figure 4D) in analyses, adjusting for sex, age at death, and TDP-43 type and after correcting for multiple testing ( $P = 0.005$ ) (Table 3). However, survival after onset did not associate with truncated *STMN2* RNA levels, after correcting for age at onset, sex, and TDP-43 subtype ( $P = 0.32$ ) (Table 3). Of interest, we observed a nominal but not statistically significant association with lower truncated *STMN2* levels in males ( $\beta$ : -0.35,  $P = 0.034$  in adjusted analyses for age and TDP-43 type) (Supplemental Table 4). In contrast, full-length *STMN2* levels failed to associate with clinical features of FTLT-DTP, including age of onset and survival (Supplemental Table 4).

## Discussion

FTLT, the neuropathological substrate of FTD, is the second most common cause of early-onset dementia (35). Although no effective treatment is available, several therapeutic approaches





**Figure 4. Truncated *STMN2* is elevated in FTLD-TDP and associates with higher p-TDP-43 burden and earlier age of FTD onset.** (A and B) Truncated and full-length *STMN2* levels were measured in RNA extracted from frontal cortex of control subjects and FTLD-TDP and PSP patients, using the NanoString PlexSet platform. (A) A significant accumulation of truncated *STMN2* RNA was detected in patients with FTLD-TDP but not in controls (see also Supplemental Table 2) or patients with PSP (see also Table 2). (B) Full-length *STMN2* levels were significantly decreased in FTLD-TDP and PSP patients (see also Supplemental Table 3). Data are presented as the median with a 95% CI, and *P* values were determined using linear regression models that were adjusted for age at death, sex, and RIN. \**P* < 0.05, \*\*\**P* < 0.005, \*\*\*\**P* < 0.001. (C and D) In our FTLD-TDP cohort (*n* = 238), significant correlations were observed between truncated *STMN2* RNA levels and (C) a higher burden of p-TDP-43 and (D) an earlier age of disease onset. See Table 3 for the correlation coefficients.

are being investigated to target TDP-43 and tau, the 2 proteins most commonly associated with the major pathological FTL subtypes. Their testing in clinical trials, however, faces a considerable barrier: our inability to determine the underlying protein pathology in living patients with FTD. Although the pathomechanisms driven by TDP-43 and tau differ, the clinical symptoms of FTD patients with TDP-43 or tau pathology overlap. Ultimately, this hinders the selection of patients with sporadic FTD who are suitable for a particular clinical trial. Since there is a clear need for biomarkers to identify FTD patients with TDP-43 pathology, we evaluated whether truncated *STMN2* could serve as a proxy for TDP-43 pathology. Using RNA-Seq and RNA quantification techniques, we show that truncated *STMN2* RNA levels were elevated in human iPSC-derived neurons depleted of TDP-43 and in bulk tissue from FTD and ALS patients with TDP-43 pathology. We did not detect truncated *STMN2* in iPSC-derived

neurons with pathogenic *TARDBP* mutations where no change in TDP-43 levels, localization, or aggregates was detected, supporting the fact that loss of TDP-43 nuclear function, and not splicing changes induced by pathogenic mutations, were driving this event. We were unable to assess postmortem tissue from patients carrying *TARDBP* mutations, which are rare in the FTD and ALS populations, as a group, but given that these patients develop florid TDP-43 pathology, we expected truncated *STMN2* to be present at levels similar to those in patients with sporadic ALS or FTLD-TDP. Additionally, we found that truncated *STMN2* RNA accumulated in the frontal cortex of patients with FTLD-TDP but not in control subjects or patients with PSP and was significantly correlated with p-TDP-43 levels and an earlier age of disease onset. Taken together, our data support the notion that truncated *STMN2* production results from TDP-43 loss of function.

**Table 2. Comparisons of truncated *STMN2* RNA in frontal cortex between patients with FTLD-TDP and patients with PSP.**

Group	n	Median (minimum, maximum) levels of truncated <i>STMN2</i> RNA	Unadjusted analysis		Adjusting for age at death, sex, and RIN		AUC (95% CI) vs. PSP cases
			Regression coefficient (95% CI)	P value	Regression coefficient (95% CI)	P value	
PSP	41	217 (22, 880)	0.00 (reference)	NA	0.00 (reference)	NA	NA
All FTLD-TDP	238	560 (8, 5210)	1.38 (0.96, 1.80)	< 0.001	1.36 (0.95, 1.77)	< 0.001	0.76 (0.69, 0.84)
FTLD-TDP type A	117	595 (8, 3495)	1.47 (0.98, 1.95)	< 0.001	1.37 (0.87, 1.86)	< 0.001	0.78 (0.70, 0.86)
FTLD-TDP type B	66	494 (62, 5210)	1.28 (0.77, 1.79)	< 0.001	1.29 (0.79, 1.78)	< 0.001	0.74 (0.65, 0.84)
FTLD-TDP type C	43	575 (106, 2071)	1.34 (0.79, 1.89)	< 0.001	1.26 (0.73, 1.78)	< 0.001	0.77 (0.66, 0.87)
FTLD-TDP type D	2	1035 (1008, 1062)	2.40 (0.25, 4.55)	0.029	3.07 (1.07, 5.07)	0.004	1.00 (1.00, 1.00)

Regression coefficients, 95% CIs, and *P* values were determined using linear regression models, where the levels of truncated *STMN2* RNA in the frontal cortex were considered on the base 2 logarithmic scale. Regression coefficients were interpreted as the difference in mean levels of truncated *STMN2* RNA in the frontal cortex (on the base 2 logarithmic scale) between the given group of patients with disease and patients with PSP. *P* values of 0.010 or less were considered statistically significant after applying a Bonferroni's correction for multiple testing.

As mentioned above, 2 independent reports recently revealed that TDP-43 regulates *STMN2* expression and that TDP-43 depletion produces a truncated, nonfunctional *STMN2* variant (7, 8). Increased truncated *STMN2* RNA levels were detected in spinal cord tissue of 17 patients with ALS but not in that of 6 controls, and ISH studies found truncated *STMN2* RNA in lower and upper motor neurons in 5 patients with ALS (8). We thus evaluated truncated *STMN2* as a proxy TDP-43 activity marker for FTD using postmortem RNA-Seq data across 11 tissues available from a large cohort of FTD, ALS, and FTLD-ALS patients. Notably, our RNA-Seq analyses detected the *STMN2* cryptic exon that gives rise to truncated *STMN2* only in FTD and ALS patients with TDP-43 pathology. In addition, the presence of abnormal *STMN2* was highly specific to brain regions characterized by TDP-43 pathology.

In addition to our analyses of RNA-Seq data, we directly measured truncated *STMN2* RNA in postmortem frontal cortex tissue from patients with FTLD-TDP, which again revealed an aberrant accumulation of truncated *STMN2* in comparison with controls and individuals with PSP. In fact, we found that truncated *STMN2* RNA levels distinguished between patients with FTLD-TDP and those with PSP, with an area under the ROC curve of 0.76. As anticipated, truncated *STMN2* accumulation in patients with FTLD-TDP was accompanied by a decrease in full-length *STMN2*, as was observed in cell culture models (7, 8). Full-length *STMN2* was also decreased in patients with PSP, which were confirmed to lack TDP-43 pathology. Given that *STMN2* is expressed by neurons and its expression is abundant in the frontal cortex, the decrease in full-length *STMN2* in patients with PSP likely reflects neurodegenerative neuronal loss. Indeed, a recent report found that full-length *STMN2* RNA levels were also decreased in the brains of patients with Parkinson's disease (36), possibly arguing for the use of *STMN2* levels as a neurodegenerative disease biomarker. It is also possible that, aside from TDP-43, various other factors regulate full-length *STMN2* expression. This could explain why full-length *STMN2* levels did not correlate with p-TDP-43 burden or with clinical features.

The accumulation of truncated *STMN2* RNA in the frontal cortex of patients with FTLD-TDP but not of patients with PSP indicates that truncated *STMN2* production is caused by patho-

logical changes in TDP-43. In line with this, we observed a significant association of truncated *STMN2* RNA with p-TDP-43 protein levels ( $P < 0.001$ , Table 3), which remained statistically significant after controlling for multiple variables (age, sex, and TDP-43 subtype;  $P < 0.001$ , Table 3). However, it is possible that p-TDP-43 levels alone may not be a perfect corollary for nuclear TDP-43 depletion. Of interest, the levels of truncated *STMN2* RNA did not differ among different TDP-43 subtypes, suggesting that truncated *STMN2* levels are elevated in all types of TDP-43 pathology.

Importantly, higher truncated *STMN2* RNA levels in the frontal cortex at death were significantly associated with an earlier age at disease onset ( $P = 0.005$ , Table 3), but we found no association with survival after disease onset. Given that truncated *STMN2* RNA serves as a proxy for TDP-43 dysfunction, this earlier age of onset in patients with higher postmortem truncated *STMN2* RNA levels could point to more severe defects in RNA metabolism resulting from aberrant TDP-43 function, including loss of *STMN2* function.

Strengths of our study include the use of samples from a clinically well-characterized, immunohistologically confirmed cohort of 238 patients with FTLD-TDP from which high-integrity RNA was extracted from the frontal cortex, and the use of quantitative measurements of truncated *STMN2* RNA and p-TDP-43 protein. Moreover, we evaluated multiple neuroanatomical regions in bulk tissue RNA-Seq from a large data set that included 1659 FTD and ALS patients with and without TDP-43 pathology. Further, our study also used 3 different approaches (RNA-Seq, NanoString, and qRT-PCR, the latter of which was also performed to validate truncated *STMN2* in patients with FTLD-TDP, as shown in Supplemental Figure 9) that have different sensitivity and specificity linked to the methodology but that all detect significant differences in truncated *STMN2*, highlighting the reproducibility of the finding. Our study also has some limitations. While we specifically designed our study to evaluate truncated *STMN2* as a marker of TDP-43 pathology — information important to establish its value as a candidate biomarker — we did not measure the translation product from this truncated variant because of a lack of commercially available antibodies. At present, it is unclear whether

**Table 3. Associations of truncated *STMN2* with p-TDP-43 and clinical characteristics in FTLD-TDP frontal cortex**

Variable	Association with frontal cortex truncated <i>STMN2</i> RNA				
	Unadjusted analysis		Multivariable analysis		Multivariable model adjustments
	Regression coefficient (95% CI)	P value	Regression coefficient (95% CI)	P value	
p-TDP-43 (doubling)	0.20 (0.12, 0.28)	< 0.001	0.19 (0.09, 0.28)	< 0.001	Age at death, sex, TDP-43 subtype
TDP-43 subtype	Overall test of difference: <i>P</i> = 0.48		Overall test of difference: <i>P</i> = 0.33		Age at death and sex
FTLD-TDP type A	0.00 (reference)	NA	0.00 (reference)	NA	
FTLD-TDP type B	-0.19 (-0.57, 0.19)	0.33	-0.31 (-0.72, 0.10)	0.14	
FTLD-TDP type C	-0.13 (-0.56, 0.31)	0.57	-0.16 (-0.60, 0.27)	0.46	
FTLD-TDP type D	0.93 (-0.91, 2.77)	0.32	0.68 (-1.17, 2.53)	0.47	
Age at onset (10-yr increase)	-0.23 (-0.41, -0.04)	0.015	-0.29 (-0.49, -0.09)	0.005	Sex and TDP-43 subtype
Survival after onset (5-yr increase)	0.12 (-0.06, 0.30)	0.19	0.10 (-0.09, 0.29)	0.32	Age at onset, sex, and TDP-43 subtype
Sex					
Male	-0.37 (-0.68, -0.06)	0.019	-0.35 (-0.67, -0.03)	0.034	Age at death and TDP-43 subtype
Female	0.00 (reference)	NA	0.00 (reference)	NA	

Regression coefficients, 95% CIs, and *P* values were determined using linear regression models, where frontal cortex truncated *STMN2* RNA was considered on the base 2 logarithmic scale. Regression coefficients were interpreted as the change in mean frontal cortex truncated *STMN2* RNA (on the base 2 logarithmic scale) corresponding to the presence of the indicated characteristic (categorical variables) or the increase shown in parentheses (continuous variables). *P* values of 0.010 or less were considered statistically significant after applying a Bonferroni's correction for multiple testing.

the predicted peptide of only 16 amino acids is translated from truncated *STMN2* RNA. Nevertheless, our findings indicate that the development of novel tools (antibodies, immunoassays) for the detection of truncated *STMN2* protein and additional studies to evaluate the role of truncated *STMN2* RNA and protein in disease pathogenesis and as biomarkers are well merited. Critical next steps to hasten the translation of this finding to the clinic include determining whether truncated *STMN2* RNA or protein is detected in the biofluids of patients with FTD (e.g., spinal fluid and blood) and, if so, whether they discriminate between FTD subtypes (i.e., with or without TDP-43). Although expression of truncated *STMN2* in affected postmortem tissues can distinguish FTLT-TDP from PSP cases, which has important implications, as it supports truncated *STMN2* as a viable proxy for the presence of TDP-43 pathology, the ability to discriminate between FTLT-TDP and PSP patients was only modest (AUC: 0.76). RNA degradation invariably occurs in postmortem brain tissues, possibly contributing to this modest AUC and potentially pointing toward the possibility of detecting truncated *STMN2* in CSF or blood, which are typically frozen immediately after collection, to discriminate between patient groups. Nonetheless, sensitive methods will be needed to detect truncated *STMN2* RNA and/or protein in human biofluids. Additionally, determining at what point during the course of disease truncated *STMN2* levels begin to be upregulated will have to be evaluated and will probably require longitudinal collections of CSF and blood first from patients with ALS, who are almost certain to develop TDP-43 pathology, and then from individuals with FTD or FTD/ALS harboring genetic variants associated with TDP-43 dysfunction.

If, as anticipated, truncated *STMN2* RNA or protein is detected in the biofluids of FTD patients with TDP-43 pathology, it may have the potential to similarly identify those patients with AD who have TDP-43 pathology. Indeed, up to 67% of patients with

AD develop TDP-43 inclusions, and, compared with AD patients without TDP-43 pathology, they are cognitively and functionally worse, suggesting that abnormal TDP-43 causes a modified AD phenotype (37–46). Our work also highlights how future means to detect truncated *STMN2* RNA or protein in biofluids may provide important information on disease activity and therefore be useful in assessing drug efficacy in clinical trials. Thus, these endeavors are expected to increase the likelihood of success in developing effective therapeutics and improving the care of patients with TDP-43 proteinopathies.

## Methods

*RNA extraction, complementary DNA synthesis, and qRT-PCR in iPSC-derived neurons.* iPSC-derived neurons were harvested, and cell pellets were lysed in TRI reagent (Zymo Research). Then, total RNA was extracted following the manufacturer's instructions using the Direct-zol RNA Miniprep Kit (Zymo Research), and 850 ng RNA was used for RT-PCR. RT-PCR of RNA samples was performed using the High Capacity cDNA Transcription Kit with random primers (Applied Biosystems), and qRT-PCR was performed using SYBR GreenER qPCR SuperMix (Invitrogen, Thermo Fisher Scientific). Samples were run in triplicate, and qRT-PCRs were run on a Prism 7900HT Fast Real-Time system (Applied Biosystems). Relative quantification was determined using the  $\Delta\Delta C_t$  method and normalized to the endogenous controls *RPLP0* and *GAPDH*. The following primers and their sequences were used: *STMN2* forward: 5'-AGCTGTCCATGCTGTCACCTG-3', *STMN2* reverse: 5'-GGTGGCTTCAAGATCAGCTC-3'; truncated *STMN2* forward: 5'-GGACTCGGCAGAACACCTTC-3', truncated *STMN2* reverse: 5'-GCAGGCTGTCTGTCTCTCTC-3'; *TARDBP* forward: 5'-AATTCTGCATGCCCCAGA-3', *TARDBP* reverse: 5'-GAAGCATCTGTCTCATCCATTTT-3'; *GAPDH* forward: 5'-GTTTCGACAGTCAGCCGCATC-3', *GAPDH* reverse: 5'-GGAATTTGCCATGGGTGGA-3'; and *RPLP0* forward: 5'-TCTACAACCCTGAAGTGCTTGAT-3', *RPLP0* reverse: 5'-CAATCTGCAGACAGACTGG-3'.

**RNA-Seq analyses from iPSC-derived neurons.** Data from Klim et al. (7) were downloaded from the NCBI's Gene Expression Omnibus (GEO) database (GEO GSE121569). All iPSC cell lines were analyzed with the same analysis pipeline. Raw FASTQ files were trimmed using fastp (47) and aligned to the hg38 build (GRCh38.primary\_assembly) of the human reference genome using STAR (2.7.2a) (48). Gene abundance for *TARDBP* and *STMN2* was calculated with featureCounts (49) using gene models from GENCODE GTF, version 31, and converted to reads per kilobase per million (RPKM) using edgeR (50). The relative abundance of truncated *STMN2* as a proportion of total *STMN2* was expressed in terms of percent spliced in (PSI) (51). This was calculated by parsing the splice junction tables from STAR and taken as the percentage of all uniquely mapped *STMN2* junction reads from exon 1 to exon 2a over the sum of all junction reads coming from exon 1.

**NYGC ALS Consortium cohort.** Patients with FTD were classified according to a pathologist's diagnosis of FTD with TDP-43 inclusions (FTLD-TDP), tau inclusions (FTLD-tau), or FUS inclusions (FTLD-FUS). ALS samples were divided into the following subcategories using the available Consortium metadata: ALS with or without reported SOD1 mutations (ALS-TDP and ALS-SOD1); ALS with frontotemporal dementia (ALS-FTLD); and ALS with AD (ALS-AD). All non-SOD1 ALS samples were grouped as "ALS-TDP" in this work for simplicity, although reporting of postmortem TDP-43 inclusions was not systematic and therefore not integrated into the metadata. Confirmed TDP-43 pathology postmortem was reported for all FTLD-TDP samples.

**NYGC ALS Consortium RNA-Seq library preparation from bulk tissue RNA.** The Consortium's sample processing has been, in part, previously described (52). In brief, RNA was extracted from flash-frozen postmortem tissue using TRIzol (Thermo Fisher Scientific) chloroform, followed by column purification (RNeasy Minikit, QIAGEN). RIN was assessed on a Bioanalyzer (Agilent Technologies). RNA-Seq libraries were prepared from 500 ng total RNA using the KAPA Stranded RNA-Seq Kit with RiboErase (KAPA Biosystems) for rRNA depletion and Illumina-compatible indexes (NEXTflex RNA-Seq Barcodes, NOVA-512915, PerkinElmer, and IDT for Illumina TruSeq UD Indexes, 20022370). Pooled libraries (average insert size: 375 bp) passing the quality criteria were sequenced either on an Illumina HiSeq 2500 (125 bp paired end) or an Illumina NovaSeq (100 bp paired end). Samples were subjected to extensive sequencing and RNA-Seq quality control metrics at the NYGC that are described below. Notably, a set of more than 250 markers was used to confirm tissue, neuroanatomical regions, and sex in the RNA-Seq data. Samples passing these metrics are available for distribution. The samples had a median sequencing depth of 42 million read pairs, with a range between 16 and 167 million read pairs.

**Analyses of bulk tissue RNA-Seq data.** Samples were uniformly processed using RAPiD-nf, an efficient RNA-Seq processing pipeline implemented in the NextFlow framework (53). Following adapter trimming with Trimmomatic (version 0.36) (54), all samples were aligned to the hg38 build (GRCh38.primary\_assembly) of the human reference genome using STAR (2.7.2a) (48), with indexes created from GENCODE, version 30 (55). Gene expression was quantified using RSEM (1.3.1) (56). Quality control was performed using SAMtools (57) and Picard Tools (58), and the results were collated using MultiQC (59). Exploratory plots were created in R, version 3.6.0 (60), using dplyr, tidyr, stringr, ggplot2, patchwork, and ggpubr packages (61–63).

RNA-Seq samples were subjected to quality control modeled on the criteria of the Genotype Tissue Expression Consortium (64). Any sample failing 1 of the following sequencing metric thresholds was removed: a unique alignment rate of less than 90%, ribosomal bases of greater than 10%, a mismatch rate of greater than 1%, a duplication rate of greater than 0.5%, intergenic bases of less than 10.5%, and ribosomal bases of greater than 0.1%. Sex was verified using XIST and UTY expression. Tissue identity was confirmed using the expression of the cerebellar marker *CBNLI* and the cortical marker *NRGN*. Spinal cord samples were confirmed by expression of the oligodendrocyte marker *MBOP*. Following quality control, 1905 samples of 1924 were retained for analysis. When comparing subgroups and tissues, only tissues with at least 50 samples and disease subgroups with at least 4 individuals were kept for analysis. This resulted in 1659 samples from 11 tissues and 439 donors.

Expression of the full-length *STMN2* isoform was estimated using RSEM, summing the estimates of the 3 annotated isoforms in GENCODE, version 30, into a gene-level estimate of expression normalized in terms of TPM. Given the large size of the NYGC ALS cohort, RNA-Seq data were produced on 2 different sequencing platforms. Exploratory analyses showed a substantial batch effect in gene expression due to the 2 sequencing platforms. We therefore split the data set by sequencing platform, along with tissue and disease status. A comparison of ALS-TDP and non-neurological controls revealed that *STMN2* had an inconsistent pattern of differential expression, with no tissue showing a marked difference in either platform (Supplemental Figure 4A). In addition, RNA degradation is known to affect the estimation of transcript abundance (65, 66), and we reasoned that systematic differences in RNA quality among samples sequenced on different platforms and between disease and control groups or tissues could confound our estimation of *STMN2* expression. We compared the RIN values for each sample between the ALS-TDP and control groups and found significant differences in RIN values in some comparisons (Supplemental Figure 4B). The small number of controls available limited further comparisons in some tissues. We then correlated the RIN with *STMN2* expression and found a positive correlation in multiple tissues and on both sequencing platforms (Supplemental Figure 4C).

**Quantifying *STMN2* splicing from RNA-Seq data.** For downstream analysis of *STMN2* splice junction reads, a bioinformatics pipeline was created using Snakemake (67, 68). Uniquely mapped reads within the *STMN2* locus were extracted from each sample using SAMtools. Any read marked as a PCR duplicate by Picard Tools was discarded. Splice junction reads were then extracted with RegTools (69) using a minimum of 8 bp as an anchor on each side of the junction and a maximum intron size of 500 kb. Junctions from each sample were then clustered together using LeafCutter (70) with relaxed junction filtering (minimum total reads per junction = 30, minimum fraction of total cluster reads = 0.0001). This produced a matrix of junction counts across all samples. Using coordinates from the Klim (7) and Melamed (8) publications, the TDP-43-associated junction of the novel truncated *STMN2* isoform was extracted from the matrix. A detection threshold of 2 or more uniquely mapped reads was applied. We found no association between truncated *STMN2* detection and RIN, sequencing platform, or total library size (Supplemental Figure 8, A–C). The relationship between expression levels of full-length and truncated *STMN2* was complex, with no overall detection of association across tissues (Supplemental Figure 8D).

*RNA extraction and NanoString analyses in postmortem tissues.* Using the RNeasy Plus Mini Kit (QIAGEN), RNA was extracted from frozen human postmortem frontal cortex tissues (33). RNA with a RIN of 7.0 or higher, as determined using an Agilent 2100 bioanalyzer (Agilent Technologies), was used to measure transcript levels using the NanoString PlexSet platform (250 ng human brain RNA, according to the manufacturer's instructions). Transcript level abundance was determined using nSolver 4.0 software (NanoString Technologies) and normalized to *HTRPI*, a housekeeping gene that was unaltered across disease subtypes in our data set and previously used to normalize human brain transcripts (71). The sequences of the NanoString probes used in this study are included in Supplemental Table 5.

*p-TDP-43 immunoassay.* The p-TDP-43 immunoassay was performed as previously described (33). Briefly, approximately 50 mg postmortem tissue from the frontal cortex of patients or unaffected individuals were homogenized in cold RIPA buffer (25 mM Tris-HCl, pH 7.6, 150 mM NaCl, 1% sodium deoxycholate, 1% Nonidet P-40, 0.1% sodium dodecyl sulfate, and protease and phosphatase inhibitors) and sonicated on ice. Homogenates were centrifuged at 100,000g for 30 minutes at 4°C. The supernatant was collected, and the pellet was resuspended in RIPA buffer, sonicated and centrifuged again to prevent carryover of soluble material. The RIPA-insoluble pellet was extracted using 7 M urea buffer and then sonicated and centrifuged at 100,000g for 30 minutes at 22°C. The protein concentration of the urea-soluble supernatant was determined by Bradford assay. p-TDP-43 levels were determined using a sandwich Meso Scale Discovery (MSD) immunoassay. The capture antibody was a mouse monoclonal antibody that detects TDP-43 phosphorylated at serines 409/410 (1:500, no. CAC-TIP-PTD-M01, Cosmo Bio USA), and the detection antibody was a sulfo-tagged rabbit polyclonal C-terminal TDP-43 antibody (2 µg/mL, 12892-1-AP, Proteintech). Urea-soluble fractions were diluted in TBS to 35 µg protein per well and tested in duplicate wells. Response values corresponding to the intensity of emitted light upon electrochemical stimulation of the assay plate using the MSD QUICKPLEX SQ120 were acquired.

*Statistics.* Comparisons of truncated *STMN2* RNA in the frontal cortex between the separate non-FTLD-TDP groups of controls and patients with PSP and various groups of patients with FTLD-TDP (all FTLD-TDP patients as well as 4 different subgroups according to the TDP-43 subtype) were made using single-variable (i.e., unadjusted) and multivariable linear regression models. Multivariable models were adjusted for age at death, sex, and RIN, and truncated *STMN2* RNA in the frontal cortex was examined on the base 2 logarithmic scale, owing to its skewed distribution. Regression coefficients (referred to as  $\beta$ ) and 95% CIs were estimated and interpreted as the difference in mean truncated *STMN2* RNA levels in the frontal cortex (on the base 2 logarithmic scale) between the given group of FTLD-TDP patients and the given reference group (cognitively normal controls or patients with PSP). Additionally, to further assess the ability of truncated *STMN2* RNA in the frontal cortex to discriminate between FTLD-TDP and non-FTLD-TDP individuals (again, separately for cognitively normal controls and patients with PSP), we estimated the area under the ROC curve (AUC); an AUC of 0.5 corresponded to the predictive ability equal to that of chance, whereas an AUC of 1.0 indicated perfect predictive ability. Bonferroni's correction was applied to adjust for the 5 different statistical tests that were

performed for the separate FTLD-TDP versus control and FTLD-TDP versus PSP analyses, after which *P* values of less than 0.01 were considered statistically significant.

In a secondary analysis, we compared truncated *STMN2* RNA levels between controls and patients with PSP and made pairwise comparisons of full-length *STMN2* RNA and p-TDP-43 (both on the base 2 logarithmic scale) between controls, patients with PSP, and patients with FTLD-TDP using the aforementioned linear regression analyses.

In the subgroup of 238 patients with disease, associations of 5 different variables (p-TDP-43 levels, TDP-43 subtype, age at onset, survival after onset, and sex) with both truncated and full-length *STMN2* RNA levels were evaluated using single-variable and multivariable linear regression models, where truncated and full-length *STMN2* RNA levels in the frontal cortex were assessed on the base 2 logarithmic scale, and multivariable models were adjusted for predefined, potentially confounding variables. The multivariable model assessing the p-TDP-43 response in frontal cortex was adjusted for age at death, sex, and TDP-43 subtype. The multivariable model assessing TDP-43 subtype was adjusted for sex and age at death. The multivariable model assessing age at onset was adjusted for sex and TDP-43 subtype. The multivariable model assessing survival after onset was adjusted for age at onset, sex, and TDP-43 subtype. Finally, the multivariable model assessing sex was adjusted for age at death, TDP-43 subtype, and genotype. Regression coefficients and 95% CIs were estimated and are interpreted as the change in the mean level of truncated or full-length *STMN2* RNA in the frontal cortex (on the base 2 logarithmic scale) corresponding to the presence of the given characteristic (categorical variables) or a specified increase (continuous variables). The p-TDP-43 response in the frontal cortex was examined on the base 2 logarithmic scale in linear regression analysis because of its skewed distribution. In order to adjust for the 5 different variables that were assessed for association with the separate truncated and full-length *STMN2* RNA outcomes, we used Bonferroni's correction for multiple testing, after which *P* values of less than 0.01 were considered statistically significant. All statistical tests were 2 sided. All statistical analyses described above were performed using R Statistical Software, version 3.6.2 (R Foundation for Statistical Computing).

Statistical analyses for iPSC studies and qRT-PCR of RNA extracted from human tissues were performed using GraphPad Prism 8 (GraphPad Software), in which 1-way ANOVA with Bonferroni's post hoc test was performed for multiple group comparisons, and an unpaired Student *t* test was performed when only 2 groups were compared. The statistical test and number of independent experiments used for each analysis are indicated in the figure legends.

*Study approval.* The NYGC ALS Consortium samples presented in this work were acquired through various IRB protocols from member sites and the Target ALS postmortem tissue core and transferred to the NYGC in accordance with all applicable foreign, domestic, federal, state, and local laws and regulations for processing, sequencing, and analyses.

Postmortem brain tissues from patients with FTLD-TDP or PSP and from cognitively normal individuals were obtained from the Mayo Clinic Florida Brain Bank. Diagnosis was independently ascertained by trained neurologists and neuropathologists upon neurological and pathological examinations, respectively. Written informed consent was given by all participants or authorized family members, and all protocols were approved by the IRB and ethics committee of the Mayo Clinic. A summary of the patients' characteristics is provided in Table 1.

Sample size was determined by tissue availability and the quality of RNA obtained from such samples, as described below.

**Data and code availability.** RNA-Seq data generated through the NYGC ALS Consortium in this study can be accessed via the NCBI's GEO database (GEO GSE137810, GSE124439, GSE116622, and GSE153960). All RNA-Seq data generated by the NYGC ALS Consortium are made immediately available to all members of the Consortium and with other consortia with whom we have a reciprocal sharing arrangement. To request immediate access to new and ongoing data generated by the NYGC ALS Consortium and for samples provided through the Target ALS Postmortem Core, complete a genetic data request form at ALSData@nygenome.org.

## Author contributions

MP, JH, and SP share co-first authorship. The order of their names was established on the basis of their individual contributions and the importance of those contributions to this study. MP led the human brain RNA and protein studies that provided the basis of the manuscript. JH led the RNA-Seq data analyses that complemented the RNA/protein studies and increased the impact of this manuscript. SP led iPSC studies and was closely involved in the RNA/protein studies as well. MP, JH, SP, TFG, PF, and LP designed the study. MP, SP, ALB, CC, YS, MY, LMD, YC, KJW, CB, AC, KT, BTS, KB, SC, and EMCF generated the iPSC lines and processed iPSC and human tissue samples for qRT-PCR and RNA-Seq. SEH and MW contributed to the experimental design for CRISPRi knockdown of *TARDBP* in iPSC-derived neurons. In the NYGC ALS Consortium, members contributed postmortem samples and clinical information that were curated deidentified by DK. and processed for RNA-Seq by IH, DC, and NP. NP and SF oversee Consortium resources and data distribution. DF and HP designed the methodology, reviewed sample and data quality, and coordinated NYGC ALS Consortium postmortem RNA research activity. JH, YCW, PS, ALB, MS, and TR performed RNA-Seq data analyses. MP, JMK, JS, EAT performed NanoString assays and analyzed the data. SP and TFG performed p-TDP-43 immunoassays. MP, MGH, MRS, ALB, and MS performed statistical analyses. MP, JH, SP, ALB, TFG, PF, and LP wrote the manuscript. CFDC, MD, SK, JN, TL, the NYGC ALS Consortium, BO, MVB, RR, NRGR, BFB,

DSK, RCP, KAJ, and DWD provided human tissue samples as well as pathological, genetic, and clinical information. All authors read and approved the manuscript.

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