A missense mutation in the CSTF2 gene that impairs the function of the RNA recognition motif and causes defects in 3’ end processing is associated with intellectual disability in humans

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ABSTRACT

CSTF2 encodes an RNA-binding protein that is essential for mRNA cleavage and polyadenylation (C/P). No disease-associated mutations have been described for this gene. Here, we report a mutation in the RNA recognition motif (RRM) of CSTF2 that changes an aspartic acid at position 50 to alanine (p.D50A), resulting in intellectual disability in male patients. In mice, this mutation was sufficient to alter polyadenylation sites in over 1300 genes critical for brain development. Using a reporter gene assay, we demonstrated that C/P efficiency of CSTF2D50A was lower than wild type. To account for this, we determined that p.D50A changed locations of amino acid side chains altering RNA binding sites in the RRM. The changes modified the electrostatic potential of the RRM leading to a greater affinity for RNA. These results highlight the significance of 3’ end mRNA processing in expression of genes important for brain plasticity and neuronal development.

INTRODUCTION

Learning, memory, and intelligence require synaptic plasticity involving persistent changes in neural gene expression. Abnormalities in brain development can result in neurodevelopmental disorders that impact intellectual functioning, behavioral deficits involving adaptive behaviors and autism spectrum disorders. Most neuronal developmental changes are accomplished by post-transcriptional processes that alter the regulation and protein coding capacity of specific mRNAs (1,2). Similarly, neurodegenerative conditions are associated with altered mRNA metabolism. Frequently, mRNAs in the brain have extremely long 3’ untranslated regions (UTRs), and altered 3’ UTRs in MECP2 (Rett syndrome), APP (Alzheimer disease) and HTT (Huntington disease) are associated with improper metabolism of each of these mRNAs leading to disease states. Though rarer, monogenic forms of intellectual deficiencies offer specific insight into neuronal plasticity and development. The most common monogenic forms of intellectual deficiency are X-linked (XLID (3,4)). For example, Fragile X Syndrome is caused by expansion of CGG repeats in the 5’ UTR of FMR1, resulting in loss-of-function of FMRP, an RNA-binding protein that promotes transport of mRNAs to dendrites for protein synthesis at synaptic sites (5,6). Similarly, mutations in mRNA processing genes encoding decapping enzymes (7,8), the polyglutamine binding protein 1 (PQBP1 (9)), spliceosomal proteins (10), hnRNA-binding proteins (11), mRNA surveillance proteins (12) and cleavage and polyadenylation factors (13,14) cause intellectual disabilities. These disorders are associated with changes in the mRNA processing landscape, especially 3’ end cleavage and polyadenylation (C/P), highlighting the importance of RNA processing in controlling neuronal function (15–18).

More than eighty different proteins are involved in mRNA C/P (19). Two core C/P factors are the cleavage and polyadenylation specificity factor (CPSF) and the cleavage stimulation factor (CstF). CPSF has six subunits...
that recognize the polyadenylation signal (AAUAAA and closely related sequences), cleaves the nascent pre-mRNA, then recruits the poly(A) polymerase, which adds up to 250 non-template adenosines to the upstream product of the cleavage reaction (20,21). CstF is the regulatory factor in C/P, consisting of three subunits, CstF-50 (gene symbol CSTF1), CstF-64 (CSTF2) and CstF-77 (CSTF3). CSTF2, the RNA-binding component of CstF, binds to U- or GU-rich sequences downstream of the cleavage site through its RNA recognition motif (RRM (22)). As such, CSTF2 regulates gene expression in immune cells (23–25), spermatogenesis (26–29), and embryonic stem cell development (30,31). Furthermore, the CSTF2 paralog, Cstf2t, has been shown to affect learning and memory in mice (32).

Here, we describe members of a family in whom a single nucleotide mutation in the RRM of CSTF2 changes an aspartic acid at amino acid 50 to an alanine (D50A). The probands presented with non-syndromic intellectual disability. The mutation, which is X-linked, co-segregated with the D50A mutation containing the D50A mutation /CSTF2D50A/ reduced C/P by 15%. However, the CSTF2D50A RRM showed an almost 2-fold increase in its affinity for RNA. Solution state nuclear magnetic resonance (NMR) studies showed that the overall backbone structure of the CSTF2D50A RRM was similar to wild type. However, repositioning of side chains in RNA binding sites and the α4-helix resulted in changes in the electrostatic potential and fast timescale protein dynamics of the RNA-bound state. Together, these changes led to an increased affinity of the CSTF2D50A RRM for RNA due to a faster $k_{on}$ rate. Differential gene expression analysis of RNA isolated from brains of male mice harboring the D50A mutation /Cstf2D50A/ identified fourteen genes important for synapse formation and neuronal development. Genome-wide 3′ end sequencing of polyadenylated mRNAs identified >1300 genes with altered C/P sites, leading to alternative polyadenylation of genes involved in neurogenesis, neuronal differentiation and development, and neuronal projection development. Our results indicate that in affected individuals, different affinity and rate of binding to the nascent mRNAs of the mutant CSTF2 RRM could misregulate expression of genes critical for brain plasticity and development.

**MATERIALS AND METHODS**

**Human subjects**

All cases had a normal karyotype, were negative for FMR1 repeat expansion, and large insertions or deletions were excluded using array Comparative Genomic Hybridization (CGH). The study was approved by all institutional review boards of the participating institutions collecting the samples, and written informed consent was obtained from all participants or their legal guardians.

**DNA isolation and X-chromosome exome sequencing and segregation analysis**

DNAs from the family members were isolated from peripheral blood using standard techniques. X-chromosome exome enrichment using DNA from the index patient, sequencing and analysis was performed as previously described (4). Segregation analysis of variants of uncertain clinical significance was performed by PCR using gene-specific primers flanking the respective variant identified followed by Sanger sequencing.

**Animal use and generation of Cstf2D50A mice**

All animal treatments and tissues obtained in the study were performed according to protocols approved by the Institutional Animal Care and Use Committee at the Texas Tech University Health Sciences Center in accordance with the National Institutes of Health animal welfare guidelines. TTEHSC’s vivarium is AAALAC-certified and has a 12-/12-h light/dark cycle with temperature and relative humidity of 20–22°C and 30–50%, respectively.

B6;Cstf2D50A/Cmca-D50A founder mice (herein Cstf2D50A; hemizygous males are designated Cstf2D50A/Y) were generated by Cyagen US Inc. (Santa Clara, CA, USA). To create C57BL/6 mice with a point mutation (D50A) in the Cstf2 locus, exon 3 in the mouse Cstf2 gene (GenBank accession number: NM_133196.6; Ensembl: ENSMUSG00000031256) located on mouse chromosome X was selected as the target site. The D50A (GAT to GCT, see Figure 1 and Supplementary Figure S1) mutation site in the donor oligo was introduced into exon 3 by homology-directed repair. A gRNA targeting vector and donor oligo (with targeting sequence, flanked by 120 bp homologous sequences combined on both sides) was designed. Cas9 mRNA, sgRNA and donor oligo were co-injected into zygotes for knock-in mouse production. The pups were genotyped by PCR followed by sequence analysis. Positive founders were bred to the next generation (F1) and subsequently genotyped by PCR and DNA sequencing analysis. Mutants were maintained as a congenic strain by backcrossing four generations to C57BL/6NCrl (Charles River) and subsequently breeding exclusively within the colony. At the time of this study, mice were bred to ~10 generations.

**Genotyping of Cstf2D50A mice by PCR and restriction enzyme digestion**

Genomic DNA was extracted from tail snips of Cstf2D50A mice by proteinase K digestion followed by isopropanol precipitation (26). PCR were performed using specific primers surrounding the D50A mutation site. The presence of the mutation converted a CGATAGG sequences to CGCATGG, thus introducing a BfaI restriction site (underlined). Digestion of the PCR products with BfaI revealed the presence of the mutation (Supplementary Figure S1, PCR-RFLP). Forward primer: 5′-GAAAGCAGTCAGAGTGGGCT-3′; reverse primer: 5′-CTGCGAAGGAAAATGAGGGT-3′.

**Cell culture, transfection, stem-loop assay for polyadenylation, immunohistochemistry, tissue protein isolation and western blots**

Culturing of HeLa cells was performed as described (22). Transfection was carried out in a 24-well plates using Lipofectamine LTX (ThermoFisher Scientific) and 250 ng of a
Figure 1. Family P167 carries a mutation in the X-linked CSTF2 gene that affects intelligence in males. (A) Pedigree of family 167 showing the index proband (III:1) who was 6 years old at the time of the study, his affected brother (III:2), and two affected uncles (II:1 and II:2). Females (I:2 and II:4) were carriers for the trait, consistent with an X-linked recessive trait. Individuals labeled Mut (II:1, II:2, III:1, III:2) or Mut/WT (II:4) were tested for co-segregation of the mutation with the clinical phenotype by PCR and Sanger sequencing of the specific products. (B) The missense mutation in exon 3 of CSTF2 substitutes an alanine codon for an aspartic acid at position 50 (p.D50A) within the RNA recognition motif (RRM). (C) CLUSTAL alignment of CSTF2 orthologs in twenty-eight species indicate that the aspartic acid (amino acid D) at position 50 (red arrow) is highly conserved. Amino acids comprising site-I (Y25 and R85) and site-II (N19 and N91) are indicated in green and blue, respectively. An * (asterisk) indicates amino acids that are conserved; a : (colon) indicates amino acids with a score greater than 0.5 in the Clustal Omega matrix; and a . (period) indicates amino acids with a score smaller or equal to 0.5 in the Clustal Omega matrix. (D) Domains of human CSTF2 (577 amino acids). Indicated are the RRM, the Hinge, the proline/glycine-rich region, the 12× MEARA repeats, and the C-terminal domain. Numbers at top indicate the amino acid positions of interest.
mixture of plasmid DNAs. Luciferase measurements were performed between 36 and 48 h after the transfection (22,33,34). Western blots to assess the abundance of the proteins were performed on the same volumes of lysates obtained from the luciferase measurements using the passive lysis buffer supplied with the Dual-Luciferase Reporter Assay System (Promega). Antibodies used for western blots were previously described (22,35). Immunohistochemistry and microscopic imaging were performed as described (22).

Protein tissue samples (wild type and Cstf2ΔD50A/Y) were obtained from four brothers from the same litter. The entire brain and pieces representing all lobes of the liver were collected from the animals and homogenized in 3 ml RIPA buffer (50 mM Tris–HCl pH 7.4, 150 mM NaCl, 1% NP-40, 0.1% SDS, 0.5% deoxycholate) supplemented with PMSF and Halt Protease Inhibitor Cocktail (ThermoFischer). Samples were homogenized with several strokes of Brinkmann polytron homogenizer and a third of the volume was spun at 16 000 × g for 10 min at 4°C. The supernatant was frozen at −80°C until use. 20 μg total protein from the brain and the liver was separated using 8% SDSPAGE, then transferred to an Immobilon-P PVDF membrane, 0.45 μm (Sigma-Aldrich). Membranes were incubated overnight at 4°C with mouse anti-CSTF2 antibody clone 3A7 (35), rabbit anti-CSTF3 antibody (Bethyl laboratories) and the mouse anti-β-tubulin clone E7 (Developmental Studies Hybridoma Bank, University of Iowa). After the addition of appropriate HRP-conjugated secondary antibody, the immunoreactivity was developed by SuperSignal West Pico Chemiluminescent Substrate (ThermoFischer) and the signal captured using an ImageQuant™ LAS 4000 imager. Images were adjusted for brightness and cropped using ImageJ software.

Plasmids and site-directed mutagenesis

The pGL3 plasmid (Promega), Renilla-luciferase construct (SL-Luc) containing a modified C/P site by the addition of two MS2 stem-loop downstream sequences was previously described (33,36). The D50A mutant was created through site-directed mutagenesis. The RRM (amino acids 1–107) and D50A mutant RRM were cloned in bacterial expression vectors as fusions with a His-tag followed by a TEV site at the amino terminal end of the RRM and was previously described (22). All plasmids were verified by sequencing before use.

Bacterial protein expression and purification

Expression and purification of the proteins over metal affinity resin and His-tag removal was as done before (22). For the NMR experiments, transfected Rosetta (DE3) pLysS cells were grown in 2× minimal M9 media using 15NH4Cl (1 g/l) and unlabeled or uniformly 13C-labeled D-glucose (3 g/l) as sole nitrogen and carbon sources, respectively (37). Induction of the transfected cells with 0.5 mM isopropyl-β-D-thiogalactopyranoside (IPTG), harvest and purification of the labeled proteins was also carried out as previously described (22).

Circular dichroism and stability assays

Circular dichroism experiments were performed on a JASCO J-815 instrument in 10 mM sodium phosphate, pH 7.25 with 10 μM of either wild type or D50A RRM protein. The spectra were scanned from 185 to 260 nm with 0.1 nm resolution. The average spectrum was obtained from three technical replicates.

Guandine–HCl experiments were performed in 5 mM HEPES pH 7.4, 50 mM NaCl with 0.5 μM protein samples. Guandine–HCl concentrations ranged from 0.5 to 3 M. Protein samples without guandine–HCl were used as reference. Spectra were collected between 205 and 230 nm with 0.1 nm resolution. Values for 216 nm and 222 nm were plotted to represent the denaturation of the secondary structure of the proteins for β-sheets and α-helices, respectively. The average spectra were obtained from three technical replicates.

3′-End fluorescent RNA labeling and fluorescence polarization/anisotropy

SVL (′-AUUUUAUGUUUCCAGGU-3′) and (GU)8 (′-GUGUGUGUGUGUGUGU-3′) RNAs were commercially synthesized (Sigma-Aldrich). 3′-end fluorescent labeling of the RNAs with fluorescein-5-thiosemicarbazide (ThermoFisher) was done as previously reported (38). The labeled RNAs were used as 3.2 nM final concentration in polarization assays.

Fluorescence polarization experiments were performed in binding buffer (16 mM HEPES pH 7.4, 40 mM NaCl, 0.008% (vol/vol) IGEPAL CA630, 5 μg/ml heparin, and 8 μg/ml yeast tRNA). Purified wild type and D50A RRMs were diluted in 20 mM HEPES pH 7.4, 50 mM NaCl, 0.01% NaN3 and 0.0001% IGEPAL CA630 to 32 μM concentration and used for 2-fold serial dilutions. The highest final concentration of the protein in the polarization assay was 16 μM. Fluorescence polarization samples were equilibrated for 2 h at room temperature in 96-well black plates (Greiner Bio) and measured on Infinite M1000 PRO instrument (Tecan Inc) with excitation set at 470 nm (5 nm bandwidth) and emission 520 nm (5 nm bandwidth). At least three technical replicates were performed. The apparent dissociation constants were calculated by fitting the data to a modified version of the Hill equation (39) using GraphPad Prism version 5.2 software for Windows (GraphPad Software). Unpaired t-test was performed using Microsoft Excel (Microsoft Corp).

Isothermal titration calorimetry (ITC)

The RRM wild type, D50A mutant RRM proteins with SVL RNAs were dialyzed overnight into 10 mM sodium phosphate, 1 mM tris(2-carboxyethyl)phosphine (TCEP), 0.05% (v/v) sodium azide, pH 6 using a 2 kDa MWCO dialysis unit (ThermoFisher). Heats of binding were measured using a MicroCal iTC200 calorimeter (GE Healthcare) with a stirring rate of 1000 rpm at 27°C. For all titrations, isotherms were corrected by subtracting the heats of RNA dilution. The concentrations of proteins and RNA were calculated by absorbance spectroscopy with extinction coefficients of ε280 = 5960 M⁻¹ cm⁻¹ and ε260 = 165.7 M⁻¹.
were generated with PyMol (49). Adaptive Poisson-Boltzmann Solver (APBS), and figures of static potential for each structure were calculated with the 6Q2I for wild type and 6TZE for D50A). The surface electrostatics are available at protein data bank (PDB codes: unlabeled SVL RNA to the 15N-labeled proteins and monomers in Pf1 bacteriophage, a concentrated stock of protein in the TITAN program (50). RNA binding on- and off-rates were calculated for both wild type and D50A mutant protein samples in the absence of a competitor and a mutation in the RNA recognition motif in CSTF2. Males in Family P167 presented with mild intellectual disability and a mutation in the RNA recognition motif in CSTF2.

**RESULTS**

Males in Family P167 presented with mild intellectual disability and a mutation in the RNA recognition motif in CSTF2. The family has four affected male individuals (II:1, II:2, III:1, and III:2, Figure 1) in two different generations con-
nected through the maternal germ line. This pattern of inheritance suggested an X-linked trait carried by females I.2 and II.4. The index proband, a 6-year-old boy (III.1) was presented to the Division of Pediatrics at Bicêtre Hospital (Kremlin-Bicêtre, France) with developmental delays. There was no parental consanguinity. He has one affected brother (II.2) and two affected maternal uncles (I.3 and I.4). All affected males were born after uneventful pregnancies and delivery. The index proband (III.1) was born at term with normal weight (3,760 g), height (50 cm) and occipitofrontal circumference (34 cm). Motor milestones, such as walking, were reached within normal limits. Developmental delay was recognized in early childhood during the first years at school and speech development was retarded. Differential scales of intellectual efficiency (Échelles Différentielles d’Efficience Intellectuelle, EDEI, 60) showed low verbal and nonverbal communication skills, with no other consistent clinical phenotype. He attended a school for children with additional speech therapy and special assistance. Brain MRI showed no specific abnormalities except for an abnormal hypersignal at the posterior thalamic nuclei due to an episode of intracranial hypertension at the age of 9.5 years (data not shown). His two uncles (ages 44- and 46-years-old, II.1 and II.2, Figure 1A) attended a special school for children with learning difficulties. They are both married and employed; one uncle has a 15-year-old unaffected girl, and the other a 7-year-old unaffected girl and a 5-year-old unaffected boy.

Initial karyotype analysis testing for Fragile X and a mutation search in a few selected XLID genes, including SLC6A4, NLGN4 and JARID1C, gave normal results. X chromosome exome sequencing of the index proband (III:1) as part of a large study of >500 unrelated males from families with likely XLID (4) identified a 3 bp deletion in STARD8, and single nucleotide variants in ALG13, COL4A5 and CSTF2. All variants co-segregated with the phenotype: they were present in all affected males and were transmitted through the maternal germ line (Figure 1A and data not shown). The single amino acid deletion identified in STARD8 is present in 180 control males in the Genome Aggregation Database (gnomAD, non-neuro control individuals, https://gnomad.broadinstitute.org/) (61,62), and was therefore interpreted as a benign polymorphism. ALG13 is an established epilepsy-associated gene in females (63), but the ALG13 missense mutation in Family P167 (chrX:110951509G>A, NCBI RefSeq NM_001099922.3:c.638G>A, p.S213N, rs374748006) has been reported in >14 000 control individuals from publicly available databases including gnomAD. The mutation was predicted as disease causing or pathogenic by several prediction tools, including MutationTaster2 (disease causing), DANN (pathogenicity score of 0.9956 with a value of 1 given to the variants predicted to the most damaging (66)) and a high CADD score of 26.9 (67). Furthermore, at the gene level, CSTF2 is highly constrained with zero known loss-of-function mutations and a Z-score of 3.43 (ratio of observed to expected = 0.37) for missense mutations in gnomAD. Conceptual translation of the cDNA demonstrated a mutation of aspartic acid (GAT) to alanine (GCT) within the RNA recognition motif (RRM) of CSTF2 (p.D50A, Figure 1B, D). The aspartic acid at position 50 is conserved in every CSTF2 ortholog surveyed (Figure 1C). We designated this allele CSTF2D50A.

**Polyadenylation efficiency is reduced by the CSTF2D50A mutation**

To determine the effectiveness of CSTF2D50A in control of C/P, we performed the Stem-Loop Assay for Polyadenylation (SLAP). SLAP permits determination of C/P efficiency of a luciferase reporter gene that contains two MS2 bacteriophage RNA stem-loops in the position of the U/GU-rich region downstream of the SV40 late polyadenylation signal; the stem-loops bind to an MS2 coat protein domain (MCP) engineered at the N-terminus of CSTF2 in transfected HeLa cells (33). Previously, using this assay we showed that mutations in the RRM of CSTF2 influenced RNA binding properties of the motif, thus altering reporter gene expression (22). Wild type MCP-CSTF2 produced a SLAP value of 4.49 ± 0.16 normalized luciferase units (NLU), whereas MCP-CSTF2D50A consistently achieved 15% lower polyadenylation efficiency (3.92 ± 0.13 NLU) than wild type MCP-CSTF2 (Figure 2A).

Mutations in the RNA binding site I and II of CSTF2 affect the synergetic function of CSTF2 and CSTF3 (22). Therefore, we co-expressed CSTF3 with either MCP-CSTF2 or MCP-CSTF2D50A to determine whether the D50A mutation affects cleavage and polyadenylation in the presence of CSTF3. CSTF3 increased SLAP for both wild type and MCP-CSTF2D50A to a similar extent, effectively eliminating the reduction observed in the MCP-CSTF2D50A construct expressed alone. Mutations in the CSTF2 RRM altered nuclear-to-cytoplasmic localization (22). Therefore, we hypothesized that the CSTF2D50A mutation might also affect the ratio between nuclear and cytoplasmic CSTF2. Immunohistochemical staining revealed that CSTF2D50A was localized more in the cytoplasm than wild type CSTF2 (Figure 2B). Co-expression of CSTF3 with CSTF2D50A increased the nuclear localization of CSTF2D50A similar to wild type CSTF2 protein (Figure 2B and (22)).

**The CSTF2D50A RRM has a greater affinity for RNA**

Because the aspartic acid (D) to alanine (A) mutation changes the charge in the loop connecting the β2 and β3-strands (Figures 3 and 4, (22,68)), we wanted...
Figure 2. CSTF2D50A is less efficient for C/P because it binds substrate RNA with a higher affinity. (A) SLAP results showing normalized luciferase units (NLU) in HeLa cells without MCP-CSTF2 (–), in cells transfected with MCP-CSTF2 or MCP-CSTF2D50A (black bars) and with CSTF3-Myc (white bars). Western blots to show the expression of FLAG-tagged MCP-CstF-64 (WB: FLAG) or Myc-tagged CSTF3-Myc (WB: Myc). β-tubulin was used as a loading control. Asterisk indicates a statistically significant P value equal to 0.0021, by student’s t-test with unequal distribution of variance. (B) Immunofluorescent images of the described constructs stained with antibodies against the FLAG tag for MCP-CSTF2 (green), Myc tag for CSTF3 (red), and counterstained with DAPI to delineate the nucleus. (C) The Kd for RNA binding was determined for the isolated RRM domain of wild type CSTF2 and CSTF2D50A mutant via the change in fluorescence polarization of a 3′-end labeled RNA substrate (C, D) and isothermal titration calorimetry (ITC; E, F). Changes in fluorescence polarization for wild type and D50A RRM binding to SVL (C) and (GU)8 (D) substrate RNAs. ITC thermograms of wild type (E) and D50A (F) RRM binding to SVL RNA. The Kd and the number of binding sites (N) are indicated on the figures along with the corresponding standard deviation from three replicates. Raw injection heats are shown in the upper panels and the corresponding integrated heat changes are shown in the bottom panels versus the molar ratio of RNA to protein. Kd's and thermodynamics, derived from the fits of the ITC data, are provided in Supplementary Table S1.

to determine whether RNA binding was altered in the CSTF2D50A RRM mutant. To measure binding via fluorescence polarization/anisotropy, bacterially expressed CSTF2 and CSTF2D50A RRMs (amino acids 1–107 (22)) were incubated with fluorescently-labeled RNA oligonucleotides from either the SV40 late transcription unit (SVL (69)) or (GU)8 (68). The wild type RRM bound to the SVL and (GU)8 RNAs with Kd, app of 1.45 ± 0.07 μM and 0.26 ± 0.02 μM, respectively (Figure 2C, D and Supplementary Table S1). The CSTF2D50A mutant RRM bound the two RNAs with significantly higher affinities (Kd, app 0.93 ± 0.13 μM and 0.17 ± 0.01 μM for the SVL and (GU)8 RNA, respectively, Figure 2C, D, Supplementary Table S1). To confirm the RNA-binding affinities, we performed isothermal titration calorimetry (ITC) titrations using the SVL RNA oligonucleotide. The average Kd was 1.52 ± 0.17 μM for wild type RRM and 0.698 ± 0.06 μM for the CSTF2D50A RRM (Figure 2E and F, Supplementary Table S1), which were in the same range as the Kd, app measured before. In both cases, binding was driven by a favorable enthalpy overcoming an unfavorable entropy. However, the enthalpy and entropy for mutant and wild type binding to SVL RNA were different (Figure 2F, Supplementary Table S1). Thus, the enthalpy-entropy compensation for the D50A mutant binding to SVL RNA was greater than that of the wild type RRM, leading to the higher observed affinity, suggesting that CSTF2D50A would bind to RNAs with a greater affinity during C/P.

The D50A mutation affects the side chain interactions and electrostatic surface of the RRM

To characterize the structural and dynamic changes that occur in the CSTF2D50A RRM upon binding to RNA, we turned to solution state nuclear magnetic resonance (NMR) spectroscopy, initially characterizing the general backbone conformation via 2D 15N,1H heteronuclear single quantum coherence (HSQC) spectra for the wild type and mutant RRMs. Predictably, the overlay of the HSQC spectra showed differences in the peak positions of the residues in the loop where the D50A mutation is located (Figure 3A and C, green bars). Otherwise, the majority of the residues in the CSTF2D50A RRM showed almost identical NMR spectra as the wild type.
Figure 3. The p.D50A mutation perturbs the environment of the β-sheet and C-terminal α-helix. The 2D $^{15}$N,$^1$H HSQC of apo (A) RRM WT (green) and RRM D50A (red) and RNA-bound (B) RRM WT•SVL (black) and RRM D50A•SVL (yellow) complexes. All structures were determined from amino acids 1–107 of human CSTF2 or CSTF2D50A. Data were collected at 600 MHz and 27°C. (C) Bar graph indicating backbone amide chemical shift perturbations (CSP) for the RNA-free (Apo) WT and D50A RRM (green) and SVL RNA-bound WT and D50A mutant RRM (red). The CSPs are calculated from the HSQC spectra from panels A and B. (D) The tertiary folds of the wild type CSTF2 and CSTF2D50A RRM domain are similar. Superimposition of the lowest energy solution structure of the WT and D50A RRM in the apo form, after PALES re-scoring against an independent set of RDC data, showing a view of site I (left panel) and the D50A loop (right view). β-strands and α-helices are labelled on the structures; the black arrows indicate the position of residue D50.
next, we determined the three-dimensional structures of the CSTF2 and CSTF2\textsuperscript{D50A} RRMs using CS-Rosetta (47,70). The CSTF2\textsuperscript{D50A} RRM was almost identical to the CSTF2 RRM structure (Figure 3D; 3.081 Å all atom root-mean-square deviation, RMSD), consistent with circular dichroism data (Supplementary Figure S2A). The major difference between the RRMs was observed in the α4-helix, which angled away to make room for the repositioned side chains of the β4-strand to interact with the α4-helix. A small difference was also noted in the relative twist of the β-sheet, which includes RNA binding sites-II and -III.

These differences in secondary structural elements resulted from repositioning of the side chains for the residues in the loop surrounding the D50A mutation and in the amino acids involved in site-I (Tyr25 and Arg85) and -II (Phe19 and Asn91; Figures 1C and 4A–C), which are two of the three sites identified as important for RNA interactions in Rna15, the yeast homolog of CSTF2 (71). Specifically, in wild type CSTF2, the carboxylic acid side chain of Asp50 formed a hydrogen bond with the hydroxyl side chain of Thr53 (Figure 4A, left). Replacing the hydrogen bond acceptor with a methyl group disrupted this interaction in CSTF2\textsuperscript{D50A} (Figure 4A, right). In addition, the Arg51 side chain formed a new ion pair interaction with the side chain carbonyl group of Glu52, instead of the backbone carbonyl of the terminal helix α4 residue Ser103, leading to re-orientation of helix α4 (Figures 3D and 4A). In the CSTF2\textsuperscript{D50A} RRM, the β1-strand had a different twist, which allows the aromatic ring of Phe19 (site-II, Figure 1C) to lift up towards the RNA binding pocket (Figure 4B), contributing to the reorientation of helix α4 in the mutant as the edge of the Phe19 aromatic ring now packs against the aliphatic portions of Asn97, Glu100, and Leu104 of helix α4. In binding site-I of the wild type RRM, the Tyr25 aromatic ring was adjacent to the amino side chain on Lys55, forming a π-cation interaction (Figure 4C). However, in CSTF2\textsuperscript{D50A}, the Tyr25 ring moved away, allowing Glu26 to form a hydrogen bond with Arg85 (site-I residue), increasing the rigidity in binding site-I (Figure 4C, right). Thus, local differences in side chain arrangement, particularly in binding sites-I and -II, contribute to the differences we observed in RNA binding.

We calculated the electrostatic potentials using the Adaptive Poisson–Boltzmann Solver (72). Changing the negatively charged Asp50 to the non-charged Ala altered the charge distribution of the loop from negative to positive in the mutant (Figure 4D). Binding site-I (Site I, Figure 4D), which forms upon RNA binding (71) in the wild type, was within a cavity with low positive electrostatic potential for RNA binding (Figure 4D, left). However, in the D50A mutant, the hydrogen bonding between Arg85 and Glu26 (Figure 4C and D) moved the positively charged side chain of Arg85 toward the inside of cavity, resulting in a higher positive electrostatic potential distribution in the binding site-I (Figure 4D, right). We hypothesize that this larger positive electrostatic potential could be the driving force for the more favorable enthalpy of binding and the tighter RNA binding affinity.

The D50A mutation affects the structure and dynamics of the RNA-bound RRM

To test effects of RNA binding, the CSTF2 and CSTF2\textsuperscript{D50A} RRMs were titrated with SVL RNA to saturation (molar ratio of 2.3:1, RNA to protein) with changes monitored in 2D \textsuperscript{15}N,\textsuperscript{1}H HSQC spectra (Figures 3B and 5A). Both proteins showed the same binding patterns (i.e. direction, magnitude and exchange regime) for most residues (e.g. Val18, Val20, Ala27, Asp90 and Leu47; Figure 5A). Gly54 located within the D50 loop was perturbed upon addition of RNA to the CSTF2\textsuperscript{D50A} RRM but not the wild type RRM (Figure 5A, left), indicating a potentially new interaction in the mutated domain. Next, we calculated amide chemical shift perturbations (CSPs) between SVL-bound CSTF2 and CSTF2\textsuperscript{D50A} RRMs (Figure 3C, red bars). The largest differences were observed in the loop containing the mutation, similar to unbound RRMs (Figure 3C, green bars), and other significant amide CSPs (i.e. above the mean, 0.032 ppm for apo and 0.048 ppm for RNA-bound) were observed in the RNA-bound domain for the N-terminus (residues 5–7) near binding site-I (25 and 85), site-II (residues 19 and 91; Figures 1C and 3C), and in the β2-strand preceding the mutation (Figure 3B and C, red bars).

The on- and off-rates of RNA binding were determined with a simple two state ligand binding model using the 2D NMR line shape analysis program (TITAN, Supplementary Figure S3 (50)). From the RNA-induced perturbations that were in the fast and intermediate exchange regimes, we calculated a \(K_d\) of 0.70 ± 0.02 μM for wild type and 0.54 ± 0.02 μM for CSTF2\textsuperscript{D50A}, which followed the trend established by fluorescence polarization and ITC (Supplementary Table S1). The off-rates (\(k_{\text{off}}\)) were similar for wild type (305.5 ± 1.4 s\(^{-1}\)) and CSTF2\textsuperscript{D50A} (312.4 ± 2 s\(^{-1}\)) RRMs. Both on-rates were the same as or exceeded the rate of diffusion, a feature seen in other protein-nucleic acid complexes where electrostatic attraction accelerates the on-rate (73,74). However, the on-rate (\(k_{\text{on}}\)) for CSTF2\textsuperscript{D50A} (5.83 ± 0.21 × 10\(^8\) M\(^{-1}\) s\(^{-1}\)) was faster than the rate for wild type (4.35 ± 0.12 × 10\(^8\) M\(^{-1}\) s\(^{-1}\)). We conclude, therefore, that the lower \(K_d\) for CSTF2\textsuperscript{D50A} was primarily due to the faster on-rate for RNA binding to CSTF2\textsuperscript{D50A} compared to the wild type.

We measured \textsuperscript{15}N R\(_1\), R\(_2\), and nuclear Overhauser effect (NOE) relaxation values at 600 MHz 27°C to derive backbone model-free order parameters (\(S^2\)), which report on the amplitude of fast timescale motion and the global correlation time (\(\tau_c\)) (53,54). In the absence of RNA, the \(S^2\) values for the CSTF2\textsuperscript{D50A} RRM domain differed from wild type only in the D50A loop, which became more rigid (Figure 5B and Supplementary Figure S4A). However, for the RNA-bound structures, the \(S^2\) values indicated different amide group flexibilities in the mutant RRM (Figure 5C and Supplementary Figure S4B). SVL RNA binding to the CSTF2 RRM reduced \(S^2\) values throughout the β-sheet (e.g. residues 18–22 and 44–48), indicating increased flexibility upon RNA binding, in agreement with earlier studies (68,75). However, SVL RNA binding to the CSTF2\textsuperscript{D50A} RRM increased \(S^2\) values for the β-sheet and the α3-helix adjacent to site-II (Phe19 and Asn91, Figure 5C). This indicated greater rigidity in the pico-to-nanosecond time scale.
Figure 4. The altered side chain interactions of CSTF2 D50A lead to different local electrostatic surface potentials. Panels A–C show the different side chain orientations and interactions present in the wild type CSTF2 RRM (green) and mutant CSTF2 D50A RRM (red). (D) Calculated electrostatic potential for wild type (left) and D50A mutant (right) RRM. The red-to-blue surface representation highlights negative-to-positive electrostatic potential from −2 to 2 \( k_B T / e \), where \( k_B \) is Boltzmann’s constant, \( T \) is the temperature in Kelvin and \( e \) is the charge of an electron.
Figure 5. The p.D50A mutation affects the RNA-binding kinetics and dynamics of the RRM. (A) Overlays of the 2D $^{15}$N,$^1$H HSQC spectra for the titration of CSTF2 (top panels) and CSTF2 D50A (bottom panels) RRM. Red-yellow-blue and red–green–blue gradients represent the 0–10% titration of wildtype and D50A RRM with SVL RNA, respectively. Titration data were acquired at 600 MHz and 27°C. All spectra are shown with the same contour base level relative to noise. The magnitude and direction of each titrated residue is shown with arrows and assignment. (B and C) Backbone amide $^{15}$N order parameter ($S^2$) for (B) apo RRM WT (green) and RRM D50A (red) and (C) SVL RNA-bound RRM WT•SVL (black), and D50A•SVL (yellow). The pink lines at an $S^2$ value of 1.0 denotes the maximum value for the order parameter.
in the RNA binding surface of the D50A mutant when bound to RNA. The order parameter is a measure of conformational entropy within a protein (76,77). Indeed, the wild type RRM, which became more flexible upon RNA binding, pays less of an entropic penalty compared to the D50A mutant (Supplementary Table S1), which maintained the same flexibility upon RNA binding.

We also determined whether the point mutation in D50A changed the stability of the secondary structure of the motif using CD spectroscopy (Supplementary Figure S2B and C). Both RRMs were thermally stable with minimal changes in ellipticity at 100°C (data not shown). However, upon chemical denaturation with guanidine, we observed that the inflection point for the wild type protein was reached at 2 M guanidine and was decreased to at 1.5 M for the CSTF2D50A RRM mutant. This result suggested that secondary structure of the mutant was less stable than the wild type.

The D50A mutation causes differential gene expression in mice

To examine effects of the D50A mutation in vivo, we developed Cstf2D50A/Y mice using CRISPR-Cas9 technology. Hemizygous male (Cstf2D50A/Y) and homozygous female (Cstf2D50A/D50A) mice are viable and fertile. Male Cstf2D50A/Y mice appear to be runted. To determine CSTF2 protein expression levels in Cstf2D50A/Y mice, we isolated total protein from brains and livers of 50-day old wild type and Cstf2D50A/Y littermates. Cstf2D50A/Y mice expressed slightly higher amounts of CSTF2 in both total brain and liver (Figure 6A). We also noted strong expression of the βCstf-64 splice variant (78,79) in brains of both wild type and Cstf2D50A/Y mice (Figure 6A, lanes 1–4, upper band).

RNA-seq (see below) did not indicate differential gene expression of the Cstf2 mRNA in the brains of Cstf2D50A/Y mice (Supplementary Table S2), suggesting that protein differences were translational or post-translational.

To further characterize the changes in the brain gene expression, we isolated whole brain RNA from 50-day old wild type and Cstf2D50A/Y littermates and performed RNA-seq. An example of last exon showing the opposite pattern (571 versus 290, Figure 6E and Supplementary Table S3). An example of last exon shortening is the RNA Binding Motif Protein 24 (Rbm24) gene (Figure 6F). Gene ontology (GO) analysis of the genes shortening the last exon showed localization of macromolecules in cells, retrograde transport, cell migration, and more (Supplementary Table S4). GO terms of the genes lengthening the last exon indicated enrichment in neurogenesis, neuron development, and more (Figure 6G).

The number of genes in which APA caused changes in the CDS favoring the pPAS in Cstf2D50A/Y animals was 3.5-fold greater than the number of genes showing the opposite pattern (396 versus 113, Figure 6H and Supplementary Table S5). For example, the mRNA for the shorter version of the Coatomer Protein Complex Subunit Gamma 1 (Copg1) gene was increased in Cstf2D50A/Y mouse brains (Figure 6I). This isoform was previously reported to be enriched in granule cells of the brain (83). GO term analysis for the 113 genes lengthening their CDSs did not reveal ontologies that were enriched (FDR < 0.05, Supplementary Table S3). However, GO terms for genes with shorter CDSs were enriched for neurogenesis, neurodevelopment and cell adhesion (Figure 6J). We propose that in the patients carrying the D50A mutation, the balance of the critical protein isoforms involved in brain development is disrupted, contributing to their cognitive disability.

We also evaluated the genes involved in the organization and long-term maintenance of the synapses. In addition to up-regulation of the pre-synaptic gene Chnl (Figure 6B), our analysis revealed that both the 3′ UTR and the CDS of the mRNA encoding the pre-synaptic cell-surface receptor Nrnx1 is shorter in brains of Cstf2D50A/Y mice (Supplementary Figure S5). Similarly, the 3′ UTR of Cntnap2 is longer in mutant mice (Supplementary Figure S5). On the post-synaptic side, 3′ UTRs of several genes involved in regulation of protein synthesis, scaffolding and signal transmission (Shank2, Fmr1, Caenog) were affected by APA (Supplementary Tables S3 and S4).

Finally, we examined hexamer frequencies within 150 nt 5′ and 3′ of the pPAS and dPAS of sites that are affected in Cstf2D50A/Y mice (Supplementary Figure S6). The hex-
Figure 6. Gene expression and 3′ APA analyses in brains of Cstf2D50A/¬ mice. (A) Protein immunoblots of CSTF2/CstF-64 (top), CSTF3/CstF-77 (middle) and β-tubulin (bottom) from two wild type (lanes 1, 2, 5, 6) and two Cstf2D50A/¬ (3, 4, 7, 8) male mice from indicated tissues (total brain and liver). Bands corresponding to CstF-64, CstF-64, CstF-77, and β-tubulin are indicated at right. Genotypes of the wild type and Cstf2D50A/¬ mice were verified by PCR-RFLP as described in the Materials and Methods. (B) Differential gene expression analysis of RNA-seq from total brain samples isolated from 50-day old male wild type and Cstf2D50A/¬ mice (three wild type and five mutants). Left, volcano plot of the differentially expressed genes. Right, list of genes that are differentially expressed, fold change (log2) and adjusted P-value (–log10) are indicated. (C) Gene ontology for biological functions of differentially expressed genes in the brains of wild type and Cstf2D50A/¬ mice. Number (n) on the right indicates the number of genes found in each functional category. (D) Schematic illustration of APA events analyzed. (Top) Genes in which APA changes the length of the last (3′-most) exon. (Bottom) Genes in which APA occurs in the coding region (CDS) changes the protein coding potential of the gene. Proximal polyadenylation site (PAS), distal PAS, start and stop codons in translation, and regions containing cis-acting RNA elements (e.g. miR and RNA-binding protein binding sites) are indicated. (E) Scatter plot showing expression change of proximal PAS isoform (x-axis) and that of the distal PAS isoform (y-axis) in total RNA. Genes with significantly shortened or lengthened 3′ ends (P < 0.05, Fisher’s exact test) in Cstf2D50A/¬ total mouse brains (three wild type and five mutant) are highlighted in red and blue, respectively. Color coded numbers indicated the number of genes shortened, lengthened, or showing no change. (F) C/P in the last exon of the Rbm24 gene in wild type (green) or Cstf2D50A/¬ (rust) mouse brains switch from the distal to the proximal poly(A) sites. Proximal and distal PASs are indicated with the maximum RPM values shown. (G) Top ten gene ontology categories (FDR < 0.05) of enriched biological functions in Cstf2D50A/¬ mice showing lengthening of their 3′-most exons. Number of genes (n) that are detected in each functional category are shown on the right. (H) Scatter plot showing expression change of proximal PAS isoform (x-axis) and that of the distal PAS isoform (y-axis) in total RNA. Genes with significantly shortened or lengthened CDSs (P < 0.05, Fisher’s exact test) in Cstf2D50A/¬ total mouse brains (three wild type and five mutant) are highlighted in red and blue, respectively. Color coded numbers indicated the number of genes shortened, lengthened, or showing no change. (I) C/P in the Copg1 gene shortens the CDS in Cstf2D50A/¬ mouse brains (rust) compared to wild type (green). Maximum RPM values are shown for each animal. Gene structure and different transcripts are shown at the bottom. (J) Top ten gene ontology categories (FDR < 0.05) of enriched biological functions in Cstf2D50A/¬ brains showing shortened transcripts in their CDS. Number of genes (n) that are detected in each functional category are shown on the right.

DISCUSSION

Many monogenic intellectual deficiencies are X-linked (XLIDs) because of hemizygosity of X-chromosomal genes in males (84–86). The X-linked CSTF2 gene is essential for embryonic growth and development (30,31). Therefore, it was surprising that a single nucleotide mutation in CSTF2 would result in structural and functional changes in 3′ end mRNA processing yet not have lethal effects. Other muta-

amer CACACA was most enriched between 50 and 150 nt upstream of the pPAS, while a similar sequence, CAACCA was enriched upstream of dPAS. Downstream sequences tended to be U-rich (pPAS) or A-rich (dPAS).

These changes in 3′ UTRs and protein coding capacity of these key synaptic genes in the excitatory synapses may lead to changes in the long-term potentiation, affecting neuronal plasticity and development, consistent with the speech delays and intellectual disability observed in humans.
tions in the RRM of CSTF2 have been reported in gnomAD but with no associated disease (62). As reported here (Figure 1), it appears that the CSTF2<sup>D50A</sup> mutation affects primarily intellectual functions and speech development in the affected males, but effects on other physiological functions were not noted, suggesting that the brain is more sensitive to this particular mutation than other organs. We propose, therefore, that the p.D50A mutation in CSTF2 results in non-syndromic intellectual deficiency in males by altering RNA binding during C/P, thus changing the expression of key genes in neurodevelopment.

Only one mutation involving a core polyadenylation protein, NUDT21 (which encodes the 25 kDa subunit of mammalian cleavage factor I), has been implicated in neuropsychiatric disorders resulting in Rett syndrome—like symptoms and intellectual disability (13,14). We did not observe altered expression of MeCP2—which is frequently associated with Rett syndrome—in our mouse model (not shown), but observed altered sites of polyadenylation in many other neurodevelopmental genes (Figure 6).

RNA-contact residues in the CSTF2 RRM play specific roles during C/P (22). Introduction of the D50A mutation into CSTF2 resulted in a small but consistent reduction in C/P efficiency (Figure 2A). Such a reduction in activity would likely result in altered C/P in vivo, favoring proximal sites over more distal sites (87,88). Analysis of genome-wide polyadenylation changes in the CSTF2<sup>D50A</sup> mouse indicated exactly that shorter RNAs (‘most exons and changes in the last exon) were enriched in brains of CSTF2<sup>D50A</sup> mice (Figure 6E, H). Residues within the CSTF2 RRM are also important for functional interactions between CSTF2 and CSTF3 during C/P (22). However, we did not observe a reduction in the ratio between MCP-CSTF2<sup>D50A</sup> alone and MCP-CSTF2<sup>D50A</sup> co-expressed with CSTF3, which we previously observed with CSTF2 RRM binding site-I and -II mutants (Figure 2). This suggests that the D50A mutation causes the phenotype independent of the interactions with CSTF3. Furthermore, the small decrease in the polyadenylation efficiency in our reporter system might suggest that the mutated protein is retained longer in the cytoplasm (Figure 2B), possibly through increased interaction with cytoplasmic RNAs (22). Indeed, we observed increased levels of CSTF2 protein (Figure 6A) in the brains of CSTF2<sup>D50A</sup> mice, which could favor proximal sites of C/P. However, we observed no change of CSTF3 protein in the same samples (Figure 6A). This suggests that the functional level of CSTF is unchanged in nuclei of neuronal and hepatic cells (22).

With reduced C/P efficiency, we observed an increased affinity of the CSTF2<sup>D50A</sup> RRM for RNA (Figures 3 and 5), probably also contributing to the retention of CSTF2<sup>D50A</sup> in the cytoplasm (Figure 2B). The <i>K<sub>c</sub></i> of the CSTF2<sup>D50A</sup> RRM for RNA was less than half that of wild type CSTF2 due to the faster <i>k<sub>off</sub></i> rate (Supplementary Table S1). Based on the on- and off-rates, the D50A mutant binds to RNA faster than wild type but releases the RNA with the same off-rate. It has been previously shown that the β2-β3 loop (D50A loop here) is important for the shape recognition of RNA in some RRM containing proteins (89,90). Our structures for the CSTF2 and CSTF2<sup>D50A</sup> RRMs allowed us to model the differences in RNA binding based on the relative orientation of the side chains of the mutant RRM, electrostatic potential, and rigidity of individual backbone atoms of the β-sheet, causing faster RNA binding in the mutant (Figure 4D) by helping to overcome the greater entropic penalty of binding resulting from the enhanced rigidity of the β-sheet in the mutant.

This mutation increases the affinity of CSTF2<sup>D50A</sup> for RNA (Figure 2C, D), which we might consider a ‘gain-of-function’ mutation at the molecular level. However, the mutation leads to reduced polyadenylation efficiency (Figure 2A), which could be considered a loss-of-function. The phenotype observed is intellectual disability (Figure 1), which many would consider to be a loss-of-function. Thus, the mutation appears to be a molecular gain-of-function that causes a loss-of-function phenotype.

To confirm that the CSTF2<sup>D50A</sup> mutation had a neuropsychological effect, we created Cstf2<sup>D50A</sup> mice with the same mutation. Several labs have examined effects of knockdowns of CSTF2 in cells in culture, but noted altered C/P in only a relatively small number of genes (87,91–93). Unlike those studies, we observed C/P site changes in 1370 genes in the brains of hemizygous CSTF2<sup>D50A</sup> Y mice (Figure 6). The majority of the changes in our study favored proximal PASs, effectively shortening the CDSs or shortening the length of the last exons (3’ UTRs). Why did the knock-down studies have fewer consequences than the D50A point mutation? Possibly, their results were confounded by the presence of tCstF-64 (gene symbol CSTF2T) compensating for decreased CSTF2 in those cells. In support of that idea, knockout of Cstf2t in mice resulted in few phenotypes in most cells (where Cstf2 was also expressed), but led to severe disruption of spermatogenesis in germ cells (where Cstf2t was expressed in the absence of Cstf2 (26)).

We noted APA in several genes involved in pre-synaptic (Nrxn1 and Nntnap2) granule cells and post-synaptic Purkinje cells (Cbln1, Shank2, Fmr1 and Caenog4) (94). Mutations in these genes have been associated with development disorders, delayed speech, and intellectual disability (6,95–97). It seems possible that these same disruptions are affecting intellectual ability in the human patients.

But it is also possible that cells in the brain (either neurons, glia, or both) are more sensitive to mutations in CSTF2. It has long been known that the brain uses alternative polyadenylation extensively to provide transcriptomic diversity and mRNA targeting signals for plasticity and behavioral adaptation (2,98–100). By extending occupancy times on the pre-mRNA, the D50A mutation likely changes the balance of APA to favor shorter mRNAs, thus reducing the effectiveness of over one thousand mRNA transcripts (Figure 6). The brain also expresses a neuron-specific alternatively-spliced isoform of CSTF2 that contains 49 extra amino acids, called βCstF-64 (Figure 6A, (78,79)). Thus, we speculate that the D50A mutation might interact with the additional domain in βCstF-64 in an as-yet undefined manner to give the brain phenotype.

How does an increase in the affinity of CSTF2 for RNA result in altered cleavage and polyadenylation? Previous work showed that formation of the C/P complex takes 10–20 seconds for a weak site; assembly on stronger sites is faster (101). We speculate that the increased affinity of CSTF2<sup>D50A</sup> alters the rate of formation of the CstF complex on the downstream sequence element of the nascent
pre-mRNA during transcription (69). We further speculate that the specificity of binding is reduced in CSTF2D50A. This combination of faster binding and reduced specificity could promote increased C/P of weaker sites, which tend to be more proximal (91), not unlike the effects of slower RNA polymerase II elongation (102,103). These C/P changes subsequently affect post-transcriptional regulation of key mRNAs by changing 3′ UTR regulation by revealing miRNA or RNA-binding protein sites, or by changing targeted localization of mRNAs to neural projections (16,17,83,104–107). Mutations in CSTF2 may have even more striking effects in other target tissues. Studying these mutation-induced changes in gene expression will be important for understanding the mechanisms of polyadenylation as well as the intricacies of neuronal development.

DATA AVAILABILITY
High-throughput sequence data has been deposited to the Gene Expression Omnibus (GEO) database repository at NCBI with the accession numbers GSE152976 (RNA-seq) and GSE152975 (3′-seq). The structures of the wild type CSTF2 (PDB 6Q2I) and CSTF2D50A (PDB 6TZE) RRM domains were deposited to the Research Collaboratory for Structural Bioinformatics Protein Data Bank.

SUPPLEMENTARY DATA
Supplementary Data are available at NAR Online.

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