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RESEARCH PAPER



## Neuronal-specific microexon splicing of *TAF1* mRNA is directly regulated by SRRM4/nSR100

Simona Capponi<sup>a</sup>, Nadja Stöffler<sup>a</sup>, Manuel Irimia<sup>b,c</sup>, Frederik M.A. Van Schaik<sup>b,d</sup>, Mercedes M. Ondik<sup>e</sup>, Martin L. Biniössek<sup>f</sup>, Lisa Lehmann<sup>f</sup>, Julia Mitschke<sup>f</sup>, Marit W. Vermunt<sup>g</sup>, Menno P. Creyghton<sup>g</sup>, Ann M. Graybiel<sup>e</sup>, Thomas Reinheckel<sup>f,h</sup>, Oliver Schilling<sup>i</sup>, Benjamin J. Blencowe<sup>j</sup>, Jill R. Crittenden<sup>e</sup>, and H. Th. Marc Timmers<sup>b,a</sup>

<sup>a</sup>German Cancer Consortium (DKTK) partner site Freiburg, German Cancer Research Center (DKFZ) and Department of Urology, Medical Center-University of Freiburg, Freiburg, Germany; <sup>b</sup>EMBL/CRG Systems Biology Research Unit, Centre for Genomic Regulation, The Barcelona Institute for Science and Technology, Barcelona, Spain; <sup>c</sup>Universitat Pompeu Fabra (UPF), Barcelona, Spain; <sup>d</sup>Molecular Cancer Research and Stem Cells, Regenerative Medicine Center and Center for Molecular Medicine, University Medical Center Utrecht, Utrecht, The Netherlands; <sup>e</sup>McGovern Institute for Brain Research, Department of Brain and Cognitive Sciences, Massachusetts Institute of Technology, Cambridge, MA, USA; <sup>f</sup>Institute of Molecular Medicine and Cell Research, Faculty of Medicine, University of Freiburg, Freiburg, Germany; <sup>g</sup>Hubrecht Institute-KNAW and University Medical Center Utrecht, Utrecht, The Netherlands; <sup>h</sup>German Cancer Research Center (DKFZ), Heidelberg, and German Cancer Consortium (DKTK) partner site Freiburg, Germany; <sup>i</sup>Institute of Surgical Pathology, Faculty of Medicine-University of Freiburg, Freiburg, Germany; <sup>j</sup>Donnelly Centre and Department of Molecular Genetics, University of Toronto, Toronto, Canada

### ABSTRACT

Neuronal microexons represent the most highly conserved class of alternative splicing events and their timed expression shapes neuronal biology, including neuronal commitment and differentiation. The six-nt microexon 34' is included in the neuronal form of *TAF1* mRNA, which encodes the largest subunit of the basal transcription factor TFIID. In this study, we investigate the tissue distribution of *TAF1*-34' mRNA and protein and the mechanism responsible for its neuronal-specific splicing. Using isoform-specific RNA probes and antibodies, we observe that canonical *TAF1* and *TAF1*-34' have different distributions in the brain, which distinguish proliferating from post-mitotic neurons. Knockdown and ectopic expression experiments demonstrate that the neuronal-specific splicing factor SRRM4/nSR100 promotes the inclusion of microexon 34' into *TAF1* mRNA, through the recognition of UGC sequences in the poly-pyrimidine tract upstream of the regulated microexon. These results show that SRRM4 regulates temporal and spatial expression of alternative *TAF1* mRNAs to generate a neuronal-specific TFIID complex.

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Alternative mRNA splicing; microexons; neuronal transcription; neurogenesis; basal transcription factors; *TAF1*; X-linked Dystonia Parkinsonism

## Introduction

Alternative microexons of 3 to 27 nt represent the most highly conserved class of pre-mRNA splicing events in vertebrates [1]. These exons are predominantly found in the nervous system and preserve the open reading frame inserting only a few amino acids in loop regions of the corresponding protein [1]. A major regulator of this “exon microsurgery” is the neuronal-specific Ser/Arg repeat splicing factor of 100 kDa (nSR100), also known as Serine/Arginine Repetitive Matrix 4 (SRRM4) [1], which recognizes UGC-containing intronic sequences of the pre-mRNA to stimulate microexon inclusion [2]. Disruption of SRRM4 results in multiple neurodevelopmental defects and has been causally linked to neurological disorders [3,4]. One of the candidate substrates of SRRM4 is *TAF1* microexon 34' [1]. The *TAF1* gene encodes the largest subunit of basal transcription factor IID (TFIID), which is the primary core promoter recognition factor and nucleates RNA polymerase II (pol II) preinitiation complex assembly for transcription [5]. While different *TAF1* isoforms are ubiquitously expressed, neuronal tissues express the *TAF1*-34' mRNA isoform (aka *N-TAF1*) [6]. This isoform results from the inclusion of the 6-nt long microexon 34', encoding two

additional amino acids (alanine and lysine), to the canonical *TAF1* (c*TAF1*) protein of 1893 residues [6]. The direct contributions of neuronal splicing factors such as SRRM4 to the formation of *TAF1* mRNAs and their expression patterns in the brain have not yet been explored.

Mutations in the *TAF1* gene have been associated with neurodevelopmental [7] and neurodegenerative conditions [6]. In particular, perturbations of *TAF1* mRNA biosynthesis have been associated with X-linked dystonia-parkinsonism (XDP, MIM: 314250) [8], an adult onset, neurodegenerative condition with progressive loss of voluntary motor control replaced by severe motor contractions (dystonia) combined with or replaced by parkinsonism features [9,10]. The neuropathology of XDP is characterized by decreased numbers of neural progenitors in the subventricular zone [11] and prominent loss of medium spiny neurons within the striatum [12], a forebrain region that controls voluntary movement. All XDP patients harbour the insertion of an SVA (SINE-VNTR-Alu) retrotransposon of the F-subclass into *TAF1* intron 32 [6], which has been proposed to affect expression and alternative splicing of *TAF1* mRNAs [6,8].

**CONTACT** H. Th. Marc Timmers  [m.timmers@dkfz-heidelberg.de](mailto:m.timmers@dkfz-heidelberg.de)  German Cancer Consortium (DKTK), partner site Freiburg German Cancer Research Center (DKFZ) and Department of Urology Medical Center-University of Freiburg, Breisacher Str. 66, 79106 Freiburg, Germany  
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Given the involvement of *TAF1* mRNA processing in human neurological disorders, we investigated the relation of SRRM4 to the brain-specific distribution of cTAF1 and TAF1-34'. Discrimination of microexon-containing mRNAs from canonical mRNAs by *in situ* methods is challenging due to the very small size of the micro-exon. We tested BaseScope™ probes in mouse brains to discriminate mRNA molecules that differ in only 6 nt. By employing this technique, we have found that *Taf1-34'* mRNAs are enriched in post-mitotic neurons, whereas *cTaf1* is more widely expressed in the brain, including cells undergoing division and post-mitotic neurons. BaseScope™ detection was validated at the protein level by using antibodies specific to TAF1 proteins that include or exclude microexon 34'. Employing mouse and human cell systems we find that SRRM4 is required and sufficient to promote microexon 34' inclusion in *TAF1* mRNAs in neuronal and non-neuronal backgrounds. The splicing event is mediated by SRRM4 recognition of two UGC motifs located in the poly-pyrimidine tract upstream of microexon 34'. Taken together, these results provide strong evidence that SRRM4 directs inclusion of microexon 34' in *TAF1* mRNAs to regulate the temporal and spatial expression of different TAF1 protein isoforms in mammalian brains.

## Results

### Analysis of cTaf1, Taf1-34' and Srrm4 expression patterns in the mouse brain

To investigate the link between the mRNA expression of *Taf1-34'* and the neuron-specific splicing factor *Srrm4*, we first explored their expression patterns in mouse brain sections at the level of the striatum. We employed *in situ* hybridization (ISH) with the BaseScope™ method, to discriminate between *cTaf1* and *Taf1-34'* mRNAs using specific probes against the 6-nt microexon 34' or against the sequence spanning the flanking exons. In adult mouse brain sections, *cTaf1* probes detected a broadly distributed expression in cerebral cortex, corpus callosum, striatum and septum (Fig. 1A). Prominent *cTaf1* expression was detected in cells along the ventricle wall and within the neurogenic sub-ventricular zone (SVZ) (Fig. 1A' and A''). Comparison of *cTaf1* and *Taf1-34'* signals indicated clear differences in their distribution patterns. The *Taf1-34'* signal was more prominent in the cerebral cortex compared to *cTaf1* and *Taf1-34'* mRNA expression was sparse in the glial-rich corpus callosum, the ventricle wall and the SVZ (Fig. 1B, B' and B''). Similarly, the expression of *Srrm4*, detected by an RNAscope® probe, which uses multiple probes to increase signal amplification [13], was abundant in neuron-rich regions relative to the low levels in the glial-rich corpus callosum (Fig. 1C and 1C'). *Srrm4*-positive cells were also present along the ventricle wall and SVZ (Fig. 1C'').

The mRNA ISH results were validated by immunohistochemical analyses (IHC) of different mouse brain regions by cTaf1-, Taf1-34'- and Srrm4-specific antibodies (Fig. 2). The affinity-purified Taf1 antibodies developed in this study were directed against the region spanning microexon 34' as the isoform-specific epitope, which is identical between mouse and human TAF1-34'. The sera displayed high specificity in the detection by IHC and

immunoblotting of both endogenous (Fig. 2A-C'') and ectopically expressed proteins (Fig. 2D-H and Supplementary Fig. 1), with very limited, if any cross-reactivity when tested against the counterpart isoform (Fig. 2D-H and Supplementary Fig. 1). As additional validation, we performed IHC and confirmed that the antibodies against Taf1-34' and Srrm4 were selectively immunoreactive with cell nuclei in neuron-rich regions of the brain, whereas the antibody against cTaf1 was ubiquitously immunoreactive across neuronal and non-neuronal cell nuclei (Supplementary Fig. 1). We next examined the distribution of cTaf1-, Taf1-34'- and Srrm4 immunoreactivity within the brain. Consistent with the mRNA results, cTaf1 protein expression was detected in the nuclei of many cells throughout the brain, including in cells of the glial-rich corpus callosum and in post-mitotic neurons within the cerebral cortex and the striatum (Fig. 2A-A''). By contrast, Taf1-34' expression was selectively enriched in neuronal nuclei (Fig. 2B and B'') and was low within the corpus callosum (Fig. 2B'). The expression of Srrm4 also appeared enriched in neuronal nuclei and at low levels in the corpus callosum (Fig. 2C-2C''). Magnification of sections at levels including the striatum showed that expression was high in the neocortex (Fig. 2C) but sparsely distributed in striatal cell nuclei (Fig. 2C'').

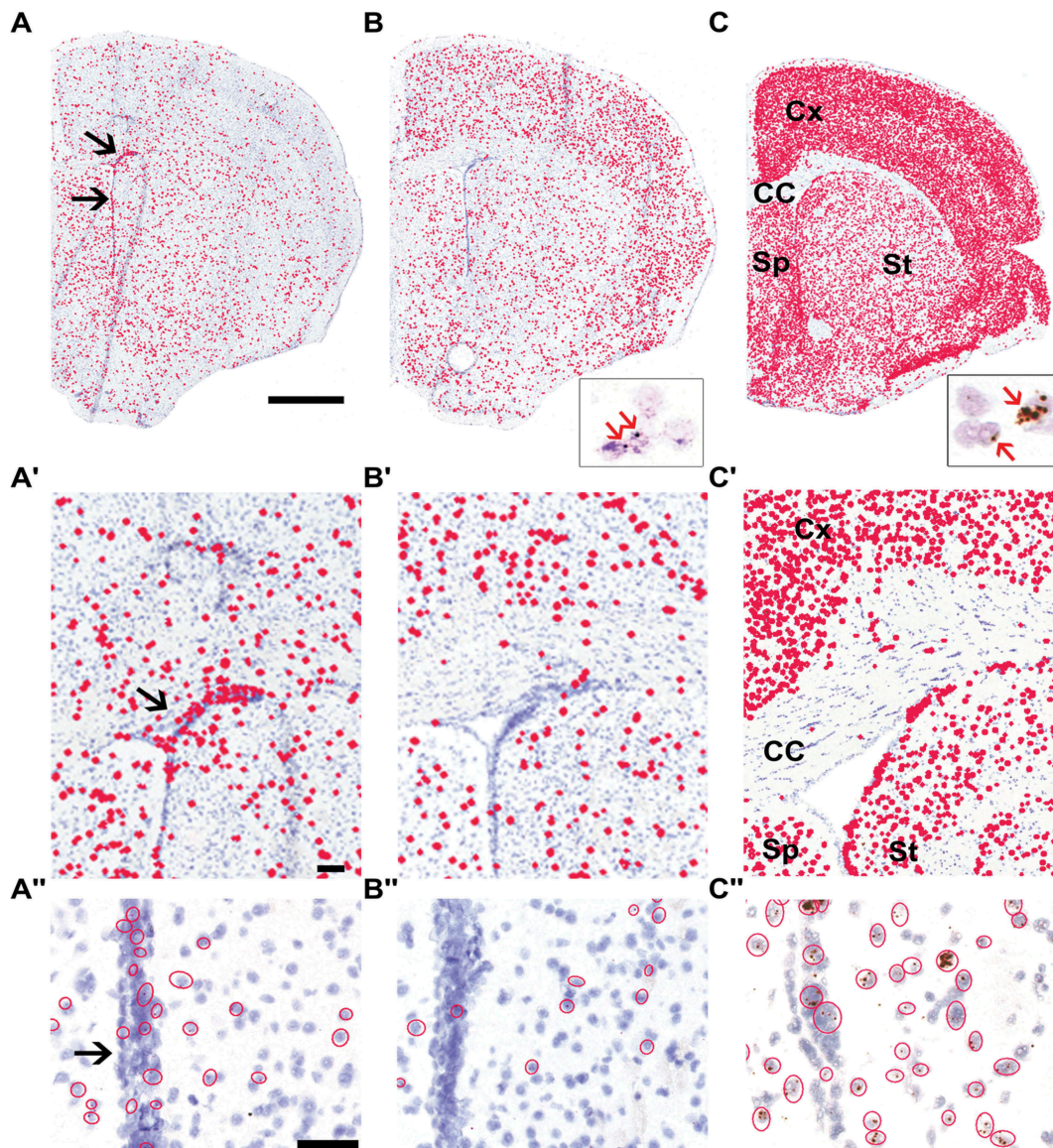
The ISH and IHC performed on mouse brains indicated that cTaf1 and Taf1-34' have different expression patterns within the forebrain. We therefore validated this differential distribution in the human brain by analysing RNA-seq data (Supplementary Fig. S2, panels A-B) and RT-PCR (Supplementary Fig. S2, panels C-D) from specimens from the striatum (caudate nucleus and putamen), neocortex, motor cortex, and thalamus. Consistent with the ISH data from the mouse brain, we confirmed that the inclusion of microexon 34' is more prominent in the neocortex than within the striatum, while not detectable in the thalamus. The expression pattern of *SRRM4* correlates with microexon 34' inclusion, supporting their inter-dependency.

In conclusion, the mRNA ISH, RNA-seq and IHC analyses show that Taf1-34' and Srrm4 are preferentially expressed in post-mitotic neurons relative to the more widespread expression of cTaf1. Importantly, the IHC analysis validated the sensitivity and accuracy of BaseScope™ technology to detect microexon 34'-containing *Taf1* mRNAs *in situ* (Fig. 1) indicating that this is a versatile approach for the *in situ* detection of microexon sequences.

### Inclusion of microexon 34' into TAF1 mRNAs occurs in the later stages of neuronal differentiation

The analysis of cTaf1 and Taf1-34' expression in the mouse brain suggested that these two isoforms display a temporal regulation. To investigate when during neural development Taf1-34' first emerges, we examined the expression profile of this isoform using RNA-seq data collected at different time points during differentiation of mouse embryonic stem cells (mESCs) into cortical glutamatergic neurons [14] (Fig. 3A). Inclusion of microexon 34' was evaluated using the Percentage-Spliced-In (PSI) metric, which determines the percentage of *Taf1* mRNAs containing microexon 34'. This study indicated that microexon 34' is incorporated into *Taf1* mRNA at the beginning of neuronal differentiation, reaching the highest value in mature post-mitotic





**Figure 1.** Differential expression of *cTaf1*, *Taf1-34'* and *Srrm4* mRNAs in the mouse brain.

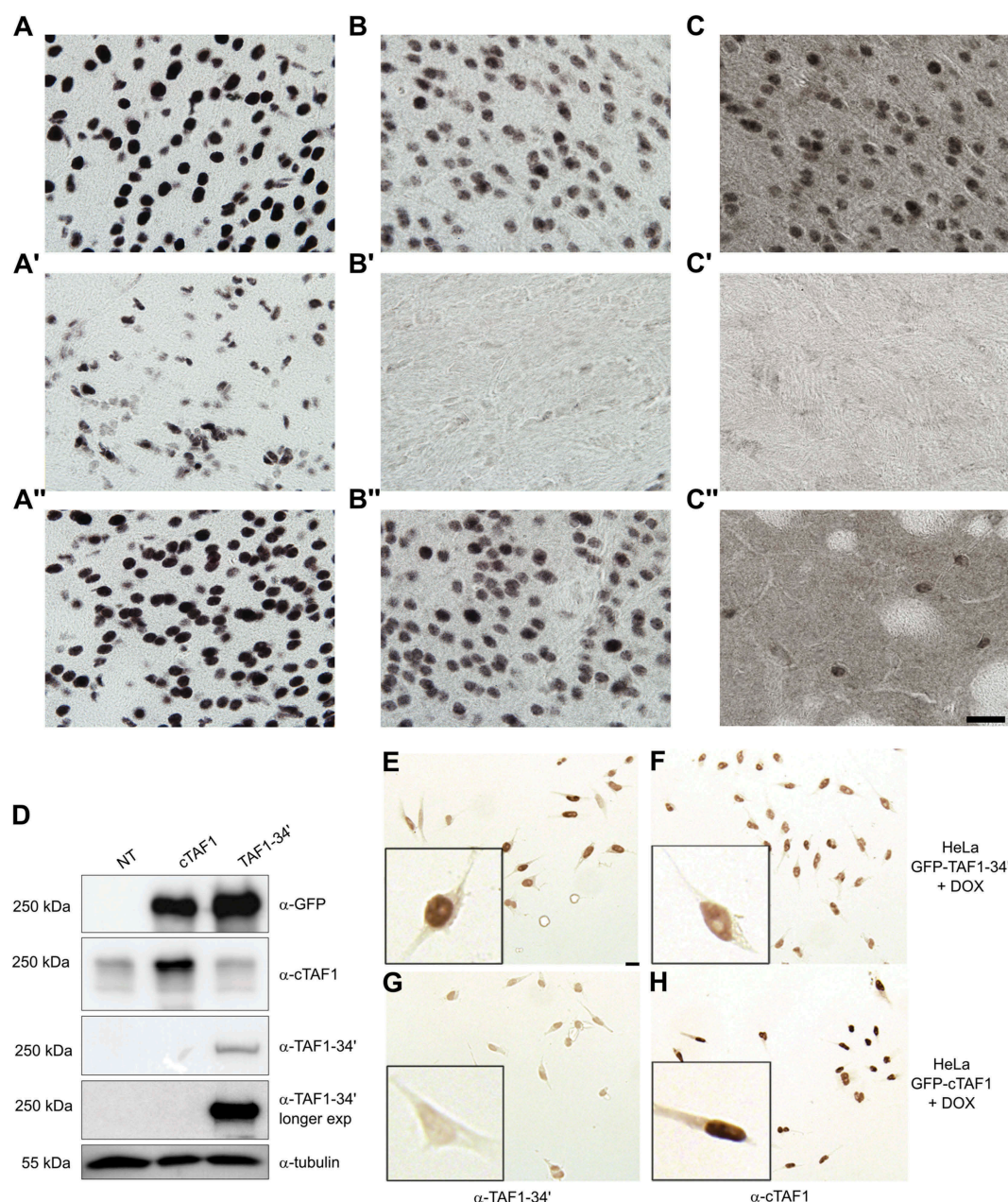
Differential expression of *cTaf1* and the splice isoform *Taf1-34'* mRNAs is detected by *in situ* hybridization and corresponds to expression of *Srrm4*. Coronal left hemisections of the adult mouse brain at the level of the striatum were analysed by BaseScope™ probes for *cTaf1* (A-A'') or *Taf1-34'* (B-B'') and an RNAscope® probe for *Srrm4* (C-C''). Whereas panels A to C display the whole brain cross section, panels A'-C' and A''-C'' show increased magnifications. Black arrows in panels A' and A'' indicate the subventricular zone. Scale bars are 1 mm in A-C and 30 µm in A'-C' and A''-C''. The distribution of the chromogenic signals is designated for illustrative purposes by filled red circles in the low magnification panels. The raw signals are visible within the open red circles in A''-C'' and in the insets of B and C as indicated by red arrows. Cx, CC, Sp and St in panels C and C' indicate respectively cerebral cortex, corpus callosum, septum and striatum regions.

neurons (Fig. 3B). At earlier stages, when neuroepithelial cells and radial glia were undergoing divisions, microexon 34' was not detectable. The expression of *Srrm4* mRNA began in glial daughters and peaked in the early phase of neuronal maturation. As neuronal differentiation proceeded, *Srrm4* expression decreased (Fig. 3C). These results indicate a switch from *cTaf1* to *Taf1-34'* expression in the early phase of neuronal differentiation with a further increase in *Taf1-34'* expression in mature neurons. Interestingly, *Srrm4* expression peaks with the earliest detection of microexon 34', suggesting that *Srrm4* initiates the transition of *cTaf1* to *Taf1-34'* mRNAs during fate determination. We next validated the temporal expression pattern of SRRM4 and the resulting shift between *cTAF1* and *TAF1-34'* in a human differentiation cell system, using the LUHMES cell line. LUHMES are

human mesencephalic cells, which have been immortalized by *v-myc*. This oncogene is repressed in a tetracycline-dependent manner, which induces LUHMES cells to exit cell cycle and differentiate into morphologically and biochemically post-mitotic dopamine-like neurons [15,16]. Immunoblot analysis showed that endogenous SRRM4 was detectable at day 2 of differentiation and its expression was decreasing in later differentiation stages (Fig. 3E). Consistent with SRRM4 expression, the incorporation of microexon 34' into *TAF1* mRNA was observed only in the latest differentiation points (day 4 and day 6) (Fig. 3D).

Taken together, this data suggests that the neuronal SRRM4 splicing factor is the driver of the switch of *cTAF1* to *TAF1-34'* expression, which coincides with the neuronal maturation of mammalian brain cells.





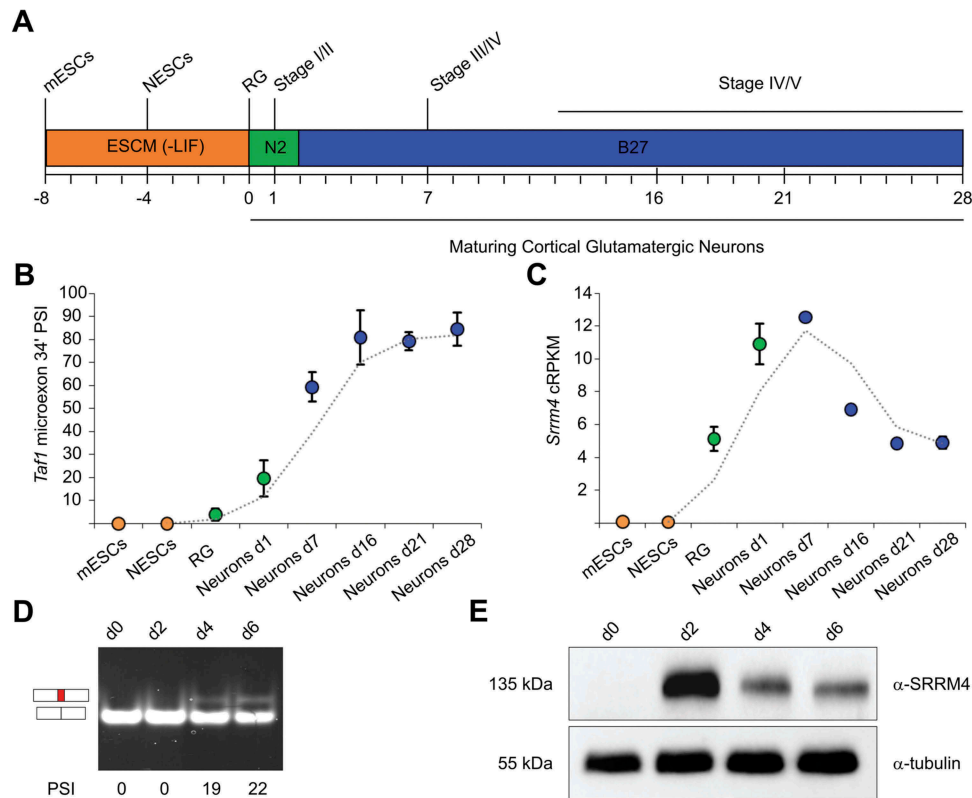
**Figure 2.** Overlapping expression of Taf1-34' and Srrm4 proteins in the mouse brain.

Immunohistochemical detection of cTaf1 (panels A-A''), Taf1-34' (panels B-B'') and Srrm4 (panels C-C'') proteins is shown for cerebral cortex (panels A-C), corpus callosum (A'-C') and striatum (A''-C''). cTaf1 immunoreactivity is evident in neurons of the cortex (A), glial cells in the corpus callosum (A') and neurons in the striatum (A''). Taf1-34' and Srrm4 immunoreactivity is observed in neurons of the cortex (B and C) but not in glial cells of the corpus callosum (B' and C'). Srrm4 expression is enriched in a subset of striatal neurons (C''). Scale bar in C'' is 10  $\mu$ m and applies to all panels. Validation of isoform-specific TAF1 antibodies was performed using 293T cells that were transiently transfected with GFP-cTAF1 or GFP-TAF1-34' cDNAs. Immunoblots of transfected cell lysates were incubated with GFP and cTAF1 monoclonal antibodies or TAF1-34' affinity-purified polyclonal antibodies.  $\beta$ -Tubulin antibodies were used as loading control. NT indicates non-transfected control lysates (panel D). HeLa cells expressing inducible GFP-fusion of cTAF1 or TAF1-34' were analysed by IHC to assess the specificity of isoform-specific TAF1 antibodies. For direct comparison of immunoreactivities, TAF1-34' (E-G) and cTAF1 (F and H) antisera were applied separately to two slides on which two types of HeLa cells had been grown in separate chambers prior to fixation with paraformaldehyde: TAF1-34' cell line induced with DOX (E and F) and cTAF1 cell line induced with DOX (G and H). Immunoreactivity for TAF1-34' was only evident upon DOX induction of the TAF1-34' expressing cell line (E). Baseline immunoreactivity for endogenous cTAF1 is evident (F) with enhanced immunoreactivity upon DOX induction (H). Inserts show magnifications of sampled cells. Images were all taken in parallel with identical camera settings. Scale bar in (E) is 10  $\mu$ m and is applicable to (E-H).

### ***SRRM4 promotes the inclusion of the alternative microexon 34' in TAF1 mRNA***

To investigate the direct involvement of SRRM4 in *TAF1* pre-mRNA splicing, we engineered HeLa cells to conditionally

express an N-terminally GFP-tagged SRRM4. DOX-inducible expression of this transgene was confirmed by immunoblotting and immunofluorescence (Fig. 4A and B). Consistent with recent results [17], the exogenous GFP-SRRM4 localized to the nuclear compartment (Fig. 4C) within the nuclear speckles,



**Figure 3.** Expression of *cTAF1*, *TAF1-34'* and *SRRM4* mRNAs during *in vitro* neuronal differentiation.

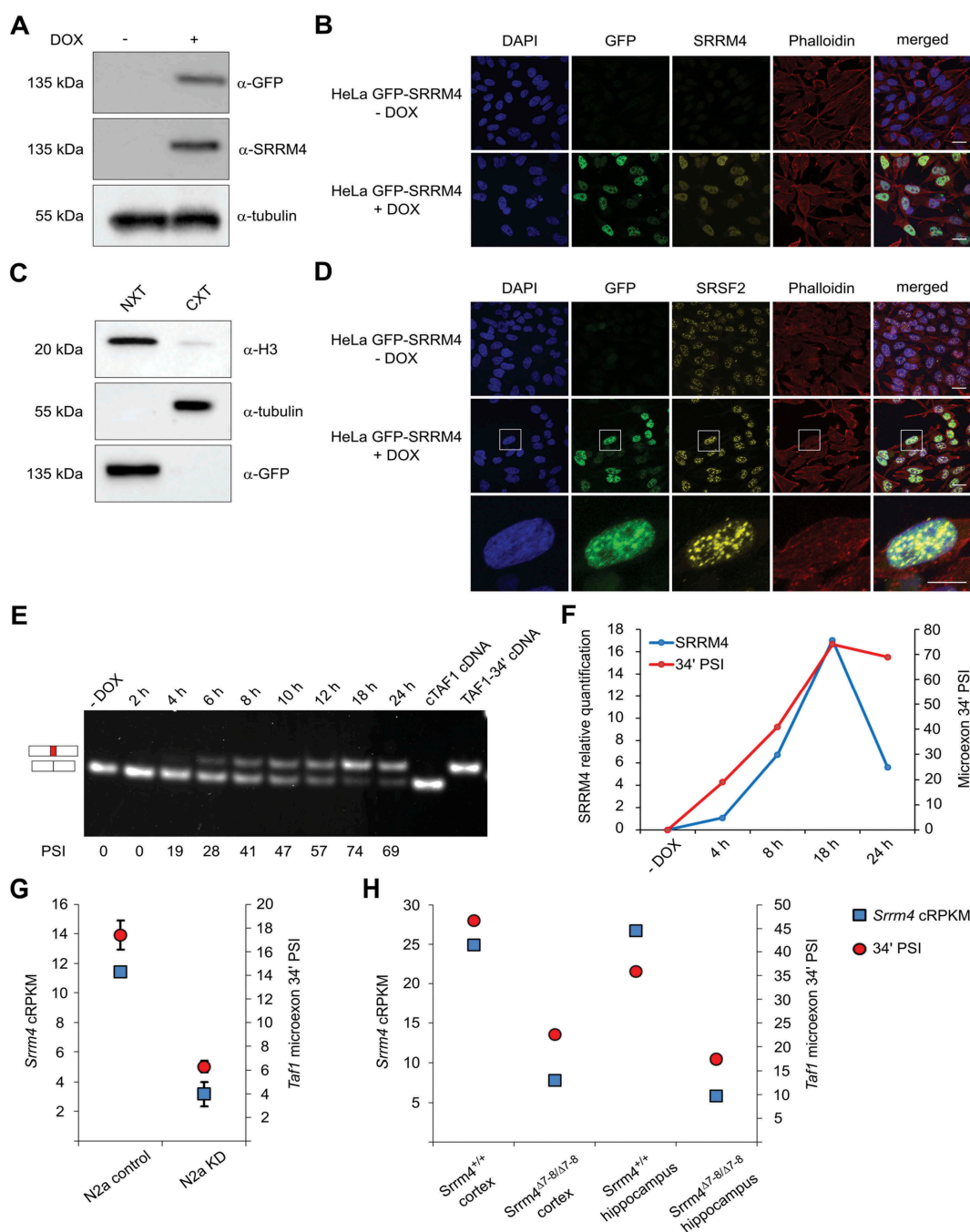
Mouse embryonic stem cells (mESCs) were differentiated towards cortical glutamatergic neurons as depicted in panel A (adapted from [15]). The percentage-spliced-in (PSI) of microexon 34' (B) and *Srrm4* expression level (C) were calculated at different time points (mESCs: day -8; neuroepithelial stem cells – NESC's: day -4; radial glia – RG: day 0; neuronal differentiation stage I-II: day 1; stage III-IV: day 7; stage IV-V: days 16, 21, 28). Dotted lines in panels B and C indicate the data trend line. cRPKM indicates corrected reads per kilo base per million mapped reads. Microexon 34' incorporation in *TAF1* mRNAs (D) correlates with *SRRM4* expression (E) in the LUHMES differentiation assay.

which was confirmed by its co-localization with the SRSF2/SC35 splicing factor (Fig. 4D).

The contribution of SRRM4 in microexon 34' incorporation in *TAF1* mRNA was examined during a time-course induction of GFP-SRRM4 expression in HeLa cells (Supplementary Fig. S3A and S3B). For this analysis, we developed a polyacrylamide gel system to resolve RT-PCR products containing the 6-nt of microexon 34' from *cTAF1*. With this system, we observed that the progressive SRRM4 expression coincided with increased microexon 34' inclusion that reached a PSI of 74% after 18 h of DOX treatment (Fig. 4E). Microexon 34' inclusion lagged only shortly behind GFP-SRRM4 induction, which indicates a rapid turnover of the pool of *TAF1* mRNAs (Fig. 4F). We examined two other microexon splicing events, the incorporation of microexon 8A in *KDM1A* and microexon 16 in *DAAM1* attributed to SRRM4 and observed inclusion of both microexons into their respective mRNAs upon SRRM4 induction in HeLa cells (Supplementary Fig. S3C and S3D).

To determine whether *Srrm4* is required for inclusion of *Taf1* microexon 34' in neuronal cells, we examined RNA-seq data from mouse neuroblastoma N2a cells after siRNA-mediated knock-down of endogenous *Srrm4* expression [2]. The PSI for microexon 34' inclusion in untreated N2a cells was ~18% and was reduced to 6% after *Srrm4* knockdown (Fig. 4G). The interdependency of microexon 34' and *Srrm4* was also confirmed using RNA-seq data from neocortical and hippocampal samples taken from

a conditional *Srrm4* knockout model [3]. These results demonstrate that the reduced expression of *Srrm4* correlates with reduced microexon 34' incorporation in *Taf1* mRNA *in vivo* (Fig. 4H). Besides microexon 34', additional alternative splicing (AS) events have been described for the downstream exons of *TAF1* [6,8,18,19]. To determine whether SRRM4 contributes to these splicing events, *TAF1* mRNAs from GFP-SRRM4-expressing HeLa cells were scanned by RT-PCR using primer pairs spanning exon 30 to the final exon 38. The primers were designed to analyse the splicing events occurring across introns or exons of this region, to detect both alternative exon inclusion and exon skipping. This scanning confirmed that *TAF1* mRNA undergoes several AS events and that only the inclusion of microexon 34' is SRRM4-dependent (Supplementary Fig. S4). In addition, this analysis confirmed that the alternative splicing of microexon 34' and exon 35' are independent events and that the two alternative exons can be part of the same *TAF1* isoform. To determine whether SRRM4 supports *TAF1* microexon 34' inclusion in different cell types, we generated GFP-SRRM4 expressing cells derived from human RPE1 retinal pigmented epithelial cells and human U-2 OS osteosarcoma cells (Fig. 5A to 5D). As with HeLa cells, DOX-induction of GFP-SRRM4 expression in these two non-neuronal backgrounds resulted in the inclusion of microexon 34' in *TAF1* mRNA (Fig. 5E). The PSI values for microexon 34' correlated with GFP-SRRM4 expression levels across different cell types. Compared to HeLa and U-2 OS cells, RPE1 cells expressed



**Figure 4.** SRRM4 promotes the inclusion of the alternative microexon 34' in *TAF1* mRNA.

Immunoblot (A) and immunofluorescence (B) analysis of Flp-In T-REx derivatives of HeLa cells expressing GFP-SRRM4. Transgene expression is verified after DOX induction for 24 h. GFP-SRRM4 is enriched in the nuclear fraction (NXT). Histone H3 was used as nuclear marker while cytoplasmic extract (CXT) was verified using  $\alpha$ -tubulin (C). GFP-SRRM4 resides within the nuclear speckles as revealed by SRSF2 co-staining (D). Scale bars in panels B and D are 10  $\mu$ m. RT-PCR analysis demonstrated the progressive incorporation of microexon 34' in *TAF1* mRNA during SRRM4 induction time curve. Plasmids containing cTAF1 and TAF1-34' were used as size-specific controls. The quantified PSI is depicted below each time lane (E). SRRM4 protein expression correlates with microexon 34' incorporation (F). RNA-seq data from N2a knockdown (KD) experiments and from the *Srm4* knockout model (*Srm4* <sup>$\Delta 7-8/\Delta 7-8$</sup> ) are depicted in panel G and H, respectively. cRPKM indicates corrected reads per kilo base per million mapped reads.

GFP-SRRM4 at lower levels and 50% of *TAF1* mRNAs included microexon 34'. By contrast, HeLa and U-2 OS cells expressed GFP-SRRM4 at similarly high levels and displayed a similar PSI (69% and 76% after 24 h of induction, respectively).

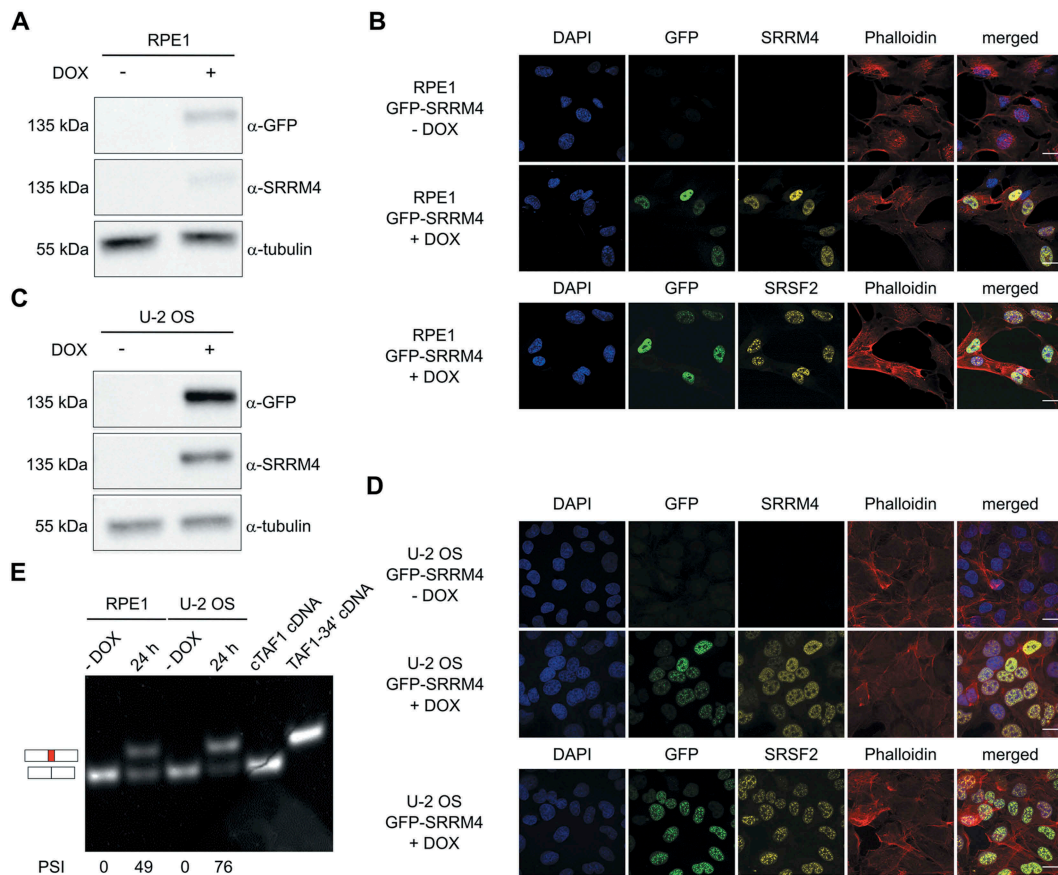
Taken together, these results show that exogenous expression of SRRM4 directs inclusion of microexon 34' in *TAF1* mRNAs. This alternative splicing event can be induced in non-neuronal cell systems, which underlines

the powerful action of SRRM4 in driving microexon 34' inclusion.

### Interactome of ectopically expressed SRRM4 supports its splicing function

Several SRRM4 interactors have been described by mass spectrometry in 293T cells [2] and N2a cells [17]. We examined the





**Figure 5.** SRRM4 promotes the alternative splicing of *TAF1-34'* in different non-neuronal cell lines.

RPE1 and U-2 OS derivative cell lines express GFP-SRRM4 as validated by immunoblot (A and C) and immunofluorescence (B and D). The expression of the transgene is verified after 24 h DOX induction. GFP-SRRM4 resides in the nucleus where it co-localizes with the nuclear speckle marker SRSF2 (B and D, lower panels). In both RPE1 and U-2 OS, the induction of SRRM4 results in microexon 34' incorporation into *TAF1* mRNA. Plasmids containing cTAF1 and TAF1-34' cDNAs were used as controls. The different PSI quantifications are depicted below each lane (E).

hierarchy of SRRM4 interactors by iBAQ-based quantitative mass spectrometry (qMS) of GFP-SRRM4 purified from nuclear extracts of DOX-induced HeLa cells (Supplementary Fig. S3E). This procedure identified U2 snRNP auxiliary splicing factor, composed of the U2AF1 (U2AF35)/U2AF2 (U2AF65), as a major interactor of SRRM4. This complex, which binds to the 3' AG dinucleotide and the polypyrimidine tract element, promotes the recruitment of U2 snRNP to adjacent branch sites [20]. Stoichiometry values indicated that ~40% of GFP-SRRM4 protein forms a stable complex with U2AF1/U2AF2. Among the other GFP-SRRM4 interactors are the RNPS1 and SRSF11 splicing factors (relative stoichiometries of 0.25 and 0.01, respectively), which have been recently identified as co-regulators of SRRM4-dependent microexon splicing [17]. Also consistent with recent results [17], core components of the Exon Junction Complex (EIF4AIII, RBM8A, and MAGOHB) and its auxiliary proteins (PNN, ACIN1, SRRM2 and SAP18) were also identified, and these account for 10-20% of the recovered SRRM4 complexes. Our analysis further confirmed the interaction with FBXW11, which has been proposed recently as a regulator of SRRM4 proteolysis [17]. The list of statistically significant interactors of GFP-SRRM4 in HeLa cell nuclear extract (with relative stoichiometry above 0.01) is provided in Supplementary Table S2.

### UGC motifs upstream of *TAF1* microexon 34' are critical for SRRM4-mediated alternative splicing

A previous study identified UGC-containing motifs as critical SRRM4 binding sites to promote neuronal microexon inclusion [2]. Sequence analysis of *TAF1* intron 34' sequences revealed the presence of two UGC motifs located in the proximity of the 3' splice site of microexon 34' (−18 and −29 nt). We investigated their involvement in microexon 34' incorporation in a minigene reporter assay. The *TAF1* minigene reporter included *TAF1* sequences spanning from exon 33 to the final codon of exon 35. This embeds the sequence of the neuron-specific microexon 34' with the flanking introns 33 and 34. The minigene reporter was designed as an in-frame fusion product with a GFP N-terminal tag to permit GFP-specific enrichment during RT-PCR analysis and to allow the discrimination from endogenous *TAF1* mRNA splicing events (Fig. 6A). *TAF1* isoform-specific antibodies also allowed to examine the cTAF1 to TAF1-34' switch at the protein level. Whereas transfection of the wild-type minigene reporter (MG) in 293T cells resulted only in cTAF1 products, co-transfection of the minigene with GFP-SRRM4 induced the incorporation of microexon 34' sequence as determined by RT-PCR analysis of mRNAs



(Fig. 6B) and by TAF1-34' specific antibodies (Fig. 6C). The two UGC motifs were subsequently mutated into UcC or UaC (Fig. 6D). We found that mutagenesis of a single UGC reduced microexon 34' incorporation by two-fold, whereas double UGC mutants displayed strongly reduced microexon 34' incorporations (average PSI wild-type mini-gene: 58.5% vs average PSI double UGC mutants: 5%) (Fig. 6E-H).

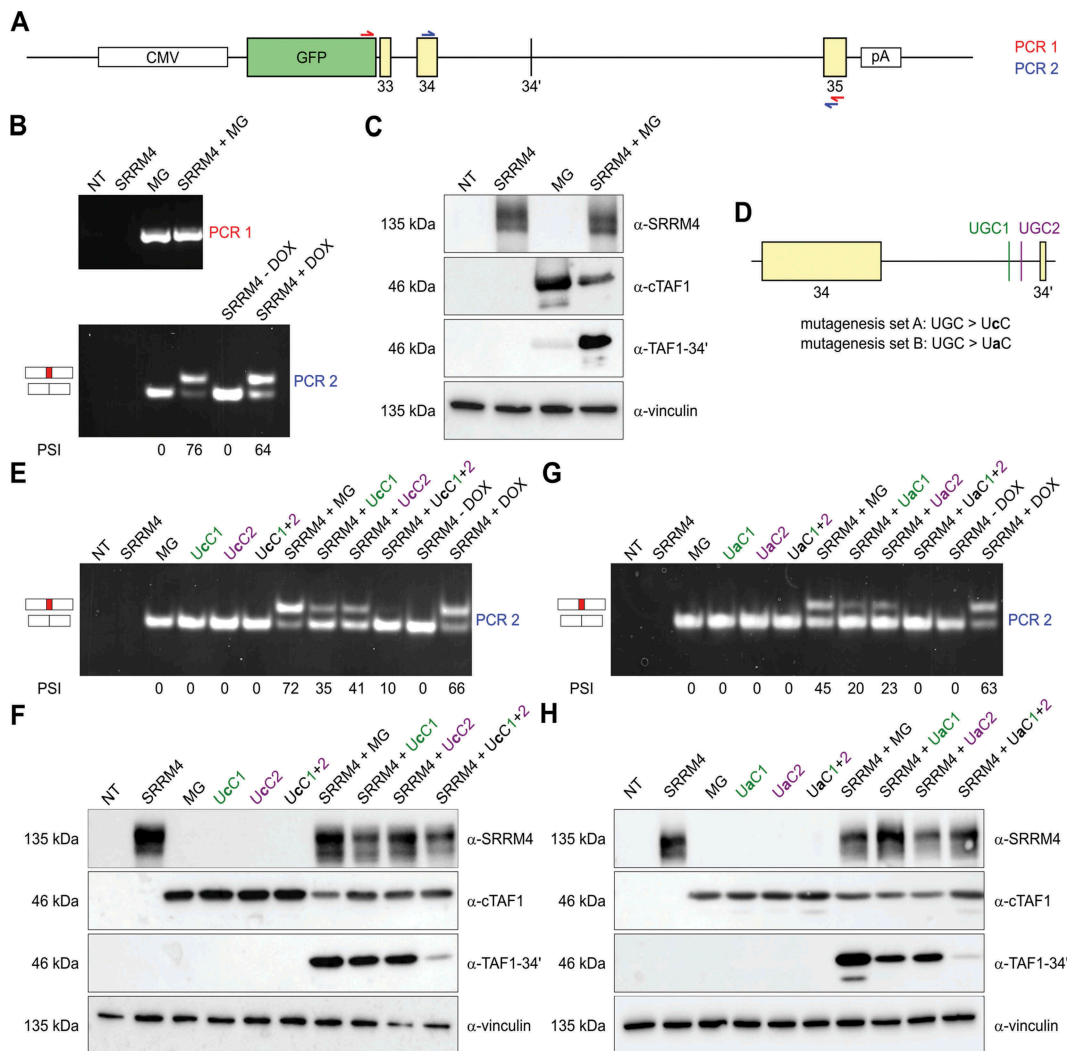
Taken together, the *TAF1* minigene transfection experiments demonstrate that expression of SRRM4 promotes microexon 34' inclusion in *TAF1* mRNA, which is dependent on the two UGC motifs located in poly-pyrimidine tract just upstream the regulated microexon 34'.

## Discussion

Alternative splicing (AS) of mRNAs is one of the central mechanisms contributing to the diversification of the coding capacity of metazoan genomes. The AS regulatory networks

are primarily directed by combinations of tissue-specific splicing factors, which enhance or silence specific splicing choices through their binding to *cis*-acting sequences in pre-mRNAs [21]. Among vertebrate tissues, the nervous system shows the most extensive and conserved AS patterns [22] required to generate protein diversity for specification and function of different cell types [23,24].

In this study, we investigated the spatial and temporal distribution of TAF1-34', the neuron-specific isoform of TAF1. The overlapping patterns of isoform-specific mRNA and protein validate the BaseScope™ method to differentiate mRNAs, which only differ by a 6-nt insertion. The specific staining patterns suggested that canonical and neuron-specific TAF1 isoforms have distinct distributions in specific regions of the brain in both mice and humans. One striking feature of these distributions is their opposing expression patterns in proliferating versus post-mitotic regions. A shift from *cTAF1* to *TAF1-34'* occurs upon neuronal maturation and this shift was confirmed by *in vitro* differentiation of mouse embryonic stem cells and



**Figure 6.** Microexon 34' incorporation is regulated by upstream UGC motifs.

The structure of the *TAF1* minigene (MG) reporter is depicted in panel A. The two sets of primers used to investigate minigene-specific RT-PCR products are depicted in red and blue, respectively. The wild-type minigene was validated using both RT-PCR (B) and immunoblot (C). Two UGC motifs were identified within poly-pyrimidine tract upstream of microexon 34'. Two sets of mutations were generated, substituting the central G of the motifs with either C or A (UGC>UcC; UGC>UaC). Six versions of the minigene reported were created, containing single or double mutations (D). RT-PCR (E and G) and immunoblot (F and H) revealed these sites as critical for microexon 34' incorporation. In panels E and G, the HeLa GFP-SRRM4 cell lines were used as positive control for the assay.

human mesencephalic cells differentiation. The observation that TAF1 and TAF1-34' are expressed in specific brain regions and at defined times of neuronal development suggest that these two isoforms exert distinct functions during development. TAF1 is part of the basal transcription factor TFIID, which represents a ~ 1.5 MDa complex composed of TBP (TATA-binding protein) and 13 TBP-associated factors (TAFs) [25]. Within the complex, TAF1 contacts TBP [26] and TAF7 [27,28] as well as downstream core promoter elements [29–31]. Although TFIID is regarded as a basal transcription factor with ubiquitous expression, tissue-specific TFIID variants have been reported [32]. An example is TAF7L, the testis-specific paralog of TAF7. In late spermatogenesis, while overall TAF7 levels are reduced, TAF7L is expressed to associate in a testis-specific TFIID [33]. An additional testis-specific TAF is TAF1L, the paralog of TAF1. TAF1L can also bind directly to TBP [34] and can replace TAF1 within TFIID (unpublished observation). Tissue-specific functions of TAFs have not only been identified in germ lines, but also in neuronal cells. TAF9B, a paralog of TAF9, promotes alternative gene expression by specifically binding to both promoters and distal enhancers of neuronal genes [35]. These observations indicate that tissue-specific forms of TFIID can display promoter-specificity driving tissue-specific transcription programmes. By analogy, we propose that the TAF1-34' isoform could exert a specific neuronal function. We have found that TAF1-34' efficiently incorporates into the TFIID complex (unpublished observations). This suggests that TAF1-34' could regulate cell-type specific transcription through mediating cell-type specific protein-protein and/or protein-DNA interactions of TFIID. This proposal is in line with observations that neuron-specific microexons encode residues that overlap surface-accessible and looped regions of proteins that often coincide with modular domains involved in protein-protein interactions [1]. This observation strengthens the concept that tissue-specific AS rewires protein-protein interaction networks [36–38]. Interestingly, the two amino acids encoded by microexon 34' are located in the unstructured C-terminal region of TAF1. The addition of an alanine-lysine dipeptide in this region might create a new binding interface or an acceptor for post-translational protein modifications. Additional studies including TAF1-34'-specific proteomics and CHIP-seq, combined with corresponding single-cell RNA-seq profiles for different brain regions, would test these hypotheses.

The alternative expression of TAF1 and TAF1-34' isoforms is restricted to specific cell types in the brain and we show that the neuronal splicing factor SRRM4 directs this switch. Using both ISH and IHC, we could confirm the overlapping expression pattern between *Srrm4* and *Taf1-34'*. Interestingly, while the cortex showed very high correlation (Fig. 3B and 3C), the striatum sample challenged this finding, showing strong immunoreactivity for *Taf1-34'* but a weak *Srrm4* staining (Fig. 2B''-C''). This data could indicate that a low expression of *Srrm4*, not detectable by IHC, is sufficient to promote *Taf1-34'* microexon inclusion. On the other hand, this could also suggest that an intricate network of AS regulation initiated by *Srrm4* is sustained by other splicing factors that have functions overlapping with *Srrm4*. This second hypothesis could be of interest in regards of detection of low levels of *Taf1-34'* in non-neuronal tissues, as striatal muscles (Fig. S1A), but additional experiments would be

required to identify these factors. The involvement of SRRM4 in cTAF1-TAF1-34' switch is demonstrated by the fact that the presence of SRRM4 is sufficient to direct microexon 34' incorporation into *TAF1* mRNA in several non-neuronal cell systems. Consistent with this, the incorporation of microexon 34' into *TAF1* mRNA is strikingly impaired when SRRM4 binding sites located just upstream of microexon 34' are mutated. Reduction of microexon 34' splicing has been linked to XDP [6,39,40]. In order to identify the molecular mechanism linking the pathogenesis of the disease with a defect in TAF1 alternative splicing, it would be important to investigate the possible SVA-induced alteration in the SRRM4-dependent 34' incorporation. In this context, the tools developed in this study will be used to investigate cTAF1, TAF1-34' and SRRM4 expression, both at mRNA and protein levels, in XDP-derived material and XDP post-mortem brains.

Several neurodegenerative diseases that show tissue-specific cell loss are known to be caused by genetic alterations in ubiquitously expressed genes. This disconnection has been a mystery that has incurred numerous hypotheses to account for the cell type-specific vulnerability. Our study combined with others [3,41,42] raises the possibility that microexons that were not previously annotated or evaluated will define tissue-specific isoforms of these 'ubiquitous' disease genes. This concept strongly applies to XDP as it affects *TAF1*, encoding a crucial component for pol II-mediated gene transcription. Our study demonstrates that the splicing factor SRRM4 directs neuron-specific *TAF1* AS profiles, which provides a mechanism to account for the restricted tissue-specificity of XDP pathology. In addition, our results suggest that the neuronal splicing factor SRRM4 shapes TFIID to drive a neuronal-specific transcription programme that is important for mature brain function in vertebrates.

## Materials and methods

### Mice

For histological experiments we used 1 to 4 months old female and male mice of C57BL/6 and FVB genetic backgrounds. Mice were group-housed by sex and kept under standard laboratory conditions. All procedures were approved by the Animal Care and Use Committee at the Massachusetts Institute of Technology (MIT), which is AALAC accredited.

### In situ hybridization on mouse tissue

Mice were deeply anesthetized with isoflurane prior to brain removal and its placement on ice. A thick block containing the striatum was dissected and placed into a cryomold containing cold Optimal Cutting Temperature (OCT) Compound (Tissue-Tek, Sakura Finetek Inc., Kyoto, Japan). The cryomold was then filled with OCT and further cooled, just until opaque, in a bath of methylbutane that was pre-cooled on dry ice. Cryomolds were sealed in plastic bags and stored at -80°C until cut into 10 µm sections on a cryostat. Sections were placed onto positively charged slides (Leica, Wetzlar, Germany) and stored at -80°C until use. For detection of *Srrm4* mRNA, single-plex chromogenic *in situ* hybridization was done as

described in the RNAscope® 2.5 manual from ACDBio (Silicon Valley, CA), with a probe directed to *Srrm4* (cat. #529,068, ACDBio). For detection of microexon 34' in *Taf1-34'*, and the exon 34 to exon 35 junction site in *cTaf1*, we used BaseScope™ (ACDBio) probes (cat. #713,761 for *Taf1-34'* and cat. #713,771 for *cTaf1*) and the provided protocol. Slides were scanned at 40X magnification with an Aperio slide scanner and images were processed in Aperio ImageScope (Leica).

### Immunohistochemistry on mouse tissue and HeLa cells

Mice were deeply anesthetized with isoflurane and trans-cardially perfused with 0.9% saline solution followed by 4% (wt/vol) freshly depolymerized paraformaldehyde in 0.1 M NaKPO<sub>4</sub> buffer as fixative. Dissected brains were post-fixed for 90 min and then were transferred to cryoprotective 25% (vol/vol) glycerol sinking solution for 12–24 hours. Brains were then frozen and cut at 30 µm on a sliding microtome, and sections were placed in 0.1% sodium azide in 0.1 M phosphate buffer solution (PBS) made from NaKPO<sub>4</sub> for storage at 4°C. Sections were treated for antigen retrieval by submerging in 95°C, pH 6.0 unmasking solution (Vector laboratories, Burlingame, CA) for 10 min. See <https://www.protocols.io/view/immunofluorescence-for-free-floating-brain-section-kracv2e> dx.doi.org/10.17504 (2017) for extra details of the immunoreactivity reactions. Briefly, sections were rinsed and treated for 10 min with 3% H<sub>2</sub>O<sub>2</sub> in PBS with 0.2% Triton X-100. Sections were incubated in TSA blocking solution (PerkinElmer, Waltham, MA) and then in primary antiserum overnight at 4°C. Sections were subsequently rinsed and incubated in a biotinylated secondary anti-rabbit antibody and followed by the Vectastain Peroxidase ABC System (Vector Laboratories) according to manufacturer's instructions. Sections were mounted and coverslipped with Eukitt (Electron Microscopy Sciences, Hatfield, PA). Immunohistochemical images were obtained on an Olympus BX61 microscope with a 100 × 1.35 NA oil immersion objective. The same procedure was adopted to perform IHC on HeLa cells.

### Preparation of human brain specimens

Flash frozen human hemispheres from unremarkable donors were acquired from the Netherlands Brain Bank. Frozen hemispheres were dissected into coronal slices in a – 20°C climate control room using a 30 cm cutting edge brain dissecting knife (Fine Scientific Tools, Foster City, CA). Further dissection of selected regions was performed using pre-cooled scalpel blades (Fine Scientific Tools) and 3 mm biopsy punches (Microtech, Fletcher, NC). Frozen tissue was pulverized using a mortar and pestle while submerged in liquid nitrogen.

### Plasmid and cell lines generation

The pcDNA5/FRT/TO/GFP-SRRM4 expression plasmid was generated using the GATEWAY cloning system according to the manufacturer's instructions (Thermo Fisher Scientific, Waltham, MA). The open reading frame of the human

*SRRM4* was amplified from the cDNA clone BC152471 (TransOMIC Technologies, Huntsville, AL) using attB-flanked primers, listed in Supplementary Table S1. The amplicon was cloned into the entry vector pDON201 (BP reaction) and subsequently recombined with the destination vector pcDNA5/FRT/TO/N-GFP (LR reaction) to generate expression clones. The gene of interest was verified using Sanger sequencing in both plasmids. The pcDNA5/FRT/TO/GFP-cTAF1 (NM\_004606) was available in the laboratory and contains two silent changes introducing an *AatII* site for cloning purposes. The pcDNA5/FRT/TO/GFP-TAF1-34' was generated via site-directed mutagenesis. Mutagenesis primers are listed in Supplementary Table S1. The *TAF1* mini-gene was generated using PCR amplification and classical cloning. The portion of the human *TAF1* locus cloned into the pcDNA5/FRT/TO/N-GFP vector has the following GRch38 coordinates: chrX:71,454,175–71,458,366.

The HeLa Flp-In T-REx parental cell line [43] was cultured in Dulbecco's modified Eagle's medium (DMEM) containing 4.5 g/litre of glucose (Lonza, Basel, Switzerland), supplemented with 10% (v/v) foetal bovine serum (Lonza), 200 µg/ml of Zeocin (Thermo Fisher Scientific) and 5 µg/ml of Blasticidin S (Invivogen, San Diego, CA), to select for the FRT site and the Tet repressor, respectively. To generate stable doxycycline (DOX)-inducible HeLa Flp-In T-REx cells expressing GFP-SRRM4, parental cells were seeded in 6-well plates and cultured for 24 h without antibiotics. Parental HeLa cells were then co-transfected with pOG44, an expression vector coding for the Flp recombinase, and the pcDNA5/FRT/TO/GFP-SRRM4, using polyethyleneimine (PEI). 48 h after transfection, the medium was supplemented with 250 µg/ml of hygromycin B (Roche Diagnostics, Basel, Switzerland) and 5 µg/ml of Blasticidin S (Invivogen), to select for the Flp-mediated recombination with the pcDNA5/FRT/TO/GFP-SRRM4 plasmid and for Tet repressor expression, respectively. The cell line was validated using immunoblot and immunofluorescence, after 18–24 h induction with 1 µg/ml DOX (MP Biomedical, Santa Ana, CA).

The procedure described above was also used to generate RPE1 and U-2 OS derivatives expressing pcDNA5/FRT/TO/GFP-SRRM4. Flp-In T-REx derivatives of human RPE1 and U-2 OS cells were kind gifts from Dr. Kops (Hubrecht Institute, Utrecht, The Netherlands) and Dr. Buhl (Harvard University, Cambridge, USA), respectively.

Neuroblastoma cell lines SH-SY5Y were a kind gift from Dr. Timmerman (University of Antwerp, Belgium). The cells were cultured in 1:1 DMEM containing 4.5 g/litre of glucose (Lonza)/F12 supplement (Lonza), supplemented with 10% (v/v) foetal bovine serum (Lonza).

Human embryonic kidney cells 293T were cultured in DMEM containing 4.5 g/litre of glucose (Lonza), supplemented with 10% (v/v) foetal bovine serum (Lonza). Transient transfection was performed using FuGENE HD Transfection Reagent (Promega) following manufacturer's instructions.

Lund human mesencephalic (LUHMES) cells were cultured and differentiated as described in [17]. Briefly, Nunclon™ flasks were pre-coated with 50 µg/ml poly-L-ornithine (Sigma, St. Louis, MO) and 1 µg/ml fibronectin (Sigma) in H<sub>2</sub>O at 37°C overnight, washed once with H<sub>2</sub>O and air-dried. For culturing



of cells proliferation medium with Advanced DMEM/F12 (Gibco, Thermo Fisher Scientific), 1x N-2 supplement (Gibco), 2 mM L-glutamine (Gibco) and 40 ng/ml recombinant basic fibroblast growth factor (R&D Systems, Minneapolis, MN) was used. Differentiation was induced by changing to differentiation medium containing DMEM/F12, 1x N-2 supplement, 2 mM L-glutamine, 1 mM dibutyl cAMP (Sigma), 2 ng/ml recombinant human GDNF (R&D Systems) and 1 µg/ml tetra-cycline (Sigma).

### Immunoblot

HeLa GFP-SRRM4 cells were seeded in 6-well plates at the density of 30,000 cells per well and were induced with 1 µg/ml DOX for 18 to 24 h. Cells were lysed in sample buffer (40 mM Tris-HCl pH 6.8, 1% SDS, 5% glycerol, 0.0125% bromophenol blue, 100 mM DTT), and equal amounts of protein were analysed by SDS-PAGE, followed by immunoblotting. After transfer to a nitrocellulose membrane, proteins were identified with the indicated antibodies and detected using Clarity ECL (Bio-Rad, Hercules, CA) and the ChemiDoc Touch system (Bio-Rad). The quantification of band intensity was performed with the 'mean pixel gray' values of the selected region of interest using ImageJ software after Gaussian blur and background subtraction.

### Immunofluorescence and confocal microscopy

GFP-SRRM4 expressing cells were seeded in 12-well plates containing 16-mm glass coverslips (Marienfeld, Lauda-Königshofen, Germany), at the density of 10,000 cells per well. After DOX induction cells were fixed in 2% paraformaldehyde (Thermo Fisher Scientific) in PBS for 20 min at room temperature. Immunofluorescence was performed using a standard protocol. Primary and secondary antibodies were incubated for 1 h at room temperature. Nuclei were stained with 2 µg/ml DAPI (4,6-diamidino-2-phenylindole, Sigma); the cytoplasm was stained with Alexa Fluor 633 Phalloidin (Thermo Fisher Scientific). Coverslips were mounted on microscopy slides (Roth, Karlsruhe, Germany) using fluorescence mounting medium (Dako, Agilent Technologies, Santa Clara, CA) and kept overnight at 4°C.

Confocal images were taken with the Zeiss LSM710 confocal microscope, using the 63x/1.4 oil plan-apochromat objective, using 405-nm violet diode, 488-nm argon and 561-nm solid state lasers.

### Antibodies

The production of cTAF1- and TAF1-34'-specific antibodies was performed in collaboration with Cambridge Research Biochemical and Abcam. To obtain antisera specific for the cTAF1 and TAF1-34' proteins, rabbits were immunized with peptides (cTAF1: [C]-TPGPYTPQPPDLY-amide or TAF1-34': [C]-TPGPYTPQAKPPDLY-amide) coupled via their N-terminal Cys residue to keyhole limpet haemocyanin (KLH) or bovine serum albumin (BSA). The selected epitopes have 100% identity between human and mouse proteins.

Rabbits were immunized by injections with KLH conjugates followed by booster injections with the BSA conjugates at regular intervals. Antibody responses were monitored by ELISA using the two peptides. Based on these ELISA results, polyclonal sera were collected from the best and most specific responders and affinity-purified using magnetic beads (Dynabeads M-280, ThermoFisher) coated with biotinylated derivatives of the immunized peptides. For immunohistochemistry on mouse tissue and HeLa cells, affinity-purified cTAF1 and TAF1-34' antibodies and SRRM4 antibody [44] were used, followed by biotinylated secondary anti-rabbit antibody (Vector Laboratories, 1:500 dilution).

For immunoblot and immunofluorescence, the following primary antibodies were used: GFP (JL-8, Clontech, Mountain View, CA), SRRM4 [44], α-tubulin (DM1A, CP06, Calbiochem, San Diego, CA), SRSF2 (SC-35, S4045, Sigma), histone H3 (ab1791, Abcam, Cambridge, UK) and vinculin (7F9, sc-73,614, Santa Cruz). Notably, the SRRM4 antibody used was raised against the first 82 amino acids of the mouse protein [44] and the identity between the mouse and the human epitope is 84%. Secondary antibodies used were: α-mouse-HRP conjugate (Bio-Rad) and α-rabbit-HRP conjugate (Bio-Rad), α-rabbit Alexa Fluor 568 (Thermo Fisher Scientific) and the α-mouse Alexa Fluor 568 (Thermo Fisher Scientific).

### Expression analysis with RT-PCR and RT-qPCR

Total RNA was isolated from the GFP-SRRM4 expressing cell lines using the RNeasy kit (Qiagen, Venlo, The Netherlands), including an over-column DNase treatment using RNase-free DNase (Qiagen), according to the manufacturer's instructions. The total RNA isolation from human brain specimens was performed using the RNeasy Lipid Tissue kit (Qiagen), with subsequent DNase treatment performed with the TURBO DNase treatment kit (Thermo Fisher Scientific). RNA integrity was analysed using Bioanalyzer (Agilent Technologies) and RNA Integrity Numbers (RINs) were above 7 for all samples.

Total RNA (1 µg) was used for cDNA synthesis (Superscript II, Thermo Fisher Scientific) using random hexamer primers. *TAF1* and *Lysine Demethylase 1A* (*KDM1A*) amplicons were analysed on a 6% acrylamide gel and *Dishevelled Associated Activator of Morphogenesis 1* (*DAAM1*) amplicon was analysed on a 2.5% agarose gel. Gels were stained with ethidium bromide. RT-PCR was performed using the primer pairs listed in Supplementary Table S1. In the *TAF1* RT-PCRs, plasmids containing *cTAF1* and *TAF1-34'* cDNAs were used as size controls. Percentage-Spliced-In (PSI) was calculated as the fraction of *TAF1* cDNA containing microexon 34' to the sum of the inclusion and exclusion *TAF1* cDNA. The quantification of band intensity was performed using ImageJ as described in the immunoblot paragraph.

RT-qPCR was performed in a CFX384 Real-Time system (Bio-Rad) using iQ SYBR Green Supermix (Bio-Rad). Primers used are listed in Supplementary Table S1. The relative expression level of *SRRM4* mRNA was evaluated with the  $\Delta\Delta C_t$  method, using *ACTB* as reference gene. The reactions were performed in technical triplicates.

## Nuclear extract and GFP affinity purification

Cytoplasmic and nuclear extracts were prepared from GFP-SRRM4 expressing HeLa cells as previously described [45]. After Bradford measurement, 1 mg of nuclear extract was used for GFP affinity purification [45], in the absence of ethidium bromide. The experiment was performed in triplicate with GFP-trap beads (gta-20, Chromotek, Planegg-Martinsried, Germany) and control agarose beads (bab-20, Chromotek). After trypsin-digestion, the peptides were desalted using C18 matrix in StageTips [46]. Peptides were eluted in 80% acetonitrile and concentrated using a SpeedVac. The lyophilized samples were diluted to the desired volume in 0.1% trifluoroacetic acid and analysed by mass spectrometry.

## Quantitative mass spectrometry (qMS)

For LC-MS/MS analysis, peptide samples were analysed by an Orbitrap Q-Exactive Plus (Thermo Fisher Scientific) mass spectrometer coupled to an Easy nano-LC 1000 (Thermo Fisher Scientific) with a flow rate of 300 nl/min. Buffer A was 0.1% (v/v) formic acid and buffer B was 0.1% (v/v) formic acid in 80% acetonitrile. For reversed phase chromatography, a gradient of increasing acetonitrile proportion was applied in combination with a C18 separating column (2 µm particle size, 100 Å pore size, length 150 mm, inner diameter 50 µm, Thermo Fisher Scientific). The mass spectrometer operated in data dependent mode with a maximum of 10 MS/MS scans following each MS1 scan. Peptide identification was performed with the Andromeda search engine using a human proteome database containing reviewed Uniprot sequences without isoforms downloaded from Uniprot 15 April 2016 (human, 20,193 entries). Peptide spectrum matching and relative protein quantitation were performed using MaxQuant v.1.5.2.8 using a false discovery rate (FDR) of 1%. Perseus v.1.6.0.7 was used to filter contaminants and reverse hits. Label-free quantification values were log<sub>2</sub> transformed and groups based on beads type were defined. Identified proteins were accepted when measured three times within at least one combination of beads, resulting in a total of 917 proteins. To determine the relative stoichiometry of the identified interactors, intensity based absolute quantification (iBAQ) values were used [47], after normalizing to the bait protein (GFP-SRRM4). FDR was set at 0.001 and the minimal fold change (s0) was set at 2.

## Data availability

The proteomics data have been deposited to the ProteomeXchange Consortium via the PRIDE [48] partner repository with the dataset identifier PXD009905.

## Keypoints

- cTAF1 and TAF1-34' mRNA and protein distinguish neurogenic from post-mitotic neurons
- Splicing factor SRRM4/nSR100 is required and sufficient to direct microexon 34' splicing into *TAF1* mRNA

- SRRM4-dependent splicing of microexon 34' relies on UGC-containing 3' sequences

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## Disclosure statement

No potential conflict of interest was reported by the authors.

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## ORCID

Frederik M.A. Van Schaik  <http://orcid.org/0000-0001-8924-2986>

H. Th. Marc Timmers  <http://orcid.org/0000-0001-7062-1417>

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