Research Report

Huntington’s Disease Protein Huntingtin Associates with its own mRNA

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Abstract

Background: The Huntington’s disease (HD) protein huntingtin (Htt) plays a role in multiple cellular pathways. Deregulation of one or more of these pathways by the mutant Htt protein has been suggested to contribute to the disease pathogenesis. Our recent discovery-based proteomics studies have uncovered RNA binding proteins and translation factors associated with the endogenous Htt protein purified from mouse brains, suggesting a potential new role for Htt in RNA transport and translation.

Objective: To investigate how Htt might affect RNA metabolism we set out to purify and analyze RNA associated with Htt.

Methods: RNA was extracted from immunopurified Htt-containing protein complexes and analyzed by microarrays and RNA-Seq.

Results: Surprisingly, the most enriched mRNA that co-purified with Htt was Htt mRNA itself. The association of Htt protein and Htt mRNA was detected independent of intact ribosomes suggesting that it is not an RNA undergoing translation. Furthermore, we identified the recently reported mis-spliced Htt mRNA encoding a truncated protein comprised of exon 1 and a portion of the downstream intron in the immunoprecipitates containing mutant Htt protein. We show that Htt protein co-localizes with Htt mRNA and that wild-type Htt reduces expression of a reporter construct harboring the Htt 3’ UTR.

Conclusions: HD protein is found in a complex with its own mRNA and RNA binding proteins and translation factors. Htt may be involved in modulating its expression through post-transcriptional pathways. It is possible that Htt shares mechanistic properties similar to RNA binding proteins such as TDP-43 and FUS implicated in other neurodegenerative diseases.

Keywords: Huntington’s disease, huntingtin exon 1, RNA-protein interaction, mRNA translation, Fmrp

INTRODUCTION

Huntington’s disease (HD) is an inherited neurodegenerative disorder caused by a CAG repeat expansion in the coding region of the Huntington’s disease gene huntingtin (HTT), which results in an increased number of glutamine residues within the N-terminus of the huntingtin protein Htt [1]. The onset of HD symptoms is temporally related to the degree of CAG repeat expansion; longer repeat expansions usually correlate with an earlier appearance of symptoms [2]. HD typically manifests during mid-life and is characterized by chorea, impaired mental abilities, altered personality, and is ultimately fatal. Post-mortem examination of the brains of
affected individuals shows massive cell loss in caudate, putamen, and to a lesser extent the cortex, indicating that these neurons are particularly sensitive to the toxic effects of mutant Htt, despite the near ubiquitous expression of the mutant allele [3]. Currently there is no cure for HD as we have an incomplete understanding of the mechanisms of the disease. We must therefore better understand Htt function if effective therapies or cures are to be developed.

Investigations into the function of wild type and mutant Htt have identified roles for the proteins in transcription, axonal transport, cytoskeletal structure/function, signal transduction, autophagy, and recently post-transcriptional gene expression [4–7]. Htt involvement in gene expression could contribute to the specific pattern of cell loss and symptoms seen in HD if particular groups of genes/messages are more adversely affected over others. These potential changes in gene expression patterns need not be large in magnitude; they are likely small to account for the delay in symptom appearance, and by all accounts apparently normal development is observed in affected individuals. Our recent works have implicated Htt in Processing (P)-body function, RNA transport, and protein synthesis. Htt associates with Ago2 and contributes to post-transcriptional gene silencing [7]. We find mRNA transport inhibited in cultured rat neurons upon knockdown of Htt [6], and Htt co-localizes with mRNAs for β-actin, BDNF, and microtubule-dependent motor proteins Kif5a and Dhc [8, 9]. Furthermore, wild type and mutant Htt co-purify with large numbers of RNA binding proteins in cytoplasmic mouse brain purifications [4]. These observations support the possibility that Htt influences gene expression at steps downstream of transcription.

Given the preponderance of evidence implicating Htt in RNA metabolism, we sought to identify mRNAs that specifically associate with Htt in the mouse brain using 3xFLAG tagged Htt knock-in mice [4]. We reasoned that differences in RNA associations could partially contribute to the tissue specific effects of HD. In our initial microarray analysis of co-purifying mRNAs from FLAG-Htt affinity purifications, we repeatedly identified the Htt mRNA as a co-purifying message. The results were also confirmed by RNA-seq. Here we show that Htt proteins co-localize with Htt mRNA and that wild-type Htt confers down regulation of a reporter gene containing the Htt 3’ UTR sequence.

MATERIALS AND METHODS

Htt affinity purifications and RNA isolation

FLAG-Htt affinity purifications were conducted essentially as described [4] with a few modifications. After the 1400× g spin, Triton X-100 was added to S1 lysates to a final concentration of 0.5%. Lysates were incubated at 4°C for 20 minutes, and centrifuged at 14,000× g for 20 minutes to give a total cytoplasmic lysate. EDTA was added to one half of the lysate at a final concentration of 20 mM or an equivalent volume of lysis buffer, and incubated at 4°C for 20 minutes. Protein concentration was measured by Bradford assay and adjusted to 6 mg/mL with lysis buffer prior to binding to 80 µL (packed bead volume) of Anti-FLAG M2 agarose beads (Sigma) for 3 hours at 4°C with gentle mixing. Beads were washed and eluted as described [4]. Endogenous Htt immunoprecipitations from Neuro-2a cells was performed also as described [4]. Eluted materials from both types of purifications were extracted with 0.5 mL of TRI-Reagent (Sigma) and processed to isolate RNA and proteins as per manufacturer’s instructions. RNA was further cleaned using a Direct-zol RNA miniprep column (Zymo Research). Eluted RNA was concentrated by ethanol precipitation in the presence of 5 µg of linear acrylamide. 20% of eluted proteins were examined for Htt recovery by western blotting and the remainder used to probe for Htt-associated proteins. Mouse brains were obtained in accordance with the guidelines established by the NIH for the care of laboratory animals, and all procedures approved by Institutional Animal Care and Use Committee (Protocol 140308-01).

cDNA synthesis, qPCR oligonucleotides

Total cDNA was generated from 1 µg of inputs and the entirety of immunoprecipitations using a cDNA synthesis kit and the 10x primer mix from USB (Affymetrix). Equal volumes of cDNA synthesis reactions were used in RT-qPCR reactions using the 2X SYBR green master mix from USB (Affymetrix). qPCR data was calculated relative to a Gapdh (Htt IP) or Htt (FLAG IP) standard curves from input cDNAs. All qPCR reactions were performed in triplicate and trends were reproduced with experimental replicates. The oligonucleotides used in the PCR reactions were as follows.
**Htt 3’ UTR**
- Forward: TAGAGCCATCGAAGGACTGCTCC
- Reverse: GCATAACGACGAAAGGC TTGGACAG

**Gapdh**
- Forward: GGTGCTGAGTATGTCGTGA
- Reverse: CGGAGATGATGACCCTTTTG

For 3xFLAG-Htt7Q/7Q mESCs, total RNA and FLAG-IP RNA were extracted with TRI-Reagent. cDNA synthesis and qPCR were carried out as above using the following oligonucleotides.

**Htt exon 1-2**
- Forward: AGGAACCGCTGCACGGA
- Reverse: CTGAGAGACTGTGCCACAAATGTT

**Htt exon 66-67**
- Forward: CCAGCCCATGGGTTCTGCGC
- Reverse: AAGCAGCCTGTGGTATGGACTTCC

**Oct4**
- Forward: GAAGCAGAAGAGGATCACCTT
- Reverse: TTCTTAAGGCTGAGCTGCAAG

**MaoA**
- Forward: GCCCAGTATCACAGGCCAC
- Reverse: CGGGCTTCCAGAACCAAGA

**Immunofluorescence microscopy**

Cells were washed 3x in pre-warmed (37°C) PBS, then fixed in pre-warmed 4% paraformaldehyde for 10 minutes at 37°C. Fixed cells were washed 3x in PBS, then blocked for 1 hour at room temperature in 5% goat serum, 0.25% saponin, in PBS. The visualization of Venus did not require the use of antibodies. All incubations in primary antibodies were carried out overnight at 4°C and secondary incubations done at room temperature for 1 hour. Coverslips were mounted in Dako fluorescent mounting medium. Images were acquired on a Zeiss META510 confocal scanning laser system on an Axiovert 200M microscope using a Plan-Apochromat 100X/1.3 oil-immersion objective lens. Brightness and contrast settings were adjusted image-wide using Adobe Photoshop.

**Antibodies**

For western blotting mouse monoclonal anti-Htt mAB2166 (Millipore) was used at 1:1000, anti-RACK1 (Gnb2l1) (BD Transduction Laboratories) at 1:2500, anti-Rps6 2217 (Cell Signaling Technology) at 1:500, anti-Myo5a (Cell Signaling Technology) at 1:1000, anti-β-tubulin (Covance) 1:1000, anti-β-actin ac15 (Santa Cruz Biotechnology) 1:400, and anti-GFP (Santa Cruz Biotechnology) at 1:500 in TBST. For immunofluorescence, mouse monoclonal anti-FLAG M2 (Sigma) was used at 1:1000, rabbit polyclonal anti-Htt (Enzo Life Sciences) at 1:1000, and anti-Fmr1 (Millipore) mAB2160 at 1:500. For immunoprecipitations 2μg of anti-Htt (Enzo Life Sciences) was used to immunoprecipitate Htt from Neuro-2a cells or 2μg of rabbit IgG (Santa Cruz Biotechnology) as the negative control.

**Plasmids**

The MS2 binding sites of the bacteriophage MS2 protein were cloned downstream of the Renilla luciferase gene in pRL-TK vector (Promega). PCR-cloned Htt 3’ UTR sequences from a mouse genomic clone containing the entire Htt 3’ UTR was inserted downstream of the MS2 binding sites to generate the MS2bs reporter constructs. pNLS-MS2-Venus was obtained from [10] and the FLAG-Ago2 plasmid was obtained from Addgene. Htt 3’ UTR sequences were also subcloned downstream of pEGFP-C1 (Clontech) to generate the GFP reporter constructs.

**Cell culture and transfections**

Mouse Neuro-2a cells were cultured in 10% FBS in DMEM (Mediatech) and transiently transfected using Lipofectamine 2000 (Invitrogen) under recommended conditions. Mouse Htt knock-in striatal precursor lines [11] were cultured in 10% FBS in DMEM and grown at 33°C. 3μg of total plasmid DNA/well of a 6-well plate was used in MS2 transfections: 3μg pNLS-Venue-MS2 in control transfections, 800 ng pNLS-MS2-Venus and 2.2μg pRL-MS2bs-Htt-long 3’ UTR, and 600 ng pNLS-MS2-Venus and 2.4μg pRL-MS2bs-Htt-long 3’ UTR. Transfections were performed for 24 hours prior to fixation. For GFP reporter assays, Neuro-2a cells were split into 6 cm plates. At 24 hours, Htt-17Q or CMV-empty control plasmid was transfected at a concentration of 3μg/plate. At 30 hours, cells were split equal into 4–25 mm plates. After 48 hours, equimolar amounts of GFP reporter were transfected: 1.18μg GFP-SV40-polyA, 2.1μg GFP-Htt-long 3’ UTR, 1.33μg GFP-Htt-short 3’ UTR. Cells were harvested after 72 hours and 20μg of soluble lysates were analyzed by SDS-PAGE followed by immunoblotting with anti-GFP and anti-Htt antibodies and visualized and quantified using Odyssey imager (LI-COR). siRNA transfections
were performed as described. Transfection of rat cortical neurons and co-localization of endogenous Htt with Htt 3’ UTR were performed as previously described [6, 12].

Experiments with mouse embryonic stem cell (mESC) lines

Three lines were used: Wild type, 3xFLAG-Htt7Q/7Q and 3xFLAG-Htt140Q/7Q. mESCs were cultured on gelatin-coated plates in standard ES medium (DMEM supplemented with 15% ES-Cult FBS, 1% sodium pyruvate, 1% NEAA, 1% penicillin-streptomycin, 0.1 mM 2-mercaptoethanol, and 1000 U/ml LIF). Cells were lysed in total lysis buffer [20 mM Tris-HCl (pH 8.0), 137 mM NaCl, 1 mM EDTA, 1% Triton X-100, 10% glycerol, 1.5 mM MgCl2, 1 mM DTT, 1 mM PMSF, and 1X protease inhibitor cocktail]. Sixty milligrams of total protein was immunoprecipitated with anti-FLAG M2 affinity beads (Sigma A2220) for 4 hours on a nutator at 4°C. The beads were first washed with total lysis buffer in a column, and then second wash with total lysis buffer w/o Triton X-100. FLAG-Htt complexes were eluted with 3X FLAG peptide (200 ng/H9262, Sigma F4799), and concentrated by TCA (trichloroacetic acid) overnight. SDS-PAGE was performed by loading 40 μg of total protein for each input sample and 5% of total eluate for each FLAG-IP sample. Membranes were probed using the following primary antibodies: Mouse anti-FLAG-M2 (Sigma F-1804), mouse anti-Htt2166 (Millipore MAB2166) and mouse anti-polyQ (clone 3B5H10, Sigma P1874). IRDye 800CW goat anti-mouse secondary antibody was used (LI-COR Biosciences). Fluorescent signal was scanned using the LI-COR Odyssey machine.

RNA-Seq analysis

RNA-Seq libraries were made using the SMARTer Ultra Low RNA kit for Illumina (Clontech, Cat # 634935 and 634936), using 1 to 2 ng of RNA. Libraries were run on an Illumina 2500, using a paired end 50 protocol. Sequencing reads were demultiplexed and converted to FASTQ format using Illumina Bcl2FastQ software. Reads were aligned to the mouse genome (build mm10/GRCm38) using the splice-aware STAR aligner [13]. PCR duplicates were removed using the Picard toolkit (http://broadinstitute.github.io/picard). HTSeq package [14] was utilized to generate counts for each gene and exon based on number of aligned reads overlapping them. These counts were used to test for differential exon usage using negative binomial generalized linear models and for visualization as implemented by the DEXSeq R package [15].

RESULTS

Purified endogenous Htt complex contains its own mRNA

In previous studies we utilized FLAG-tagged wild type (F7Q/F7Q) and mutant (F140Q/+ Htt) knock-in mice [16] to isolate Htt-containing protein complexes from juvenile mouse brains. Co-purifying proteins identified by mass spectrometry revealed many RNA binding proteins and translation factors, suggesting a new role for Htt in post-transcriptional gene regulatory pathways [4]. The finding prompted us to analyze RNA that might be present in the Htt protein complexes. Surprisingly, microarray and RNA-Seq analyses of co-purifying mRNAs consistently identified the Htt mRNA as the most enriched RNA in our FLAG-Htt purifications from mouse brains compared with the negative control (FLAG-IP eluates of normal mouse brains expressing no FLAG-tagged proteins). Multiple RT-qPCR experiments on co-eluted RNA from FLAG affinity purifications confirmed the drastic enrichment of Htt mRNA in F7Q and F140Q/+ purifications compared with the control. Comparison of the recovery of Htt mRNA to other mRNAs (Oct 4, MaoA, as examples) in the FLAG-IP eluates clearly demonstrated selective retention of the Htt mRNA in the FLAG immunoprecipitates (Supplementary Figure 1). No other mRNAs showed enrichment above the levels found in the control immunoprecipitate, which represented non-specific binding. We wondered whether this interaction might be an artifact of the purification scheme used to purify Htt as the FLAG tag is positioned at the N-terminus of wild type and mutant alleles. It is possible that our Htt purifications recovered Htt mRNA co-translationally; i.e., incompletely translated FLAG-Htt was immunoprecipitated and as such was associated with the Htt mRNA solely through the whole translational apparatus. To rule out this possibility, we dissociated ribosomes prior to FLAG affinity purification by the addition of EDTA to total cytoplasmic lysates [17]. Addition of EDTA did not significantly affect the amount of FLAG-Htt recovered from F7Q affinity purifications, but reduced F140Q/+ recovery by approximately 50% (Fig. 1a, lanes 13–16). However, EDTA pre-treatment nearly
completely abolished the wild type and mutant Htt interactions with ribosome-associated proteins Rps6 and Gnb2l1 (Rack1) (Fig. 1a, lanes 13–16). In contrast, the Htt-Myo5a interaction was not completely abolished by this treatment. These data show that EDTA addition does not prevent FLAG-mediated purification of FLAG-Htt and that interactions of Htt with ribosomal proteins is largely dependent upon the integrity of the whole ribosome.

We next measured the levels of Htt mRNA present in FLAG-Htt purifications by quantifying it relative to the amount of Htt mRNA present in 1 μg of total input RNA, which is expressed as a numerical ratio (Fig. 1b). FLAG IP of both wild type (F7Q/F7Q) and mutant (F140Q/+) Htt protein recovered significant amounts of Htt mRNA compared with negative control. Although addition of EDTA drastically reduced the enrichment ratio of Htt mRNA for both wild type and mutant, these ratios were still 12 and 24 times greater than that of the negative control, for wild type and mutant, respectively (Fig. 1b, EDTA-treated). These data show that a pool of the Htt protein-Htt mRNA association occurs independently of ribosomal integrity.
Furthermore, the enrichment of Htt mRNA in EDTA-treated samples compared with the negative control argues against the possibility that the Htt protein-Htt mRNA association is an experimental artifact. To further substantiate the Htt protein-Htt mRNA interaction, we immunoprecipitated endogenous Htt from Neuro-2a cells and measured the amount of Htt 3’ UTR present within control (IgG) and Htt immunoprecipitates (Htt) by RT-qPCR. Htt immunoprecipitates contained 3 times as much Htt 3’ UTR as the negative control, indicating that Htt protein-Htt mRNA association does not occur through the FLAG tag.

**Htt protein co-localizes with Htt mRNA in neuronal cells**

We have previously shown that Htt co-localizes with the 3’ UTR of β-actin and Bdnf mRNA in cultured rat neurons [8]. We hypothesized that the 3’ UTR of Htt is likely to contain element(s) responsible for associating with Htt protein. To confirm this possibility we examined the location of Htt with respect to a reporter construct harboring the short and long versions of the Htt 3’ UTR in a mouse immortalized striatal precursor cell lines expressing wild-type Htt (STHdhQ/7Q) or mutant Htt (STHdh111Q/111Q) [11]. Our assays employed the bipartite MS2 reporter system [10, 18] to visualize Htt protein with respect to a transfected mRNA containing the short and long versions of the Htt 3’ UTR (Fig. 2a), preceded by the Renilla luciferase coding sequence and 24 tandem copies of the bacteriophage MS2 binding sequence. Control experiments indicated that Venus signal was primarily confined to the nucleus in wild-type and mutant STHdh cells transfected with the NLS-MS2-Venus reporter alone (Fig. 2b, Venus panels), indicating that the presence of the NLS on the Venus reporter confined Venus to the nucleus. In contrast, wild-type cells (Fig. 2b, STSdhQ/7Q) expressing both the NLS-MS2-Venus and MS2 binding site-tagged Htt short 3’ UTR displayed numerous large and small cytoplasmic puncta. Many of these cytoplasmic puncta co-localized with wild-type Htt (Fig. 2b, short UTR, arrowheads), evidenced by the appearance of yellow spots in the merged image. A recent cross-linking immunoprecipitation study of fragile-X mental retardation protein (Fmrp) identified Htt RNA as a target of Fmrp binding in mouse brains [17]. Consistent with this finding, we found co-localization of Fmrp with the Htt 3’ UTR in this assay (Fig. 2b, short UTR, blue-green spots). Furthermore, we observed several Htt short 3’ UTR puncta, which were labeled for both Htt and Fmrp, suggesting that both proteins are capable of associating with the Htt short 3’ UTR (Fig. 2b, upper panel, short UTR, arrow). Not surprisingly, wild-type cells expressing NLS-MS2-Venus and MS2 binding site-tagged Htt long 3’ UTR constructs also displayed numerous cytoplasmic puncta, which also stained positively for Htt and Fmrp (Fig. 2b, long UTR, arrow and arrowhead). These data demonstrate that the 3’ UTR of Htt is sufficient to attract Htt protein to an exogenous mRNA in wild-type cells.

Mutant cells (Fig. 2b, STSdh111Q/111Q) expressing NLS-MS2-Venus and MS2 binding site-tagged Htt short 3’ UTR also displayed cytoplasmic distribution of Venus signal, although the puncta were often larger and more concentrated than those observed in wild-type cells. Like wild-type cells, Htt short 3’ UTR as reported by Venus signal was frequently found to co-localize with both Fmrp and Htt, particularly at the periphery of the cell. This peripheral localization of all three elements was also observed in mutant cells in which the localization of the long version of the Htt 3’ UTR was monitored (Fig. 2b, long UTR). Finally, we transfected MS2 binding site-tagged Htt long 3’ UTR into rat cortical neurons and demonstrated co-localization with endogenous rat Htt by immunofluorescence (Fig. 2c).

**Htt represses gene expression through its 3’ UTR sequence**

Next we sought to determine the effect that the 3’ UTR of Htt might have on the expression of a linked GFP reporter. To do this, we generated GFP expression constructs containing either the SV40 polyadenylation signal alone, the short Htt 3’ UTR, or the long Htt 3’ UTR (Fig. 3a). These constructs were transfected into Neuro-2a cells at equal molar ratios and GFP expression levels monitored by western blotting. Addition of the Htt short 3’ UTR to the GFP expression construct led to a 57% reduction in GFP protein (Fig. 3b), while the long 3’ UTR had an even larger inhibitory effect, reducing GFP expression by 95% compared with the empty control. These data agree with the results from [19] and support an inhibitory role for the 3’ UTR of Htt on protein expression.

Given that Htt likely associates with its own mRNA and that much of this association is dependent upon the presence of intact ribosomes (Fig. 1a, b), we...
Fig. 2. MS2-tagged Htt 3’ UTR co-localizes with Htt in wild type and mutant Htt striatal precursor cell lines. a) Schematic shows the elements of the bipartite MS2 reporter system containing short or long Htt 3’ UTR sequence. Control experiments express NLS-MS2-Venus alone. b) Htt and Fmrp co-localize with Htt short and long reporters in wild-type (STHdh7Q/7Q) and mutant (STHdh111Q/111Q) striatal precursor cells. Merged images and individual channels for enlarged regions (boxes) are shown for each experimental combination. The arrows highlight instances of all three co-localization events, while the arrowhead shows Htt and Htt 3’ UTR co-localization. Scale bars are 10 μm. c) Endogenous Htt co-localizes with long Htt 3’ UTR in primary rat cortical neurons.

wanted to determine the effect that overexpression of Htt had on the levels of GFP expression in the context of the different Htt 3’ UTRs. Overexpression of full-length wild-type Htt (Htt17Q) led to a nearly 50% decrease in the levels of GFP-Htt short when compared with the control without exogenous
Fig. 3. The Htt 3' UTR reduces the expression of a GFP reporter in cis and Htt expression affects the levels of the GFP-Htt 3' UTR in trans. 

(a) Schematic showing the design of GFP reporter constructs. Control (empty) GFP reporter contains the SV40 polyadenylation element alone that is present in pEGFP-C1. The “short” Htt 3’ UTR GFP reporter contains the short version of the 3’ UTR and similarly the “long” Htt 3’ UTR GFP reporter contains the long version of the 3’ UTR. Putative polyadenylation signals are labeled in red text and the engineered stop codon immediately after GFP coding sequence is also shown.

(b) Addition of the long Htt 3’ UTR to the GFP reporter substantially reduces expression of GFP while the short version expresses at half the level of the control. Western blots show N2a cell experimental triplicates transfected with the indicated reporter constructs probed with antibodies to GFP and actin (loading control). The average intensity of GFP bands relative to the empty control are quantified in the bar graph and normalized to actin intensity. *P < 0.05, t-test.

(c) Overexpression of wild-type Htt (Htt17Q) acts to further reduce the levels of the GFP reporter harboring the Htt 3’ UTRs. Western blots show N2a cells with (+Htt17Q) or without (Control) transfected wild-type Htt expression in the context of the expression of the indicated GFP reporter constructs. Blots were probed with antibodies to the indicated proteins and done in experimental triplicate. The graph shows the levels of GFP reporters relative to the sample transfected with vector alone (Control). *P = 0.05, t-test.

(d) Depletion of Htt with siRNA enhances GFP reporter harboring the Htt 3’ UTR. Experiments were performed similar to c) except endogenous Htt was depleted by transfecting a mixture of siRNA to Htt. The graph shows the levels of GFP reporters relative to the sample transfected with a mixture of siRNA containing non-targeting sequences. *P = 0.1, t-test.
Htt expression (Fig. 3c). The already low levels of GFP-Htt long did not change with Htt overexpression.

To further test the possibility that Htt acts to negatively regulate the levels of protein produced from mRNA containing the Htt 3’ UTR, we examined the effects of Htt knockdown on the expression of the GFP reporter constructs. siRNA-mediated knockdown of Htt led to a nearly 10-fold decrease in the levels of Htt compared with the control knockdown (Fig. 3d, Htt level), and a 50% increase in GFP-Htt short, and a 40% increase in GFP-Htt long expression levels compared with their levels in the control knockdown (Fig. 3d, GFP level). These results suggest that Htt has a role in the maintenance of its protein levels through the 3’ UTR.

RNA-Seq analysis of Htt co-purifying RNA identifies mis-spliced Htt exon 1 mRNA

We performed RNA-Seq analysis of the same set of FLAG-immunoprecipitated samples from mouse brains previously subjected to microarray analysis. We indeed detected Htt mRNA sequences across the entire length of the gene at levels orders of magnitude greater than in the control immunoprecipitate (from normal mouse brains expressing no FLAG-tagged proteins to control for non-specific binding to the FLAG antibody) in both wild type and mutant Htt immunoprecipitates (Fig. 4a). Furthermore, analysis of sequence reads in FLAG immunoprecipitates from F140Q/+ brains showed presence of nearly equal number of reads corresponding to endogenous wild-type mouse allele and FLAG-tagged knock-in allele, which can be distinguished by a short human sequence present in the knock-in allele (data not shown). This provides further compelling evidence that Htt protein-Htt mRNA association does not occur through the FLAG tag or through pull down of nascent FLAG-Htt polypeptide. A recent report identified the presence of mis-spliced Htt mRNA containing the sequence of Htt exon 1 and beginning of intron 1 specifically in cells and tissues expressing mutant Htt protein [20]. This prompted us to look for mRNA corresponding to Htt intron 1. We indeed detected sequence reads from intron 1 only in Htt immunoprecipitates obtained from 14Q-expressing mouse brains (Fig. 4b, c). These results indicate that Htt protein complex contains both full-length and mis-spliced Htt mRNA.

Consistent with the above results, RNA-Seq analysis of FLAG-Htt140Q immunoprecipitates from mouse embryonic stem cells expressing the FLAG-tagged protein were also found to contain full-length and exon1 Htt mRNA (data not shown). Furthermore, immunoblotting of the FLAG-Htt immunoprecipitates demonstrated the presence of a polypeptide migrating at ~80kDa (aberrant migration due to expanded polyQ sequence) detected by α-FLAG and α-polyQ antibodies (Fig. 5a, c), but not α-Htt (2166) monoclonal antibody whose epitope is located C-terminal to the region encoded by exon 1 (Fig. 5b). This polypeptide was found only in cells expressing mutant Htt, confirming the existence of the N-terminal mutant Htt fragment encoded by the mis-spliced Htt mRNA.

DISCUSSION

Here we have shown that Htt associates with its own mRNA, possibly through the 3’ UTR. We do not yet know whether the interaction is direct or through other RNA binding proteins with which Htt co-purifies. This finding has interesting parallels to the discovery that the RNA binding protein TDP-43 regulates the levels of its own mRNA through the 3’ UTR to keep them within a certain acceptable threshold [21], thereby preventing the toxic accumulation of the protein [22]. TDP-43 aggregation is found in amyotrophic lateral sclerosis (ALS) and Huntington’s disease [23, 24]. Mutations in TDP-43 are associated with both familial and sporadic ALS [25, 26]. The 3’ UTR of TDP-43 is conspicuously long and conserved [21], and overexpression of wild-type TDP-43 is sufficient to kill cells [22]. Like TDP-43, mutant Htt causes tissue-specific cell death and dysfunction, mutant Htt forms intracellular aggregates [3], the 3’ UTR of Htt can be quite long in neurons [27, 28], and Htt seems to have some role in controlling the levels of its own mRNA.

The Htt 3’ UTR contains many putative miRNA binding sites and as one would predict, the longer version contains more miRNA binding sites than the shorter version. Exogenous expression of miRNAs capable of binding to the Htt 3’ UTR reduces the expression of a luciferase reporter fused to said UTR [19]. We found that a reporter mRNA harboring the Htt 3’ UTR, could localize to cytoplasmic concentrations of Ago2 (data not shown) and Fmrp, which likely represent stress granules or P-bodies [29]. Similarly, we found that the fusion of the Htt 3’ UTR to a GFP reporter acted to substantially reduce the levels of GFP protein. Overexpression of wild-type Htt protein further decreased the GFP levels. We also found
Fig. 4. RNA-Seq analysis of Htt immunoprecipitates reveals full-length Htt mRNA and mutant Htt-specific aberrantly spliced Htt mRNA associated with Htt protein. 

a) Differential expression of Htt exons. The plot shows the fitted expression values of each of the exons of Htt for each of the three conditions based on estimated dispersions as calculated by the statistical method DEXSeq. 

b) Read coverage for Htt exon 1, intron 1, and exon 2. Coverage was normalized relative to the total number of mapped reads for each sample. Same scale was used for graphing all samples to illustrate relative expression levels. 

c) Read coverage for Htt exon 1, and the beginning of intron 1. Coverage was normalized relative to the total number of mapped reads for each sample. Same scale was used for graphing all samples to illustrate relative expression levels.
that reduced Htt protein levels (by siRNA) acted to enhance the protein levels of a GFP reporter fused to the Htt 3’ UTR. It is possible that Htt negatively influences the translation of mRNAs containing the Htt 3’ UTR. However, given the complexity of biological processes, it is most likely that like TDP-43, multiple mechanisms exist to keep Htt expression at an appropriate level.

It was recently reported that a mis-spliced mRNA encoding the N-terminal Htt exon 1 fragment is present specifically in cells and tissues expressing mutant Htt protein [20]. This prompted us to search for mRNAs mapping to introns of the Htt gene in our immunoprecipitates containing Htt in addition to full-length Htt mRNA that we first identified. Indeed, we detected both the mis-spliced mRNA containing the exon 1 sequence and its encoded polypeptide only in the immunoprecipitates containing the mutant Htt protein, confirming the above report. This finding raises the question of how Htt complex associates with both mis-spliced and full-length Htt mRNA since the two messages differ in their 3’ UTR sequence. A motif search program (MEME) identified several short sequences in the 3’ UTR from both mRNAs that included a putative binding site for CELF4, a neuronal RNA binding protein reported to bind to 15%–20% of the brain transcriptome [30]. CELF4 binds specifically in the 3’ UTR of target mRNAs enriched for synaptic functions. Like Htt, CELF4 is found in RNA granules suggesting a potential link to the Htt complex. Since the full-length and mis-spliced mRNAs have a common 5’ UTR it is possible that Htt protein associates with both mRNAs through this region. We observed co-purification of wild type Htt protein with the full-length wild type Htt mRNA. We also found mutant Htt protein to co-purify with the full-length and mis-spliced Htt mRNA. The two mRNAs might be differentially regulated by wild type or mutant Htt protein leading to potential changes in their levels thus contributing to HD pathogenesis.

The new discovery has raised a potentially significant role of Htt exon 1 in the pathogenesis of HD. Transgenic mice expressing Htt exon 1 with expanded polyQ sequence were made decades ago and have served as important mouse models of HD [31]. These animals show rapid progression to neurodegeneration and die within 3 months suggesting that mutant Htt exon 1 encoded polypeptide is highly toxic. By contrast, knock-in animals or BAC transgenic ani-
mals expressing full-length mutant Htt show slower progression to disease and live nearly 2 years [32]. The exon 1 Htt model has been thought of as an acute case of HD; however, given the new finding, toxicity associated with the N-terminal polypeptide albeit at low levels could contribute to the disease progression in HD. Potential role for mutant exon 1 mRNA associated with the Htt complex remains to be investigated.

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CONFLICT OF INTEREST

The authors have no conflict of interest to report.

SUPPLEMENTARY MATERIAL

The supplementary table and figure are available in the electronic version of this article: http://dx.doi.org/10.3233/JHD-150177.

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