Global Patterns of Tissue-Specific Alternative Polyadenylation in Drosophila

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SUMMARY

We analyzed the usage and consequences of alternative cleavage and polyadenylation (APA) in Drosophila melanogaster by using >1 billion reads of stranded mRNA-seq across a variety of dissected tissues. Beyond demonstrating that a majority of fly transcripts are subject to APA, we observed broad trends for 3’ untranslated region (UTR) shortening in the testis and lengthening in the central nervous system (CNS); the latter included hundreds of unannotated extensions ranging up to 18 kb. Extensive northern analyses validated the accumulation of full-length neural extended transcripts, and in situ hybridization indicated their spatial restriction to the CNS. Genes encoding RNA binding proteins (RBPs) and transcription factors were preferentially subject to 3’ UTR extensions. Motif analysis indicated enrichment of miRNA and RBP sites in the neuronal extensions, and their termini were enriched in canonical cis elements that promote cleavage and polyadenylation. Altogether, we reveal broad tissue-specific patterns of APA in Drosophila and transcripts with unprecedented 3’ UTR length in the nervous system.

INTRODUCTION

Alternative cleavage and polyadenylation (APA) has substantial impact on transcript diversity and function (Di Giammartino et al., 2011; Licatalosi and Darnell, 2010). APA can affect coding exons and protein sequence, but it more commonly affects the extent of 3’ untranslated region (UTR) sequence. 3’ UTRs harbor much of the cis-regulatory information for posttranscriptional regulation, including binding sites for microRNAs (miRNAs) and diverse RNA binding proteins (RBPs) (Flynt and Lai, 2008; St Johnston, 2005). Collectively, these can either positively or negatively regulate transcript stability or translational efficiency, as well as influence transcript localization. Consequently, shortening or lengthening of 3’ UTRs can substantially alter gene function across isoforms. For example, loss of distal 3’ UTR sequences allows certain oncogenes to evade repression by miRNAs, thereby potentiating their activity (May and Bartel, 2009).

APA has recently been appreciated as a global phenomenon that can be broadly modulated under different cell conditions. This was originally inferred from analysis of cDNA libraries (Tian et al., 2005; Zhang et al., 2005), and examined much more deeply with the use of genome-wide techniques such as tiling microarrays (Ji et al., 2009; Ji and Tian, 2009; Sandberg et al., 2008), mRNA sequencing (RNA-seq) (Mangone et al., 2010; Ozsolak et al., 2010), and sequencing of transcript 3’ ends (Jan et al., 2011; Mangone et al., 2010; Shepard et al., 2011). Several trends emerged from such studies, including that cells proliferating upon T cell activation express shorter 3’ UTRs (Sandberg et al., 2008); that cell transformation may be correlated with 3’ UTR shortening independently of proliferation rate (Fu et al., 2011; May and Bartel, 2009); that a general transition to shorter APA isoforms is observed during reprogramming of somatic cells into iPSCs (Ji and Tian, 2009); that global lengthening of 3’ UTRs occurs during mouse embryonic development and differentiation of C2C12 myocytes (Ji et al., 2009); reciprocally, that 3’ UTRs generally shorten during Caenorhabditis elegans development (Mangone et al., 2010); and that 3’ UTRs in mammalian neurons exhibit a broad trend for lengthening (Shepard et al., 2011).

To date, 3’ UTR diversity at the genome-wide scale has received relatively little attention in Drosophila melanogaster, as compared to other well-studied eukaryotes. In this study, we use strand-specific RNA-seq across a panel of dissected tissues to reveal broad usage of APA in Drosophila. Notably,
we identify large cohorts of genes that exhibit 3' UTR shortening in the testis and 3' UTR lengthening in the central nervous system (CNS), and we provide extensive experimental confirmation of these APA events by using qPCR, northern analysis, and in situ hybridization. The usage of distal APA sites expands the transcript models of hundreds of neural genes and frequently generates extremely long 3' UTRs, raising unanticipated complexity in their posttranscriptional regulation.

RESULTS

Analysis of D. melanogaster 3' UTRs through Stranded mRNA Sequencing

As part of our efforts to annotate the function of every base in the Drosophila genome (modENCODE, 2010), we have conducted extensive tiling microarray and mRNA-sequencing studies (Cherbas et al., 2011; Graveley et al., 2011). Although powerful, these strategies were limited in that the transcribed strand of origin was not captured. In the current phase of the project, we switched to a stranded mRNA-seq protocol that preserves this information, and we generated libraries from 29 dissected tissues and 25 tissue culture cell lines (J.B.B. et al., unpublished information, and we generated libraries from 29 dissected organs (e.g., brain, larval CNS, and eye). In addition, data from poly(A)-spanning reads showed that these various neural tissues exhibited 3' UTRs with median lengths that were 25%–40% longer than FlyBase gene models (Figure 1C, red asterisks).

Table 1. Stranded RNA-Seq Library Read and Mapping Summaries

<table>
<thead>
<tr>
<th>Sample</th>
<th>Uniquely Mapped Reads:</th>
<th>Uniquely Mapped Reads:</th>
<th>Total Uniquely Mapped Reads</th>
<th>modENCODE/NCBI-SRA Accession Numbers</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Replicate 1</td>
<td>Replicate 2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Third instar larvae, CNS</td>
<td>12,921,055</td>
<td>40,022,667</td>
<td>52,943,722</td>
<td>modENCODE_3466 SRR070409, SRR070410</td>
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<tr>
<td>Pupae, white prepupae + 2 days, CNS</td>
<td>44,833,331</td>
<td>37,635,655</td>
<td>82,468,986</td>
<td>modENCODE_3469 SRR100271, SRR070412</td>
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<td>Mated female, eclosion + 4 days, heads</td>
<td>59,341,425</td>
<td>60,448,967</td>
<td>119,790,392</td>
<td>modENCODE_3448 SRR070414, SRR070415</td>
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<td>Mated male, eclosion + 4 days, heads</td>
<td>54,635,860</td>
<td>54,555,328</td>
<td>109,191,188</td>
<td>modENCODE_3449 SRR070416, SRR070400</td>
</tr>
<tr>
<td>Mated female, eclosion + 4 days, ovaries</td>
<td>62,102,229</td>
<td>168,912,884</td>
<td>231,015,113</td>
<td>modENCODE_3451 SRR070431, SRR100277, SRR100283</td>
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<td>Mated male, eclosion + 4 days, testes</td>
<td>249,972,580</td>
<td>171,471,971</td>
<td>421,444,551</td>
<td>modENCODE_3452 SRR070422, SRR350980, SRR070423, SRR100276, SRR350961</td>
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<tr>
<td>S2-R+</td>
<td>55,121,051</td>
<td>N/A</td>
<td>55,121,051</td>
<td>modENCODE_3487 SRR070279, SRR124149, SRR070286</td>
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<tr>
<td>Total</td>
<td>538,927,531</td>
<td>533,047,472</td>
<td>1,071,975,003</td>
<td></td>
</tr>
</tbody>
</table>

Details on RNA-seq data sets produced for this study. All the raw sequences reads and mapping files can be obtained under the designated accession numbers from the modENCODE Data Coordination Center (http://www.modencode.org/publications/dcc/index.shtml) and the NCBI Short Read Archive (http://www.ncbi.nlm.nih.gov/sra/).
Thus, neural 3’ UTRs are in fact substantially longer than currently appreciated. Reciprocally, testis-expressed genes had the shortest median 3’ UTRs across all tissues (Figure 1C, green asterisk); ovaries expressed 3’ UTRs of intermediate length (Figure 1C, black asterisk). We selected neural tissues and testis for detailed experimental and computational analysis of tissue-specific alternative 3’ UTR patterns.

The Testis Transcriptome Is Strongly Biased for Usage of Proximal Poly(A) Sites
As the 3’ UTRs of testis-expressed transcripts had the shortest median length (using two or more poly(A)-spanning reads), we were interested to identify cases of APA in which a proximal site was utilized in testis. Analysis of the stranded RNA-seq data recovered 100 genes exhibiting 3’ transcript ends that were clearly shorter in testis compared to ovaries (Figure S1 and Table S1 available online). From the poly(A)-spanning reads pooled from all the tissue libraries, 47 of these genes had poly(A) support for the proximal 3’ end and 63 had poly(A) support for the distal 3’ end. Figure 2 illustrates typical examples of this phenomenon.

To confirm differential expression of 3’ UTR isoforms in gonads, we performed RT-PCR with the use of a common proximal primer and two unique distal primers for each gene.
assays indicated preferred (Figures 2A–2C) or exclusive (Figure 2D) expression of transcripts using distal poly(A) sites in ovaries compared to testis. Additional examples can be found in Figure S1. The black regions correspond to mapped RNA-seq reads from each tissue, green indicates poly(A)-spanning reads pooled from 29 poly(A)-enriched RNA-seq libraries corresponding to the strand on which the gene of interest is expressed. The gray regions in (B) correspond to reads from the neighboring gene transcribed on the opposite strand. In (A–D), RT-PCR was performed using a common forward primer (black arrows) and UTR-specific reverse primers (blue arrows). To the right of the RNA-seq data are ethidium bromide stained gels, in which lower molecular weight bands derive from both long and short 3' UTR isoforms, whereas higher molecular weight bands are specific to long 3' UTR isoforms. Note the reduced levels of distal APA isoforms in testis.

(F) qRT-PCR demonstrates 3' UTR shortening in testis compared to ovaries for oaf, polo, slmb, e(r), and lack. Relative RNA levels are presented as ratio of total/long. Data are represented as mean ± SEM.

Figure 2. The Testis Transcriptome Is Biased toward Proximal Poly(A) Site Usage

(A–D) RNA-seq tracks for representative genes expressed in both male and female gonads, that exhibit evidence for truncated 3' UTRs in testis compared to ovary. Additional examples can be found in Figure S1. The black regions correspond to mapped RNA-seq reads from each tissue, green indicates poly(A)-spanning reads pooled from 29 poly(A)-enriched RNA-seq libraries corresponding to the strand on which the gene of interest is expressed. The gray regions in (B) correspond to reads from the neighboring gene transcribed on the opposite strand. In (A–D), RT-PCR was performed using a common forward primer (black arrows) and UTR-specific reverse primers (blue arrows). To the right of the RNA-seq data are ethidium bromide stained gels, in which lower molecular weight bands derive from both long and short 3' UTR isoforms, whereas higher molecular weight bands are specific to long 3' UTR isoforms. Note the reduced levels of distal APA isoforms in testis.

(E) Schematic for quantification of alternative length 3' UTRs in ovary and testis by qRT-PCR.

(F) qRT-PCR demonstrates 3' UTR shortening in testis compared to ovaries for oaf, polo, slmb, e(r), and lack. Relative RNA levels are presented as ratio of total/long. Data are represented as mean ± SEM.

The Neural Transcriptome Is Strongly Biased for Usage of Novel Distal Poly(A) Sites

We observed a strikingly converse trend in RNA-seq data from larval and pupal CNS and adult head, whose median 3' UTR lengths were much longer than for genes expressed in all other tissues (Figure 1C). We focused our analysis on 3' UTR extensions resulting from APA events in the 3' UTR (UTR-APA) and not from alternative cleavage events located in internal introns or exons. We identified 66 transcripts, contained within FlyBase 5.32 gene models, for which CNS/head RNA-seq evidence clearly demonstrated usage of distal PAS relative to other tissues.

Manual browsing revealed extensive transcribed regions downstream of current gene annotations in neural tissues but not in non-neural tissues. We therefore systematically searched libraries from larval and pupal CNS and adult head for 3' UTR extensions supported by continuous RNA-seq evidence, distal to annotated FlyBase models. This yielded 317 additional genes exhibiting UTR-APA, with longer 3' UTR species in the nervous system (Table S1). These extensions had a significant impact on the catalog of exonic sequence in Drosophila, collectively adding >760 kb of novel sequence to the transcriptome.
In total, we recognized 383 genes exhibiting neural-specific 3’ UTR extensions. Comparison of the 3’ UTR lengths of these 383 neural extended transcripts with FlyBase 5.32 annotations highlighted that this set of genes was rich in exceptionally long 3’ UTRs (Figures 3A and 3B). Indeed, only a handful of known *Drosophila* transcripts exhibit 3’ UTRs in excess of 5 kb (disregarding artifactual annotations, see Extended Experimental Procedures), whereas 51 of our newly annotated 3’ UTRs surpass this limit. We therefore sought experimental confirmation of these unusually extended 3’ UTRs.

The TRIM-NHL family member *brat* exhibited extensive transcription downstream of the annotated gene model in neural tissues, comprising an ~8.5 kb 3’ UTR (Figures 3C and 3D). In addition, distinctive accumulation of poly(A)-spanning reads was observed in the larval CNS compared to ovary, with the former terminating at intermediate and distal sites and the latter at a proximal site (Figure 3C). During embryogenesis, the universal *brat* probe detected maternal deposition of *brat* and two broad stripes of zygotic expression, whereas later stage expression was confined to the brain and ventral nerve cord (VNC) (Figure 3E). In contrast, the extended isoform-specific probe failed to detect maternal or early zygotic transcripts, and hybridized exclusively to CNS transcripts at later stages (Figure 3E). Therefore, the proximal APA isoform of *brat* exhibits an expression pattern that is distinct from its distal APA isoform.

A more extreme example of an extended 3’ UTR is illustrated by *mei-P26*, which also encodes a TRIM-NHL protein. Stranded RNA-seq showed tremendous variance in tissue-specific 3’ UTR lengths, with a short 3’ UTR in testis, an intermediate 3’ UTR length in ovary and an ~18.5 kb UTR in neural samples (Figures 3F and 3G). RT-PCR analysis confirmed that the extended isoform is strongly expressed in the head samples relative to body, ovary, and testis (Figure 3H). In addition, the extended isoform appeared only in embryos 12 hr of age and older (Figure 3H). Interpretation of this temporal pattern required spatial analysis (Figure 3I). A probe detecting *mei-P26* coding sequence revealed maternally deposited transcripts and posterior accumulation at stage 5. At stage 14, staining was largely ubiquitous, with enrichment in the developing brain and VNC becoming apparent by stage 17. In contrast, a probe 13 kb distal lacked maternal or early embryonic staining, and showed exclusive CNS expression in late stage embryos. Additional in situ probes designed against proximal or distal regions of the *mei-P26* 3’ UTR confirmed these distinctive spatial patterns (Figures S2A and S2B). Thus, the temporal accumulation of the extended *mei-P26* APA isoform was a consequence of its spatial expression in the maturing nervous system. Similar results were observed with several other genes with apparent developmental lengthening (e.g., *shep*, *heph*, and *msi*, Figures S2C–S2J), for which a temporal trend of 3’ UTR lengthening was attributable to the expression of 3’ UTR extended isoforms in the nervous system. These data highlight the need to coordinate expression patterns determined in whole animals with knowledge of tissue-specific gene expression.

**Northern Analysis Confirms Exceptionally Long APA Isoforms as Bona Fide Transcripts**

The RNA-seq, RT-PCR, and in situ data do not formally prove 3’ UTR extensions of stable transcripts, as opposed to transient or unstable RNA species, products of runaway transcription or improper termination, or distinct transcripts downstream of protein-coding genes (Mercer et al., 2011; Ponjavic et al., 2009). We distinguished these possibilities using northern analysis, which is also uniquely suited for assessing the relative accumulation of different full-length isoforms. We first compared the signals of universal probes (that should hybridize to all 3’ UTR isoforms) with extension probes specific to distal APA isoforms for *cam*, *shep*, *cut*, and *brat* (Figure 4A) and *mei-P26* (Figure 4B). In all cases tested, the extension probes detected a subset of species detected by the universal probe, and these always comprised longer isoforms that were strongly enriched in heads. These data provide strict evidence of 3’ UTR extension isoforms that are contiguous with their neighboring protein-coding mRNA annotations. Many of these exceeded the longest RNA size standards, including the existence of full-length *mei-P26* transcripts (Figure 4B) estimated from RNA-seq data to be some 23 kb in length, of which >18 kb was 3’ UTR.

Additional northern assays using universal probes provide broad support for extended APA isoforms that are enriched or indeed restricted to heads (Figures 4C and 4D). Interestingly, several genes exhibited both 3’ UTR lengthening in head and shortening in testis, relative to intermediate-sized isoforms expressed in body and/or ovary (e.g., *mei-P26*, *bol*, *sm*, *orb*, and *orb2*, Figures 4B and 4C). Perusal of our APA lists revealed 23 transcripts that exhibit such dual patterns of CNS and testis APA (Table S1C). Altogether, these extensive northern analyses provide a compelling view of APA dynamics leading to the accumulation of extraordinarily long APA isoforms in the *Drosophila* nervous system.

**Predominant CNS Restriction of Neural Distal APA Isoforms**

We followed up these northern analyses with additional in situ hybridization studies. As before, we compared proximal probes that would detect all 3’ UTR mRNA isoforms (universal) with extension-specific probes specific to distal APA regions. In some cases, both probes detected similar expression patterns in the CNS (e.g., *khc*-73, Figure 5A; CG4612 and *fas1*, Figure S3). However, we also observed spatially discrepant patterns of expression of paired probes. Consistent with the RNA-seq data, the extended probe often detected transcripts in a subset of the tissues for which expression was also detected for the universal probe. For example, the universal and extended probes for *bru-3* detected strong expression in the brain and VNC but strong visceral muscle primordium staining was only observed with the universal probe (Figure 5B).

Curiously, in every case the distal probes detected transcripts in the CNS, and staining was often exclusive to the CNS (Figure 5 and Figure S3). This was particularly notable for a set of known pan-neural (CNS and PNS) transcripts, for which we observed distal APA isoforms largely or completely restricted to the CNS. Examples of these genes included *fne*, *sct*, *elav* (Figures 5C–5E), and *cut* (Figure S3). In the case of the CNS-expressed gene *mub*, we observed that its extended 3’ UTR isoform was expressed in a subdomain of the brain (Figure 5F). The repeated observation of pan-neural APA transcripts with distinct CNS-restricted distal APA isoforms suggests that the mechanism underlying these unusually long 3’ UTR extensions is not simply...
Figure 3. Unusually Long 3' UTR Extensions in Neural Genes

(A) Distribution of 3' UTR lengths of 383 transcripts that exhibit neural lengthening, relative to the longest annotated 3' UTRs of all FlyBase 5.32 genes.

(B) Focusing on the tail of the distribution, the neural 3' UTR extensions comprise most of the longest 3' UTRs in the Drosophila transcriptome.
neural-specific, but biased to the CNS relative to the PNS (Figure 5).

**Enrichment of Nucleic Acid-Binding Proteins among Neural Genes Exhibiting Distal APA**

We tested for enrichments of gene ontology annotations among the 383 neural-extended transcripts. Perhaps not surprisingly, this set of genes was highly enriched for biological processes relating to neural development or neural function (Table S2A). Strikingly though, among molecular function terms, the highest statistical enrichments observed (not including the generic categories “binding” and “protein binding”) concerned various classes of nucleic acid binding proteins. In particular, sequence-specific transcription factors were enriched at

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(C and D) RNA-seq tracks from indicated tissues (C) and transcript models for brat (D). A splice variant that alters the last few amino acids of Brat was apparent from RNA-seq data (gray), but this isoform was not differentially expressed.

(E) In situ hybridization revealed that both proximal and distal probes detect CNS expression in germband retracted embryos and late stage embryos, but that the proximal probe alone detects maternal deposition and blastoderm expression in two stripes along the anterior-posterior axis.

(F and G) RNA-seq tracks from indicated tissues (F) and gene models for mei-P26 (G).

(H) Semiquantitative RT-PCR indicates that RNA corresponding to the 3' end of the coding sequence and proximal 3' UTR can be readily detected in all tissues and embryonic time-points examined, whereas an amplicon ∼8 kb into the 3' UTR was predominantly detected in heads and late stage embryos.

(i) In situ hybridization reveals that both proximal and distal probes detect late stage expression in the brain and ventral nerve cord, but that the proximal probe alone detects maternal deposition at stage 2, posterior expression at stage 5, and ubiquitous expression at stage 14.
a p value of 2.68E-08, and mRNA binding was enriched at a p value of 5.77E-06. Other categories of strongly enriched molecular functions included kinases (1.96E-06) and signal transduction components (3.31E-05) (Table S2A). We did not observe a reciprocal coherence of the genes subject to utilization of proximal APA sites in testis, because no molecular function terms were enriched among the 100 transcripts in this cohort (Table S2B). In summary, we observed several classes of genes with regulatory functions preferentially subject to 3' UTR extensions in the nervous system.

Quality of Poly(A) Sites and Local Conservation of Alternative Transcript Termini Analysis

We performed motif analysis on sequences surrounding alternative 3' ends of CNS APA (Figure 6A) and testis APA (Figure 6B) transcripts, for which poly(A)-spanning reads pooled from all

Figure 5. In Situ Hybridization Reveals Distinctive CNS Expression of Distal APA Isoforms
Schematics to the left include gene models with 3' UTR extensions in head RNA-seq data relative to another tissue. H, head; O, ovary; 10–12 and 12–14 hr embryos. The locations of proximal (black) and distal (red) in situ probes are indicated. The gene of interest is displayed in a plus strand orientation, and the name and orientation of the neighboring downstream gene is indicated. Panels with single prime labels show in situ hybridization patterns detected with a universal probe designed to detect all transcripts from the gene of interest. Panels with double prime labels show in situ hybridization patterns of probes specific for the extended isoforms.

(A) khc-73 illustrates a gene with CNS staining detected with both universal and extended probes.
(B) The universal bru-3 probe stains visceral muscle (VM) and CNS, whereas its extension probe stains only CNS.
(C–E) fne (C), scrt (D), and elav (E) are pan-neuronally expressed in the peripheral nervous system (PNS) and CNS, yet their distal APA isoforms are specific to the CNS.
(F) mub is an example of a neural transcript where the extension probe specifically detects expression in subregions of the embryonic brain (arrow) compared to the universal probe that detects expression throughout the embryonic brain and CNS.
Figure 6. Quality and Conservation of Alternative Poly(A) Sites and miRNA Sites

(A and B) Average 3' UTR lengths of the proximal and distal transcript isoforms for genes exhibiting CNS 3' UTR lengthening (A) and testis 3' UTR shortening (B).

(C and D) Sequence motifs involved in polyadenylation around poly(A) sites identified by poly(A)-spanning reads: Canonical PAS (in blue), GU-rich DSE (in green) and variant PAS (in gray). (C) For CNS extended genes, distal sites are enriched for the canonical PAS compared to proximal sites (Fisher's exact test p value = 4.9E-16), whereas the frequency at the intermediate poly(A) sites lies in between. The distal poly(A) sites are also enriched for the DSE motif, compared to the proximal and intermediate poly(A) sites (Fisher's exact test p value = 7.2E-4). All poly(A) sites show similar collective levels of variant PAS; see also Figure S5 for analysis of individual PAS variants. (D) Testis APA transcripts display similar patterns of motif occurrence and conservation as the CNS transcripts (enrichment of AAUAAA in distal versus proximal sites, Fisher’s exact test p value = 2.6E-4).

(E) PhastCons conservation scores in the vicinity of poly(A) sites. Proximal and intermediate CNS poly(A) sites have intermediate conservation levels whereas distal sites have a region of high conservation just upstream of the poly(A) site and a region of low conservation downstream of the poly(A) site.

(F) Proximal poly(A) sites in testis transcripts have intermediate conservation levels whereas the distal sites have a region of high conservation just upstream of the poly(A) site and region of low conservation downstream of the poly(A) site.

(G) Cumulative distribution of numbers of miRNA target sites across all Drosophila 3' UTRs, compared with the proximal and distal APA 3' UTR variants of the 383 transcripts exhibiting neural lengthening. The distal extensions bear significantly more miRNA binding sites than Drosophila 3' UTRs do in general.

(H) Cumulative distribution of miRNA target site numbers in the 100 testis APA transcripts.

(I) Motifs conserved above background in CNS APA extensions. Shown are the top 6-mer motifs conserved above background in greater than or equal to seven Drosophilid species among the 383 3' UTR extensions of neural transcripts. Many sites correspond to miRNA seeds or neural RBP sites (e.g., Pumilio and U-rich sequences that may potentially include Elav binding sites). A complete list of 6-mer and 7-mer motifs conserved above background are shown in Table S3. S/N, signal/noise ratio.

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29 samples described above definitively marked the site of cleavage and polyadenylation. The precise demarcation of transcript ends using the poly(A)-spanning reads enabled us to search for cis elements potentially interacting with the poly(A) machinery. De-novo searches identified the canonical AUAAA PAS upstream of poly(A) sites, and a G/U rich sequence downstream of (the distal) poly(A) sites (Figure S4) that resembled the GU-rich downstream sequence element (DSE), known to increase the efficiency of 3’ end processing in mammalian cells via interaction with CstF-64 (MacDonald et al., 1994). We also observed a degenerate A-rich motif enriched mostly upstream of poly(A) sites (Figure S4). Because no motifs were found that were clearly distinct from known poly(A)-associated elements (Hu et al., 2005; Ozsolak et al., 2010), we proceeded to analyze the canonical and variant polyadenylation signals and the inferred DSE motif in more detail.

Upon comparing proximal, intermediate, and distal isoforms of our collection of genes with neural extensions, we observed a progressively increasing fraction bearing the canonical AUAAA polyadenylation signal just upstream of poly(A) sites (Figure 6C). In contrast, variant PAS (Figure S5) were collectively equally represented upstream of these various cohorts of transcript ends (Figure 6C). We observed that distal neural poly(A) sites contained substantially higher frequency of DSE motifs, relative to intermediate or proximal isoforms. We observed a similar trend in the 3′ termini of genes that exhibit testis shortening, with the distal poly(A) sites exhibiting a much higher frequency of canonical AUAAA PAS and DSE motifs (Figure 6D).

We also assessed the levels of conservation at the various categories of poly(A) sites, using PhastCons scores (http://genome.ucsc.edu). This revealed an intermediate level of conservation (0.5–0.6) in the proximity of proximal poly(A) sites, that was relatively similar upstream and downstream of the poly(A) sites. In contrast, the levels of conservation in the vicinity of distal neural poly(A) sites rose sharply in the preceding ~50 bp, peaking at ~0.8 at the poly(A) sites, and then rapidly dropped to background levels of 0.2 (Figure 6E). This suggests that the regions in the immediate upstream vicinity of distal neural poly(A) sites are selected for sequences that mediate highly efficient cleavage and polyadenylation. A similar trend was observed in the vicinity of distal poly(A) sites of transcripts that were preferentially shortened in testis (Figure 6F).

Impact of Neural 3’ UTR Extensions on Posttranscriptional Regulation

We sought to infer the functional impact of tissue-biased patterns of APA in Drosophila. One strategy was to investigate the frequency of miRNA binding sites in the proximal versus distal portions of APA regulated transcripts. For each category of 3′ UTR we cataloged the number of conserved seed matches (two to eight 7-mer sites) for miRNAs conserved between D. melanogaster and D. pseudoobscura (Ruby et al., 2007). Cumulative distribution plots showed that proximal regions of CNS APA transcripts carried higher numbers of conserved miRNA sites than observed with all Drosophila 3′ UTRs (Figure 6G). Kolmogorov-Smirnov (KS) tests demonstrated significantly higher numbers of miRNA binding sites in distal versus proximal regions of the CNS APA transcripts. Analysis of the testis APA transcripts showed that the proximal 3′ UTRs utilized in testis had similar numbers of miRNA binding sites relative to all Drosophila 3′ UTRs, but the distal regions of these APA transcripts exhibited significantly more miRNA binding sites (Figure 6H). A simple interpretation is that APA brings neural transcripts with extended 3′ UTRs under posttranscriptional control unique to the CNS, whereas APA spares testis transcripts with shortened 3′ UTRs from posttranscriptional control that applies to longer isoforms expressed outside the testis.

We next employed an unbiased strategy to ask what motifs are most preferentially conserved in the extended portions of neural distal APA transcripts. We assessed all 6-mers or 7-mers for conservation in the 383 neural 3′ UTR extensions, above a background binomial distribution of control motifs with similar occurrence and GC content. Interestingly, many of the most-conserved motifs corresponded to miRNA binding sites, including those of miR-190, K box miRNAs (miR-2/11/13, etc.), and Brd box miRNAs (miR-4/79). Another highly conserved motif corresponds to the binding site for Pumilio, and many conserved U-rich motifs potentially including Elav binding sites (Figure 6I). Additional miRNA seeds were observed among less-conserved (but still significantly-conserved) motifs (Table S3).

Overall, the logic of our observed UTR-APA examples appears to bring neural extended transcripts under the control of neural post-transcriptional regulatory machinery, because miR-190 is strongly enriched in adult head relative to body (Ruby et al., 2007), a cluster of miR-2/13 miRNAs is specifically expressed in the CNS (Aboobaker et al., 2005), Brd box miRNAs are known to regulate nervous system development (Lai and Posakony, 1997; Lai et al., 2005), and Pumilio and Elav are well known as regulatory RNA binding proteins in the nervous system (Menon et al., 2004; Soller and White, 2003).

DISCUSSION

Toward a More Complete Description of the Drosophila Transcriptome

One of the charges of the modENCODE project is the comprehensive characterization of the fly and worm transcriptomes. Although we recently analyzed 3.5 billion RNA-seq reads spanning Drosophila development (Graveley et al., 2011), our new libraries add substantially to the catalog of genic regions in this species. We confidently identify 317 transcripts with previously unannotated 3′ UTR extensions in the nervous system, comprising >760 kb of novel 3′ UTR sequence. The large number of transcript models affected was surprising, given that the nervous system is historically one of the more well-studied tissues in Drosophila. Equally unexpected was the sheer length of many neural distal APA isoforms. A number of them extend more than 10 kb, longer than virtually all other known Drosophila transcripts, and we validated many of these as stable full-length transcripts using northern analysis.

These comprise a conservative annotation of 3′ UTR variation, as there were additional instances of extended transcription for which the bounds and continuity could not be confidently judged. In addition, we observed orphan poly(A) sites downstream of 3′ UTRs that were not clearly associated with RNA-seq
evidence. These may include transcripts with low or restricted expression in the nervous system. Reciprocally, our analysis of 3’ UTR shortening focused on APA that differed between male and female gonads, and did not include genes that were not expressed in both testis and ovary.

Beyond these APA isoforms, the datasets described herein comprise a valuable resource for furthering the annotation of the transcriptome. In particular, the dissected tissues enrich for transcripts that are rare in whole animals, and the stranded nature of the data help distinguish closely apposed transcription units, especially when they are produced from opposite strands. A fuller accounting of novel transcribed regions revealed by these data will be reported elsewhere (J.B.B. et al., unpublished data).

Tissue-Biased Features of Drosophila APA and Comparison with Other Model Systems

Our collection of >1 million poly(A)-spanning reads provides the largest resource of alternative 3’ ends in Drosophila to date, and provides direct evidence for APA events in more than 50% of detected Drosophila genes. Among this broad palette of APA transcripts, we discovered trends for shortening in the testis to 383 genes with 3’ UTR extensions, and several of which were shown to be neural-specific. Our tissue-specific RNA-seq data broadly extend these findings to other vertebrates. For example, analysis of murine ESTs showed a trend of shortened 3’ UTRs in spermatogenesis that correlated with reduced usage of canonical AAUAAA signals (Liu et al., 2007). Reciprocally, microarray analysis and deep sequencing of 3’ ends in mouse and human tissues showed that brain and nervous system were among the tissues that tended to favor distal PAS usage (Sandberg et al., 2008; Shepard et al., 2011; Zhang et al., 2005). The mechanistic bases of these tissue-specific APA trends are poorly understood, but our finding that these trends are conserved in Drosophila indicates that this genetic system will be valuable for dissecting these processes.

Biological Significance of Tissue-Specific APA in Drosophila

In the nervous system, the expression of 3’ UTR lengthened isoforms subjects these genes to spatially restricted posttranscriptional control. We show this to apply to many hundreds of transcripts, and find that binding sites for neural miRNAs and neural RNA binding proteins are among the most highly conserved motifs within these extensions. Certainly miRNA-mediated control might serve to restrict transcript function, potentially in the context of local translation or transcript recycling in response to environmental cues and neural activity. However, neural 3’ UTR extensions may not solely confer downregulation. A distal APA variant of mammalian brain-derived neurotrophic factor (BDNF), but not its proximal APA variant, localizes to dendrites where it plays a role in long term potentiation (An et al., 2008). Moreover, Drosophila polo undergoes APA, in which the distal isoform is required to support efficient Polo translation (Pinto et al., 2011); thus, 3’ UTR extensions can promote translation. Finally, it is worth considering whether the considerable real estate within these neural 3’ UTR extensions may serve structural or scaffolding functions, or perhaps act as “sponges” that attract miRNA or RBP complexes.

We are intrigued by the fact that among the network of Drosophila genes exhibiting 3’ UTR lengthening in the nervous system, the top-enriched molecular functions are transcription factors and RBPs. This may imply special needs to regulate these key regulatory molecules in the nervous system. Interestingly, several of the RBPs that are subject to extraorordinarily lengthened 3’ UTRs are themselves involved in 3’ UTR determination and/or miRNA-mediated regulation (e.g., elav, mei-P26, brat, pumilio, ago1). This raises the possibility of a complex network of auto- and cross-regulating posttranscriptional regulatory factors in the CNS. Recent technical advances in identifying RBP interactions with RNA elements such as PAR-CLIP and HITS-CLIP (Hafner et al., 2010; Licatalosi et al., 2008) enable genome-wide examination of RBP-target interactions, and should prove useful for elucidating these complex networks.
EXPERIMENTAL PROCEDURES

Sample Preparation and RNA-seq

Total RNA was isolated from tissues dissected from Oregon R animals in biological duplicates or from cultured S2R+ cells. Strand-specific RNA-seq libraries were prepared using pre-release Directional mRNA-seq Library Kits (Illumina). Briefly, poly(A)+ RNA was isolated by oligo-dT selection, fragmented, and treated with phosphatase and polynucleotide kinase to repair the ends. RNA adapters (3' and 5') were then serially ligated to the RNA fragments and reverse transcribed using a primer complementary to the 3' linker. The libraries were then PCR amplified and sequenced on either an Illumina GAIIx using paired-end 76 bp chemistry or a HiSeq2000 using paired-end 100 bp reads. Reads were simultaneously aligned to the genome and splice junctions using Bowtie (Langmead, 2010) and SGA to report uniquely aligned reads as described (Graveley et al., 2011). The raw fastq RNA-seq data were deposited at the NCBI Short Read Archive, and the processed bam files were deposited at the modENCODE Data Coordination Center; the accession numbers are summarized in Table 1.

Poly(A) Enrichment

One nanogram of the strand-specific RNA-Seq libraries were reamplified by PCR using a primer complementary to the 5' adaptor and a second primer complementary to the 3' adaptor with six T residues at the 3' end. After 10 rounds of amplification, the 3' primer with the T extension was replaced with a 3' primer complementary to the adaptor with a 5' extension containing a 6 nt index sequence and a sequence complementary to the flow cell primer. After an additional 15 rounds of amplification, the libraries were quantitated, 10–12 libraries were pooled together and sequenced on an Illumina HiSeq2000 using paired-end 100 bp and 6 bp index read chemistry. Reads were split into the respective samples using the index sequence and aligned as described above. All of the raw fastq data and alignments of poly(A)-spanning reads from the poly(A)-enriched libraries were deposited at NCBI Gene Expression Omnibus under Series GSE3390.

Experimental Analysis of Gene Expression

In situ hybridization was performed on mixed stage Canton S embryos according to the BDGP 96-well plate in situ hybridization procedure (Tomancak et al., 2002), with modifications to utilize 1.7 ml tubes. Quantitative RT-PCR was performed using SYBR green PCR mastermix (QIAGEN) on a CFX96 real-time system (BioRad). Northern analysis of was performed as previously described (May and Bartel, 2009) using 2–3 μg of poly(A)+ RNA or 8–12 μg of total RNA per lane. Detailed experimental procedures are provided in the Extended Experimental Procedures. Oligo sequences used to generate in situ probes, northern probes and qPCR amplicons are listed in Table S4.

Analysis of PAS Defined by Poly(A)-Spanning Reads

Read pairs that failed to align to the genome were examined to identify cases where the first read contained ≥10 A residues at the 3' end or the second read contained ≥10 T residues at the 5' end. Terminal A or T residues were trimmed from the reads and uniquely aligned poly(A)-spanning reads identified. These were filtered for instances of oligo-dT priming to potentially genomically encoded poly(A) stretches by removing reads where at least eight out of the ten nucleotides downstream of the matching regions were adenines. Remaining reads were clustered so that all reads mapping to the same strand and ending within ten nucleotides were collapsed into a cluster. Clusters supported by at least two reads were considered for further analysis.

3' UTR Analysis

A combination of bioinformatic searches and manual browsing was performed to identify genes with neural 3' UTR extensions. This was necessitated by the discontinuous nature of stranded RNA-seq data. The computational scan was based on the number of reads proximal and distal to annotated 3' ends in head and ovary libraries, assuming that genes that displayed a higher ratio of distal/proximal reads in head samples compared to ovary samples could be mRNAs with neural 3' UTR extensions. To identify genes with 3' UTR truncations in tests, we performed a computational search for an increased ratio of proximal/distal reads between tests and ovary. Manual browsing to validate transcript models was performed using Jbrowse (browse.org) and Integrated Genome Viewer (www.broadinstitute.org/gv) for the entire Drosophila genome with tracks loaded from multiple Drosophila tissues.

Motif Analysis

Regions around the polyadenylation sites (±50 nt) were scanned for sequence motifs using MEME (Bailey et al., 2006) and Weeder (Pavesi et al., 2001). We also performed directed searches for the canonical PAS AAUAAA and known variants (Retelska et al., 2006). The neural extended and tests shortened portions of the 3' UTRs were scanned for conserved 6-mers and 7-mers using the approach described in (Xie et al., 2005). We also searched for known miRNA seeds (Ruby et al., 2007) within annotated and extended 3' UTRs.

ACCESS NUMBERS

Raw fastq RNA-seq data have been deposited at the NCBI Short Read Archive, and the processed .bam files have been deposited at the modENCODE Data Coordination Center. Accession numbers for these data are summarized in Table 1.

SUPPLEMENTAL INFORMATION

Supplemental Information includes Extended Experimental Procedures, six figures, four tables, and one data set and can be found with this article online at doi:10.1016/j.cell.2012.01.001.

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**Extended Experimental Procedures**

**In Situ Hybridization**
In situ probes were prepared using two-step amplification starting with genomic DNA or cDNA templates. The first reaction consisted of gene-specific oligonucleotides designed to amplify 500-700 nucleotide products and include linker regions at either end; sequences are listed in Table S4. A second round of PCR with primers designed against the linker regions was performed on these products to append T7 and T3 recognition sites. Purified products were used as template for RNA probe synthesis using T7 Digoxigenin RNA Labeling Kit (Roche). In situ hybridization was carried out on mixed stage Canton S embryos according to the Berkeley Drosophila Genome Project 96 well plate in situ hybridization procedure (Tomanack et al., 2002), with modifications to utilize 1.7 ml tubes. RNA probes were detected with anti-Digoxigenin-AP and BM purple (Roche). Oligonucleotide sequences used to generate in situ probes are listed in Table S4.

**First Strand Synthesis and RT-PCR**
For cDNA preparation, reverse transcription was performed using Superscript III reverse transcriptase (Invitrogen) on total RNA treated with DNase I (Ambion). qPCR was performed using SYBR green PCR mastermix (QIAGEN) on a CFX96 real-time system (Biorad). We used the delta delta Ct method to quantify levels. Oligonucleotide sequences of qPCR amplicons are listed in Table S4.

**Northern Analysis**
Total RNA was extracted from Canton S tissues using Trizol (Invitrogen), and poly(A)+ RNA was prepared from total RNA using oligotex mRNA mini kit (QIAGEN). Ovary and testis were manually dissected. Body and head samples were obtained by separation using liquid nitrogen to freeze, followed by vortexing and separation of heads from bodies using a sieve. Northern analysis was performed essentially as previously described (May and Bartel, 2009) using 2-3 μg of poly(A)+ RNA or 8-12 μg of total RNA per lane. Oligonucleotide sequences used to generate northern probes are listed in Table S4.

**Analysis of PAS Defined by Poly(A)-Spanning Reads**
To identify poly(A)-spanning reads from the paired-end sequencing data, read pairs that failed to align to the genome were examined to identify cases where the first read contained ≥10 A residues at the 3’ end or the second read contained ≥10 T residues at the 5’ end. The terminal A or T residues were trimmed from the reads and uniquely aligned poly(A)-spanning reads identified using Bowtie. These were further filtered to remove instances of oligo dT priming to potentially genomically encoded poly(A) stretches. Reads where at least 8 out of the 10 nucleotides downstream of the matching regions were adenines were filtered out. The poly(A)-spanning reads are available as a BED file in the Data Set S1. Remaining reads were clustered so that all reads mapping to the same strand and ending within 10 nucleotides were collapsed into a cluster. Clusters supported by at least 2 reads were considered for further analysis. To assess differences in 3’ UTR length between tissues, gene expression data from the FlyAtlas (Chintapalli et al., 2007) were used. For each tissue, all genes with enrichment scores of at least 2 were selected. The longest and shortest 3’ UTRs supported by clusters of at least 2 reads were computed, along with longest and shortest 3’ UTRs based on FlyBase 5.32 annotations of the same genes (Tweedie et al., 2009). For each tissue, median 3’ UTR lengths were reported.

**3’ UTR Analysis**
We used FlyBase 5.32 as our reference for current 3’ UTR annotations. For analysis of long FlyBase 3’ UTR lengths (Figure 3A, B), we disregarded a handful of annotations that were likely artifacts. These included atypically long annotations that lacked RNA-seq support and/or had RNA-seq evidence for 3’ UTR splicing that caused mature 3’ UTR to be much shorter than annotated. To identify genes with neural 3’ UTR extension, we performed a combination of computational searching and manual browsing. This was necessary as we found that discontinuities in the stranded RNA-seq data frequently caused automatic exon calls to terminate in the midst of a transcribed region that was visually evident. Briefly, we determined the number of reads proximal and distal to annotated 3’ ends in head libraries compared to ovary libraries. We reasoned that the genes that displayed a higher ratio of distal/proximal reads when comparing between head and ovary samples could be mRNAs with neural 3’ UTR extensions. Manual browsing to validate transcript models was performed using JBrowse (jbrowse.org) for the entire Drosophila genome with tracks loaded from multiple Drosophila tissues. Candidates were then further screened using Integrated Genomics Viewer (IGV) 2.0 (Robinson et al., 2011). To identify genes with 3’ UTR truncations in testsi, we performed a computational search for an increased ratio of proximal/distal reads between testsi and ovary, and confirmed candidates using manual browsing.

**GO Analysis**
Gene Ontology term enrichment was calculated with AmiGO (Ashburner et al., 2000), using the hypergeometric distribution and the Bonferroni correction to correct for multiple hypothesis testing.

**Motif Analysis**
Regions around the polyadenylation sites (± 50 nt) were scanned for sequence motifs using MEME (Bailey et al., 2006) and Weeder (Pavesi et al., 2001). We also performed directed searches for the canonical PAS AUAUAAA and the known variants AUUAAA,
UAUAAA, AGUAAA, UUUAAA, AAUAUA, CAUAAA, AAUACA, GAUAAA, AAGAAA, AAAAAG, AAAACA and AAUAGA (Retelska et al., 2006). The neural extended and testis shortened portions of the 3’ UTRs were scanned for conserved 6-mers and 7-mers (above a background of sequences with similar frequency and CG content) using the approach described in (Xie et al., 2005). Only sequences that occurred at least 5 times and had at least 50 sequences in the control set were reported. P-values were computed using the binomial distribution, and corrected for multiple hypothesis testing using the FDR method (Benjamini and Hochberg, 1995).

SUPPLEMENTAL REFERENCES


Figure S1. Additional Examples of Genes Exhibiting 3’ UTR Shortening in Testis Compared to Ovary, Related to Figure 2

Fourteen examples are illustrated showing representative RNA-seq traces from ovary, testis, and accumulated poly(A)-spanning reads pooled from 29 RNA-seq libraries. The y axis for the RNA-seq traces indicates the read coverage. Gray regions in gene models and RNA-seq plots indicate neighboring genes, and RNA-seq reads associated with them.
Figure S2. Additional Examples of Developmental Expression of APA Isoforms of Genes with Neural 3' UTR Extensions, Related to Figure 3

(A) RNA-seq traces, transcript models and additional probe locations for mei-P26 (partially reproduced from Figure 3 for clarity). (B) Additional in situ hybridizations for mei-P26 using the indicated probes not shown in the main figure. (C) RNA-seq traces, transcript models and probe locations for shep. (D) RT-PCR amplification of the indicated regions showing that the distal-most amplicon (55) is detected strongly in 12-16 hr and older embryos and in adult heads, but only weakly or not at all in other tissues examined. (E) qRT-PCR of the indicated amplicons showing strong enrichment of the long shep isoform in head versus body samples. (F) in situ hybridization for shep transcripts using the indicated probes reveals absence of head mesoderm staining for the probe specific for the extended isoform. (G) RNA-seq traces, transcript models and probe locations for heph. (H) The relative abundance of the extended isoform of heph increases with embryonic age, and is higher in head than body (left panel). Northern blot showing that the extended isoform of heph is expressed in the adult head (right panel). (I) RNA-seq traces, transcript models and probe locations for msi. The relative abundance of the extended isoform of msi increases with embryonic age, and is higher in head than body (left panel). (J) Northern blot indicating that several extended isoforms of msi are expressed in the adult head (right panel). Note that the northern blots in (H) and (J) are also presented in Figure 2 and are re-produced here for clarity. Data are represented as mean +/- SEM.
**Figure S3. Additional Examples of Embryonic Expression of APA Isoforms of Genes with Neural 3' UTR Extensions, Related to Figure 5**

(A) *wech*, which by RNA-seq data expresses a short isoform only in early embryogenesis, shows no difference in expression patterns between the universal and extended probes. (B) RNA-seq analysis of *cut* suggests APA during embryogenesis. Staining in the developing PNS in stage 11 embryos is observed for the universal probe, but not for the extended probe. At stage 14, the extended probe staining is CNS restricted, while the universal probe detects expression in the CNS and PNS. Expression is seen for universal and extended forms in the CNS at stage 17. (C) No clear differences were observed in the staining of the universal and extended probes for *CG4612*. Staining was at low intensity and ubiquitous with enrichment in the brain and VNC at stage 14, and restricted to the brain and VNC at stage 17. (D) Both the universal and extended probes for *fas1* detected brain, CNS and PNS staining at stage 14, becoming restricted to the brain and VNC at stage 17.
MEME was used to search for motifs within ±50 bp of pA sites for 3' end regions supported by poly(A)-spanning RNA seq reads. All motifs with MEME p-values below 1e-3 are shown for proximal and distal pA sites for the neural extension and testis shortening genes. mean pos = mean position of motif; SD = standard deviation.

**CNS 3'UTR lengthening genes**

- Proximal neural pA sites: p-val: 6.4e-50, mean pos=-9, SD=22
- Distal neural pA sites: p-val: 4.2e-194, mean pos=-20, SD=15

**Testis 3'UTR shortening genes**

- Proximal testis pA sites: p-val: 2.2e-10, mean pos=-9, SD=21
- Distal testis pA sites: p-val: 2.5e-34, mean pos=-18

Figure S4. De Novo Motif Search of Regions Surrounding Polyadenylation Sites, Related to Figure 6

MEME was used to search for motifs within ±50 bp of pA sites for 3' end regions supported by poly(A)-spanning RNA seq reads. All motifs with MEME p-values below 1e-3 are shown for proximal and distal pA sites for the neural extension and testis shortening genes. mean pos = mean position of motif; SD = standard deviation.
Figure S5. Enrichment of Variant PAS in the Vicinity of APA Sites, Related to Figure 6
Shown are a series of variant PAS analyzed for the transcripts exhibiting lengthening in the CNS and for the transcripts exhibiting shortening in the testis.
Figure S6. Genes with 3’ UTR CAGE Peaks Are Not Associated with Accumulation of Distinct 3’ UTR Transcripts, Related to Figure 4

We present four genes with neural 3’ UTR extensions and broad 3’ UTR CAGE peaks (cam) or punctate 3’ UTR CAGE peaks (shep, cut, mei-P26). However, Northern analysis of proximal and distal extension probes (re-produced from Figure 4) show evidence for accumulation of the full length 3’ UTR extended transcripts. We did not observe shorter bands detected by the extension-specific probes that would correspond to distinct 3’ UTR-only transcripts.