dTIS11 Protein-dependent Polysomal Deadenylation Is the Key Step in AU-rich Element-mediated mRNA Decay in Drosophila Cells*

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Background: TIS11 proteins control the degradation of mRNA containing AU-rich elements (ARE).

Results: Drosophila dTIS11 activates the polysomal deadenylation of the ARE mRNA Cecropin A1.

Conclusion: Drosophila dTIS11 controls fewer aspects of ARE mRNA decay as compared with its mammalian homologs.

Significance: TIS11 proteins may have acquired additional functions to control mRNA decay during evolution from invertebrates to mammals.

The destabilization of AU-rich element (ARE)-containing mRNAs mediated by proteins of the TIS11 family is conserved among eukaryotes including Drosophila. Previous studies have demonstrated that Tristetraprolin, a human protein of the TIS11 family, induces the degradation of ARE-containing mRNAs through a large variety of mechanisms including deadenylation, decapping, and P-body targeting. We have previously shown that the degradation of the mRNA encoding the antimicrobial peptide Cecropin A1 (CecA1) is controlled by the TIS11 protein (dTIS11) in Drosophila cells. In this study, we used CecA1 mRNA as a model to investigate the molecular mechanism of dTIS11-mediated mRNA decay. We observed that during the biphasic deadenylation and decay process of this mRNA, dTIS11 enhances deadenylation performed by the CCR4-CAF-NOT complex while the mRNA is still associated with ribosomes. Sequencing of mRNA degradation intermediates revealed that the complete deadenylation of the mRNA triggers its decapping and decay in both the 5′-3′ and the 3′-5′ directions. Contrary to the observations made for its mammalian homologs, overexpression of dTIS11 does not promote the localization of ARE-containing mRNAs in P-bodies but rather decreases the accumulation of CecA1 mRNA in these structures by enhancing the degradation process. Therefore, our results suggest that proteins of the TIS11 family may have acquired additional functions in the course of evolution from invertebrates to mammals.

The dynamic control of mRNA stability is one of the key strategies used by eukaryotic cells to regulate gene expression.

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5 The abbreviations used are: ARE, AU-rich element; TTP, Tristetraprolin; AMD, ARE-mediated decay; cRACE, circularized rapid amplification of cDNA ends; CCR4, carbon catabolite repression 4; CAF1, CCR4 associated factor 1.
phosphorylation of TTP (10, 11). Although deadenylation seems to play a major role in TTP-mediated mRNA decay in mammals, the observation that TTP can promote AMD when the poly(A) tail is artificially replaced by a histone 3' end-processing sequence suggests that TTP can exert its effects through alternative mechanisms (13).

The detection of the ARE-binding proteins TTP and BRF1 in complex with the decapping machinery and the observation that decapping of mammalian ARE mRNAs can be a limiting step in the degradation process suggested a mechanism in which TTP could also induce AMD by promoting mRNA decapping (14). This proposed mechanism is further supported by the fact that TTP activates DCP2 decapping activity in vitro (15).

In eukaryotes, RNA degradation enzymes are found concentrated in P-bodies, which are considered as "factories for mRNA decay" (see Ref. 16 for review). The mammalian members of the TIS11 protein family are all found at least partially localized in P-bodies (17). The observation that ARE mRNAs are specifically targeted to P-bodies (18) and that in some cases inhibition of P-body formation prevents AMD (19) suggested that an additional function of TTP and of related mammalian proteins was to deliver ARE mRNAs to degradation in P-bodies. Alternatively, P-bodies could function as reservoirs to sequester ARE mRNAs from the translational machinery when mRNA decay is delayed (18). ARE mRNA degradation can occur through the 5'-3' exosome-dependent degradation pathway (20, 21). However, in mammalian cells, the observation that both ARE mRNAs and TTP proteins accumulate in P-bodies when the 5'-3' decapping pathway is inhibited suggested that this mode of degradation is favored in TTP-mediated ARE mRNA decay (18, 19). Altogether, these observations indicate that in mammals, TTP and the other members of the TIS11 family are multipotent proteins, which favor AMD by enhancing several crucial steps in the mRNA degradation process.

AMD is a conserved mechanism among eukaryotes. In yeast, CTH2p, the homolog of TTP, regulates the stability of mRNAs important for iron metabolism through an ARE-dependent process (22). Recently, we and others have shown that AMD is conserved in Drosophila. In this organism, dTIS11 regulates the stability of ARE-containing mRNAs including messenger RNAs involved in the immune response, such as antimicrobial peptide-coding mRNAs (23–25). We have observed that AMD requires the CAF1 protein in Drosophila. However, the molecular mechanism of mRNA decay mediated by proteins of the TIS11 family remains so far mostly uncharacterized in invertebrates. By using the mRNA encoding the Cecropin A1 (CecA1) antimicrobial peptide as a model substrate of AMD, we explored the involvement of dTIS11 in several steps of mRNA degradation. We observed that the deadenylation and decay of CecA1 mRNA is a biphasic process. The initial deadenylation occurs in the absence of mRNA decay, and the final deadenylation occurs while the messenger is still loaded on polysomes and is the key step controlled by dTIS11 to target the mRNA for degradation. Interestingly, in Drosophila cells, ARE-containing mRNAs are neither specifically addressed to P-bodies nor do they seem to be preferentially targeted for decapping. Our results suggest that the activity of dTIS11 is mainly based on its capacity to enhance deadenylation in Drosophila cells and that proteins of this family may have acquired additional functions later in the course of evolution from invertebrates to mammals.

**EXPERIMENTAL PROCEDURES**

**Reagents**—Actinomycin D and DNA oligonucleotides were purchased from Sigma-Aldrich. Hygromycin B and peptidoglycan (Escherichia coli K12) were purchased from InvivoGen. Tobacco acid pyrophosphatase was purchased from Epicenter, and T4 RNA ligase was from New England Biolabs.

**Plasmids and dsRNAs**—Expression vectors coding for dTIS11 and GFP were prepared by cloning corresponding RT-PCR products in the pMT/V5-His vector (Invitrogen). CAF1 cDNA was amplified with a 3' primer replacing the natural stop codon by a HA tag and cloned in the pMT vector. The CAF1-D53A/E55A dominant negative form was obtained by site-directed mutagenesis to replace aspartate 53 and glutamate 55 by alanines. dsRNAs were prepared by in vitro transcription (MEGAscript RNAi, Ambion) of either PCR products or cDNAs cloned in the pBluescript vector. Amplified sequences were selected with the E-RNAi software (26).

**Cell Culture, Transfection, and RNAi Experiments**—S2 cells were maintained at 25 °C in Schneider medium (Invitrogen) supplemented with 10% fetal bovine serum. Highly transfectable S2 subclone (kind gift of N. Silverman, University of Massachusetts, Worcester, MA) were stably transfected with FuGENE® HD (Roche Applied Science) and maintained in Schneider medium supplemented with 300 μg/ml hygromycin B. For RNAi experiments, cells were grown in the presence of dsRNA (5 μg/ml) for 7 days.

**RNA Analysis**—In Drosophila cells, negative cross-talks operating in the immunodeficiency (Imd) pathway result in transient activation of the Cecropin A1 promoter in response to peptidoglycan stimulation (Ref. 27 and references therein). This process allowed us to evaluate the stability of CecA1 mRNA for several hours after stimulation of S2 cells with peptidoglycan and without the addition of potentially toxic pharmacological inhibitors of transcription. However, as this kinetic of transcriptional shutdown cannot be precisely controlled, evaluations of CecA1 mRNA stability over short periods of time were performed in the presence of actinomycin D to achieve rapid transcriptional inhibition. Northern blot and RT-PCR were performed as described elsewhere (23). For high resolution Northern blot, RNA samples were separated by electrophoresis on 3% agarose gels. Molecular weight markers were prepared by in vitro transcription of defined size cDNAs using 32P-labeled UTP. In vitro deadenylated CecA1 mRNA was prepared by hybridization of total RNA with oligo(dT) (18–21-mer) at 37 °C for 30 min in the presence of 1 unit of RNase H. Circularized rapid amplification of cDNA ends (cRACE) analysis was performed according to the adaptation of Rissland and Norbury based on previously described methods (Ref. 28 and references therein). Briefly, total RNA from S2 cells was purified with TRI Reagent® (Molecular Research Center). Samples were treated or not with alkaline phosphatase and tobacco acidic phosphatase to remove the cap structure prior to circularization with T4 RNA ligase. cDNA clones obtained by RT-PCR amplification of the CecA1 mRNA using divergent primers (forward, GGCCAGAATGAGAGCGACG;
Mechanism of ARE-mediated Decay in Drosophila

Overexpression of dTIS11 Accelerates the Degradation of CecA1 mRNA but Does Not Alter the Biphasic Profile of mRNA Deadenylation and Decay—General mRNA deadenylation and decay is a biphasic process in eukaryotes. The first slow and synchronous deadenylation shortens the poly(A) tail without detectable decay of the mRNA. The second phase of deadenylation is less synchronous and is accompanied by mRNA decay (recently reviewed in Ref. 29). In Drosophila S2 cells, the mRNA encoding the antimicrobial peptide Cecropin A1 is rapidly accumulated in response to the activation of the Imd pathway by Gram-negative peptidoglycan. Due to the ARE present in its 3′-UTR, this messenger RNA is also rapidly degraded. The analysis of the deadenylation process of the CecA1 mRNA is facilitated by its small RNA body size (353 nucleotides) and can be directly evaluated by Northern blot. Therefore, the degradation of CecA1 mRNA is a useful model to study the mechanism of AMD in Drosophila cells. We have previously shown that this mRNA is deadenylated by a two-step mechanism (23). Our RNAi experiments indicated that both dTIS11 and CAF1 were necessary for the second phase of deadenylation, whereas depletion of these proteins alone had no detectable influence on the first phase of deadenylation (23).

To better delineate the influence of dTIS11 on the deadenylation and decay mechanism of Drosophila ARE-containing mRNAs, we established S2 cell lines stably transfected with an expression vector where dTIS11 or GFP coding sequences were placed under the control of the Metallothionein gene promoter. S2 cells were treated with copper sulfate to induce the accumulation of GFP or dTIS11 (Fig. 1C) and stimulated with peptidoglycan to transiently activate the Imd pathway (27). We observed that CecA1 mRNA accumulation over time is significantly reduced in cells overexpressing dTIS11 as compared with cells expressing GFP (Fig. 1A, compare left and right panels and Fig. 1B), demonstrating that the accumulation of dTIS11 is sufficient to enhance CecA1 mRNA degradation. In GFP control cells, the average length of the CecA1 mRNA is markedly decreased during the initial 2 h of stimulation (Fig. 1A, lanes 2–4), suggesting a rapid deadenylation of the mRNA after synthesis, whereas a decrease in the total amount of mRNA is initiated only after 2 h of stimulation (Fig. 1B). As described previously (23), this degradation phase of the mRNA is concomitant with a further decrease in length of the remaining pool of messenger RNA (Fig. 1A, lanes 4–8). In dTIS11-overexpressing cells, CecA1 mRNA degradation is detectable as soon as 1 h after stimulation (Fig. 1B), whereas this mRNA is already partially deadenylated as indicated by the smear detectable in Fig. 1A (lane 11). Transcriptional blockade by actinomycin D during peptidoglycan stimulation further confirmed the absence of mRNA degradation during the initial phase of CecA1 mRNA deadenylation in control cells (Fig. 1D, lanes 2–8, and Fig. 1E). Interestingly, in dTIS11-overexpressing cells, this initial deadenylation of ~70–80 nucleotides occurs more rapidly as it is completed after 15–30 min of actinomycin D treatment (Fig. 1D, compare lane 8 with lanes 11–12). However, the total amount of CecA1 mRNA remains constant and not significantly different from the level observed in control cells during this period of time (0–30 min after actinomycin D addition, Fig. 1E). After 30 min of actinomycin D treatment, we observed a processive deadenylation and decay of the CecA1 mRNA pool (Fig. 1D, lanes 12–16). Therefore, although dTIS11 is able to accelerate the deadenylation and the decay process of CecA1 mRNA, our observations suggest that this factor is only able to trigger mRNA decay when the poly(A) tail has been markedly reduced from its initial size.

The Initial Deadenylation of CecA1 mRNA Is Indifferently Performed by the PAN2-PAN3 and CCR4-CAF1-NOT Complexes, whereas the Late Deadenylation Phase Relies Solely on the CCR4-CAF1-NOT Complex and on CAF1 Catalytic Activity—To identify the enzymatic activities involved in the different...
deadenylation steps of CecA1 mRNA, S2 Drosophila cells were treated with dsRNA targeting messenger RNAs encoding all the Drosophila deadenylation complexes described so far (5), including the PAN2-PAN3 complex, which was reported in mammals to be responsible for the initiation of general mRNA deadenylation (30). Following incubation with dsRNA, cells were stimulated with peptidoglycan. After 30 min, actinomycin D was added for short periods of time to observe the initial phase of CecA1 mRNA deadenylation. As shown in Fig. 2A (left panel), the size of CecA1 mRNA is rapidly reduced after actinomycin D treatment (compare lanes 2 and 4). Pretreatment of cells with dsRNAs targeting mRNAs encoding CAF1 or PAN2 and PAN3 proteins efficiently reduces the accumulation of these mRNAs (Fig. 2B) but has no effect on the shortening of CecA1 mRNA (Fig. 2A, second and third panels). Similarly, knockdown of mRNAs encoding CCR4, Nocturnin, and Angel deadenylases does not affect the initial deadenylation of the mRNA (not shown). However, the combined knockdown of
PAN2, PAN3, and CAF1 leads to the accumulation of the full-size CecA1 mRNA (Fig. 2A, right panel), thereby indicating that the PAN2-PAN3 complex and CAF1 protein play redundant functions in controlling the initial phase of CecA1 mRNA deadenylation.

The CAF1 protein belongs to the CCR4-CAF1-NOT complex involved in general mRNA deadenylation in eukaryotes. Both the CCR4 and the CAF1 components of this complex contain deadenylase domains. Although only CCR4 seems to be catalytically involved in deadenylation in yeast, both proteins are active in mammals (9). In Drosophila, CAF1 has been identified as the main enzyme involved in the general deadenylation of mRNAs (Ref. 31 and references therein). Because we previously described that CAF1 was necessary for the second deadenylation phase and for the decay of ARE mRNAs in Drosophila (23), we determined whether the intrinsic catalytic activity of CAF1 was required for the deadenylation of CecA1 mRNA. S2 cell lines were transfected with vectors encoding the GFP or a
dominant negative form of CAF1 (Fig. 2D). We compared the accumulation of CecA1 mRNA in response to prolonged peptidoglycan stimulation of these cell lines. The expression of the catalytic mutant form of CAF1 has little effect on the accumulation of CecA1 during the initial 3 h of peptidoglycan stimulation corresponding to the initial phase of mRNA deadenylation but significantly increases the amount of CecA1 mRNA detected during the later phase of deadenylation and decay occurring after 3 h of stimulation (Fig. 2, C and E). The ability of the dominant negative form of CAF1 to inhibit the secondary deadenylation of the mRNA was further confirmed by the broader band detected on Northern blot after 4 h of stimulation with peptidoglycan (Fig. 2C).

We tested whether subunits of the CCR4-CAF1-NOT complex were required for the second phase of CecA1 mRNA deadenylation. We confirmed that the depletion of CAF1 inhibits the last deadenylation step of CecA1 mRNA (Fig. 2F). Inhibition of Not1 and Not3 expression also induces the accumulation of a partially deadenylated form of CecA1 mRNA. In contrast, knockdown of Not4 and Caf40 had no effect on CecA1 mRNA deadenylation (Fig. 2G). Altogether, our results show that the catalytic activities of CAF1 as well as the NOT1 and NOT3 subunits are required to induce the second phase of CecA1 deadenylation and to target this mRNA to degradation, suggesting that CAF1 deadenylase is active on CecA1 mRNA in the context of the CCR4-CAF1-NOT complex. Because total depletion of NOT subunits cannot be achieved by RNAi, we cannot determine at this point whether all subunits of this complex are required for the deadenylation of this ARE-containing mRNA, but our results suggest that a decrease in the abundance of NOT4 and CAF40 does not affect the deadenylating activity of the complex toward ARE mRNAs.

Decapping Conditions CecA1 mRNA Deposition and Occurs Only on Fully Deadenylated Messenger RNA Molecules—In mammals, proteins of the TIS11 family are able to enhance the decay of ARE mRNAs by stimulating mRNA decapping. TTP and BRF-1 can interact with the Dcp1-Dcp2 decapping complex. TTP has also been shown to activate decapping in vitro (15), and the decapping activator Lsm1 is essential for AMD in mammalian cells (19). Alternatively, decapping of mammalian ARE mRNAs can also be achieved by the Nudt16 decapping factor (32). Analysis of the general mRNA degradation pathway in mammals indicates that decapping can take place after the initial phase of deadenylation or later when the mRNA has reached a fully deadenylated state (30).

To analyze the influence of decapping on AMD in Drosophila cells, we assessed the degradation of CecA1 mRNA in S2 cells where decapping has been inhibited by depletion of the DCP1/2 components of the decapping complex and of the Me31b decapping co-activator (Fig. 3B). Cells were stimulated with peptidoglycan for increasing periods of time, and CecA1 mRNA decay was evaluated by Northern blot (Fig. 3A). As
The ability of dTIS11 to stimulate the decapping mechanism of ARE mRNAs could not be evaluated by this approach because the presence of long poly(A) tails on CecA1 mRNA molecules from dTIS11-depleted cells disrupts the CRACE procedure. However, because we previously observed that depletion of dTIS11 blocks the secondary deadenylation and decay phase of ARE-containing mRNAs (23), we can conclude that dTIS11 is active on ARE-containing mRNA before the decapping step.

**Mechanism of ARE-mediated Decay in Drosophila**

We have previously observed that dTIS11 does not localize in P-bodies in Drosophila S2 cells (23). However, to determine whether dTIS11 could favor the accumulation of CecA1 mRNA in P-bodies as described in mammals, we analyzed the localization of CecA1 mRNA in P-bodies (18). Indeed, in human cells, the accumulation of ARE-mRNAs in P-bodies correlates with cellular levels of TIS11 proteins. Depleting TTP and BRF1 by siRNA reduces the accumulation of ARE-mRNAs in P-bodies, whereas overexpression of TTP enhances the accumulation of an ARE mRNA to P-bodies (18). These observations suggest that one function of TIS11 proteins in mammals is to stimulate mRNA decay by enhancing the contact between ARE mRNAs and degradation factors concentrated in P-bodies. Alternatively, these factors could silence gene expression by localizing ARE mRNAs away from the translation machinery (18).

To determine whether this function of TIS11 proteins is conserved in invertebrates, we evaluated the localization of CecA1 mRNA in the course of degradation in *Drosophila* cells by RNA FISH experiments. As shown in Fig. 4A, CecA1 mRNA is barely detectable in the cytoplasm of unstimulated S2 cells. After 30 min of stimulation by peptidoglycan, CecA1 mRNA accumulates in the cytoplasm. The localization of the messenger remains unchanged upon prolonged exposure of the cells to peptidoglycan. CecA1 mRNA is homogeneously distributed in the cytoplasm during the deadenylation and decay process and does not concentrate in cytoplasmic foci (Fig. 4A, pictures 2–5, and Fig. 4B, lanes 2–5). Interestingly, in cells treated with actinomycin D, CecA1 mRNA is detectable in cytoplasmic granules in ~15% of the cells (Fig. 4A, pictures 6–7, and Fig. 4C) concomitantly with the accumulation of a fully deadenylated form of this molecule (Fig. 4B, lanes 6–7). Co-localization of these granules with overexpressed DCp1 proteins identifies them as P-bodies (Fig. 4D). In mammals, the formation of P-bodies is stimulated by depletion of the mRNA degradation machinery (18). Therefore, accumulation of CecA1 mRNA in P-bodies could result from a depletion of the RNA degradation machinery induced by actinomycin D in S2 cells.

The presence of a short poly(A) tail on all cDNA clones generated from capped mRNA and the absence of poly(A) tail on all clones generated from decapped mRNA suggest that decapping occurs after the last phase of deadenylation when the mRNA is fully deadenylated. Moreover, these data show that the degrada-tion of decapped CecA1 mRNA molecules can occur in the 3’ to 5’ direction as demonstrated by the 3’ truncation of 50% of cDNA clones derived from decapped mRNA. A parallel analysis of the 5’ sequence shows that decapped CecA1 mRNA can also be degraded in the 5’ to 3’ direction, whereas as expected, capped mRNA are protected from 5’ exonucleolytic degrada-tion (Fig. 3C, right panel). An analysis of the 5’ region by the RNAfold algorithm (33) predicted two hairpin structures centered at position 48 and 83 (not shown). These structures could contribute to the enrichment of intermediate degradation products observed around these positions in the mRNA (Fig. 3C, right panel).
Mechanism of ARE-mediated Decay in Drosophila

FIGURE 4. Differential localization of CecA1 mRNA in S2 cells treated or not with actinomycin D. A, S2 cells were stimulated with peptidoglycan (PEP) alone or peptidoglycan with the addition of actinomycin D (ActD) for the indicated periods of time. Cells were fixed and hybridized with an in vitro transcribed antisense biotinylated CecA1 riboprobe. Fluorescent labeling was performed with the TSA-Oregon green labeling kit (Molecular Probes). Samples were mounted in Dako fluorescent mounting medium in the presence of DAPI and observed at 1000× magnification to count CecA1 mRNA-positive granules. D, cells were treated as in A. For each experimental condition, 300 cells were observed individually at 630× magnification to count CecA1 mRNA-positive granules.

and after 4 h, this accumulation was approximately four times lower in cells treated with a dsRNA directed against dTTS11 as compared with untreated cells (Fig. 4F).

Similarly, depletion of CAF1 decreased CecA1 mRNA accumulation in P-bodies. In mammals, it has been reported that the formation of P-bodies requires active deadenylation (9). Because we have shown that dTTS11 induces CecA1 mRNA deadenylation, our observations that dTTS11 and CAF1 depletion inhibits the accumulation of CecA1 mRNA in P-bodies could be the consequence of the inhibition of the deadenylation process rather than the ability of dTTS11 to actively target ARE mRNAs to P-bodies.

We further tested the ability of dTTS11 to directly address CecA1 mRNA to P-bodies by comparing the localization of CecA1 mRNA in cells overexpressing a GFP-dTTS11 fusion protein or GFP alone. In both conditions, no CecA1 mRNA granule is observed in the absence of actinomycin D (Fig. 4G, pictures 1–4 and 8–11). Upon treatment with actinomycin D after peptidoglycan stimulation, the number of cells containing detectable CecA1 mRNA in cytoplasmic foci is markedly reduced in cells transfected with GFP-dTTS11 as compared with cells transfected with GFP alone (Fig. 4G and H), correlating with the highest degradation rate of the CecA1 mRNA in the presence of high amount of dTTS11 (Fig. 1A). Therefore, contrary to the observations made for TTP in mammalian cells (18), overexpression of dTTS11 in Drosophila cells is not able to induce the accumulation of an ARE mRNA in P-bodies. The ability of ARE mRNAs to localize in P-bodies in the presence of actinomycin D in S2 cells could be a consequence of the deadenylation induced by dTTS11 protein and the inhibition of degradation potentially induced by actinomycin D.

dTTS11 Stimulates the Polysomal Deadenylation of ARE-containing mRNA—Because dTTS11 does not induce CecA1 mRNA accumulation in specialized degradation structures such as P-bodies, we determined whether it promotes mRNA deadenylation once these are released in the cytoplasm or whether this protein targets polysome-associated mRNA molecules. Therefore, we analyzed the polysome distribution of CecA1 mRNA in S2 cells transfected with GFP or dTTS11 expression constructs. Cells were stimulated with peptidoglycan for 1 or 3 h, and cytoplasmic extracts were separated by ultracentrifugation on sucrose gradients. In cells overexpressing a GFP control construct and stimulated with peptidoglycan for 1 h, CecA1 mRNA is mainly distributed in 80S and polysomal fractions, whereas partially degraded mRNAs are found in the ribosome-free fraction (Fig. 5A, left panel). Interestingly, a minor shortened form of CecA1 mRNA is also present in the ribosomal fractions. This minor form co-migrates with an in...
vitro deadenylated CecA1 mRNA, suggesting that the deadenylated form of CecA1 mRNA can be found in association with the translational machinery. The ectopic expression of dTIS11 strongly enhances the proportion of this fully deadenylated form of CecA1 mRNA in the 80 S and polysomal fractions (Fig. 5, right panel, and Fig. 5D). Upon dissociation of polysomes by puromycin, all CecA1 mRNA molecules were shifted toward the top of the gradient (data not shown), further confirming that both adenylated and deadenylated CecA1 mRNA molecules can be found in association with polysomes. Stimulation of the cells with peptidoglycan for 3 h further enhances the deadenylated form of CecA1 mRNA associated with polysomes (Fig. 5, B and C, compare left panels). As described in Fig. 1, dTIS11 overexpression greatly increases the degradation of CecA1 mRNA after 3 h of stimulation with peptidoglycan, and this mRNA is no more detectable in polysomes in such experimental conditions (not shown). Altogether, our data indicate that dTIS11 promotes the deadenylation of CecA1 mRNA while still associated into polysomes and accelerates its degradation.

**DISCUSSION**

The destabilization of ARE-containing mRNAs mediated by proteins of the TIS11 family is conserved among eukaryotes. Previous studies have demonstrated that TTP, a human protein of the TIS11 family, induces the degradation of ARE-containing mRNAs through a large variety of mechanisms including deenylation, decapping, and P-body targeting. Sequence alignment of TIS11 family members reveals a high degree of conservation in the CCCH zing finger RNA-binding region (34), whereas other regions of the protein are poorly conserved between invertebrates and mammals, suggesting that the molecular mechanism controlling the activity of TIS11 proteins may have evolved in the course of eukaryotic evolution.

We have investigated the molecular mechanism mediating the action of the *Drosophila* dTIS11 protein on the ARE-containing mRNA coding for the antimicrobial peptide CecA1. We observed that the degradation of CecA1 mRNA is initiated by a two-step deadenylation mechanism. We have shown that CAF1 is required for the late deadenylation of the CecA1 mRNA.
Moreover, CAF1 requires the presence of at least the NOT1 and NOT3 proteins to deadenylate this mRNA. Our data also suggest that the PAN2-PAN3 and CCR4-CAF1-NOT deadenylation complexes play a redundant role at the beginning of the deadenylation process because the initial deadenylation phase is inhibited solely when proteins from both complexes are depleted. Our results corroborate previous data indicating that both the PAN2-PAN3 and the CCR4-CAF-NOT complexes were involved in mRNA deadenylation in Drosophila (35).

The ability of TIS11 family members to enhance the deadenylation of ARE-containing mRNAs has been clearly established in mammals (6, 9). TTP is able to recruit CAF1 deadenylase, and this interaction relies on the recruitment of NOT1 by the C-terminal region of TTP (10–12). We have also observed a co-immunoprecipitation of dTIS11 with the CCR4-CAF1-NOT complex (data not shown), suggesting that the ability of TIS11 proteins to recruit the CCR4-CAF1-NOT complex on ARE-containing mRNAs has been conserved from invertebrates to mammals. By depletion of CCR4-CAF1-NOT complex subunits, we observed that NOT1 and NOT3 are required for the second deadenylation phase of CecA1 mRNA, whereas depletion of NOT4 and CAF40 does not alter this process. Although we cannot exclude the requirement of NOT4 and CAF40 on the basis of RNAi depletion experiments, our results confirm previous observations that CAF1, NOT1, NOT2, and NOT3 are required for the general deadenylation mechanism of mRNA in Drosophila and that NOT4 and CAF40 do not seem necessary for this process. Moreover, NOT4 was not detected as a stable component of the CCR4-CAF1-NOT complex (31).

We observed that overexpression of TIS11 accelerates the decay of CecA1 mRNA (Fig. 1), whereas we have previously shown that dTIS11 is essential for the late, CAF1-dependent deadenylation but is not required for the early deadenylation of ARE mRNAs (23). Therefore, the ability of TIS11 to stimulate the CAF1-dependent deadenylation of ARE mRNAs might be timely controlled. Phosphorylation events could transiently inactivate the ability of TIS11 to stimulate CAF1-induced deadenylation as observed with the TTP protein in mammals (10, 11). Alternatively, when both deadenylation complexes are present, early deadenylation of ARE mRNAs could be preferentially carried out by PAN2-PAN3 and therefore would prevent TIS11 from modulating the early steps of deadenylation in a CAF1-dependent manner. Our observations that a dominant negative form of CAF1 preferentially inhibits the late deadenylation of the CecA1 mRNA but does not alter the early deadenylation step (Fig. 2C) are in accordance with this hypothesis. However, these models are not mutually exclusive, and both mechanisms could coexist in vivo. Overall, our results suggest that dTIS11 enhances a general mechanism of mRNA deadenylation.

In mammals, evidence suggests that other deadenylasates could mediate TTP function. Poly-A specific ribonuclease (PARN) deadenylase has been involved in AMD and was shown to interact with TTP in vitro (7, 8), although this interaction is not detected in cells overexpressing both factors (11). TTP is also able to recruit PAN2 deadenylase in a phosphorylation-dependent manner, suggesting that this enzyme could also participate in TTP-induced mRNA deadenylation (11). In Drosophila, the absence of PARN ortholog and the implication of the PAN2-PAN3 complex in the initial dTIS11-independent deadenylation phase of CecA1 mRNA suggest that the activity of dTIS11 protein is concentrated on the second deadenylation phase of ARE-containing mRNAs, where it enhances the CAF1-induced deadenylation of ARE-containing mRNA.

In eukaryotes, the removal of the cap structure is believed to occur on transcripts with short poly(A) tails but not necessarily on fully deadenylated mRNAs (30). In our cRACE analysis, the absence of residual poly(A) tail on decapped CecA1 mRNA and the systematic presence of a poly(A) tail on capped mRNA suggest that decapping occurs only on fully deadenylated CecA1 mRNA molecules. These observations could reflect the fact that the accelerated deadenylation induced by dTIS11 reduces the probability for the decapping machinery to trap an mRNA molecule before completion of the deadenylation process. Alternatively, partially deadenylated CecA1 mRNA could remain inaccessible to the decapping machinery. Because depletion of dTIS11 leads to the accumulation of partially deadenylated CecA1 mRNA (23), whereas decapping occurs on fully deadenylated messenger RNAs, it is unlikely that dTIS11 influences the decapping of ARE-containing mRNA in Drosophila, contrary to what has been observed with TTP (15).

By depleting components of the decapping machinery, we have observed the accumulation of a fully deadenylated form of CecA1 mRNA. Moreover, by the cRACE approach, we have observed the degradation of CecA1 mRNA core only of decapped mRNA, suggesting that decapping is required for mRNA degradation. Sequencing of cRACE cDNA clones also revealed that the core of CecA1 mRNA can be degraded both from the 5’ and from the 3’ ends. In mammals, ARE mRNA silencing involves the delivery of transcripts to P-bodies by TTP or by the related protein BRF-1 (18). In S2 cells, we detected CecA1 mRNA in P-bodies at late stages of mRNA degradation only when cells were treated with the transcription inhibitor actinomycin D. Moreover, this actinomycin D-dependent localization of CecA1 mRNA in P bodies is markedly reduced by dTIS11 overexpression (Fig. 4, G and H), suggesting that in contrast to its mammalian counterparts, dTIS11 is devoid of the ability to address ARE-containing mRNA to P-bodies. We also observed that preserving CecA1 mRNA from deadenylation by depletion of dTIS11 or CAF1 also abolished its migration into P-bodies in response to actinomycin D (Fig. 4E). Therefore, the migration of CecA1 mRNA into P-bodies in these conditions might be caused by the accumulation of deadenylated CecA1 mRNA in waiting for further degradation due to the saturation of the degradation machinery. In mammals, ARE mRNAs migrate into P-bodies in normal degradation conditions, but the amount of ARE mRNA localized in these structures is enhanced by depleting components of the RNA degradation machinery. Altogether, these observations could reflect the fact that the degradation machinery is more rapidly saturated in mammalian cells than in Drosophila S2 cells.

The general model of mRNA degradation suggests that ribosome dissociation is necessary for mRNA decapping and degradation (36). However, recent studies suggest that in yeast, mRNA degradation can occur in polysomes (37). In the major
eukaryotic mRNA degradation pathway, mRNA decay is concomitant with the second phase of mRNA deadenylation (reviewed in Ref. 29). Our observation that dTIS11 enhances the second phase of Ceca1 mRNA deadenylation before decapping without further addressing to P-bodies suggests that dTIS11 could target Ceca1 mRNA in polysomes. This hypothesis is confirmed by our observation that the second phase of Ceca1 mRNA deadenylation occurs in polysomes and is accelerated by overexpressing dTIS11. In agreement with this observation, both the CCR4-CAF1-NOT complex and TTP have been detected in polysomes (38, 39). Therefore, the polysomal deadenylation of ARE-containing mRNAs might represent the ancestral function of TIS11 family members, whereas new functions could have emerged in the course of evolution leading to the optimization of AMD.

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