The Ccr4-Not complex is a key regulator of eukaryotic gene expression

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The Ccr4-Not complex is a multisubunit complex present in all eukaryotes that contributes to regulate gene expression at all steps, from production of messenger RNAs (mRNAs) in the nucleus to their degradation in the cytoplasm. In the nucleus it influences the post-translational modifications of the chromatin template that has to be remodeled for transcription, it is present at sites of transcription and associates with transcription factors as well as with the elongating polymerase, it interacts with the factors that prepare the new transcript for export to the cytoplasm and finally is important for nuclear quality control and influences mRNA export. In the cytoplasm it is present in polysomes where mRNAs are translated and in RNA granules where mRNAs will be redirected upon inhibition of translation. It influences mRNA translatability, and is needed during translation, on one hand for co-translational protein interactions and on the other hand to preserve translation that stalls. It is one of the relevant players during co-translational quality control. It also interacts with factors that will repress translation or induce mRNA decapping when recruited to the translating template. Finally, Ccr4-Not carries deadenylating enzymes and is a key player in mRNA decay, generic mRNA decay that follows normal translation termination, co-translational mRNA decay of transcripts on which the ribosomes stall durably or which carry a non-sense mutation and finally mRNA decay that is induced by external signaling for a change in genetic programming. Ccr4-Not is a master regulator of eukaryotic gene expression. © 2016 The Authors. WIREs RNA published by Wiley Periodicals, Inc.

INTRODUCTION

Cells must continuously adapt to changing environmental conditions. They face the challenge of being able to respond rapidly to dramatic alterations in the environment while buffering responses to mild environmental changes. They must achieve homeostasis in all growth conditions at a minimal energy cost. Regulation of gene expression must respond to this fundamental need of the cell. While this is evident in single cell organisms, which will strive to survive in different growth conditions, it is equally critical in multicellular organisms for appropriate development and health. For a very long time it was believed that the major control of gene expression occurred at the first step, namely transcription. However, it has become increasingly evident that gene expression control occurs also in a major way at all subsequent post-transcriptional steps. Studies on messenger RNA (mRNA) export from the nucleus to the cytoplasm and on the post-translational modifications of the chromatin template that has to be remodeled for transcription have provided important insights into the mechanisms that regulate gene expression. The Ccr4-Not complex is a key regulator of eukaryotic gene expression.© 2016 The Authors. WIREs RNA published by Wiley Periodicals, Inc.
nucleus, localization of mRNAs, control of mRNA decay, and translational control have all risen to the forefront of the gene expression field.

The Ccr4-Not complex is a multisubunit protein complex that is conserved in all eukaryotes and contributes to regulate RNA metabolism at all steps, from synthesis to decay. It has gradually emerged as an essential regulator of gene expression homeostasis in eukaryotes. After the three first reviews more than 10 years ago, many new reviews have been published on different aspects of this complex in the last 2 years. In this review, after a brief biochemical and genetic description of the complex I will summarize the evidence for Ccr4-Not roles at various steps of the gene expression pathway.

BIOCHEMICAL AND GENETIC DESCRIPTION OF THE Ccr4-Not COMPLEX

The Ccr4-Not complex consists of a scaffold protein, Not1, and a number of evolutionarily conserved core proteins that dock onto Not1 (Table 1 and Figure 1). The main core proteins are the Not2 and Not5 heterodimer, the Caf40 subunit and the Ccr4 and Caf1 catalytic subunits. The Not4 RING E3 ubiquitin ligase is a conserved protein, but it is only a stable subunit of the complex in yeast. However, its function is conserved since the human protein complements a deletion of NOT4 in yeast. Yeast has an additional subunit, Caf130, and it carries Not3, a protein that is homologous to Not5 and most likely originates from a gene duplication event. It is unfortunate that the ortholog in higher eukaryotes was named after Not3 rather than after Not5, since it seems that Not3 is functionally much less relevant than Not5 in yeast. In Drosophila, there is no easily recognizable ortholog of Caf130. Instead there are two additional subunits called Not10 and Not11. The human complex is similar in composition to the one in flies, but there are alternatively used genes encoding Caf1 and Ccr4 orthologs (Table 1). Additionally there is a mammalian-specific protein TAB182. The subunits are known under different alternatively used names (Table 1) so I will use the core names defined in Table 1 throughout the review.

The docking of Ccr4-Not subunits onto the Not1 scaffold has been well characterized, the most extensively in Drosophila, but also in yeast and human (Figure 1A). Several crystal structures of domains of Not1 with their interaction partners are available (Ref 2 and reviewed in Ref 8). We lack structural information about species-specific subunits and some domains are not included in the available structures. Our current understanding of the overall architectural organization of the Ccr4-Not complex is a low-resolution electron microscopy reconstruction of the complex purified via Not1 from yeast after chemical cross-linking. It indicates an L-shaped structure with two arms of similar lengths of 180–190 Ångström (Figure 1(b)).

The structural data available for the Ccr4-Not complex has greatly increased very recently but we are still far from a comprehensive structural understanding of this complex. Moreover, we still do not know whether there are multiple functional forms of the complex. Core complexes of about 1 MDa, and larger

| Table 1 | Name of Ccr4-Not Genes in Yeast, Flies, and Human |
|-------|----------------------------------|
| Core Name | Drosophila Gene Name | Saccharomyces cerevisiae Gene Name | Homo sapiens Gene Name |
| Not1 | Not1 | NOT1/CDC39 | CNOT1 |
| Not2 | Regena (Rga) | NOT2/CDC36 | CNOT2 |
| Not5 | Not3 | NOT5 | CNOT3 |
| Caf1 | Pop2 | CAF1/POP2 | CAF1a/CNOT7/CAF1 and CAF1b/CNOT8/POP2/CALIF |
| Ccr4 | twin | CCR4 | CCR4a/CNOT6C and CCR4b/CNOT6L |
| Caf40 | Rcd1 | CAF40 | CNOT9/Rcd1/CAF40/RQCD1 |
| Not4 | Not4 | NOT4/MOT2/SIG1 | CNOT4 |
| Not10 | Not10 | |
| Not11 | Not11 | |
| Caf130 | CAF130 | |
| Not3 | NOT3 | |
| TAB182 | TAB182 | |

I chose Not5 as the core name for fly Not3, human CNOT3, and yeast NOT5 because yeast Not5 rather than Not3 is certainly the yeast functional ortholog of the fly and human proteins.
complexes with all core subunits, are found in cells of different eukaryotes, but the subunits are additionally detectable in smaller complexes. It is not known whether this reflects partial dissociation of Ccr4-Not complexes during in vitro work or instead partial complexes that truly exist and have functional relevance in vivo.

Genetic Description of the Ccr4-Not Components

Ccr4 on one hand, and the five Not proteins on the other, were first identified by different genetic selections in the yeast *S. cerevisiae*. The Caf proteins were isolated by virtue of their interaction with Ccr4. Caf1 has the same mutant phenotypes as Ccr4, but neither do Caf40 nor Caf130. Deletions of Ccr4 or Caf proteins do not pass the genetic selection that led to the identification of the Not proteins and deletions of the Not proteins do not pass the selection that led to the identification of Ccr4 (Refs 24,26 and unpublished observations). This was early genetic evidence that the Ccr4-Not complex is multifunctional.

Not1 is the only essential protein in yeast, but the deletion of Not2, Not4, or Not5 leads to a very...
slow growth phenotype at permissive temperature and the proteins are essential for growth at high temperature.21 The deletions of Caf1 and Ccr4 display only mild phenotypes for yeast grown on rich media in comparison to the deletions of the Not proteins (Ref 21 and unpublished observations). Deletion of Caf40 or Caf130 does not display any identified growth phenotype (unpublished observations). However mild increase or decrease of sensitivity to a variety of different conditions have been reported from global studies and are summarized on the Saccharomyces genome database (www.yeastgenome.org).

As we progress in our characterization of the Ccr4-Not complex functionalities, the early genetic experiments in yeast should not be forgotten, in particular the phenotypes of double deletion mutants. A double mutant that grows no worse than the single mutants indicates components likely to work together. This is true for Ccr4 and Caf1, as well Not2, Not3, and Not5, but deleting either Ccr4 or Caf1 with Not2 or Not5 is lethal.21 The deletion of Not4 is lethal when combined with that of either Ccr4 or Not3, indicating a different functionality, and this is very consistent with the fact that Not4 is not a stable subunit of the complex in flies or human. This also indicates that Not3 has maintained some functional relevance in yeast.21

Cellular Localization of the Ccr4-Not Complex
Ccr4-Not components have been detected both in the cytoplasm and in the nucleus.26,30–32 One study showed that the localization of a Ccr4-Not subunit was regulated according to the stage of the cell cycle, with accumulation in the nucleus in G0 or G1, but mostly cytoplasmic distribution by the time cells are in S phase.33 In the cytoplasm, Ccr4-Not subunits are present in polysomes where mRNAs are being translated,34,35 and they have been detected in processing bodies (P-bodies),36 sites to which mRNAs that undergo inhibition of translation, decapping and/or initiation of degradation, delocalize.

FUNCTIONS OF THE Ccr4-Not COMPLEX ALONG THE GENE EXPRESSION PATHWAY
mRNA Production in the Nucleus

Chromatin Status
DNA packaging into nucleosomes and organization into chromatin fibers generates constraints for transcription. Chromatin remodelers are required for RNA polymerase II (RNAPII) to access the DNA. Post-translational modifications of histone tails constitute a code that contributes to transcription by serving as a landing platform for protein complexes such as ATP-dependent chromatin remodeling complexes. Two such histone marks are under the control of the Ccr4-Not complex. Histones H3 and H4 are globally hypoacetylated in yeast cells lacking Not4 or Not5.37 Gcn5 is the histone acetyltransferase (HAT) that acetylates histones within nucleosomes when associated with other proteins in HAT complexes. Gcn5 purified from cells lacking Not5 can acetylate free histones but it fails to acetylate nucleosomes in vitro. Another chromatin mark that is affected by the Ccr4-Not complex is histone H3 lysine four tri-methylation. This mark is very much reduced in cells lacking Not4 or Not538,39 probably because Not4 is responsible for the turnover of the Jhd2 demethylase that demethylates histone H3.40

Transcription Initiation
Ccr4-Not subunits are present at sites of transcription and can be cross-linked to promoters in yeast and in higher eukaryotes.41–46 Tethering of Ccr4-Not complex subunits to promoters can activate or repress transcription.47,48 In higher eukaryotes, a couple of examples have suggested that Ccr4-Not subunits can interact with transcription factors and repress their ability to activate transcription. For instance Not1 interacts in a ligand dependent manner with the retinoid acid X receptor49 and Caf40 binds transcriptional activators such as c-Myb and AP-1.50 Several transcription factors or transcriptional regulators are targets of the Not4 ubiquitin ligase in yeast or mammals. Examples are the DNA binding activator Yap1 in yeast51 or the Paf1 subunit of the PAF complex in mammalian cells.52 This suggests that transcription factors recruit Ccr4-Not subunits to promoters but can also be regulated by Not4.

Besides its effect on specific transcription factors, the Ccr4-Not complex is important for the relative presence of TFIID and SAGA at promoters.33,54 TFIID and SAGA are general transcription complexes that can both recruit the TATA binding protein to promoters, but TFIID is generally present at constitutive housekeeping genes, while SAGA is mostly present at highly inducible genes. Ccr4-Not subunit crosslinking is enriched at SAGA-dependent genes43 and genome-wide gene expression analyses of ccr4-not mutants indicate that SAGA-dependent genes are mostly affected.55 Numerous studies describe both genetic and physical interactions of Ccr4-Not subunits with TFIID or SAGA. However,
we still do not know whether presence of Ccr4-Not at the sites of transcription is relevant for SAGA-related phenotypes.

The connections between Ccr4-Not and transcription initiation are summarized in Figure 2.

**Transcription Elongation**

Strains harboring mutations in CCR4-NOT genes are sensitive to inhibitors of transcription elongation, namely 6-azauracil and mycophenolic acid. In addition the deletion of CCR4-NOT genes displays a synthetically enhanced slow growth phenotype when combined mutations in transcription elongation factors. *In vivo* evidence for a role of the Ccr4-Not complex in transcriptional elongation is based upon an assay, which monitors the last wave of RNAPII across a long gene after its initiation is repressed. This wave clears the gene more slowly in ccr4-not mutants compared to the wild-type. In another assay the *in vivo* role of Ccr4-Not on transcription elongation was evaluated by measuring gene-length dependent accumulation of mRNA (GLAM). Changes in GLAM were identified in ccr4-not mutants. *In vitro* Ccr4-Not can interact with transcription elongation complexes prepared from highly purified yeast RNAPII. This requires the separable Rpb4/7 module of the polymerase, in accord with a reported interaction between Rpb4 and both Not3 and Not5. Ccr4-Not directly affects transcription elongation by rescuing backtracked RNAPII. It has no effect on the transcription rate of un-arrested RNAPII. Interestingly, the ability of Ccr4-Not to rescue backtracked polymerase depends upon the length of the transcript suggesting that an association of Ccr4-Not with the new transcript contributes to promote transcription elongation. Another transcription factor that directly promotes transcription elongation and rescues backtracked RNAPII is TFIIS. It does so by stimulating intrinsic nucleolytic activity of RNA-P II and transcript cleavage (reviewed in Ref 60). This mechanism is not shared by Ccr4-Not, but Ccr4-Not has the additional effect of improving association of TFIIS to the elongating polymerase. The effects of Ccr4-Not on transcription elongation are summarized in Figure 3.

**Quality Control and mRNA Export**

Processing and maturation of newly produced transcripts depends upon many factors that assemble into heterogenous ribo-nucleoprotein particles (hnRNPs)
to prepare the mRNA for its export to the cytoplasm. The mRNPs are sensed in a quality control process to ensure that only functional mRNAs are exported. When export components are defective a number of phenotypes are observed, some of which depend upon the nuclear exosome subunit Rrp6 indicating that this exonuclease is important for the mRNA quality control process. The Ccr4-Not complex interacts with proteins that participate in maturation of mRNPs coupled to their export, such as the nuclear poly(A) binding protein Nab2 or Hrp1 of the THO complex, or Mlp1 of the inner basket of the nuclear pore complex (NPC). The deletion of Ccr4-Not subunits displays synthetic growth phenotypes when combined with the deletion of export proteins such as Mft1 of the THO complex, or when combined with the deletion of Rrp6. Yet Rrp6-dependent quality control defects observed in mft1 mutants are rescued by the co-deletion of Ccr4-Not subunits. This might indicate that the double mutants export defective mRNAs and hence suggests a role for Ccr4-Not in the nuclear quality control mechanism. In line with this idea, overexpression of Not4 exacerbates the poly(A) mRNA export defect observed in an NPC mutant. Moreover polyadenylated snRNAs indicative of defective Rrp6-dependent nuclear quality control accumulate in ccr4-not mutants. These interactions of Ccr4-Not related to mRNA export and nuclear quality control are depicted in Figure 4.

**Cytoplasmic mRNA Fate**

Stability and translation of mRNAs in eukaryotes depend upon the 7-methylguanylate cap at the 5’ end of the mRNA and the poly(A) tail at the 3’ end of the mRNA, the former added co-transcriptionally to the new message and the latter added after transcript cleavage. In mammalian cells the mRNA cap is first bound in the nucleus by the cap binding protein (CBP) that will support a pioneer round of translation in the cytoplasm. The eIF4E translation initiation factor replaces subsequently CBP and will direct steady state rounds of translation. CBP and eIF4E particles are associated with nuclear and cytoplasmic poly(A) binding proteins (PABP), respectively. CBP is not essential in yeast and newly produced mRNAs can be bound directly by eIF4E.
Cytoplasmic mRNAs cycle between poly-somes and RNA granules. The mechanisms and directionality of these movements are not fully understood. Translationally repressed mRNAs will accumulate in granules and RNA granules can release transcripts to allow their translation. Functional mRNAs arriving in the cytoplasm are not necessarily recruited immediately into polysomes to be translated, nor are they necessarily immediately degraded after leaving polysomes.

The best-characterized granules are the P-bodies (PBs) and stress granules (SGs). They have features in common and specific features. PBs contain the mRNA decay machinery whereas SGs contain many translation initiation factors. SGs form in the absence of new translation. They harbor mRNAs from pre-existing polysomes and arise in response to inhibition of translation initiation, followed by polysome elongation and a progressive loss of ribosomes. PBs are also formed from translationally repressed mRNPs onto which complexes of decapping components are recruited. mRNAs that exit translation can first go to PBs but then remodel to acquire translation initiation factors and move to SGs where translation is repaired and mRNAs can be released when they are ready to translate again. SGs may also form independently of PBs, and mRNPs may enter SGs prior to entering polysomes when produced under conditions of translation inhibition. Globally, the role of the RNA granules is still not entirely clear. They might induce a local concentration of proteins favorable for mRNA decay (PBs) or reinitiation of translation (SGs), or provide a buffering system to maintain a proper ratio of translational capacity to the pool of mRNAs that are translating (for instance to avoid titrating limiting factors), or finally serve as a reservoir of translatable mRNAs (for review see Ref 66).

The Roles of Ccr4-Not in the Translation Process

Ccr4-Not and mRNA Translatability
Recent data has indicated that translatability of several mRNAs, defined as the presence of
polyadenylated forms of those specific mRNA in polysome fractions relative to total levels, is altered in yeast lacking Not5, some increased, others decreased.\textsuperscript{59} The mechanism is unknown. Not5 might be altering the cycling of specific mRNAs between RNA granules and polysomes, or regulating the distribution of specific newly produced mRNAs to polysomes or RNA granules, or both. Not5 has RNA binding activity\textsuperscript{67} and it is also important for integrity of the Ccr4-Not complex that contains other proteins with RNA binding activity.

**Ccr4-Not and Translation**

The level of total polysomes is reduced in yeast cells lacking Not2, Not4 or Not5.\textsuperscript{35} These strains grow slowly, so it is difficult to definitively conclude that reduced polysomes are not a consequence of slow growth. However several observations in yeast are consistent with a role of the Ccr4-Not proteins during normal translation. All Ccr4-Not subunits are detected in polysomes\textsuperscript{35} and newly produced proteins aggregate massively in \textit{not2}, \textit{not4}, and \textit{not5} mutants. This suggests that folding or protein interactions of newly produced proteins are defective.

The importance of Not5 during translation is exemplified by the need for Not5 during \textit{de novo} synthesis of the largest subunit of RNAPII, Rpb1.\textsuperscript{59} Rpb1 tends to aggregate and must associate with Hsp90 and a co-chaperone called R2TP to form a soluble assembly-competent intermediate.\textsuperscript{68} Not5 is needed for the presence of R2TP at ribosomes producing Rpb1 (Figure 5(a)), and as a consequence it is needed for efficient assembly of RNAPII. Assembly of another multisubunit complex, the proteasome, is also dependent upon the Ccr4-Not complex, in particular upon Not4.\textsuperscript{69} Not4 is needed for assembly of the proteasome lid, and it is important for a proteasomal chaperone, Ecm29, to associate with proteasome subunits. Interestingly in fission yeast the ortholog of Ecm29 is present in polysomes producing proteasomal subunits, suggesting that Not4 may be contributing to co-translational assembly of the proteasome with one of its chaperones (Figure 5(b)).

The Not4 E3 ligase ubiquitinates a ribosomal protein Rps7A and a ribosome associated chaperone called nascent polypeptide associated complex (NAC).\textsuperscript{35,70} This observation has fostered interest in Not4 as a possible translational regulator. Such a role for Not4 has been studied with artificial mRNA templates encoding two open reading frames (ORFs) with an intervening stretch of RNA coding for 12 subsequent lysines known to provoke ribosome stalling.\textsuperscript{34,71,72} Initially Not4 was reported to ubiquitinate the stalled peptides,\textsuperscript{34} but subsequently described instead to have an impact on levels of the mRNAs with stalled ribosomes.\textsuperscript{72} A third study suggested that in the absence of Not4 translation levels were reduced\textsuperscript{71} in contrast to a fourth study indicating that Not4 was needed to repress translation.\textsuperscript{73} Not4 was also described to participate in global translational repression after nutrient withdrawal in yeast, when reduced concentrations of amino acyl tRNAs are thought to promote generalized ribosome stalling.\textsuperscript{73} These different possible effects of Not4 in co-translation quality control are summarized in Figure 5(c).

A role for the Ccr4-Not complex in translational repression upon non-sense mediated mRNA decay (NMD) or upon its’ tethering to mRNAs by RNA binding proteins (RBPs) or by the micro-RNA machinery has been characterized. This translational repression ability is independent of its deadenylase function. Indeed, tethering of the Ccr4-Not complex to mRNAs lacking a poly(A) tail \textit{in vitro}\textsuperscript{74} and \textit{in vivo}\textsuperscript{75} can induce translation arrest, and in yeast repression of translation by the Mpt5 RBP occurs via Caf1 in cells that lack the active deadenylase subunit Ccr4.\textsuperscript{76} A mechanism for translation repression has recently emerged from studies in \textit{Xenopus}\textsuperscript{77} and \textit{Drosophila}.\textsuperscript{78} The elf4E transporter protein (4E-T), a member of the elf4E binding proteins, can mediate translational repression of tethered Ccr4-Not. 4E-T interacts with elf4E and with the mRNA decay machinery, both with components interacting with the 3’ ends of the mRNAs (such as the LSM1-7-PAT1 complex) and with the 5’ cap (such as the dead box protein DDX6 known as Dhh1 in yeast). 4E-T also interacts with the Ccr4-Not complex itself\textsuperscript{77-79} and in yeast it was described instead to have an impact on levels of the mRNAs with stalled ribosomes.\textsuperscript{72} A third study suggested that in the absence of Not4 translation levels were reduced\textsuperscript{71} in contrast to a fourth study indicating that Not4 was needed to repress translation.\textsuperscript{73} Not4 was also described to participate in global translational repression after nutrient withdrawal in yeast, when reduced concentrations of amino acyl tRNAs are thought to promote generalized ribosome stalling.\textsuperscript{73} These different possible effects of Not4 in co-translation quality control are summarized in Figure 5(c).

The interaction between elf4E and 4E-T is necessary for deadenylation-independent translational repression by the Ccr4-Not complex\textsuperscript{77} and for mRNA decay.\textsuperscript{27} The proposed chain of events is that RBPs recruit Caf1 that in turn recruits Not1 to the mRNA that will then recruit 4E-T. 4E-T will physically link the 3’ terminal mRNA decay machinery to the 5’ cap (Figure 5(d)).

Not1 also directly interacts with DDX6 via its MIF4G domain (see Figure 1).\textsuperscript{80-82} In yeast it was shown that Dhh1 is an activator of the decapping complex, but that it also functions in translation repression.\textsuperscript{83} It accompanies slowly translocating ribosomes and when tethered to an mRNA leads to an accumulation of ribosomes on the mRNA.\textsuperscript{84} It has been suggested that translational repression after nutrient withdrawal needs Dhh1 like Not4.\textsuperscript{73} However overexpression of Dhh1 rather than deletion of Dhh1 passes the genetic selection that led to identification of the NOT genes,\textsuperscript{85} indicating that Dhh1 and...
Not4 might have opposite rather than similar effects on gene expression.

The described positive and negative roles of the Ccr4-Not complex during translation seem contradictory, but they may in fact be reconcilable. The necessity for proteins to interact with partners cotranslationally could cause ribosome pausing. If partners fail to appear in a timely manner, the Ccr4-Not complex might contribute to translational repression and mRNA decay. However, the Ccr4-Not complex might initially preserve the ribosome-mRNA complex to give time for partners to interact co-translationally, and only if the delay is too long allow the co-translational responses to proceed (Figure 5(e)). An imbalance in this mechanism would lead to the production of mis-folded proteins.

Since the Ccr4-Not complex is important for control of poly(A) tail length, it is to be expected that...
it will play an important role in the movement of mRNAs from the translating pool to the pool directed to RNA granules. In HeLa cells the depletion of Not1 suppresses largely the formation of PBs. Ccr4-Not subunits localize to PBs in stressed yeast but they do not seem to be a major component of PBs in flies.

The Roles of Ccr4-Not in mRNA Degradation

Since stability and translation of mRNAs depend upon the cap and the poly(A) tail of the mRNA, translation and decay are intimately connected and generally in competition with each other. The major mRNA decay pathway in eukaryotes involves removal of the cap followed by $5'$→$3'$ exonuclease degradation. The initial and rate-limiting step in this pathway is deadenylation performed mainly by Caf1 and Ccr4, the major eukaryotic deadenylases.

Initiation of mRNA degradation is intimately connected to the process of translation. Different types of mRNA decay can be distinguished: generic decay of mRNAs that have undergone normal cycles of translation, quality control induced mRNA decay, which occurs upon ribosomal stalling without regular translational termination, and finally induced mRNA decay brought about by tethering of the Ccr4-Not complex to specific mRNAs in response to specific signaling.

Generic mRNA Deadenylation

Not much is known about how the Ccr4-Not complex participates in generic decay of all mRNAs upon translation termination. Either Ccr4-Not is present during translation of all mRNAs in an inactive form, and then is modified upon translation termination to induce deadenylation, or it is recruited upon translation termination to initiate shortening of the poly(A) tail. One study has described a series of coupled events starting with the recruitment of the translation termination factors eRF1 and eRF3, recruitment of the minor deadenylation complex Pan2-Pan3, and finally recruitment of Ccr4-Not by the BTG/TOB family of proteins via their interaction with PABP. BTG/TOB proteins are absent in budding and fission yeast and they are not constitutively expressed in mammalian cells (for reviews see Ref 89). Thus, while this mechanism (Figure 6(a)) might be a response that leads to a general mRNA deadenylation response in metazoans, it cannot be the mechanism for generic mRNA decay.

Non-sense Mediated mRNA Decay (NMD)

NMD results from the presence of an aberrant stop codon within the open reading frame. The recruitment of the Ccr4-Not complex to initiate deadenylation of NMD targets has been characterized in vertebrates (Figure 6(b)). During the pioneer round of translation, CBP interacts with the conserved Upf1 protein that will become phosphorylated. This promotes recruitment of the Smg 5, 6, and 7 proteins. Smg6 is an endonuclease and the Smg5-Smg7 heterodimer recruits the Ccr4-Not complex via interactions between Smg7 and Pop2/CNOT8 (but not its homolog Caf1/CNOT7). The recruitment of Ccr4-Not by Smg7 is critical for NMD in cells lacking Smg6, indicating that Smg6 and the Ccr4-Not complex act redundantly to promote mRNA decay of NMD substrates. In yeast, Upf1 is not phosphorylated and there are no orthologs of Smg5, 6, and 7. NMD targets are rapidly decapped without prior need for deadenylation. A specific role for Ccr4-Not in NMD in yeast is uncertain.

Targeted mRNA Decay

Certainly the most well established function of the Ccr4-Not complex is its ability to repress gene expression upon its targeted recruitment to the 3' untranslated regions (UTRs) of mRNAs. The complex is recruited by RBPs or the micro-RNA machinery that associate with specific sequences at the 3' end of the mRNAs. Experiments in yeast have demonstrated that the tethering of the Ccr4-Not complex is sufficient to destabilize an mRNA in vivo. This has established that Ccr4-Not is constitutively active to deadenylate an mRNA provided that the deadenylase subunits are recruited to the mRNA (Figure 6(c)).

Many sequence specific RBPs that recruit Ccr4-Not have been described. They either bind Not1 directly, or they recruit Not1 by binding to other subunits of the Ccr4-Not complex. They can also interact with proteins such as the BTG/TOB proteins that in turn associate with a Ccr4-Not subunit. Three binding partners have been characterized structurally. Tristetraprolin (TTP) interacts with Not1 and associates with ARE elements in the 3' UTRs of genes such as TNFa, c-Fos, and c-Myc. Nanos1 is important for germ cell line maintenance and survival and interacts also with Not1. Finally, BTG/TOB proteins interact with Caf1 homologs to mediate their antiproliferative activity.

The interaction of the miRNA machinery with the Ccr4-Not complex and mRNA targets has been well studied. It is a multilayered process starting with
base pairing of the Argonaute (Ago1) miRNPs to the target mRNA. This leads to recruitment of the GW182 protein that in turn will recruit deadenylases including Ccr4-Not. This recruitment involves multiple regions in GW182, a 400 amino acid C-terminal region called CED being the most important one. It can recruit Not1 by interaction with a central domain of Not1 in three different ways, one being mediated by Caf40 (Figure 6(d)). These interactions involve tryptophan residues in GW182 and Trp-binding pockets in Not1 and Caf40. In turn Not1 is associated with Caf1 and Ccr4 that will deacylenlate the mRNA, and it can bind DDX6 that has the effects discussed above (reviewed in Ref 94).

Cross-Talk Between Different Levels of Gene Expression Via Ccr4-Not

The different functions of the Ccr4-Not complex at different stages of gene expression described above are compatible with the presence of the subunits both in the nucleus and in the cytoplasm. Moreover the reported change in localization of a Ccr4-Not subunit in response to the stage of the cell cycle is compatible with the ability of the Ccr4-Not complex to differentially modify its role along the gene expression pathway in response to the need of the cell. In addition the Ccr4-Not subunits harbor a multitude of post-transcriptional modifications that have not yet been functionally characterized.

One can imagine that the Ccr4-Not complex acts independently at different steps along the gene expression pathway, or instead that its functions at the different steps are physically connected (Figure 7). It has been proposed that gene-specific regulators could recruit specific Ccr4-Not subunits to promoters for subsequent post-transcriptional regulation. Moreover the Ccr4-Not complex can interact with the transcription elongation complex and with

Deadenylation-Independent Decapping

Deadenylation-independent decapping of certain mRNAs has been reported for Not2, Not4, and Not5 in yeast. As mentioned above, Dh1 associates with the Not1 MIF4G domain and can function as a decapping activator. It could be that specific mRNAs need Ccr4-Not to be efficiently decapped because Ccr4-Not interacts with Dh1.
the nascent transcripts. Hence the Ccr4-Not complex may associate with some mRNAs as they are synthesized and then remain bound within the mRNPs exported to the cytoplasm. This may not concern all mRNAs, but only those whose synthesis depends upon TFs that can recruit Ccr4-Not or those synthesized from genes whose transcription elongation is particularly prone to mobilize Ccr4-Not. If mRNAs are bound in the nucleus by Ccr4-Not, probably this is a form of the complex that is not immediately active for induction of deadenylation, translational repression, and decapping.

The Rpb4 subunit of RNAPII was reported to associate with newly produced mRNAs under stress and to be exported with them out to the cytoplasm, where it will contribute to their translation, targeting to PBs and/or decay. Interestingly, Rpb4 is essential for association of Ccr4-Not with elongating RNAPII and Rpb4 export to the cytoplasm depends upon Not5. Hence an intriguing possibility is that Rpb4 could be necessary in the nucleus for the Ccr4-Not complex to associate with new transcripts, allowing Ccr4-Not to module their function post-transcriptionally (Figure 7).

What might be the advantage of imprinting mRNAs in the nucleus with Ccr4-Not? The post-transcriptional fate of such mRNAs may be more immediately determined. Moreover, it provides a mean for the eukaryotic cell to connect physically the nuclear and cytoplasmic stages of gene expression. Changes in transcription levels may be compensated for by opposite changes in mRNA degradation via the Ccr4-Not complex to maintain similar steady state levels of mRNAs, but also by changes in translation to produce similar levels of protein. Such gene expression buffering is known to exist. Rpb4 and the Ccr4-Not complex have been connected to this forward buffering (reviewed in Ref 98). Reciprocally, Ccr4-Not is important during translation for production of the transcription machine (RNAPII), and this provides Ccr4-Not with the ability to communicate the translation status of the cell back to the nucleus.

CONCLUSION

The Ccr4-Not complex intervenes at all stages of the gene expression pathway, helping to produce mRNAs and contributing positively to translation, but also initiating translation repression and mRNA degradation. A major challenge facing our understanding of this master regulator is to determine in
which forms the Ccr4-Not complex acts for each of these functions and how it is modified to change roles. Characterizing the flow of Ccr4-Not between the nucleus and the cytoplasm is an essential part of these questions.

Finally, one has to wonder why such a complex regulator as Ccr4-Not was conserved across the eukaryotic kingdom. The most obvious explanation is that a regulator such as this one is the most efficient to coordinate the different steps of gene expression. It is ideally suited to mediate changes in programs of gene expression, but also ideally designed to buffer undesirable changes in a given step along the gene expression pathway. Not surprisingly, evidence for essential roles of Ccr4-Not in cellular programming and gene expression buffering is increasing.

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FURTHER READING

Many reviews have been written on the Ccr4-Not complex, the first ones about 10 years ago,\textsuperscript{1–3} and recently many more, some of which cover aspects not discussed in this review, such as for instance more detailed structural information on the complex or the physiological roles of Ccr4-Not in different species or genetic programming, that are exciting additional reading.\textsuperscript{4–18}

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