Synchronicity: policing multiple aspects of gene expression by Ctk1

Michael Hampsey¹ and Terri Goss Kinzy²,³

¹Department of Biochemistry, Division of Nucleic Acid Enzymology, University of Medicine and Dentistry of New Jersey-Robert Wood Johnson Medical School, Piscataway, New Jersey 08854, USA; ²Department of Molecular Genetics, Microbiology and Immunology, University of Medicine and Dentistry of New Jersey-Robert Wood Johnson Medical School, Piscataway, New Jersey 08854, USA

Transcription and translation are coordinated events in all organisms. In prokaryotes, the process that couples these two events is clear: The ribosome begins translation of the nascent mRNA while the DNA template is still being transcribed. Indeed, cotranscriptional protein synthesis underlies key regulatory mechanisms in bacteria, including attenuation, the mechanism that regulates RNA polymerase processivity in response to ribosome movement along the mRNA. But how is transcription coordinated with translation in eukaryotic organisms, where mRNA is synthesized in the nucleus and protein synthesis occurs in the cytoplasm? Although these two events are spatially distinct, separated by the nuclear envelope, efficient control of gene expression necessarily requires that transcription and translation be regulated in a coordinated manner. As an example, dFOXO-mediated transcriptional activation produces both an inhibitor of cap-dependent translation, eukaryotic translation initiation factor 4E (eIF4E)-BP, and a form of the insulin receptor mRNA that is translated by a cap-independent mechanism [Marr et al. 2007]. In addition, translation requires a fully and accurately processed mRNA, and has mechanisms to help sense that appropriate processing has occurred.

In the absence of physical coupling of transcription and translation, how are these two processes coordinated in eukaryotes? In this issue of Genes & Development, Röther and Sträßer [2007] report that the Ctk1 kinase, a key enzyme that facilitates passage of RNA polymerase II [Pol II] through specific stages of the transcription cycle, is also found in the cytoplasm associated with ribosomes actively engaged in protein synthesis [Fig. 1]. Their work defines a physiological role for Ctk1 in translation by showing that cellular depletion of Ctk1 decreases total protein synthesis as well as the fidelity of translation elongation. These effects are likely to be a direct effect of Ctk1, since they identified the small ribosomal subunit protein rpS2 as the specific target of the kinase. Moreover, site-directed replacement of rpS2 Ser238, which they identified as the target of Ctk1, results in the same translational defects as depletion of Ctk1. Thus, Röther and Sträßer [2007] propose that Ctk1 piggybacks with either the mRNP particle or with the pre-40S ribosomal subunit to transit the nuclear envelope to regulate protein synthesis in the cytoplasm.

Ctk1 was first identified as a subunit of the yeast CTDK-I complex that catalyzes phosphorylation of the Pol II C-terminal domain [CTD] [Sterner et al. 1995], a reiterated heptapeptide sequence [Tyr1–Ser2–Pro3–Thr4–Ser5–Pro6–Ser7] present at the C terminus of Rpb1 [Kobor and Greenblatt 2002]. The CTD couples Pol II transcription with RNA processing, apparently forming a platform for the association and exchange of transcription and RNA processing factors [Hirose and Manley 2000; Orphanides and Reinberg 2002; Proudfoot et al. 2002; Bentley 2005; Meinhart et al. 2005]. These factors include the 5′-capping enzymes, the splicing machinery, the 3′-end processing complex, and the transcription export complex [TREX] that facilitates mRNA translocation to the cytoplasm. Progression of Pol II through the transcription cycle is accompanied by changes in the phosphorylation status of the CTD. Pol II is recruited to the promoter in an unphosphorylated form [Pol IIA] that becomes extensively phosphorylated at Ser2 and Ser5 during different stages of the transcription cycle. Differential CTD phosphorylation promotes the exchange of initiation and elongation factors at promoter clearance [Pokholok et al. 2002] and the exchange of elongation and 3′-end processing factors at termination [Kim et al. 2004].

CTD phosphorylation is catalyzed by C-type cyclin-dependent kinases [Prelich 2002]. The first of these complexes to be identified is Kin28–Ccl1, which functions as a subcomplex of the general transcription factor TFIIH. Kin28 catalyzes Ser5 phosphorylation coincident with transcription initiation and as a prerequisite for capping [Hengartner et al. 1998; Rodriguez et al. 2000]. The second complex, CTDK-I, is composed of three subunits, the Ctk1 kinase, Ctk2 cyclin, and Ctk3 accessory protein that forms a regulatory complex with Ctk2 [Sterner et al. 1995; Hautbergue and Goguel 2001]. CTDK-I catalyzes Ser2 phosphorylation during elongation, coinci-
the 5' binding protein (Pab) provides communication between mRNAs and 3' polyadenylation, and nuclear export, occurs cotranscriptionally in a manner coordinated by the Pol II CTD. Progression of Pol II through the transcription cycle is regulated by CTD phosphorylation, catalyzed by the Kin28 and Ctk1 cyclin-dependent protein kinases. The processed mRNA is transported to the cytoplasm, where it becomes a substrate for the translational apparatus. The Ctk1 kinase (shaded oval) phosphorylates (P) both the CTD of Pol II on Ser2 in the nucleus and Ser238 of the small ribosomal subunit protein rpS2 in the cytoplasm.

As with every stage of gene expression, accuracy must be maintained. During translation, this includes recognition of the appropriate AUG and the use of the Met-tRNA\textsuperscript{Met} at initiation, cognate aa-tRNA selection and the maintenance of the correct reading frame during elongation, and recognition of one of the three stop codons during termination. Examples of conditions where modifications alter the accuracy of translation are not common.

Post-transcriptional and post-translational modifications of components of the translational apparatus also affect the efficiency and/or accuracy of protein synthesis. The target of rapamycin (TOR) pathway plays a key role in signaling that sufficient nutrients and energy are available for protein synthesis, leading to phosphorylation of rpS6, the eEF2 kinase, elF4E-BP, and other translation factors [Hay and Sonenberg 2004]. Effects on the elongation phase of translation are seen as a result of altered modifications of translation components. Failure to form pseudouridine within the tRNA at the peptidyl transferase center of the ribosome results in reduced translation [King et al. 2003], and loss of other rRNA-modifying proteins affects several forms of translational fidelity [Baxter-Roshek et al. 2007]. Loss of the diphtheria toxin gene expression

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\caption{Coordination of eukaryotic Pol II transcription, mRNA processing, and protein synthesis. RNA processing, which includes 5'-cap addition, splicing, 3' polyadenylation, and nuclear export, occurs cotranscriptionally in a manner coordinated by the Pol II CTD. Progression of Pol II through the transcription cycle is regulated by CTD phosphorylation, catalyzed by the Kin28 and Ctk1 cyclin-dependent protein kinases. The processed mRNA is transported to the cytoplasm, where it becomes a substrate for the translational apparatus. The Ctk1 kinase (shaded oval) phosphorylates (P) both the CTD of Pol II on Ser2 in the nucleus and Ser238 of the small ribosomal subunit protein rpS2 in the cytoplasm.}
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\caption{Ctk1 and gene expression}
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\caption{Figure 1. Coordination of eukaryotic Pol II transcription, mRNA processing, and protein synthesis. RNA processing, which includes 5'-cap addition, splicing, 3' polyadenylation, and nuclear export, occurs cotranscriptionally in a manner coordinated by the Pol II CTD. Progression of Pol II through the transcription cycle is regulated by CTD phosphorylation, catalyzed by the Kin28 and Ctk1 cyclin-dependent protein kinases. The processed mRNA is transported to the cytoplasm, where it becomes a substrate for the translational apparatus. The Ctk1 kinase (shaded oval) phosphorylates (P) both the CTD of Pol II on Ser2 in the nucleus and Ser238 of the small ribosomal subunit protein rpS2 in the cytoplasm.}
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\caption{Ctk1 and gene expression}
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lation components, differ from Ctk1p phosphorylation of rpS2 demonstrated in the work of Röther and Sträßer [2007] in several ways. First, diminished rpS2 phosphorylation associated with either Ctk1 depletion or the Ser238Ala replacement results in reduced translational fidelity at the A site as well as reduced total translation. The increased missense and nonsense errors are not an effect of reduced elongation, as mutants in translation elongation factors that reduce total translation do not always result in altered fidelity. In addition, a phosphorylation event that results in altered translational fidelity is novel. Second, while coordinate regulation of multiple components of the translational apparatus is seen in the TOR pathway, the Ctk1 kinase is not specific to the translational apparatus but also impacts an earlier stage of gene expression. Thus, instead of coordinating a single step of gene expression, Ctk1 might coordinate distinct events in gene expression. The specific effect of Ser238 phosphorylation of rpS2 on ribosome function remains unknown. The fact that the sup4-41 allele of RP52, which encodes a Ser200Tyr replacement near Ser238, is a paromomycin-sensitive omnipotent suppressor, which encodes a Ser200Tyr replacement near Ser238 [Pipperberg et al. 1980]. It is possible that the fidelity is manifest through the A-site-binding elongation factors, especially eEF1A, which is known to affect missense, frameshift, and nonsense suppression [Sandbaken and Culbertson 1988; Dinman and Kinzy 1997; Cavallius and Merrick 1998]. While the location of rpS2 is not within the A site, its location could modulate the accuracy of aa-tRNA selection, consistent with the effect of Ctk1 depletion or mutation of Ser238.

The effects of Ctk1 on capping, splicing, 3’-end processing, and mRNA export can all be explained by the coupling of Pol II transcription with RNA processing, a connection that is mediated by differential phosphorylation of the CTD. But the discovery that Ctk1 catalyzes phosphorylation of a substrate other than Ser2 of the CTD is not the first hint of a broader role played by Ctk1 in gene expression. In an effort to identify additional Ctk1 substrates, Goguel and colleagues [Bouchoux et al. 2004] found that Ctk1 is partially localized to the nucleolus and affects Pol I transcription at the rDNA locus. These results were recently extended by showing that Ctk1 affects the stability of the rDNA tandem repeats [Grenetier et al. 2006]. It seems unlikely that the effect of Ctk1p dephosphorylation on fidelity is related to changes in ribosome synthesis, based both on the results of Röther and Sträßer [2007] and a large bank of ribosome assembly mutants that have been identified but not linked to reduced translational fidelity. Thus, Ctk1 might serve to link or coordinate the path of mRNA from its site of synthesis to its use as a substrate.

How does Ctk1 traverse the nuclear membrane to regulate both transcription and translation? One of the most well-characterized transcription factors in yeast, Pho4, is regulated by nuclear–cytoplasmic translocation by the Msn5 exportin and Pse1 importin [Komeil and O’Shea 2000]. Are there comparable proteins that regulate subcellular Ctk1 localization and under what physiological conditions? The CTD kinase activity of Ctk1 is regulated by the Ctk2–Ctk3 heterodimer [Hautbergue and Goguel 2001]. Do Ctk2 and Ctk3 affect translation? Progression of Pol II through the transcription cycle and the exchange of RNA processing factors also involves CTD dephosphorylation, catalyzed by the Fcp1 and Ssu72 phosphatases [Cho et al. 2001; Mandal et al. 2002, Reyes-Reyes and Hampsey 2007]. Is there a cytoplasmic phosphatase that catalyzes rpS2 dephosphorylation? Finally, in light of the association of mammalian Ctk9 kinase with polyribosomes [Röther and Sträßer 2007] and a large bank of ribosome assembly mutants that have been identified but not linked to reduced translational fidelity, it will be especially interesting to learn whether Ctk9 in the P-TEFb complex also affects protein synthesis.

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