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Making ends meet: coordination between RNA 3'end processing and transcription initiation

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Abstract

RNA polymerase II (RNAPII)-mediated gene transcription initiates at promoters and ends at terminators. Transcription termination is intimately connected to 3’end processing of the produced RNA and already when loaded at the promoter, RNAPII starts to become configured for

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7 this downstream event. Conversely, RNAPII is 'reset' as part of the 3'end processing/termination
8 event, thus preparing the enzyme for its next round of transcription – possibly on the same gene.
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10 There is both direct and circumstantial evidence for preferential recycling of RNAPII from the gene
11 terminator back to its own promoter, which supposedly increases the efficiency of the
12 transcription process under conditions where RNAPII levels are rate limiting. Here, we review
13 differences and commonalities between initiation and 3'end processing/termination processes on
14 various types of RNAPII transcribed genes. In doing so, we discuss the requirements for efficient
15 3'end processing/termination and how these may relate to proper recycling of RNAPII.]
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27 The conversion of genetic information into functional molecules begins with gene transcription. In
28 eukaryotes, multisubunit RNA polymerases (RNAPs) execute the transcription of nuclear-encoded
29 genes. There are five eukaryotic RNAPs - RNAPI, II, III, IV and V, where the latter two only exist in
30 plants^{1,2}. RNAPII appears to be the most versatile of these enzymes, both with respect to the types
31 of transcription units it engages and in terms of the cellular levels, sizes and physical properties of
32 the generated RNA molecules, which include both protein-coding mRNA as well as long- and short
33 non-coding (nc)RNA.
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41 Common to all genes is that they are linear entities confined by promoters and terminators that
42 dictate where transcription starts and ends. Where studied, transcription termination is tightly
43 coupled to the 3'end processing of the produced RNA molecule and the combined process is
44 necessary for release of the transcript and the RNA polymerase from the gene template as well as
45 from each other³⁻⁵. In addition to this crucial step in gene expression, recent studies from several
46 laboratories suggest that RNA 3'end processing/transcription termination can both positively and
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negatively influence initiation of the next round of transcription⁶⁻¹². Such functional interaction between its start and end suggests that a gene functions, at least partially, as a closed circuit.]

RNAPII transcription – from beginning to end

Transcription initiation

RNAPII transcription initiates at the gene promoter, which is usually positioned upstream, but in some cases immediately downstream, of the transcription start site (TSS)^{13,14}. Although the exact mechanism by which transcription is initiated is far from uniform among all promoters, the first step of RNAPII transcription initiation involves the formation of a pre-initiation complex (PIC) consisting of the promoter DNA, general transcription factors (GTFs) and RNAPII – a process that is stimulated by, or perhaps even dependent on, transcriptional activators, chromatin remodeling factors and the mediator complex. *De novo* PIC assembly takes place through the sequential binding of GTFs at the promoter core elements, ending with the recruitment of RNAPII¹⁵⁻¹⁸. After RNAPII is released from the promoter and starts elongating, a sub-complex of the PIC called the re-initiation complex can remain behind and facilitate transcription re-initiation¹⁹.

Transcription termination

After transcribing the gene body, RNAPII reaches the terminator region downstream of the RNA 3'end processing site. It is generally perceived that transcriptional termination is tightly coordinated with RNA 3'end processing, but no unifying model exists for termination of transcription on all types of RNAPII genes. The process is best described for genes encoding polyadenylated RNA (hereafter referred to as 'polyA genes'), for which two models have been proposed. The 'allosteric' model suggests that RNAPII undergoes conformational changes due to dissociation of elongation factors and/or association of termination factors upon passage of the 3'end processing signal, which

ultimately leads to pausing and termination of RNAPII²⁰⁻²³. Conversely, the 'torpedo'-model, is based on the fact that pre-mRNAs are endonucleolytically cleaved as part of the 3' end processing reaction and that the resulting RNAPII-associated downstream fragment is degraded from its 5' end by the nuclear 5'-3' exonuclease XRN2 (Rat1p in *S. cerevisiae*). The model states that XRN2 catches up with the transcribing RNAPII and, in a poorly understood manner, triggers transcription termination^{24,25} (Figure 1). Additional endonucleolytic cleavage sites downstream of the polyA signal (the so-called co-transcriptional cleavage (CoTC) sites), specific DNA sequence elements that slow down or pause RNAPII, as well as enzymatic activities destabilizing RNA:DNA hybrids, including the one covered by RNAPII itself, have also been implicated in the termination process^{24,26-31}. Evidence exists in support of both transcription termination models and it is likely that both contribute to varying degrees on different RNAPII-transcribed genes^{3-5,32}.

Setting and resetting RNAPII

During the transcription process the RNAPII complex is subject to extensive changes in modification. A central platform for this modulation is the C-terminal domain (CTD) of the largest subunit of RNAPII, which acts as a scaffold for interactions with multiple factors involved in transcription initiation, elongation and termination as well as with factors important for co-transcriptional maturation of the RNA product. The CTD contains a conserved heptad repeat sequence (consensus: Tyr₁Ser₂Pro₃Thr₄Ser₅Pro₆Ser₇) and most of its amino acid residues can be post-translationally modified (Figure 1). This dynamic property of the CTD is of great importance for the transcription cycle as it assists the coordination between the many facets of transcription and RNA processing³³⁻³⁷. The patterns of phosphorylation of serines 2 (Ser2P) and 5 (Ser5P) and the consequences thereof have been the focus of particular intense investigations, but recent attention has also been directed towards phosphorylation of Tyr1 (Tyr1P), Thr4 (Thr4P) and Ser7 (Ser7P) that appear to be important for specific types of 3' end processing reactions³⁸⁻⁴³.

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Although the phosphorylation pattern of the CTD is gene specific some overall features relate to the distance travelled by RNAPII away from the promoter (see Figure 1). In general, RNAPII is loaded on the PIC with a hypophosphorylated CTD, which then becomes Ser5 phosphorylated during the initial phases of transcription. As RNAPII travels through the gene Ser5P gradually declines and Ser2P levels rise. In yeast, the switch from dominating Ser5P to Ser2P is estimated to occur ~450 bp downstream of the TSS^{44,45}, after which high levels of Ser2P persist until a terminator is reached. Ser7P and Thr4P display roughly similar patterns to Ser5P and Ser2P, respectively (Figure 1)^{42,45,46}. Thus, when RNAPII encounters a terminator on short genes (<450 bp) the CTD is mainly in a Ser5P/Ser7P state, whereas on longer genes (>450 bp) Ser2P/Thr4P will predominate. Accordingly, the associated 3'end processing/termination processes on short and long genes have evolved to be differentially stimulated by the specific modification of the CTD matching the time of its usage (see below and Figure 2A). An interesting new discovery here is the observation from yeast that although the profile of Tyr1P follows that of Ser2P it declines earlier than Ser2P and before RNAPII reaches the 3'end processing signal (Figure 1)³⁸. This allows for the dismantling of specific elongation factors as well as the binding of specific termination factors and Tyr1P thereby 'shields' the CTD from becoming prematurely termination-prone before reaching the gene end. It is likely that such coordinated departure of elongation factors and entry of termination factors constitutes a checkpoint for efficient transcription termination.

Some distance into the terminator region, the CTD is eventually dephosphorylated^{45,47-49}, and it is generally observed that RNAPII which is not bound to the DNA template is hypophosphorylated^{50,51}. It therefore appears that RNAPII can be 'reset', at least partially, in preparation for its next round of transcription either prior to, or concomitant with its termination. Another important, albeit less well studied, aspect of such RNAPII resetting is its release of the produced RNA. This can occur while RNAPII is still attached to, or after it has been dismantled from, the chromatin template⁵²⁻⁵⁵. Interestingly, coupled *in vitro* transcription/3'end formation assays performed in HeLa cell nuclear extracts have revealed that a nascent mRNA can remain attached to the RNAPII CTD even after

cleavage at its polyadenylation site and is only released upon productive 3'end polyadenylation⁵⁶. Thus, at least in some situations, all stages of RNA 3'end processing would have to be completed before RNAPII can be recycled to perform another round of initiation.

RNAPII-transcribed genes - the long and the short of it

Above RNAPII transcription has been generalized, but in reality there are variations for each individual gene type. RNAPII-transcribed genes can broadly be placed into three categories depending on the nature of the produced mature RNA: (1) protein-coding genes, which in many metazoans fall in two different groups, namely (1a) those encoding polyadenylated mRNA and (1b) those encoding replication-dependent non-adenylated histone mRNA, (2) short ncRNA genes, such as those encoding a subset of sn/snoRNAs and (3) long non-coding (lnc)RNA genes. We will not discuss the latter class further here, since only little general knowledge has been obtained on features of its transcription initiation and 3'end processing/termination.

Even though promoters vary between and within these categories in terms of which complement of transcription factors is bound, a common (sub)set of GTFs are believed to be positioned at the core of committing RNAPII for transcription^{13,57-61}. Additionally, the machineries that mediate RNA 3'end processing and RNAPII termination on these types of genes display commonalities as well as major differences. In the remainder of the review we will mainly focus on knowledge from metazoan organisms except where indicated.

Protein-coding genes with a polyA terminator

The vast majority of protein-coding genes produce polyadenylated mRNA with 5'- and 3'- untranslated regions (UTRs), an open reading frame (ORF), and in most cases several introns. The

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position of the 3' end of the mature transcript is dictated by the RNA polyA site, however, after its passage, RNAPII continues into the terminator region where transcription eventually ceases. The distance from the TSSs of polyA genes to their polyA sites (i.e. the length of the gene body) varies in size from ~500 to several millions of base pairs (bp). Furthermore, the distance covered by RNAPII from the polyA site and until it terminates differs between, and within, genes from a few hundred to several thousand bp (Figure 2A)^{46,49}. 3' end processing of polyA gene products is driven by two enzymatic reactions – pre-mRNA cleavage and 3' end polyadenylation of the upstream cleavage fragment (Figure 2B). A polyA signal consists of several RNA elements of which the best-defined is the polyA-hexamer AWUAAA (W is either A or U) found 1-40 nt upstream of the cleavage site in the majority of metazoan RNAPII transcribed protein coding polyA genes⁶²⁻⁶⁵. Additionally, a polyA signal consists of a less well-defined 'downstream sequence element' (DSE) as well as 'auxiliary sequences' that can be positioned up- and downstream of the cleavage site. The factors and enzymes that recognize these sequences and finally mediate the cleavage and polyadenylation reaction are in tight association in a major 3' end processing complex, which broadly defined contains >85 proteins^{4,66,67}. A core of these can be divided into four major multisubunit complexes (so-called 'factors'; Figure 2B)⁶⁸: **(1)** CPSF (cleavage and polyadenylation specificity factor), which recognizes the polyA-hexamer (or functionally equivalent sequences) and contains the CPSF73 enzyme responsible for the endonucleolytic cleavage reaction⁶⁹, **(2)** CstF (cleavage stimulatory factor) that binds the DSE and stimulates the cleavage reaction⁷⁰, and **(3,4)** cleavage factors I and II (CFI_M and CFII_M), where CFI_M recognizes an auxiliary sequence upstream the cleavage site⁷¹. After transcript cleavage, CPSF directs the polyadenosine polymerase (PAP) to add the polyA-tail, a coordination of cleavage and polyadenylation, which ensures that the 3' end is rapidly protected from 3'-5' exonucleases. During polyA-tail synthesis, nuclear polyA-binding protein (PABPN1) binds to the nascent stretch of PAP-produced A's and strongly stimulates further polyadenylation by retaining CPSF at the polyadenylation signal^{72,73}. The interaction between PABPN1 and CPSF with the polyA tail persists until it has reached ~250 A's after which the stimulation by CPSF is disrupted⁷³.

Evidence exists for several functional connections between transcription initiation, elongation and 3' end processing/termination on polyA genes. The CPSF73 and CstF64 proteins can be detected by chromatin immunoprecipitation (ChIP) not only at the gene 3' end but also at the promoter and in case of CPSF throughout the entire transcription unit^{49,74}. At the promoter, this recruitment is believed to be aided by GTFs, since several CPSF components (CPSF 73/100/160) interact with TFIID and CstF 64 interacts with TFIIB as well as the transcription factor PC4⁷⁴⁻⁷⁶. CPSF (CPSF 73/100/160) and CstF (CstF 50/64/77) also bind the CTD of RNAPII⁷⁷ and their juxtaposition at the promoter is likely to mediate a smooth transfer to RNAPII when transcription commences. The interaction with the CTD is highly stimulatory for 3' end formation since it positions the factors in proximity to the newly synthesized pre-mRNA upon transcription of the polyA signal⁷⁸. Although the mentioned interactions appear to be independent of the phosphorylation status of the RNAPII CTD, there are also examples of 3' end processing/termination factors that depend on CTD modification. Most prominently, it has been shown in yeast that Pcf11p, a factor important for both 3' end processing and RNAPII termination^{22,79}, selectively recognizes CTD Ser2P^{20,80,81}. In addition, Tyr1P, which drops immediately upstream of the polyA signal, antagonizes Pcf11p interaction and indeed, Pcf11p only binds to the Ser2P CTD if Tyr1 and Ser5 are unphosphorylated³⁸. Thus, effectively RNAPII is prepared for polyadenylation/termination already at the promoter, but it will not be fully competent before it has undergone specific modification changes and passed/produced specific sequence elements.

Replication-dependent histone genes

Unique among metazoan protein-coding genes are those encoding the replication-dependent histones (hereafter referred to as 'histone genes'). At least three features set this class of genes apart from the bulk of protein-coding genes: (1) they are located in clusters in the genome, (2) they are intronless, and (3) the encoded mRNA is 3' end processed in a specialized manner⁸²⁻⁸⁴. Furthermore, histone genes are generally shorter than their polyA gene counterparts with gene

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body lengths from ~350 to ~800 bp (Based on data from ref. 83). In contrast to polyA gene terminators, RNAPII is detectable on histone genes as a relatively narrow peak within a ~1 kb region downstream of the histone 3' end processing signal, indicating that transcription termination occurs in a more efficient and homogenous manner (Figure 2A)^{46,85}. This is potentially aided by the presence of a strong RNAPII arrest/pause site immediately downstream of the 3' end processing signal⁸⁶. Two sequence elements are required for proper 3' end processing of histone mRNAs, a stem-loop and a purine-rich 'histone downstream element' (HDE) located 15-20 nt downstream of the stem-loop. The histone pre-mRNA is endonucleolytically cleaved between these two elements^{87,88}, resulting in a mature transcript containing a 3' stem-loop potentially protecting against 3'-5' degradation (Figure 2B). Interestingly, as for polyA site cleavage the responsible endoribonuclease is CPSF73^{89,90}. In addition, cleavage/polyadenylation factors CPSF, CstF77, CstF64, Fip1 and Symplekin are also involved in histone mRNA 3' end processing^{91,92}. Besides this overlap in factor usage, the implicated complexes are different. The histone stem-loop is bound by 'stem loop binding protein' (SLBP), which facilitates the binding of U7 snRNA to the HDE by basepairing⁹³. SLBP bound to the stem-loop interacts with ZFP100⁹⁴, which in turn associates with the U7 snRNA-associated protein Lsm11 thus bridging the stem-loop and the HDE^{95,96}. How CPSF73 is recruited to the cleavage site is still not known, but it has been suggested that FLASH, another essential component of the histone mRNA 3' end formation machinery, mediates the contact between CPSF73 and the pre-mRNA (Figure 2B)⁹⁷.

Any coupling between transcription and RNA 3' end processing on histone genes is only poorly understood currently. FLASH, SLBP, CPSF73 and CstF77 are present at the histone promoter^{43,49,98} and the former two proteins co-localize with the histone gene transcription factor NPAT, which may imply pre-loading of 3' end processing factors⁹⁸. Interestingly, CTD-Thr4P is required for histone mRNA 3' end processing whereas it is dispensable for polyadenylation of polyA genes⁴³. In line with this, Thr4P is essential in vertebrates⁴², but not in budding and fission yeasts^{99,100}, consistent with the fact that no specialized 3' end processing machinery exists for histone genes in these species. There are conflicting observations about the involvement of Ser2P in histone mRNA 3' end processing^{101,102}.

However, these studies suffer from the problem that the drugs used to inhibit the Ser2 kinase likely also inhibit phosphorylation of Thr4. It has yet to be revealed how Thr4P is mechanistically coupled to histone 3'end processing.

U snRNA genes

RNAPII transcribes a subset of autonomous small ncRNA genes, including some snRNA and snoRNA loci¹⁰³. The resulting RNAs are usually involved in the processing/modification of other RNA molecules. Besides being short (~150-350 nt; Figure 2A), these RNAs are characterized by their lack of introns and their production from genes having specialized promoters and terminators^{57,104}. The human U1 and U2 snRNA-encoding genes are most comprehensively described with respect to 3'end processing and transcription termination and will be discussed in the following paragraphs.

U1 and U2 snRNA gene promoters contain two important regions; the proximal and distal sequence elements (PSE and DSE), respectively¹⁰⁵. Lack of either of these reduces RNA levels by more than 100-fold¹⁰⁶. Like other RNAPII-genes, U1 transcription relies on GTFs such as TFIIA, TFIIB, TFIIF, TFIIE⁶⁰ together with TBP¹⁰⁷. However, TBP and a subset of its interacting factors (TAFs) are recruited to the U1 promoter in complex with the snRNA-activating protein complex (SNAPc) and not in context of the usual TFIID GTF^{61,107}. The general involvement of U snRNAs (and snoRNAs) in RNA metabolism requires their robust expression, which is probably governed by a specialized mode of transcription as well as the long half-lives of these molecules. 3'end processing of U snRNAs is accomplished by a dedicated machinery, which is different between metazoans and yeast. In animals, a protein complex called Integrator carries out 3'end formation involving an endonucleolytic cleavage event upstream of the 3'end-positioned so-called 3'box (Figure 2B)¹⁰⁸. The endonuclease, Int11, and another Integrator subunit, Int9, are homologs of CPSF73 and CPSF100, respectively¹⁰⁸. Apart from these factors, the snRNA 3'end processing complex is completely

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different from the described mRNA 3'end processing complexes and no polyadenylation of the naked snRNA 3'end occurs. Instead, it appears that the U snRNA is protected against degradation by upstream secondary structures or bound proteins. How processing at the 3'box connects with termination of RNAPII is virtually unknown. Nuclear run-on (NRO) analyses suggested that the primary transcript produced from the U2 gene can extend at least ~800 nt from the TSS, which means that RNAPII, similar to histone genes, may continue to transcribe ~600 bp past the 3'box¹⁰⁹. In contrast, termination on the U1 snRNA gene was found to occur immediately downstream of the 3'box^{109,110}. However, this observation is probably affected by the many genomic copies of U1, which diverge in their sequences downstream of the gene body. Indeed, unpublished data imply that RNAPII transcription here can also continue further downstream (D. O'Reilly and S. Murphy, personal communication).

The CTD of RNAPII is essential for proper 3'end formation of U1 and U2 snRNA¹¹¹⁻¹¹³ and its phosphorylation status is central for this dependency³⁹⁻⁴¹. In particular, Ser7P appears to be essential for the process³⁹. The Integrator subunit Int11 interacts with the CTD, but it is most efficiently recruited to Ser2P/Ser7P configured heptad repeats⁴⁰, which implies that Integrator-mediated 3'end processing is only efficient within a rather narrow window of distance from the TSS where both these modifications are present at the same time. Since Ser7 generally starts to become dephosphorylated when RNAPII travels further than ~450 bp this may explain why increasing the distance between the promoter and the 3'box of the U1 snRNA gene decreases the efficiency of 3'end processing¹¹⁴.

Similar to the early recruitment of 3'end processing factors on polyA genes, the Integrator complex (Int2 and Int9-11) is found at the promoter of snRNA genes¹⁰⁸. It is not known whether initial recruitment occurs independently of RNAPII, but since several studies have shown that elements in snRNA promoters are needed for efficient U snRNA 3'end processing^{104,115,116}, this would be in accordance with promoter-specific Integrator recruitment and its later handover to RNAPII.

Short RNA genes in *S. cerevisiae*

In *S. cerevisiae*, yet another means of transcription termination is employed. Unlike in higher eukaryotes, most short *S. cerevisiae* ncRNAs derive from independently-transcribed loci where 3' end processing/termination depends on the Nrd1p/Nab3p/Sen1p (NNS) complex (Figure 2B), which is not homologous to Integrator¹¹⁷⁻¹¹⁹. Indeed, a functional equivalent of NNS has not yet been identified in higher eukaryotes. NNS consists of the RNA binding proteins Nrd1p and Nab3p as well as the helicase Sen1p^{117,120,121}. Sen1p (and its human ortholog Senataxin) is also involved in 3' end processing/termination of some polyA genes^{27,122,123}. RNAPII-bound NNS targets specific sequence motifs in the nascent RNA and by an uncharacterized mechanism, likely involving Sen1p, terminates RNAPII transcription^{117,121,124-127}. NNS also recruits the 'Trf4-Air2-Mtr4 polyadenylation' (TRAMP) and RNA exosome complexes that, depending on the nature of the target RNA 3' end, mediate either trimming/processing or the complete degradation of the affected transcript (Figure 2B)¹²⁰. Via its CTD interacting domain (CID), Nrd1p interacts directly and selectively with the Ser5P modification, restricting this type of termination to occur mainly within a short distance from the TSS¹²⁸⁻¹³⁰.

Functional communication between terminator and promoter

Where most of the above described interactions between nascent RNA 3' end processing factors and transcription take place within the same transcription cycle, evidence for a functional link between the 3' end processing/termination reaction back to the promoter is emerging. A first indication was the discovery of a transcription-dependent physical interaction between terminator and promoter via so-called gene loops. In yeast, loop formation is mediated by an interaction between TFIIIB and the Ser5-phosphatase, 3' end processing and transcription elongation factor Ssu72p^{7,9,131-133}. Gene loops result in close contact between the promoter and terminator and are suggested to facilitate

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transcription re-initiation and to impose directionality onto the promoter^{10,11,133,134}. Moreover, several reports imply that impairment of 3' end processing and/or transcription termination leads to decreased transcription initiation: In a genome-wide study on yeast cells, it was observed that termination was impeded on both protein coding and small non-coding genes upon inhibition of the phosphatase activity of Ssu72p⁴⁷. In parallel, a general drop in RNAPII levels at the promoters and in the gene bodies of the termination-impeded genes was detected. Similar effects were reported by a series of study examples using individual reporter genes. For instance, inactivation of Rat1p (yeast homolog of XRN2) was shown to decrease levels of RNAPII at the promoter and inside the gene body of the assayed *ADH1* gene¹³⁵ and depletion of human Senataxin led to elevated levels of RNAPII in the terminator region of the *β-actin* gene accompanied by a significant decrease at the promoter²⁷. These examples do not *per se* imply recycling within the same gene, since lowered levels of RNAPII at the investigated promoters could be a consequence of global sequestration in terminator regions. However, in two recent studies similar phenomena were detected on genes with 3' end processing/termination defects imposed in *cis*^{6,8}. In the first, compromised 3' end processing/termination caused by a single point mutation of a polyA signal yielded a lower transcription initiation rate as measured by NRO as well as promoter-ChIP of RNAPII and the two GTFs TBP and TFIIB⁸. Surprisingly, complementary ChIP analysis revealed that the CTD of RNAPII present downstream of the mutated polyA site, and supposedly still attached to the nascent uncleaved RNA, was largely devoid of Ser2P and Ser5P and an equally curious elevated level of both TBP and TFIIB could be measured at the same positions⁸. While these phenomena were consequences of the impaired 3' end processing/termination, they may also reflect natural processes that just take place with faster kinetics at a functional terminator. Thus, perhaps dephosphorylated RNAPII can bind some GTFs already before disengaging from the chromatin template, which is in agreement with the suggestion that RNAPII can be prepared for the next round of transcription before or concomitant with termination. Furthermore, it was recently shown that the described 3' end processing/termination defect correlated with the disappearance of a detectable promoter-

terminator loop, which implies that gene looping could be responsible for efficient recycling of RNAPII back to the promoter¹¹. In a second study, polyA genes of different lengths were analyzed and it was found that a short <450 bp gene was transcriptionally repressed at the initiation level compared to its longer (>450bp) counterparts. Transcription repression required that the gene harboured a polyA terminator; i.e. repression was relieved upon its replacement by a replication dependent histone gene-terminator⁶. This implies a strong context-specific requirement for the efficient communication between 3' end processing/termination signals and the promoter. Assuming that the switch from dominating Ser5P to Ser2P also happens ~450 bp downstream of the TSS in mammalian species, there is a striking correlation between the predicted RNAPII-CTD phosphorylation status and the efficiency by which the different 3' end processing signals stimulate transcriptional re-initiation (see *Conclusion* for further elaboration). A somewhat surprising addition to these observations is that the actual RNA polyadenylation event also appears to affect transcription initiation as cells treated with cordycepin, a drug that inhibits the addition of adenosines to the elongating polyA tail, displayed decreased promoter RNAPII occupancy at polyA genes concomitant with increased RNAPII signals downstream of the corresponding polyA sites. In contrast, RNAPII occupancy in the terminator region of histone genes were not affected by cordycepin⁴⁶. The mechanism underlying these observations is not understood, but taken together with the finding that the actual polyadenylation event is needed for release of the mRNA from RNAPII⁵⁶, it is possible that proper transcription termination in some cases depends on polyadenylation, which in turn affects re-initiation (Figure 1).

Sidebar title

[Please include sidebars in the body of the text where appropriate]

Comment [A5]: You are encouraged to include sidebars ("boxed" information that is relevant to but separate from the main text) especially to highlight contemporary interdisciplinary themes. Each sidebar should be a maximum of 250 words. Do not include more than two sidebars.

Conclusion

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Although not formally demonstrated for all gene classes, it is generally assumed that transcription termination is intricately coupled to RNA 3'end processing. The 'strength' of the 3'end processing signal and its surrounding terminator region determine how rapidly the RNA is processed and how efficiently RNAPII terminates. While a major determinant for 3'end processing strength is the composition of the signal itself in terms of elements and their sequences, it is also clear that the distance to the TSS plays a decisive role in how well a given signal is utilized. Due to CTD-modification changes as a function of RNAPII transcriptional progress, a promoter proximal polyA signal is not recognized as efficiently as a distal one and, conversely, a distal snRNA 3'end processing signal functions worse than a proximal one. In addition, RNA 3'end processing efficiency is influenced by upstream RNA processing events; e.g. splicing of the last intron is stimulatory for the cleavage/polyadenylation reaction as well as for release of the polyadenylated mRNA from RNAPII^{56,136-139}. Similarly, 3'end processing of snRNAs is promoted by 7-methyl-G capping of the RNA 5'end¹⁴⁰. Finally, as many 3'end processing factors are already recruited at the promoter, it is possible that critical decisions can be made very early in the transcription process.

The accumulating body of evidence highlighting the importance of productive 3'end processing/termination for transcriptional re-initiation of the same gene calls for mechanism(s) to recycle RNAPII from the terminator back to the promoter. Gene-looping from 3'end processing sites to promoters may be one way to maintain sufficient local concentrations of RNAPII within each individual gene circuit. Alternatively, the compartmentalization of genes into a spatially confined environment (aka 'gene factories'), could increase the chances for an RNAPII complex to re-initiate on the same recently transcribed gene, simply by being in proximity^{141,142}. This was shown for gene loci on polytene chromosomes in *D. melanogaster* salivary glands¹², and it may be an important feature for clustered genes. As the availability of RNAPII varies between different conditions and cell types there may not always be a need for intragenic recycling of RNAPII¹⁴³. However, in favor of a general importance of recycling, it was recently shown that RNAPII is present in surprisingly few copies per cell in proliferating *S. pombe* (~1 molecule per gene)¹⁴⁴.

If the lowered transcription initiation activity of genes undergoing defective/weak 3'end processing/termination is indeed caused by inefficient RNAPII recycling, what is then causing this? When RNAPII encounters a weak/defective 3'end processing signal, it may become somewhat prone for termination while still being partially fit for continued elongation. Such an 'upset' polymerase may terminate erroneously at a more or less random position without being properly reset – i.e. without proper CTD dephosphorylation and/or without release of the RNA (both the gene-body encoded RNA and the polymerase-associated RNA) – and therefore not be prepared for re-initiation (Figure 1). In fact, the major determinant for efficient recycling may be the ability of RNAPII to be reset during termination. Such resetting would then depend on the kinetics of 3'end processing and termination, which in turn are governed by the strength of the encountered processing signal.

What then is an efficient terminator? In order to avoid transcriptional interference into a downstream gene, termination is required, especially in cases where genes are closely spaced. Nonetheless, even when the 3'end processing signal is supposedly strong, RNAPII often travels far into the terminator – up to 1 kb on histone and snRNA genes and up to several kb on polyA genes. The rather broad distribution of RNAPII molecules in the terminator region – especially on polyA genes – implies that completion of the transcription process is somewhat stochastic and depends on limiting events/factors. This may reflect an inherent disinclination for an elongating RNAPII to terminate, ensuring that it only happens when the transcription and RNA processing events have been completed. Although the resetting and recycling processes of RNAPII need to be studied in much greater detail, the fact that the CTD can be dephosphorylated a short distance into the terminator region and that the unprotected 3'RNA trailing from the RNAPII is normally rapidly degraded implies that resetting of RNAPII can take place on the chromatin template. Intuitively, this appears beneficial if recycling is to be tightly controlled. Perhaps the final release from the terminator is even determined by the encounter with a re-initiation complex bound promoter – possibly from the same gene.

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Notes

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Comment [A6]:
Notes are generally discouraged in scientific reviews, but authors writing from a humanities or social sciences perspective may wish to employ them as necessary. Notes should be indicated by superscript letters, both in the text and in the notes list.

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Figure captions

Fig. 1: The RNAPII transcription cycle

Illustration showing the changes of RNAPII during transcription and recycling based on transcription of a polyA gene. Colored circles of the respective residue indicate phosphorylation of the CTD: Tyrosine 1 (Y1) in yellow, Serine 2 (S2) in green, Threonine 4 (T4) in purple, Serine 5 (S5) in red and Serine 7 (S7) in blue. For S2 and S7, intense colors illustrate high levels of phosphorylation whereas less intense coloration indicates lower levels of phosphorylation. XRN2 is depicted with a yellow pacman. For further details see text.

Fig 2: Overview of different types of RNAPII transcribed genes and their 3'end processing machineries.

(A) Schematic illustration of the general lengths of polyA (green), histone (red) and snRNA (purple) genes (terminators included). See text for details. (B) Overview of systems engaged with 3'end processing of precursor RNAs arising from polyA genes, histone genes and snRNA genes as well as

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3'end processing by the NNS complex. Top panels show sequence-elements and complexes involved with an arrow emphasizing the endoribonuclease catalyzing the cleavage reaction. Lower panels show the processed RNA product, which in the case of NNS-mediated processing is further processed by TRAMP and the RNA exosome. Complexes involved in 3'end processing of polyA mRNAs are depicted in shades of green, for histone mRNAs in shades of red and for snRNA in purple. The machineries responsible for 3'end processing of histone and polyA mRNAs share several factors, which are in shades of green (see text for further details). The proteins of the NNS complex are depicted in shades of grey.

Tables

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151	Polyadenylation and transcription termination
136	Non mRNA 3' end formation

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