Divergent RNA transcription
A role in promoter unwinding?

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New approaches using biotinylated-psoralen as a probe for investigating DNA structure have revealed new insights into the relationship between DNA supercoiling, transcription and chromatin compaction. We explore a hypothesis that divergent RNA transcription generates negative supercoiling at promoters facilitating initiation complex formation and subsequent promoter clearance.

Packaging DNA into chromatin creates a repressive environment. This is thought to reduce transcriptional noise and provide additional levels of gene regulation. Many processes have been described that alter chromatin architecture and facilitate or repress transcription. These include transcription factor binding, recruitment of RNA polymerases, ATP-dependent chromatin remodeling and changes in histone modifications. The level of DNA supercoiling is also known to be important for transcription but remains poorly understood, as few techniques are available for directly investigating it. This is about to change: new approaches have recently been used for analyzing DNA topology in cells and show transcription dependent changes in DNA supercoiling that impact on higher levels of chromatin structure. Furthermore, new data indicates that DNA supercoiling and divergent RNA transcription are linked providing new ideas for understanding the role of DNA topology in regulating gene transcription.

The structure of DNA within mammalian cells is not known, although it is often thought to be in a B-form configuration. However, this will depend on the base composition, extent of supercoiling and DNA binding proteins, such as HMGs. As DNA moves through the RNA polymerase positive (over-wound) supercoils are generated ahead of the polymerase and negative (under-wound) supercoils behind, described as the twin supercoiled domain model. Therefore, transcription generates significant torsion in the DNA, and the observation that topoisomerase I, an enzyme that relaxes supercoils, is associated with the transcription machinery and localizes to actively transcribed chromatin led to the suggestion that it might relieve topological strain. However, early experiments show that genes are more efficiently transcribed if they are encoded on negatively supercoiled plasmids. In addition, superhelical tension is also a prerequisite for TFIID binding promoting the formation of a preinitiation complex, transcription factor binding and promoter clearance. Consequently, cell based experiments using psoralen, as a probe for DNA supercoiling, show that promoter regions are under-wound and supercoiling may prime specific genes for transcription. Together this suggests that DNA supercoiling must be precisely controlled and is an additional level of transcriptional regulation. However, how DNA supercoiling is introduced at TSS prior to transcription remains unanswered.

A major clue arose from an in vivo study demonstrating that local domains of supercoiling can affect transcription. In this work, a promoter is activated due to transcription generated negative supercoils from a nearby divergent promoter and inhibition of endogenous topoisomerase I further increases transcription. While this

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study exploited in vitro generated plasmid templates; divergent transcription, whereby transcription is initiated in both sense and anti-sense directions from a promoter, has been shown to be a feature of many active mammalian genes.\textsuperscript{16-18} The high resolution mapping of short RNA transcripts reveals extensive sense and anti-sense transcript peaks at approximately 50 bp and 250 bp respectively from the TSS. Antisense RNAs are capped\textsuperscript{19} but are present at a relatively low abundance as they are substrates for the exosome.\textsuperscript{18} Histone modifications reflective of transcription initiation (H3K4me3) are found both in the sense and antisense direction while histone marks indicative of transcriptional elongation (H3K79me2) are found in the sense direction.\textsuperscript{17,20} Consequently divergent transcription is more appropriately described as divergent initiation as only productive elongation occurs in the direction of the gene.

Divergent initiation is very prevalent (approximately 80\% of transcribed mammalian genes) but does it have a regulatory function or can it just represent “sloppy” transcription initiation events resulting from an open promoter architecture? Several functions have been proposed\textsuperscript{21} but, in particular, Seila et al.\textsuperscript{22} suggest divergent initiation may under-wind promoter regions facilitating productive transcription initiation and elongation. To better investigate DNA supercoiling we have developed an approach for directly analyzing it in cells.\textsuperscript{1} This technique builds on previous studies that have exploited the cell permeable drug tri-methyl psoralen (TMP), which preferentially intercalates and can be cross-linked into negatively supercoiled (under-wound) DNA.\textsuperscript{23} The addition of a biotin group, attached via a linker, to the psoralen (bTMP) enables the selective enrichment of under-wound DNA fragments that can be mapped by hybridization to high-resolution genomic microarrays. In RPE1 cells, bTMP binding analysis demonstrates that TSSs are under topological strain and negatively supercoiled in a region that extends ~20 kb into the body of the gene and 10 kb upstream (Fig. 1A). Transcription inhibition by α-amanitin or DRB\textsuperscript{1} causes significant remodeling of TSS DNA to a more positively supercoiled state and subsequent α-amanitin washout promotes a remodeling of the TSS back to its under-wound level. Treatment of cells with bleomycin (introducing DNA nicks) relaxes DNA supercoiling demonstrating that promoter regions are under topological tension, which is also regulated by topoisomerase I and II.\textsuperscript{12}

As transcription clearly introduced negative DNA supercoils under-winding promoter regions, we investigated further the role of divergent initiation in this process. In our experimental approach, the elongating form of RNA polymerase II (RNAPII) is very sensitive to α-amanitin and is rapidly degraded in a proteasome dependent manner (Fig. 1B). In contrast, the initiating form of the polymerase is more resistant to degradation suggesting that changes in supercoiling upon α-amanitin washout are manifest by the initiating form of the polymerase. This was additionally confirmed as blocking transcription elongation specifically with flavopiridol rapidly reduced long RNA production, but there was a lag before a drop in the synthesis of short RNAs, consistent with initiating RNA polymerase producing short RNAs.\textsuperscript{1} Thus, we reasoned that elongating polymerase synthesizes long RNAs while the initiating form produces shorter RNA species.\textsuperscript{24} To test this, we trace labeled cells with 3H-Uridine and showed that the production of long RNAs (> 200 nt) was substantially reduced after 5 h α-amanitin treatment and continued to decrease after α-amanitin washout (Fig. 1C). In contrast, short RNAs (< 200 nt) were produced in abundance by 1 h recovery and continued to increase at 2 h (Fig. 1C), concomitant with remodeling of DNA supercoiling (Fig. 1A). We hypothesized that these short RNAs are the products of divergent transcription.\textsuperscript{16-18} Using data from Core et al.,\textsuperscript{16} we investigated the expression of several short antisense or sense RNA transcripts around the promoters of the expressed IGFB1 (Xq13.1) and LDHA (11p15.1) genes (Fig. 2A).

Long RNAs produced within the gene body were inhibited by α-amanitin and did not recover within 2 h (Fig. 2B). However, in agreement to what is seen globally (Fig. 1C), short antisense and sense RNAs are produced upstream and downstream of genes a short period of time after α-amanitin washout and recovery (Fig. 2C). Our data suggests that the initiating form of RNA polymerase, functioning both upstream through divergent transcription and downstream through abortive sense transcription, generates short transcripts and negatively supercoils TSSs creating a permissive environment for subsequent transcription. Furthermore, as DNA supercoiling impacts on higher levels of chromatin organization,\textsuperscript{1} there is a possible spreading of supercoiling from one locus to another, which could facilitate transcription of surrounding genes and might provide a rationale for gene clustering in the human genome.

This model harmonizes recent advances in our understanding of transcriptional regulation. Traditionally, RNAPII recruitment is thought to be the rate limiting step and, thus, the key regulatory step in eukaryotic transcription; however, genome wide profiling of RNAPII indicates that it is bound and initiated at both active and inactive genes.\textsuperscript{25} Indeed, for a large proportion of metazoan genes (20–30\%), RNAPII density is enriched downstream of many TSSs and this has been described as RNAPII promoter proximal pausing.\textsuperscript{26,27} Pausing is now thought to be a widespread regulatory mechanism with the Negative Elongation Factor (NELF) and DRB-Sensitivity Inducing Factor (DSIF) protein complexes binding to and arresting RNAPII 60 nts downstream of the TSS. Subsequent recruitment of Positive Transcription Elongation Factor b (P-TEFb) to this paused RNAPII complex and phosphorylation by DSIF, NELF and Ser2 on the RNAPII C-terminal results in dissociation of NELF and productive transcriptional elongation. Both sense and antisense RNAPII complexes are involved in RNAPII pausing and both depend on PTEF-b recruitment.\textsuperscript{19}

The purpose of pausing is not known but it is frequently found at developmental control genes and stimulus-responsive pathways and is thought to allow their rapid and synchronous induction in response to extracellular signals.\textsuperscript{27} Consequently, loss of pausing through knockdown of the pause-inducing factor NELF leads to broadly attenuated immune gene activation.\textsuperscript{28} One function of paused RNAPII is to establish a permissive chromatin environment\textsuperscript{29} and paused polymerase has
been shown to block nucleosome assembly at promoters, thus maintaining an open chromatin architecture. In support of this, we propose that promoter proximal pausing permits time for divergent transcription to produce short RNAs and, concomitantly, negatively supercoiled promoter regions to facilitate transcription. After transcriptional pause-release, polymerases in collaboration with topoisomerases and helicases can then maintain the supercoiling state of the locus in a regulated manner. This is consistent with pausing being more frequently associated with developmentally regulated genes, while constitutively expressed genes are maintained with optimal levels of DNA supercoiling. However, anti-sense RNA transcription is significantly less prevalent in Drosophila, but there is pronounced promoter pausing. This may suggest that the purpose of pausing may be different in mammalian and Drosophila genomes, or that abortive sense transcription is sufficient to negatively supercoil the promoter to facilitate transcription factor binding and subsequent processive transcription. The fact that no function has yet been assigned to the short sense and antisense RNAs further substantiates our idea that they are by products of a critical process necessary to create a transcriptionally friendly chromatin environment. Future work combining targeted RNAi of RNA polymerase II pausing and elongation factors with DNA supercoiling analysis will elucidate the mechanism by which chromatin structure influences transcription.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.
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Figure 2. Short RNA synthesis at TSSs. (A) Diagram showing IGBP1 and LDHA gene loci with sense and anti-sense RNA transcripts. From Core et al.15 (B) Graph showing transcription elongation in the gene-body measured by RT-PCR. Long RNAs (> 200 nt) were reverse transcribed (Superscript II, Invitrogen) using random primers and quantified by qPCR (Fast start SYBR green, Roche). Primer sequences are:
- IGBP1 Exon1-Intron1: Fwd: ATCTTCAAAC CGTGAGGAGT
- IGBP1 Exon1-Intron1: Rev: AAAACCTTAG CGGCCTTTTT
- IGBP1 Intron2-Exon3: Fwd: TTTCATGCT CTTTTTGGC
- IGBP1 Intron2-Exon3: Rev: GCTAAACTCT TGCCACATGA
- LDHA Intron3-Exon4: Fwd: CAAGAAGGTT TTGAGGAGCA
- LDHA Intron3-Exon4: Rev: CTTCCTCCCT TTGGTGAAGC
- LDHA Intron2-Exon3: Fwd: AATGGGAGTC CCTCTACTTT
- LDHA Intron2-Exon3: Rev: AGGCTGCGAT GTGGAGAT

(C) Graph showing short RNA transcription measured by miRT-PCR. ± values show distance from TSS. Short RNAs (< 200 nt) were detected by first polyadenylating and then reverse transcribed using tagged oligo-dT and random primers (miScript kit, Qiagen). They were then quantified using qPCR with a specific forward primer and universal reverse primer (miScript primer assay, Qiagen). Primer sequences are:
- IGBP1 −277 TTGTCTCTCT ACCTGGCTCT
- IGBP1 −17 GAAGATCCGG TCGCTTGAG
- LDHA +193 CGATTCGGT CTTGACTT
- LDHA +279 AGGATGAGC GGGTACAG

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