

Nascent transcript sequencing visualizes transcription at nucleotide resolution

L. Stirling Churchman¹ & Jonathan S. Weissman¹

Recent studies of transcription have revealed a level of complexity not previously appreciated even a few years ago, both in the intricate use of post-initiation control and the mass production of rapidly degraded transcripts. Dissection of these pathways requires strategies for precisely following transcripts as they are being produced. Here we present an approach (native elongating transcript sequencing, NET-seq), based on deep sequencing of 3' ends of nascent transcripts associated with RNA polymerase, to monitor transcription at nucleotide resolution. Application of NET-seq in *Saccharomyces cerevisiae* reveals that although promoters are generally capable of divergent transcription, the Rpd3S deacetylation complex enforces strong directionality to most promoters by suppressing antisense transcript initiation. Our studies also reveal pervasive polymerase pausing and backtracking throughout the body of transcripts. Average pause density shows prominent peaks at each of the first four nucleosomes, with the peak location occurring in good agreement with *in vitro* biophysical measurements. Thus, nucleosome-induced pausing represents a major barrier to transcriptional elongation *in vivo*.

Accumulating evidence now reveals that transcription elongation is not a straightforward read-out of the downstream DNA sequence. Co-transcriptional processing events dictate the covalent nature and fate of RNA transcripts¹. Indeed many transcripts are targeted co-transcriptionally for rapid degradation and hence are effectively invisible to approaches that monitor mature messenger RNAs^{2–4}. In addition to these processing events, the strong propensity of RNA polymerase (RNAP) to pause creates barriers to elongation and provides an opportunity for regulation and coordination of co-transcriptional events^{5,6}. *In vitro*, RNAP pausing is found to be ubiquitous⁷. Biophysical approaches have provided a structural and energetic understanding of RNAP pausing which results from both intrinsic properties of the polymerase itself as well as interactions with its DNA template, including the presence of bound proteins (for example, histones)^{8–12}. In the cell, elongation factors probably alter the energetic landscape of transcription, but the extent and mechanism of RNAP pausing in eukaryotic cells remain largely unknown. Bridging the divide between *in vivo* and *in vitro* transcriptional views requires approaches that visualize transcription with comparable precision afforded by *in vitro* transcriptional assays. More generally, the ability to monitor quantitatively nascent transcripts would provide broad insights into the roles and regulation of transcription initiation, elongation and termination in gene expression.

Historically, two strategies have been used to provide snapshots of transcriptional activity *in vivo*. In the first approach, RNAP is crosslinked to DNA and RNAP-bound DNA elements are identified by microarrays or deep sequencing^{13,14}. Although providing a global view of RNAP binding sites, these measurements are of limited spatial and temporal resolution and do not reveal the identity of the transcribed strand or even whether RNAP molecules are engaged in transcription. In the second approach, transcription is halted *in vivo* and then reinitiated in isolated nuclei under conditions that allow labelling of nascent chains, thereby enabling them to be distinguished from bulk RNA^{15,16}. Such 'nuclear run-on' strategies reveal actively transcribed DNA regions but require extensive manipulations that limit resolution and depend on the efficient re-initiation of transcription under non-physiological conditions.

To monitor the transcriptional states of unperturbed cells, we sought to determine the precise *in vivo* position of all active RNAP complexes. Here we present an approach (native elongating transcript sequencing, NET-seq) that accomplishes this goal by exploiting the extraordinary stability of the DNA–RNA–RNAP ternary complex¹⁷ to capture nascent transcripts directly from live cells without crosslinking. The identity and abundance of the 3' end of purified transcripts are revealed by deep sequencing¹⁸, thus providing a quantitative measure of RNAP density with single nucleotide precision. Using NET-seq, we expose rapidly degraded transcription products, locate the position of RNAP pauses, and identify factors and chromatin structure that regulate these transcription events.

Quantifying transcription at nucleotide resolution

We focused on the transcription by RNAPII of protein-coding genes in the budding yeast *Saccharomyces cerevisiae*, although the NET-seq approach should be readily adaptable to other systems. To facilitate purification, we worked with a strain that endogenously expressed a functional variant of RNAPII with a 3×-Flag epitope attached to its third subunit (Rpb3). Log-phase cultures were collected by filtration and flash frozen in liquid nitrogen (Fig. 1a). After cryogenic lysis, RNAPII was efficiently immunoprecipitated (Supplementary Fig. 1a). We prepared the co-purified RNA for deep sequencing using a protocol that allows efficient RNA capture while minimizing bias¹⁹, and sequenced 40 bases from the 3' end. The alignment of these sequences to the yeast genome identified the final nucleotide that was incorporated by RNAPII, and the number of sequencing reads at each position along the genome indicated the density of transcriptionally active RNA polymerases at that site (Fig. 1b, alignment statistics displayed in Supplementary Table 1). A metagene analysis of RNAPII distribution across transcription units shows higher RNAPII density for the first 700 base pairs (bp) from the 5' end (Fig. 1c), consistent with lower resolution observations seen using a global run-on approach¹⁶.

Several observations indicate that we are detecting nascent transcription. First, we robustly capture transcripts from introns and regions

¹Department of Cellular and Molecular Pharmacology, Howard Hughes Medical Institute, University of California, San Francisco and California Institute for Quantitative Biosciences, San Francisco, California 94158, USA.

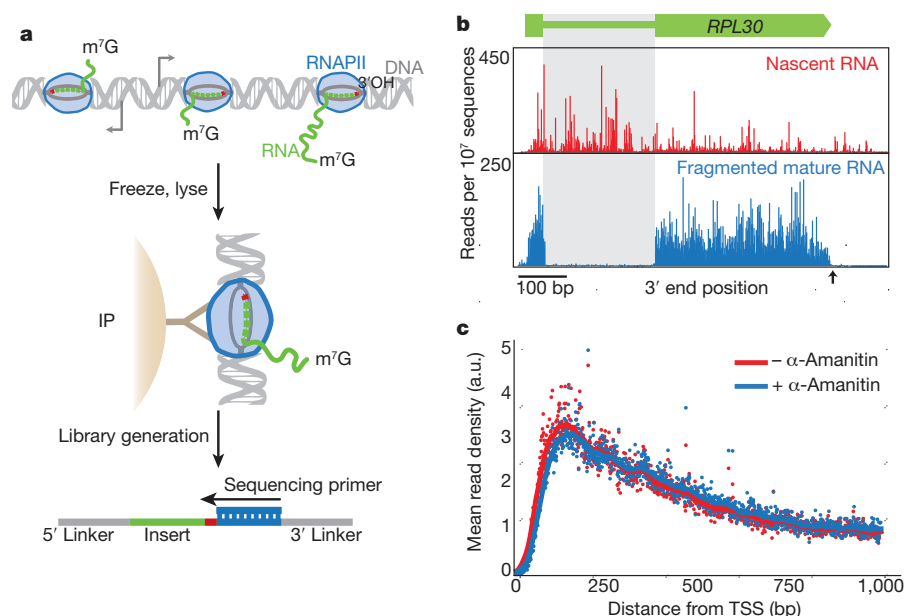


Figure 1 | NET-seq visualizes active transcription via capture of 3' RNA termini. **a**, Schematic diagram of NET-seq protocol. A yeast culture is flash frozen and cryogenically lysed. Nascent RNA is co-purified via an immunoprecipitation (IP) of the RNAPII elongation complex. Conversion of RNA into DNA results in a DNA library with the RNA as an insert between DNA sequencing linkers. The sequencing primer is positioned such that the 3' end of the insert is sequenced. m⁷G refers to the 7-methylguanosine cap structure at the 5' end of nascent transcripts. **b**, The 3' end of each sequence is

after polyadenylation sites; areas that are present in nascent transcripts but absent from mature messenger RNAs (Fig. 1b). Second, we verified that transcripts do not associate with RNAPII after cell lysis (Supplementary Table 2). Third, we saw negligible degradation of RNA under the immunoprecipitation conditions. Nevertheless, our library generation protocols prevent detection of co-purified degradation products by requiring that input RNAs have 3' hydroxyl termini, as hydrolysis and degradation products primarily have terminal phosphates²⁰. Finally, we saw that transcription did not proceed during processing of lysates as addition of the transcription inhibitor α -amanitin to the lysis buffer did not change the RNAPII density (Fig. 1c).

In addition to nascent transcripts, the RNAPII immunoprecipitation captures splicing intermediates (that is, the 5' exon and the excised lariat). Their 3' hydroxyl termini allow them to appear in our data at the 3' ends of exons and introns (Supplementary Fig. 5). These observations indicate the widespread existence of co-transcriptional splicing in yeast and establish NET-seq as a powerful tool for studying such events.

Direct observation of transcription of unstable RNA

NET-seq monitors transcripts regardless of their stability, making it ideally suited to the analysis of unstable transcripts. Recent studies have revealed a class of cryptic unstable transcripts (CUTs) that are short (less than ~700 nucleotides), upstream and antisense to an annotated gene and rapidly degraded by the exosome^{2-4,15,21}. Divergent transcription, yielding the production of antisense CUTs and mature messenger RNAs from the sense direction, is seen at many promoters in both yeast and metazoans. The observation of widespread divergent transcription was surprising and it remains unclear how antisense transcripts initiate and what biological function they may have. It is likely that the nucleosome-free region associated with promoters facilitates antisense transcription. Additionally, it has been suggested that antisense and sense transcription levels are co-dependent^{15,21}, as transcription in the sense direction could promote upstream antisense transcription (and vice versa) by creating negatively supercoiled DNA and recruiting factors that set permissive histone marks²². Critical evaluation of these

hypotheses has been limited by the difficulty in quantitatively monitoring the levels of unstable antisense transcripts.

As NET-seq directly monitors the production of transcripts, we were able to quantify the relative amounts of nascent sense and antisense transcripts (Fig. 2a, b). We focused our analysis on promoters between genes encoded on the same strand (tandem genes), because in those instances, antisense transcripts can be clearly differentiated from the stable upstream transcript. To quantify divergent transcription, we integrated the transcript levels for the first 500 bp of transcribed DNA in each direction. Although we clearly observed divergent promoters, the large majority of promoters had much less antisense transcription than sense transcription; for more than half of the promoters, sense transcription was at least eight times higher than antisense transcription, and for 80% of the promoters the sense-to-antisense transcription ratio exceeded threefold (Fig. 2b). Notably, a comparison between the levels of sense and antisense transcription showed only modest correlation (Spearman correlation coefficient, $r_s = 0.34$) (Fig. 2c).

The above analysis establishes that antisense transcription is not an obligatory consequence of having an active promoter. What then dictates whether a promoter is directional? Transcription initiation is known to occur in nucleosome-free regions; however, we failed to see a correlation when we compared antisense transcription levels with published data²³ reporting on promoter nucleosome-free-region size and promoter average nucleosome occupancy (Supplementary Fig. 2a, b). We also investigated whether histone modifications associated with active promoters correlated with antisense transcription, as it was observed that H3 acetylation peaks in regions of antisense transcription in human fibroblasts¹⁵. Notably, we found a strong positive correlation ($r_s = 0.65$) between antisense transcription levels and earlier measurements of the levels of H4 (and to a lesser extent H3) acetylation enrichment²⁴ (Fig. 2d and Supplementary Fig. 2c, d).

Rpd3S promotes promoter directionality

The strong correlation between antisense transcription and H4 acetylation indicates that H4 acetylation may have a causative role in

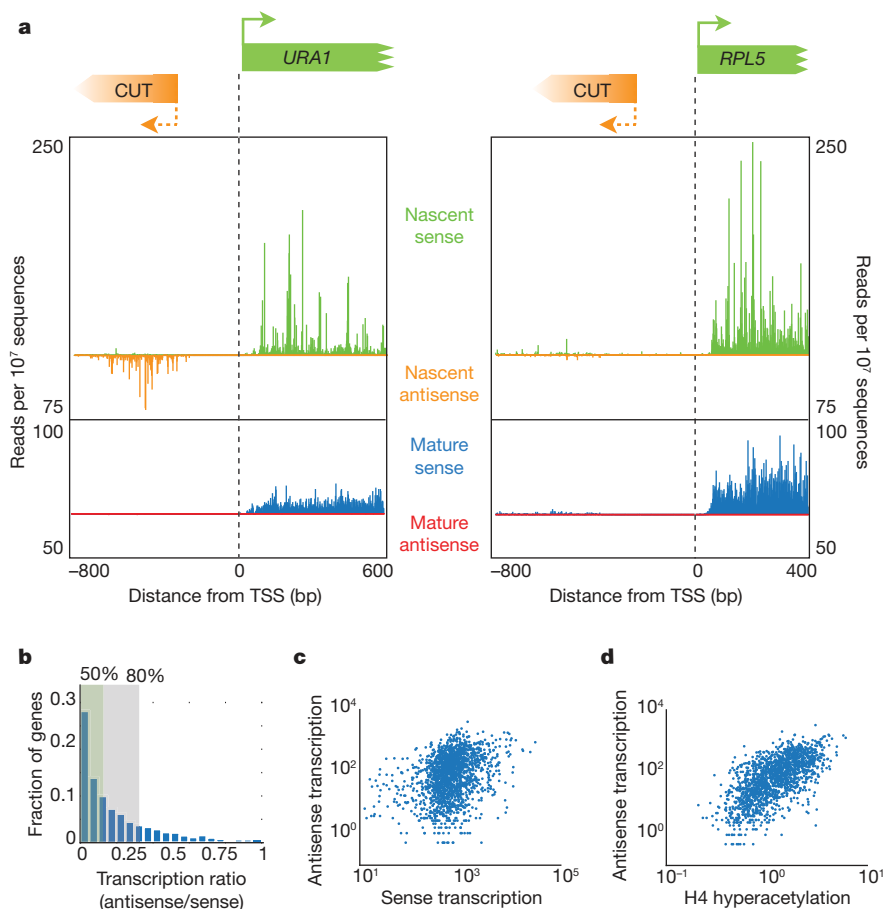


Figure 2 | Observation of divergent transcripts reveals strong directionality at most promoters. **a**, Nascent and mature transcripts initiating from *URA1* and *RPL5* promoters in the sense and antisense directions. Note that there are cryptic unstable transcripts (CUTs) in the antisense direction for *URA1* but not *RPL5*. **b**, A histogram of the transcription ratio (antisense/sense transcription levels) for 1,875 genes. The green and grey boxes indicate the subset of genes with a ratio of less than 1:8 and less than 1:3, respectively. **c**, Antisense transcription levels are plotted versus sense transcription for each tandem gene (Spearman correlation coefficient, $r_s = 0.34$). **d**, The level of antisense transcription for each promoter is plotted versus the local enrichment for H4 hyperacetylation using available data²⁴ ($r_s = 0.65$).

facilitating antisense transcription. To test this, we examined the effect on antisense transcription of loss of *RCO1*, a required and dedicated subunit of the Rpd3 small (Rpd3S) H4 deacetylation complex^{25,26}. We focused on Rpd3S, as earlier studies had shown that it contributes to deacetylation of H4 in the 3' region of transcripts and the large majority of antisense transcripts overlap the 3' ends of upstream genes. Previous global studies of Rpd3S monitored accumulation of mature stable RNAs and so would not detect the effects of Rco1 on transient RNA species^{25,26}. Our analysis revealed a pervasive increase (average fourfold) in unstable antisense transcription (Fig. 3a, b). This effect was the dominant transcriptional phenotype that we observed and was specific to antisense transcription: we found no systematic increase in RNAPII density at the beginning of sense transcripts (Supplementary Fig. 3). Importantly, antisense transcripts seen in the *rco1Δ* strain have the same transcription start sites and the same lengths as the wild-type transcripts, indicating that Rco1 is acting at the initiation stage of antisense transcription and does not affect termination (Fig. 3c). Additionally, we observed that deletion of *EAF3*, another subunit of Rpd3S, mimicked the increases seen in the *rco1Δ* data ($r_s = 0.88$, Supplementary Fig. 4). Thus, the primary function of the Rpd3S histone deacetylase complex seems to be to enforce promoter directionality.

This raises the question of how Rpd3S is recruited to positions designated for suppression of antisense transcription. The Rco1 and Eaf3 components of the Rpd3S complex bind H3 lysine 36 methylation marks made by Set2 and that binding activates the deacetylase activity of Rpd3S (refs 25–27). However, a distinct RNAPII-associated methyltransferase, Set1, has also been implicated in Rpd3S-dependent repression²⁸. Moreover, even in the absence of methylation, RNAPII is capable of recruiting Rpd3S to gene bodies during transcription²⁹.

To investigate how Rpd3S is localized to suppress antisense transcription, we analysed nascent transcripts in cells lacking Set1 or Set2. *SET1* deletion caused a weak increase in antisense transcription in a

manner that correlated only modestly with the *rco1Δ* and *eaf3Δ* data ($r_s = 0.36$ and $r_s = 0.38$ respectively; Supplementary Fig. 5). In contrast, deletion of *SET2* led to a pronounced increase in antisense transcription that was highly correlated with the *rco1Δ* and *eaf3Δ* data ($r_s = 0.88$ and $r_s = 0.89$ respectively; Supplementary Fig. 5). These data together with earlier work on the Set2/Rpd3S pathway indicate that the major mechanism for Rpd3S action on antisense transcription involves Set2 recruitment to elongating RNAPII via Ser 2 phosphorylation on its carboxy-terminal domain³⁰. This in turn, through the Set2 methylation activity, allows recruitment of Rpd3S to the 3' ends of genes, suppressing antisense transcription from downstream nucleosome-free regions. Future challenges will be to explain how histone acetylation in the body of antisense transcripts can affect transcription initiation, and to determine other mechanisms that localize Rpd3S, particularly for the handful of antisense transcripts that do not overlap the 3' ends of genes.

Pausing occurs throughout transcription elongation

The ability of NET-seq to map the density of nascent transcripts enables in-depth investigation of the extent and sources of RNAP pausing *in vivo*. Our data revealed strong and highly reproducible spikes in the density of 3' ends of nascent transcripts along a given gene indicative of RNAPII pause sites (for example, *GPM1*; Fig. 4a). We developed an algorithm to identify RNAPII pause positions that finds points where the read density is at least three standard deviations above the mean in a local 200-bp window. We found that pauses occur frequently throughout the body of RNA messages and are evenly distributed after the first ~700 bp (Fig. 4b and Supplementary Fig. 6). The high density of pauses was not an artefact of library generation and sequencing biases, as we detected tenfold fewer spikes in data from messenger RNA lightly fragmented by alkaline hydrolysis (Supplementary Fig. 7). Notably, 70% of the more than 2×10^5 pause sites that we identified had an A at the 3' end of the transcript.

Additionally, there was a preference for the pause to be followed immediately by a T and then G (Fig. 4c). None of these biases was seen in the control sample of fragmented mRNA (Supplementary Fig. 7a).

Largely from *in vitro* studies, one mechanism of RNAP pausing has been shown to involve backtracking: after encountering a blockage, RNAP reverses direction and moves upstream³¹. In the backtracked state, the 3' end of the RNA transcript is no longer aligned with the active site and RNAP must either return to the initial pause site or cleave the transcript. The latter option is aided by the presence of the elongation factor TFIIS (Dst1 in yeast) that enhances RNAP's intrinsic RNA cleavage activity (Fig. 5a)^{32,33}. Although the role of TFIIS is well established *in vitro*, its mechanism *in vivo* has been less explored^{34–36}.

To investigate the role that backtracking has in pausing *in vivo*, we deleted *DST1* and repeated the NET-seq assay. Notably, we saw a large-scale downstream shift in the position of the pauses, an average of 5–18 bp (Fig. 5b, c). This shift was observed for ~75% of the pauses (Supplementary Fig. 8) and was accompanied by a global change in the sequences surrounding pause sites; the preference for A at the pause was lost and instead there was a strong preference for T immediately downstream of the pause (Fig. 5d). These observations confirm that the observed spikes in NET-seq data result from RNAPII pausing, and indicate that pausing followed by backtracking—which previously had been observed at promoter-proximal pauses³⁵—is prevalent throughout the body of transcripts. Additionally, our studies indicate that Dst1-stimulated RNA cleavage has a strong sequence bias and that a slow step follows cleavage before transcription resumes.

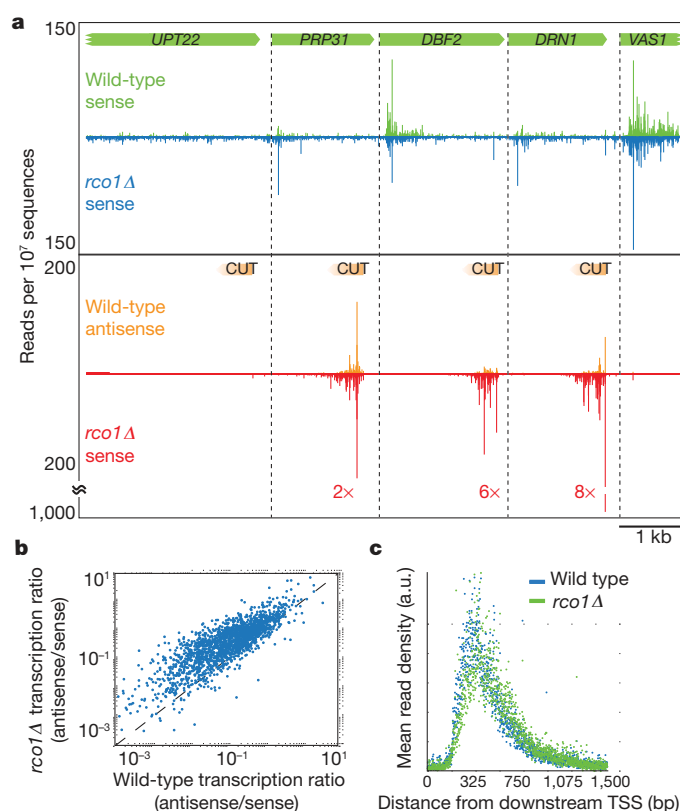


Figure 3 | Rco1 suppresses antisense transcription at divergent promoters. **a**, Examples of cryptic unstable transcripts (CUTs, orange data) upstream and antisense of *DBF2*, *DRN1* and *VAS1* promoters. The fold increase of CUT transcription in the *rco1Δ* strain is marked. **b**, The transcription ratio (antisense/sense) in the *rco1Δ* strain is plotted against the transcription ratio in the wild-type strain for each gene. **c**, A metagenome analysis of well-expressed antisense transcription ($n = 171$, >1 read per bp).

RNAPII pause density peaks before the nucleosome dyad

The pauses observed in the *dst1Δ* strain reveal positions where RNAPII began to backtrack and, therefore, represent the primary point of transcriptional blockage. By analysing these pause positions, we can evaluate what induced RNAPII to backtrack. *In vitro*, nucleosomes induce RNAPII backtracking and TFIIS aids the progression of RNAPII through them^{10,12}. *In vivo*, it is unknown whether nucleosomes interfere with transcription, as chromatin remodelling factors could greatly diminish the nucleosome barrier or remove nucleosomes before RNAPII arrival^{37,38}. Global high-resolution measurements of steady-state nucleosome occupancy revealed that the first few nucleosomes after the transcription start site are phased and well positioned^{23,39}. Thus, by correlating the relative density of RNAPII pauses with nucleosome positions, we can evaluate whether nucleosomes promote RNAPII pausing *in vivo*.

We compared the pause positions in the *dst1Δ* strain to the centre positions of nucleosomes using previously published data²³. Notably, we saw marked peaks of mean pause density at each of the first four nucleosomes (Fig. 6). The precise position of the point of maximal RNAPII pausing at the +1 nucleosome is obscured because it is located just after the transcription start site where many nascent transcripts are too short for unique alignment to the genome. For the +2, +3 and +4 nucleosomes, however, the pause density peaks just before the nucleosome dyad axis (Fig. 6). As would be expected from RNAPII backtracking, the excess pause density before the nucleosome dyad in the wild-type strain is spread out over the upstream region (Supplementary Fig. 9).

Our finding that the peak in pause density occurs just before the nucleosome dyad is particularly remarkable as it is in excellent agreement with earlier biophysical measurements. Specifically, optical trapping studies that physically unwrapped the DNA of a nucleosome off the histone core observed that the dyad is the point where the strongest DNA–histone contacts are found⁴⁰. Moreover, high-resolution optical trapping experiments that followed RNAPII transcribing through a nucleosome found that the RNAPII pause density peaked before the nucleosome dyad¹⁰. Taken together, the above observations provide strong evidence that nucleosomes do indeed present a barrier to elongating polymerases *in vivo* and that this barrier leads to polymerase pausing and backtracking.

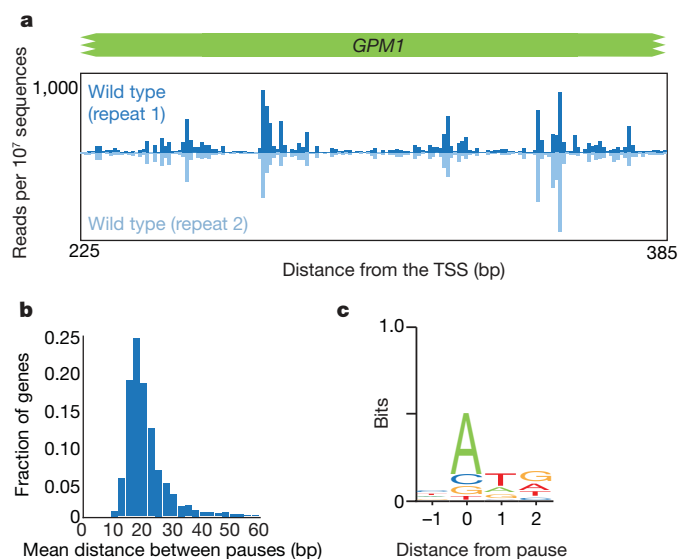


Figure 4 | Frequent RNAPII pausing throughout gene bodies. **a**, NET-seq data at the *GPM1* gene for biological replicates. **b**, A histogram of the mean distance between pauses for each well-expressed gene ($n = 1,006$, >2 reads per bp). **c**, The consensus sequence of the DNA coding strand surrounding pause sites found from all genes.

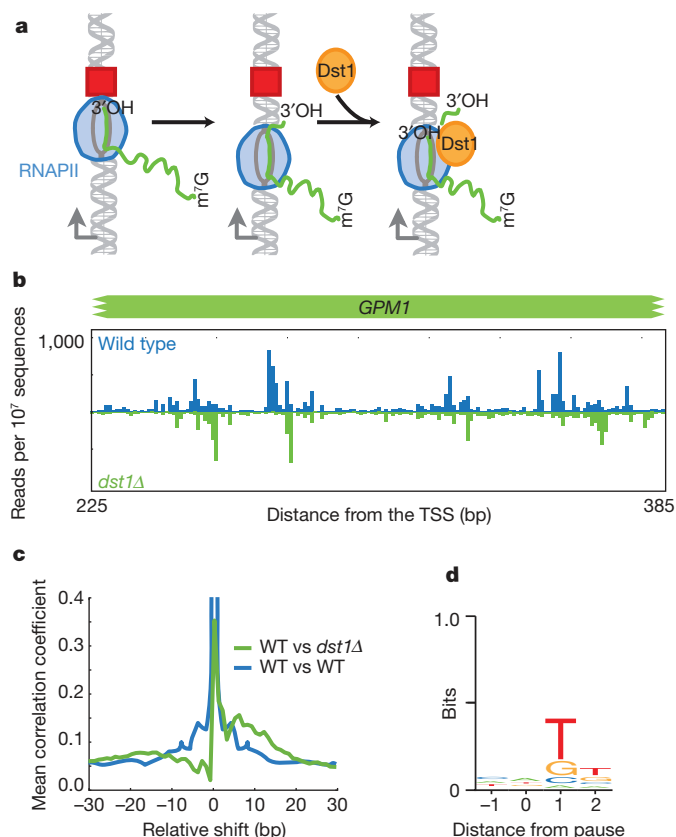


Figure 5 | Dst1 relieves RNAPII pausing after backtracking. **a**, A schematic describing an existing model for how RNAPII pauses at an obstacle (red square), backtracks and is induced to cleave its transcript through binding to Dst1 (refs 32, 33). **b**, A comparison of NET-seq data for wild-type and *dst1Δ* strains at the *GPM1* gene. **c**, Mean cross-correlation between the *dst1Δ* and wild-type data of well transcribed genes ($n = 770$, >2 reads per bp) (green line) was calculated by determining the Pearson's correlation coefficient at each gene between fixed *dst1Δ* data and shifted wild-type data followed by averaging over all genes. This analysis is compared to the mean autocorrelation of the wild-type data for well transcribed genes (blue line). **d**, The consensus sequence for all pauses observed in the *dst1Δ* strain.

Perspective

One of the major surprises in the transcription field in recent years has been the widespread observation of divergent transcription, revealing that the majority of promoters engage in canonical transcription in the sense direction along with the production of unstable transcripts in the antisense direction^{2–4,15,21}. NET-seq provides an ideal tool to look at this phenomenon and uncovers several fundamental properties of divergent transcription. First, most promoters show a strong directionality favouring the sense transcript. Second, suppression of antisense transcripts is enforced by two distinct mechanisms: Rpd3S-mediated deacetylation that prevents antisense initiation, and an independent mechanism, previously characterized to involve the Nrd1–Nab3–Sen1 complex⁴¹, that terminates antisense transcripts and shuttles them to the exosome for degradation. Interestingly, sense transcription may also use this termination mechanism, as our data showed an enrichment for transcripts at the 5' end of genes that mirrors what we observed for antisense transcripts and complements observations that Nrd1 localizes to the 5' end of genes⁴². Third, our observations indicate independence between the initiation of the sense and antisense transcripts. Specifically, we found only modest correlation between sense and antisense transcription levels. Moreover, even among the set of antisense transcripts that increased when *RCO1* is deleted, no increase in sense transcription levels was seen. These findings argue against models in which antisense transcription serves to promote sense

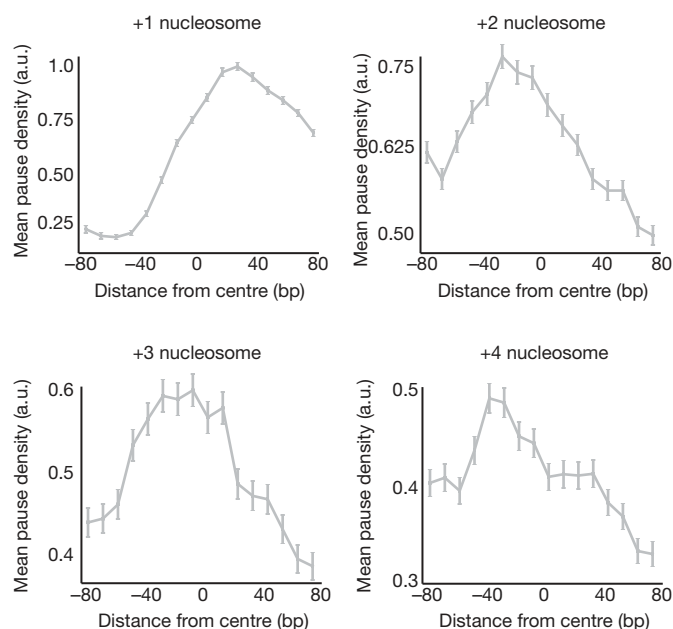


Figure 6 | Nucleosomes are a major barrier to transcription. Plot of mean pause densities in *dst1Δ* data relative to the first four nucleosomes after the transcription start site using available nucleosome positioning data²³. Error bars represent one standard deviation.

transcription (for example, by unwinding DNA supercoils or by removing nucleosomes).

The potential for RNAP to pause has been apparent for decades, motivating interest in the mechanisms and regulatory roles of pausing in the process of transcription^{7,9,11,12,15}. NET-seq provides the first in-depth view of pausing in a eukaryotic cell, revealing that transcription is punctuated by pauses throughout the body of all RNA messages. Taking into account both the abundance and magnitude of the pauses, we conclude that RNAPII spends comparable time in a paused state and moving forwards (Supplementary Fig. 10). We establish that nucleosomes induce pausing *in vivo*, and may be the major source of pausing considering that the increase in pause density at nucleosomes is comparable to the increase in nucleosome occupancy²³. Our observation that pausing peaks at the nucleosome dyad reveal a striking similarity between our measurements and optical trap measurements, indicating that the physical forces observed in purified *in vitro* systems are at play in the cell. NET-seq's ability to follow the physical basis of transcription *in vivo*, allowing direct comparison with high-resolution *in vitro* measurements, may prove to be the most transformative aspect of this approach.

METHODS SUMMARY

Nascent RNA purification. All experiments were conducted using derivatives of yeast strain BY4741. Epitope-tagged Rpb3 (C-terminal $3 \times$ -Flag) was expressed from its endogenous locus. Deletion strains were made by standard PCR-based methods. Litres of log phase culture in YEPD were harvested by filtration and flash frozen by plunging into liquid nitrogen. Frozen cells were lysed cryogenically via six cycles of pulverization using a mixer mill.

Clarified and DNase-I-digested lysate was added to washed anti-Flag M2 affinity gel (Sigma Aldrich), incubated at 4 °C and nutated for 2.5 h. After washing, bound proteins were eluted twice with 2 mg ml⁻¹ $3 \times$ -Flag peptide (Sigma Aldrich). RNA from the eluates was purified using the miRNeasy kit (Qiagen).

RNA linker ligation, cDNA synthesis and PCR. An RNA linker that was 5' adenylated and 3'-end blocked with a dideoxy-C base (5'-CTGTAGGCACC ATCAAT, Integrated DNA Technologies) was ligated onto the 3' end of the immunoprecipitated RNA based on a previously described strategy⁴³. Ligation conditions (see Methods) were systematically optimized to maximize ligation efficiency to ~90% to ensure that the majority of the input RNA was ligated and thus avoiding any bottleneck biases.

cDNA synthesis and sequencing was performed as described with a few modifications¹⁹. The sequencing primer binding site was positioned so that sequencing would start at the 3' end.

Comparing pause densities to nucleosome positions. Nucleosome positions²³ were assigned as +1, +2, +3 etc according to their position relative to transcription start sites. The mean pause density (MPD) relative to a particular nucleosome was determined by the number of pauses observed at that position (N_p) divided by the total number of opportunities it could be observed there (N_o):

$$\text{MPD}_k(x) = \left(\frac{N_p}{N_o} \right)_x = \frac{\sum_i^{\text{all genes}} g_i(y)}{\sum_i^{\text{genes with TSS} < y} 1}$$

$$y = n_i^k + x$$

where k is the nucleosome number, $g(y)$ is the binary function indicating whether a pause occurs at y , and n_i^k are the centre nucleosome positions. The error of the pause density was calculated via the standard deviation of the binomial distribution:

$$\frac{\sqrt{N_p \left(1 - \frac{N_p}{N_o} \right)}}{N_o}$$

Full Methods and any associated references are available in the online version of the paper at www.nature.com/nature.

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Supplementary Information is linked to the online version of the paper at www.nature.com/nature.

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Author Information Raw sequencing data and processed data are available for download at <http://www.ncbi.nlm.nih.gov/geo/> via GEO accession number GSE25107. Reprints and permissions information is available at www.nature.com/reprints. The authors declare no competing financial interests. Readers are welcome to comment on the online version of this article at www.nature.com/nature. Correspondence and requests for materials should be addressed to J.S.W. (weissman@cmap.ucsf.edu).

METHODS

Strain construction. All experiments were conducted using derivatives of yeast strain BY4741. Epitope-tagged Rpb3 (C-terminal 3×-Flag) was expressed from its endogenous locus. Deletion strains were made by standard PCR-based methods.

Extract and total RNA preparation. Yeast strains were grown in YEPD at 30 °C with shaking from an initial optical density (OD) of 0.1 to mid-log phase with an OD of 0.6–0.8. Two litres of yeast culture were harvested in turn by filtration onto 0.45-µm-pore-size nitrocellulose filters (Whatman). The culture was scrapped off the filter with a spatula pre-chilled by liquid nitrogen and flash frozen by plunging into liquid nitrogen. Frozen cells were pulverized for six cycles, each of 3 min at 15 Hz, on a Retsch MM301 mixer mill. Sample chambers were pre-chilled in liquid nitrogen and re-chilled between each pulverization cycle.

One gram of ground cells (~1 l at 0.7 OD) was added to 5 ml of ice-cold lysis buffer (20 mM HEPES, pH 7.4, 110 mM KOAc, 0.5% Triton X-100, 0.1% Tween 20, 10 mM MnCl₂, 50 U ml⁻¹ SUPERase•In (Ambion)) supplemented with protease inhibitor cocktail (1× Complete, EDTA-free, Roche). The experiment using α-amanitin included 10 µg ml⁻¹ α-amanitin (Sigma Aldrich) in the lysis buffer. After re-suspending the lysate by pipetting, 660 units of DNase I (Promega, RQ1 RNase-Free DNase) was added and incubated for 20 min on ice. The lysate was then clarified by centrifugation at 4 °C at 20,000g for 10 min. The supernatant is reserved for immunoprecipitation.

Two-hundred microlitres of clarified lysate is reserved for total RNA purification which was done by the hot acid phenol method. Typical yields were 20 µg. **Native affinity purifications of RNAPII.** 0.5 ml of Anti-Flag M2 Affinity Gel (Sigma Aldrich) was washed twice with lysis buffer. The clarified lysate was added to the washed gel, incubated at 4 °C and nutated for 2.5 h. The immunoprecipitation was washed 4 × 10 ml with wash buffer (20 mM HEPES, pH 7.4, 110 mM KOAc, 0.5% Triton X-100, 0.1% Tween 20, 50 U ml⁻¹ SUPERase•In (Ambion), 1 mM EDTA). Bound proteins were eluted twice with 150 µl elution buffer (20 mM HEPES, pH 7.4, 110 mM KOAc, 0.5% Triton X-100, 0.1% Tween 20) with 2 mg ml⁻¹ 3×-Flag peptide (Sigma Aldrich). RNA from the combined eluates was purified using the miRNeasy kit (Qiagen, 217004). A typical yield from approximately one litre of log-phase yeast culture was 3 µg.

mRNA purification and fragmentation. Polyadenylated mRNA was purified from 50 µg total RNA using magnetic oligo-dT DynaBeads (Invitrogen). Purified RNA was eluted in 20 µl 10 mM Tris, pH 7.0. The purified mRNA was mixed with an equal volume of 2× alkaline fragmentation solution (2 mM EDTA, 10 mM Na₂CO₃, 90 mM NaHCO₃, pH ≈ 9.3) and incubated for 5 min at 95 °C. These conditions yielded lightly fragmented RNA of size distribution similar to that of the nascent RNA. The fragmentation reaction was stopped by the addition 0.56 ml of ice-cold precipitation solution (final 300 mM NaOAc pH 5.5, plus GlycoBlue (Ambion) as a co-precipitant) and RNA was purified by a standard isopropanol precipitation, as follows: after adding 650 µl of isopropanol, samples were placed at -30 °C for at least 30 min. Precipitated RNA was pelleted by centrifugation at 4 °C at 20,000g for 30 min. The pellet was air dried after a quick wash with 80% ethanol and then re-suspended in 10 mM Tris pH 7.0.

A total of 6.4 µg of fragmented mRNA was dephosphorylated in a 50 µl reaction with 1× T4 polynucleotide kinase buffer without ATP, 0.5 U SUPERase•In (Ambion) and 22.5 units T4 polynucleotide kinase (NEB). The dephosphorylation reaction was incubated at 37 °C for 1 h followed by 10 min at 75 °C for enzyme heat inactivation. RNA was precipitated with GlycoBlue by standard methods (see above).

RNA linker ligation, fragmentation and size selection. An RNA linker that was 5' adenylated and 3'-end blocked with a dideoxy-C base (5'-CTGTAGGCACCA TCAAT, Integrated DNA Technologies) was ligated onto the 3' end of the immunoprecipitated RNA, the fragmented mRNA and a synthetic 28-base RNA oligonucleotide (oNTI199, 5'-AUGUACACGGAGUCAGCCGCAACG CGA) similarly to what has been described⁴³. Specifically, 3 µg of each RNA sample was broken into three reactions and diluted to 10 µl with 10 mM Tris, pH 7.0. After a brief denaturation the reactions were brought to 20 µl with a buffer that gave final concentrations of 12% PEG8000, 50 ng µl⁻¹ linker, 1× T4 Rnl2, truncated reaction buffer and 2 units µl⁻¹ of T4 Rnl2, truncated (NEB). The reaction was incubated at 37 °C for 3 h. Ligation conditions were systematically optimized to maximize ligation efficiency to ~90% to ensure that the majority of the input RNA was ligated.

Fragmentation of the ligated samples allowed for the final DNA library to contain inserts of a narrow range to reduce any length biases of downstream enzymatic reactions. EDTA was added to all reactions for a final concentration of 17 mM. 20 µl of 2× alkaline fragmentation solution (2 mM EDTA, 10 mM Na₂CO₃, 90 mM NaHCO₃, pH ≈ 9.3) was added to each reaction and incubated at 95 °C for 30 min. The reactions were stopped by the addition of 0.56 ml of ice-cold precipitation solution (final 300 mM NaOAc pH 5.5, plus GlycoBlue (Ambion) as a co-precipitant), followed by a standard isopropanol precipitation (see above).

The ligated and fragmented samples were size-selected by gel electrophoresis. The purified reactions along with the oNTI199 RNA oligonucleotide was mixed with 2× Novex TBE-Urea sample prep buffer (Invitrogen) and briefly denatured, then loaded on a Novex denaturing 15% polyacrylamide TBE-urea gel (Invitrogen) and run according to the manufacturer's instructions. The gel was stained with SYBR Gold (Invitrogen) and the 35–85-nucleotide region was excised. The gel was physically disrupted and either allowed to soak overnight in gel elution buffer (300 mM NaOAc pH 5.5, 1 mM EDTA, 0.1 U µl⁻¹ SUPERase•In) or incubated in 200 µl of water treated with diethylpyrocarbonate (DEPC) for 10 min at 70 °C. The gel debris was removed from the water or buffer using a Spin-X column (Corning) and RNA was precipitated with GlycoBlue as a co-precipitant using standard methods.

cDNA synthesis. cDNA synthesis was performed as described with a few modifications¹⁹. The primer used for reverse transcription was oLSC003 (5'-pTCG TATGCCGCTCTTCTGCTTG•AATGATACGGCGACCACCGATCCGACGAT CATTGATGTGCCTACAG) where the initial 'p' indicates 5' phosphorylation and '•' indicates the following spacer added for increased flexibility, 18 carbon spacer molecule-CACTCA-18 carbon spacer molecule. Efficient circularization of the RT product was performed as described¹⁹ with CircLigase (Epicentre) according to the manufacturer's directions. Any ligation bias at this step is averaged out as the random fragmentation leaves a range of 5' ends for each 3' end. The PCR was performed directly on the circularized product as described¹⁹, resulting in DNA with Illumina cluster generation sequences on each end and a sequencing primer binding site positioned so that sequencing would start at the 3' end. DNA was purified from a PCR reaction that had not reached saturation and was quantified using the Agilent BioAnalyser High Sensitivity DNA assay. DNA was then sequenced on the Illumina Genome Analyser 2 according to the manufacturer's instructions, using 4–6 pM template for cluster generation and sequencing primer oLSC006 (5'-TCCGACGATCATTGATGTGCCTACAG).

Data analysis. Data analysis was performed using scripts written in Python 2.6 that are available upon request.

Sequencing analysis. Image data obtained by the Illumina Genome Analyser 2 was analysed using the GAPIipeline to extract raw sequences. Matrix and phasing parameters were estimated from a φX control lane.

Sequence alignment. Raw sequences 40 bases long were composed of the cDNA of the fragmented RNA sequence. For RNA fragments smaller than 40 bases, the sequence is followed by part of the 5' Illumina linker sequence which was removed *in silico*. Alignments to the yeast genome were performed by the alignment program, Bowtie 0.12.0⁴⁴ (<http://bowtie-bio.sourceforge.net/>). Bowtie settings were chosen so that three mismatches were allowed and alignments were required to be unique. The shortest sequenced fragments were approximately 18 nucleotides due to the RNA size selection step after ligation and random fragmentation. Eighteen-base-pair sequences would occur by chance every 6.9×10^{10} bp, which is sufficiently rare for 18-bp sequences to be generally uniquely aligned to the 1.2×10^7 bp yeast genome. Alignments were first performed against tRNA and rRNA sequences to remove them. The remaining sequences were aligned against a recent version of the yeast genome downloaded from the Saccharomyces Genome Database (SGD, <http://www.yeastgenome.org/>) on 11 October 2009. Statistics on sequence alignments are reported in Supplementary Table 1.

Quantifying antisense and sense transcription levels. At tandem promoters sense transcription was determined using available annotated transcription start sites³. To allow for the error involved in these transcription start site measurements, we calculated the sum of the read density in 500-nucleotide windows for the first 700 bases after the transcription start site and chose the highest sum. The antisense transcription was determined by starting 100 bases upstream of the transcription start site and the read density sum in 500-nucleotide-wide windows was calculated for the subsequent 1,000 bases. The highest sum was used for downstream analysis.

Metagene analysis. Each gene included in the analysis is normalized by the mean number of reads in a 400-bp window beginning 100 bases downstream from the transcription start site. A mean read density (MRD) is then calculated for each position over all genes as described below.

$$MRD(i) = \frac{\sum_j^{\text{all genes}} \left(\frac{r_j^i}{\sum_{i=100}^{500} r_j^i / 400} \right)}{\sum_j^{\text{all genes}} 1}$$

where r_j^i are the reads for the j th gene at the i th position after the transcription start site.

Extracting pause positions. Pauses were identified in previously annotated transcription units³ of well-expressed genes. Pauses were defined as having reads

higher than three standard deviations above the mean of the surrounding 200 nucleotides which do not contain pauses. Pauses were required to have at least four reads regardless of the gene's sequencing coverage. Sequence consensus was calculated by WebLogo 3 (<http://weblogo.threeplusone.com/>)⁴⁵.

Comparing pause densities to nucleosome positions. Nucleosome positions²³ were assigned as +1, +2, +3 etc according to their position relative to transcription start sites. The mean pause density (MPD) relative to a particular nucleosome was determined by the number of pauses observed at that position (N_p) divided by the total number of opportunities it could be observed there (N_o):

$$\text{MPD}_k(x) = \left(\frac{N_p}{N_o} \right)_x = \frac{\sum_i^{\text{all genes}} g_i(y)}{\sum_i \text{genes with TSS} < y} 1$$

$$y = n_i^k + x$$

where k is the nucleosome number, $g(y)$ is the binary function indicating whether a pause occurs at y , and n_i^k are the centre nucleosome positions. For the +2 and

+3 nucleosomes, the number of pause opportunities was uniform at every position and was simply the number of genes included in the analysis. The +1 nucleosome analysis required that the number of pause opportunities at each position represent the number of genes where that position occurs after the transcription start site. The error of the pause density was calculated via the standard deviation of the binomial distribution

$$\frac{\sqrt{N_p \left(1 - N_p / N_o \right)}}{N_o}$$

The densities were then binned by averaging across windows ten nucleotides wide. The error for each bin was calculated by computing the sum of the variances of the binned measurements and calculating the square root.

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