Asymmetry between Activation and Deactivation during a Transcriptional Pulse

Highlights
- Gene transcription switches between variable rates
- Single-cell microscopy and mathematical modeling quantifies switch dynamics
- We observe an asymmetry in the activation/deactivation of transcriptional bursts

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In Brief
Dunham et al. apply mathematical tools to single-cell imaging data to quantify gene expression dynamics, revealing a pattern of all-or-nothing activation and a graded deactivation in the regulation of transcriptional bursts.
Asymmetry between Activation and Deactivation during a Transcriptional Pulse

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INTRODUCTION

Transcription in eukaryotic cells occurs in gene-specific bursts or pulses of activity. Recent studies identified a spectrum of transcriptionally active “on-states,” interspersed with periods of inactivity, but these “off-states” and the process of transcriptional deactivation are poorly understood. To examine what occurs during deactivation, we investigate the dynamics of switching between variable rates. We measured live single-cell expression of luciferase reporters from human growth hormone or human prolactin promoters in a pituitary cell line. Subsequently, we applied a statistical variable-rate model of transcription, validated by single-molecule FISH, to estimate switching between transcriptional rates. Under the assumption that transcription can switch to any rate at any time, we found that transcriptional activation occurs predominantly as a single switch, whereas deactivation occurs with graded, stepwise decreases in transcription rate. Experimentally altering cAMP signalling with forskolin or chromatin remodelling with histone deacetylase inhibitor modifies the duration of defined transcriptional states. Our findings reveal transcriptional activation and deactivation as mechanistically independent, asymmetrical processes.

SUMMARY

Quantitative analysis of optical reporter genes, such as the firefly luciferase gene, provides real-time measurements of transcription dynamics from individual living cells. This enables the development of mathematical models and the formulation of subsequent hypotheses regarding the nature of bursts of promoter activity in a noisy molecular system. Previously applied “random telegraph” models assumed that transcription operated as a binary function, identifying defined on/off states of activity (Raj and Oudenaarden, 2009; Harper et al., 2011; Suter et al., 2011). However, observation of transcription profiles from large numbers of individual cells in either a basal or stimulated state suggested that variable rates may in fact occur (Harper et al., 2011; Molina et al., 2013; Corrigan and Chubb, 2014), and therefore an on/off model is too simplistic to describe dynamic transcription. While several studies have applied multi-state models of transcription, many categorize transcriptional phases into only a few discrete states (Neuert et al., 2013; Zhang and Zhou, 2014; Zoller et al., 2015; Bintu et al., 2016). Hey et al. (2015) developed a stochastic switch model (SSM) assuming probabilistic interactions between low numbers of intracellular molecules, and extrinsic noise, which is due to more global variations in cell states (Elowitz et al., 2002; Carey et al., 2013; Dadiani et al., 2013). Production of mRNA transcripts occurs in transcript-specific bursts, reliant on promoter architecture (Zoller et al., 2015; Suter et al., 2011; Hocine et al., 2015), chromatin status (Bintu et al., 2016; Noordermeer et al., 2011), and exogenous stimuli (Harper et al., 2011; Molina et al., 2013; Featherstone et al., 2016). These bursts of activity are separated by intermittent “off” periods of transcriptional inactivity (Chubb et al., 2006; Zenklusen et al., 2008; Noordermeer et al., 2011). It has been suggested that this inactive phase comprises multiple independent states (Zoller et al., 2015). This phase includes a refractory period, during which a new round of transcriptional activation cannot be initiated (Harper et al., 2011; Suter et al., 2011). The rate-limiting duration of this state is assumed to be a result of a combination of processes including signaling, chromatin remodeling, transcription factor complex formation, and the recruitment and activation of RNA polymerase II (Larson et al., 2011; Sorre et al., 2014; Bintu et al., 2016).

INTRODUCTION

The expression of many genes has been shown to be highly dynamic and heterogeneous in individual living cells (Raj et al., 2006; Harper et al., 2011). Transcription is an inherently noisy process contributing to intercellular variation, where low molecular numbers amplify this variation (Becskei et al., 2005). An individual cell is subject to both intrinsic noise, which arises from
that transcription could occur at any rate, and could switch in any direction at any time. When applied to the transcription dynamics of the human prolactin (hPrl) gene within the pituitary gland, the model provided a more graded or analog view of transcription (Featherstone et al., 2016). By not limiting the process to discrete transcriptional on-states, the model also provides a more quantitative insight into dynamic and complex transcriptional states, supporting recent work suggesting the occurrence of transcription along a continuum of rates (Corrigan et al., 2016; Sepúlveda et al., 2016).

RESULTS
Promoter Structure and Associated Chromatin Remodeling Confer Expression Dynamics

To test the importance of promoter complexity on the pattern of pulsatile expression, we created two reporter cell lines by stably transfecting one copy of a −840/+1 bp or −3,348/+1 bp hGH proximal promoter-luciferase construct into the GH3 rat pituitary cell line (Figures 1A, 1B, S1A, and S1B). Both stable transfectant cell lines exhibited dynamic and heterogeneous luminescence activity in the presence or absence of serum or stimuli (Figures 1A–1F and S2). The luminescence produced by either hGH construct was approximately tripled following stimulation of the cyclic AMP (cAMP) signaling pathway with forskolin (Fsk) (Figures 1D and 1F). However, while the −840/+1 bp promoter showed no response to the inhibition of histone deacetylase (HDAC) by trichostatin A (TSA), the larger −3,348/+1 bp promoter greatly increased luminescence output. The combination of Fsk and TSA has a synergistic effect, further increasing −3,348/+1 bp luminescence production. (−840/+1 bp: BSA, n=75; Fsk, n = 41; TSA, n=33; Fsk + TSA, n=46; −3,348/+1 bp: BSA, n = 97; Fsk, n = 59; TSA, n = 42; Fsk + TSA, n = 40).

Figure 1. Heterogeneous Promoter-Specific Activity in Single Cells
(A and B) Example images of single-cell bioluminescence in GH3 pituitary cells expressing either an −840/+1 bp (A) or −3,348/+1 bp (B) hGH-luciferase transgene in serum-starved conditions (BSA) for 0–48 hr.
(C and E) Real-time luminescence plots demonstrate heterogeneous expression dynamics (shades of green/blue) but an overall stable population output (black).
(D and F) Mean population luminescence following stimulation shows promoter-specific responses. While the luminescence of both promoters is tripled within 4 hr upon cAMP stimulation with forskolin (Fsk), only the larger −3,348/+1 bp construct responds with increased output following HDAC inhibition by trichostatin A (TSA). The combination of Fsk and TSA has a synergistic effect, further increasing −3,348/+1 bp luminescence production. (−840/+1 bp: BSA, n=75; Fsk, n = 41; TSA, n=33; Fsk + TSA, n=46; −3,348/+1 bp: BSA, n = 97; Fsk, n = 59; TSA, n = 42; Fsk + TSA, n = 40).

Previous single-cell bioluminescence analysis of luciferase activity driven by a short human growth hormone (hGH) proximal promoter (−496/+1 bp) has highlighted the pulsatile dynamics of this gene (Norris et al., 2003). The cell-type-specific expression of this pituitary hormone is controlled by an extensive sequence that regulates chromatin remodeling and proximal promoter access (Ho et al., 2002, 2011, 2015; Shawchuk et al., 2002, 2006; Bodner et al., 1988; Lipkin et al., 1993; Cohen et al., 1999; Alonso et al., 1998). We have taken advantage of the pulsatile nature of hGH gene expression to assess the influence of different types of regulatory element on gene transcription dynamics. Furthermore, we compared these dynamics with those of the hPrl gene, an independently regulated pituitary hormone gene (Niall et al., 1971). The statistical modeling approach was validated by single-molecule fluorescence in situ hybridization (smFISH), quantitatively confirming the modeled estimates of the distribution of numbers of mRNA molecules. These data indicate that for each promoter there is an all-or-nothing “on-switch” in transcriptional activation, whereas transcriptional inactivation involves a decaying series of “off-switches.” Our data suggest mechanistic hypotheses for the way in which promoters may fully engage with transcriptional machinery and then may switch off through a series of distinct states.
correlation of luminescence patterns between individual cells (Figures S1C and S1D) suggests that this synergistic increase was not due to increased activity of an already active subpopulation but to a reduction in cell-cell heterogeneity. We and others (Harper et al., 2011; Suter et al., 2011) have proposed that the "off-state" involves a refractory period that originates from a period of chromatin remodeling, during which a new round of transcription cannot be initiated. For the larger promoter this chromatin-remodeling step may promote population heterogeneity, and therefore protect against noisy activation.

Estimation of Transcriptional States Using a Stochastic Switch Model Suggests a Continuum of Rates
An SSM was previously developed and used here for inference on switches in transcription rate from the observed reporter gene activity (Figure 2A) through the fitting of various parameter...
prior distributions (Figure 2B) (Hey et al., 2015). Through 30,000 iterations of this reverse jump-Markov chain Monte Carlo-based algorithm, the locations of significant changes in transcription rate were identified. These estimations produced several potential switch profiles for each single-cell luminescence trace, each with varying probability (Figure 2C). These probable scenarios inform the generation of a single likely transcription profile (Figure 2D).

The SSM was evaluated using smFISH (Figures 2E and 2F) as an independent and directly quantitative assay to measure the distribution of mRNA molecule numbers. This showed close agreement between observed and modeled mRNA numbers (Figure 2G). Starting with the assumption that any transcription rate is possible, and that switching between variable rates is independent of the previous rate, we observed a significant inverse relationship between the rate and duration (\( \tau \)) of a transcriptional phase when fitted to the single-cell luminescence data produced by each \( hGH \)- and \( hPrl \)-luciferase promoter constructs (Figures 2H–2J). High rates of transcriptional activity lasted for the shortest periods of time, while low rates of transcription could be maintained for longer. The continuous distribution of rates as modeled by the SSM approach supported the view that rather than this being a discrete binary on/off system, transcription occurs along a spectrum of possible rates (Molina et al., 2013; Zhang and Zhou, 2014; Featherstone et al., 2016; Corrigan et al., 2016; McNamara et al., 2016).

Asymmetrical Transcriptional Control

Unlike previous binary models of transcriptional activity, the SSM used here estimates a switch in rate in any direction, independent of any previous switch direction or rate. Using such assumptions, we observed an unequal number of rate-increasing and rate-decreasing switches, with approximately twice as many “down-switches” identified per cell for all \( hGH \)- and \( hPrl \) promoter constructs throughout a 48-hr period (Figure 3A). Despite this asymmetry in the number of these deactivating switches, the overall transcription rate across the entire cell population was maintained (Figures S3A and S3B), indicating a constant and stable population output. Therefore the greater number of down-switches was not due to a general systematic reduction in transcriptional activity throughout the experimental period, but instead indicates the nature of the underlying process. A slight increase in the number of deactivating switches is observed following Fsk stimulation of the \(-840/+1 \) bp fragment (Figure 3B), but the number of down-switches remains unaffected by TSA treatment. A significant decrease in the number of these deactivating steps is seen of the \(-3,348/+1 \) bp promoter when HDAC is inhibited, reducing the number of switches by half.
Figure 3C plots the amplitude of two consecutive switches allowing for the identification of any consistent switch patterns. We identify predictable patterns for each scenario when a Down switch is the secondary switch, with Up-Down and Down-Down switch amplitudes being highly correlated (Table S3). This predictability suggests a system memory controlling the transcription-deactivating mechanism. Over 50% of each cell line consistently exhibits a graded pattern of transcriptional switching, with more cells containing the larger constructs producing the asymmetrical graded deactivation pattern (Figures 3D, S3C, and S3D).

While previous research has suggested a transcriptional pulse to be either binary or graded (Biggar and Crabtree, 2001; Giorgetti et al., 2010; Bintu et al., 2016; Ochab-Marcinek and Tabaka, 2015), we propose that a single transcriptional burst is predominantly a combination of an all-or-nothing activation, followed by a graded reduction, tuning transcriptional output along a continuum to the post-stimulatory environment.

Quantifying Mechanistic Influences on Transcription Phase Duration
Noting the characteristic differences between the regulation of transcriptional activation and deactivation (Figures 2 and 3), we investigated a modification to the duration of phases following a deactivating switch (Figure 4). Observing the duration distribution of such phases (Figures 4A–4C), we identified promoter-specific variations in response to treatment. While the short hGH promoter-produced periods remain unchanged when treated, the durations produced by the larger hGH and hPrl promoter constructs are significantly altered. The median phase duration of both the −3,348/+1 bp hGH and hPrl promoter is increased by HDAC inhibition. Interestingly, all three promoters produce an absolute minimum deactivated phase duration of 3 hr when unstimulated in BSA medium, suggesting a state during which transcription cannot be activated (Figure S4). For the larger hGH and hPrl promoter constructs, this period is reduced upon removal of HDAC activity. To statistically quantify this observation we developed a three-state model of transcription, refractory model (Figure 4D). This model infers the duration of two independent states (“off” and “primed”), which comprise the duration of the phase following a deactivating switch (Figures 4A–4C). Statistical modeling identified significant reductions in the duration of the refractory state for each promoter following specific treatments (Figures 4E–4G). Most notably, the median refractory phase of the shorter hGH promoter is reduced from 4.3 hr (in BSA) to 1.9 hr with Fsk treatment, while the larger...
gene expression control is far more dynamic than was once thought. Recent work describing transcription dynamics in living cells and tissues has shown that at all levels from signaling to protein synthesis, gene expression control is far more dynamic than was once thought (Harper et al., 2011; Molina et al., 2013). The general view has been that these dynamics arise from stochastic switching between on- and off-states of transcription. The observation of the importance of a refractory phase in the pulsatility of gene expression (Harper et al., 2011; Suter et al., 2011; Cesbron et al., 2015; Zoller et al., 2015) has suggested the idea that the timing may involve a key period of chromatin remodeling that determines characteristic transcription dynamics for a given gene. Here, we show the non-intuitive result that there is a consistent asymmetry between the probability of switching between activation and deactivation of transcription for different promoters, with different integration sites. While the occasional stepwise increase in transcriptional rates is possible, we predominantly observe a single, all-or-nothing activating switch followed by several steps of graded reductions of decreasing amplitude. These results arise from unbiased statistical analysis of time-lapse reporter gene expression. The picture that emerges is that genes tend to achieve the highest transcriptional state of the current cycle within a single switch, but then switch down through a series of different levels of transcription with the first switch often being the largest (Figure 2). Thus in a given period we observe more down-switches than up-switches, even in a population of cells that are maintaining a stable level of gene expression. This phenomenon was observed for two different hGH promoter constructs (Figure S1), and also for the prolactin promoter. Although these genes are evolutionarily related and are both regulated by the Pit-1 transcription factor, they display different structural characteristics and organization, and cell-type and development-specific expression.

Transitions between binary and graded processes may allow for the tuning of transcriptional output according to the initial activating stimulus. The binary-like activating switch is likely to be a function of several cumulative transcription-initiating interactions breaching a threshold of activation. Above this threshold, the increasing promoter and transcription factor interaction may increase the probability and opportunity of establishing a stable transcription complex, inhibiting further graded increases and recruiting polymerase II (Pol II) to its current maximal capability. Following activation, these promoter interactions may degrade and dissociate at varying rates, along with Pol II pausing, which will therefore gradually reduce the overall rate of transcription in a stepwise manner until a level of activity is met that reflects the new cell state (Saccani et al., 2004; Selia et al., 2009; Hammar et al., 2014; Jonkers and Lis, 2015). Larger promoter fragments driving the luciferase reporter appeared to direct an increased number of down-switches during the stepwise decay, likely due to more extensive and maintained chromatin status and a greater number of promoter-protein interactions.

The stochastic bursting of transcriptional activity is believed to be a general phenomenon, with temporal dynamics being highly promoter specific (Raj and Oudenaarden, 2009; Suter et al., 2011; Coulon et al., 2013), and individual transcription events for specific mRNAs occurring on a time scale of minutes (Muramoto et al., 2012). We show here that longer-scale maintenance of chromatin status is associated with defined characteristics of timing in the transcriptional cycles that we observe. Gene-specific phases of transcription and refractory periods are thought to contribute to the cyclic bursts, and their variability results in heterogeneous transcriptional states across a population of cells or tissue (Harper et al., 2010; Featherstone et al., 2011, 2016). In the present study, removal of chromatin-remodeling processes through HDAC inhibition removes this heterogeneity and alters the defined bursting pattern. Evidence here suggests that simpler promoters may be more subject to fluctuations in extrinsic noise through modification of transcriptional phases upon stimulation of signaling pathways. It is important to note that differing promoters with different architecture and complexity here display essentially the same patterns of activation and inactivation. In the same way that bursting behavior has been seen as a common feature for multiple regulated promoters (Suter et al., 2011), it may therefore be true that binary activation and graded inactivation are a common transcriptional phenotype. The potential transcription factor binding sites in the two GH promoters are shown in Figure S1, but the exact role of specific response elements and factor binding sites remains to be evaluated in detail.

Unbiased variable-rate modeling provides a robust inference of the underlying dynamics of transcription. This modeling approach has previously been evaluated using biological and synthetic data (Hey et al., 2013). Here we have tested its inferences regarding mRNA molecule number by direct assay using smFISH. Our statistical analysis has shown that transcription is likely to occupy a distribution of active rates. Although we still categorize transcription into discrete phases, these phases display a spectrum of rates (Molina et al., 2013; Featherstone et al., 2016; Corrigan et al., 2016). In addition to the previously identified gene-specific transcriptional burst frequency, amplitude, and duration, we also observe promoter-specific graded switching patterns. Greater promoter complexity is likely to support a rich variety of epigenetic control mechanisms to generate accurate regulation across different time domains in cells and tissues.

**STAR METHODS**

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Supplemental Information includes four figures and three tables and can be found with this article online at https://doi.org/10.1016/j.cels.2017.10.013.

AUTHOR CONTRIBUTIONS
L.S.S.D., H.M., M.R.H.W., and J.R.E.D. designed research; L.S.S.D., C.V.H., D.G.S., A.M., and K.F. contributed new reagents/analytic tools; L.S.S.D. and H.M. analyzed data; D.A.R., B.F., M.R.H.W., and J.R.E.D. supervised research; L.S.S.D., H.M., and J.R.E.D. wrote the paper.

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# STAR★METHODS

## KEY RESOURCES TABLE

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| **Bacterial and Virus Strains** | | |
| DH5a Competent Cells | Invitrogen | 18265017 |
| **Chemicals, Peptides, and Recombinant Proteins** | | |
| Forskolin | Sigma-Aldrich | F3917 |
| Trichostatin A | Sigma-Aldrich | T1952 |
| Cycloheximide | Sigma-Aldrich | C104450 |
| Actinomycin D | Sigma-Aldrich | 01815 |
| **Critical Commercial Assays** | | |
| FuGENE Transfection reagent | Promega | E2311 |
| Copy number assay kit | ThermoFisher | 4442487 |
| PureLink Genomic DNA kit | ThermoFisher | K182001 |
| Stellaris FISH | Biosearch Technologies | VSMF-1007-5 |
| Miniprep | Qiagen | 27104 |
| Maxiprep | Qiagen | 10023 |
| **Deposited Data** | | |
| Raw data | This paper | N/A |
| **Experimental Models: Cell Lines** | | |
| GH3 | Tashjian et al., 1968 | N/A |
| GH3/hGH3348 | This paper | N/A |
| GH3/hGH840 | This paper | N/A |
| GH3/hPrl | This paper | N/A |
| **Oligonucleotides** | | |
| Copy number Taqman probe | FAM-AATCGCTCAACAGTATGAG-MGB | N/A |
| Copy number luciferase forward primer | CCGCGAAGCAGCATTTATAATGA | N/A |
| Copy number luciferase reverse primer | CCACGGTAGGCTCGGAAA | N/A |
| GGT1 probe | Cy5-CCGAGAAGCAGCCACACCTACCTAC-GHG | N/A |
| GGT1 forward primer | CCACCGCTCTCCTACTCTAC | N/A |
| GGT1 reverse primer | GCCCAGACAGCTGTTGTC | N/A |
| **Recombinant DNA** | | |
| pGL3-hGH3348/+1-luciferase | This paper | hGH-3348/+1bp |
| pGL3-hGH -840/+1-luciferase | This paper | hGH-840/+1bp |
| pGL3-hGH96/+1-luciferase | Norris et al., 2003 | hGH-96/+1bp |
| Cosmid K2B | Jones et al., 1995 | N/A |
| pGL3-5kbPrl- luciferase | Takasuka et al., 1998 | hPrl |
| pGL3-basic | Promega | E1751 |
| pVITRO2-hyg-mcs | Invivogen | pvitro1-mcs |
| **Software and Algorithms** | | |
| Micro-Manager (Version 1.4) | Edelstein et al., 2014 | https://micro-manager.org/wiki/Micro-Manager_Version_Archive |
| UCSC Genome Browser | Kent et al., 2002 | https://genome.ucsc.edu/ |
| TRANSFAC 7.0 Public 2005 | Matys et al., 2006 | http://gene-regulation.com/pub/databases.html |
| AQM6 Image Analysis Software (Kinetic) | Andor | |

(Continued on next page)
CONTACT FOR REAGENT AND RESOURCE SHARING
Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact Julian Davis (julian.davis@manchester.ac.uk).

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Cell Lines
Female rat pituitary GH3 cells were used as a model in this study (Tashjian et al., 1968). These cells require growth in phenol-red free DMEM with pyruvate (Gibco) supplemented with 1% glutamine, and 10% fetal calf serum (Gibco). All imaging was performed in serum-starved conditions in phenol-red free DMEM with pyruvate, 1% glutamine, and 0.25% bovine serum albumin (Gibco). Cells were maintained at 37°C and 5% CO₂.

METHOD DETAILS

Plasmid Preparation and Transfection
Human growth hormone promoter-luciferase constructs were generated via amplification from the cosmid K2B (Jones et al., 1995). The amplified fragments were digested with SacI and ligated into the SacI digested pGL3-basic plasmid (Promega) 5’ to the luciferase reporter gene coding sequence. Generation of the 5kb prolactin-luciferase construct and cell line has been previously described (Takasuka et al., 1998). 10μg of either plasmid was isolated via miniprep (Qiagen) and was mixed with 3μg pVITRO2-hyg-mcs in 1ml of 150mM NaCl to allow for co-transfection into 1 x 10⁵ GH3 cells using FuGENE transfection reagent (Promega). Media was changed every 3 days post transfection, and was supplemented with 500μg/ml hygromycin selection antibiotic. Positively transformed cells formed colonies after 2-3 weeks and were ring cloned into individual wells of a 48 well plate and cultured. The clones were screened for luciferase expression and response to stimuli (5nM forskolin, 100nM dexamethasone, and 50μM T₃), prior to establishing gene copy number.

Copy Number Validation
The copy number of GH3 transfected luciferase constructs was quantified using a customisable copy number assay kit (Life Technologies), comparing qRT-PCR amplification of the luciferase gene with known controls.

Genomic DNA Extraction
The PureLink Genomic DNA kit (Life Technologies) was used to extract all genomic DNA (gDNA) from the transformed GH3 cell lines following the manufacturer’s instructions. gDNA concentration and purity was quantified using a Nanodrop 2000 UV-Vis spectrophotometer (Thermo Scientific). The absorbance was measured between 260 and 280nm. All gDNA were of sufficient purity with a A₂₆₀/A₂₈₀ ratio between 1.9-2.1.

qRT-PCR
The gDNA was amplified using the copy number assay kit (Life Technologies). 20ng of gDNA was mixed with manufacturer instructed amounts of TaqMan genotyping master mix, copy number assay, copy number reference assay and nuclease-free water. The copy number assay contains a TaqMan probe and quencher directed to the luciferase using customised luciferase oligonucleotides (Forward- CCGCAGACACATTATGAA; Reverse- CCACGTTAGGCTGCGAAA; Probe- FAM-AATTGCACTAGTATGGG-MGB) (Semprini et al., 2009). Amplification of the luciferase target was compared directly to the amplification of the GGT1
housekeeping gene using primers and a probe previously described by Pawlak et al. (1988) (Fwd- CCACCCCTTCCCTACTCCTAC; Rev- GGCCACAGAGCTGGTTGTC; Probe- Cy5-CCGAGAAGCAGCCACAGCCATACCT-BHQ2). qRT-PCR was performed using a StepOnePlus (Applied Biosystems) with cycling parameters as follows: initial denaturation at 95°C for 10 min; 40 cycles of 95°C 15 secs, 60°C 60 secs; hold at 72°C for 5 min.

**Real-Time Luminescence Imaging**

**Image Acquisition**

1 x 10^5 transformed GH3 cells were plated onto a CELLview 35mm glass-bottomed cell culture dish in FCS-containing media. 24h prior to imaging cells were washed with PBS and serum starved in media containing 1mM luciferin. Application of stimuli was performed immediately prior to transfer to the stage of a Zeiss Axiovert 200 equipped with an XL incubator to maintain the cells at 37°C, 5% CO2. Luminescence images were obtained using a Fluar 10x, 0.5NA objective (Zeiss) and captured using an ImagEM EM-CCD cooled camera (Hamamatsu photonics). Images were integrated over a 15m period using a 1 x 1 binning and acquired using Micro-Manager software (Version 1.4). In treated conditions, forskolin (50μM) and trichostatin A (50nM) were added immediately prior to imaging.

**Protein and mRNA Inhibition**

For the back-calculation from observed luminescence to the estimated transcriptional dynamics we required the calculation of mRNA and protein degradation rates. Whilst both hGH-luciferase constructs have identical 3' UTRs, their respective genome insertion sites may vary and therefore affect post-translational modification and mRNA and protein half-lives. We imaged single-cell bioluminescence of transformed GH3 cells when stimulated with 5nM forskolin followed by the application of cycloheximide (10μg/ml) or actinomycin D (3μg/ml) (Sigma) to block translation and transcription, respectively.

**Single Molecule RNA-In Situ Hybridisation**

**Sample Preparation**

4 x 10^4 transformed GH3 cells were plated in FCS-containing media onto glass coverslips pre-treated with poly-L-lysine. Cells were cultured for two days, treated with 5μM forskolin or left untreated, and fixed four hours later. Coverslips were fixed then hybridised with the Stellaris FISH probe set against firefly luciferase conjugated with Quasar 670 dye (VSMF-1007-5, Biosearch Technologies), using the manufacturer’s protocol for adherent cells.

**Image Acquisition**

Images were obtained using a Delta Vision Core restoration microscope (Applied Precision) using a 60x/NA 1.42 Plan Apo objective and Sedat Quad filter set (Chroma Technology), and collected using a Coolsnap HQ2 camera (photometrics)(Mueller et al., 2013).

**Quantification and Statistical Analysis**

**Bioinformatic Promoter Analysis**

hGH promoter sequence was obtained using the UCSC Genome Browser (http://genome.ucsc.edu/), locating the hGH start site and exporting the upstream promoter sequence. Transfac analysis was performed on the -3348bp of promoter sequence using weight limits: matrix = 0.9, core = 0.95 to identify likely transcription factor binding sites (TRANSFAC 7.0 Public 2005).

**Analysis of Image Data**

Individual cells were tracked using AQM6 image analysis software (Kinetic). Regions of interest were drawn around each individual cell, and mean intensity data collected. Monadic noise between individual frames was removed, and background noise was subtracted from the luminescence signal. Analysis of an individual cell ceased at the point of cell division.

**qRT-PCR Analysis**

Relative fluorescence was then analysed using Copy Caller Software (ThermoFisher) generating a relative quantification value for number of gene copies within each clone.

**FISH Data Processing**

Raw images were deconvolved using the Softworx software then converted to .tif stacks using a custom ImageJ script. Transcript counts for each cell were determined using FISH-quant (Jin et al., 1999).

**Correlation Coefficient Calculation**

To analyse the temporal correlation of luminescence dynamics in response to specific treatments we calculated the correlation coefficient for sequential 1h pooled periods. The correlation of luminescence throughout an hour period was calculated between each and every single cell under a particular treatment. The median and distribution of these hour correlation coefficients were plotted for a 20h period demonstrating the degree of similarity between individual cells. The significance of each pool was compared using an unpaired t-test.

**Stochastic Switch Model**

To estimate switches between variable rates of transcription, we applied a previously developed stochastic switch model (Hey et al., 2015). This reversible jump Markov chain Monte Carlo algorithm estimates the temporal location of a significant change in transcriptional activity through the back-calculation of protein and mRNA degradation rates (which remain unchanged by stimulation, Table
The below stochastic reaction network is incorporated within this model, allowing for the identification of luminescence above stochastic noise within the system:

\[ \emptyset \xrightarrow{\beta(t)} \text{mRNA} \]

\[ \text{mRNA} \xrightarrow{\delta_m} \emptyset \]

\[ \text{mRNA} \xrightarrow{\delta_m} \text{mRNA} + \text{Protein} \]

\[ \text{Protein} \xrightarrow{\delta_p} \emptyset \]

The degradation rates of reporter mRNA and protein are denoted by \(\delta_m\) and \(\delta_p\), respectively, whilst \(\alpha\) denotes the rate of translation and \(\beta(t)\) denotes the time varying rate of transcription. Our transcription function is given by:

\[ \beta(t) = \beta_i \text{ for } t \in [s_{i-1}, s_i] \text{ for } i = 1, \ldots, K, \]

where \(K\) is the number of transcriptional switches, occurring at times \(s_1, s_2, \ldots, s_K\) and \(\beta_1, \beta_2, \ldots, \beta_K\) are the corresponding transcriptional rates. We impose no restriction to the form of the transcriptional levels but note that the conventional binary switch behaviour can be seen as a specific example where \(\beta_i = \beta_{LOW}\) if the gene is inactive in the time period \([s_{i-1}, s_i]\) or \(\beta_i = \beta_{HIGH}\) if the gene is active.

Assuming light intensity measurements are related to reporter protein levels by the equation,

\[ Y(t) = \alpha P(t) + \varepsilon(t), \]

\[ \varepsilon(t) \sim N(0, \sigma^2), \]

inference is performed through the linear noise approximation to the stochastic reaction network coupled with this measurement equation to obtain the posterior transcriptional function for each single cell. To ensure model identifiability, we impose informative prior distributions about the degradation parameters, obtained from independent half-life experiments. In addition, we specify a hierarchical framework over each dataset, as individual parameters are unlikely to change substantially.

In order to estimate both the number and positioning of transcriptional switches, we employ a reversible jump Markov Chain Monte Carlo (MCMC) algorithm (Green, 1995). Consequently, the posterior distribution consists of all possible transcriptional profiles. In order to extract the information regarding the estimated transcriptional dynamics, the posterior samples go through a post-processing procedure outlined below.

A parametric model is fitted to the marginal posterior switch distribution (as described by Jenkins et al., 2013). Specifically, a Gaussian mixture model is fitted to the marginal posterior distribution of the possible switch times. All possible sub-models are extracted, to take into account the co-occurrence of switches. For example, if the marginal posterior has two possible switch positions, the sub-models will consist of a zero switch model, two mutually exclusive one switch models, and the two switch model. Counting the frequency with which each of the sub-models was sampled in the MCMC, we can associate a weight or probability to each sub-model.

Therefore, this post-processing procedure associates each single cell to a set of mutually exclusive transcriptional profiles. The analysis presented in the main paper has been calculated from the set of all possible transcriptional profiles, weighted by their probability of occurrence.

**Protein and mRNA Half-Life Estimation**

**Half-life Calculation**

We used these luminescence decay assays to first calculate the protein degradation rate \(\delta_p\) using the function:

\[ \frac{dP}{dt} = c_p - \delta_p P(t) \]

where \(c_p\) is a small non-negative constant.

This then allowed the estimation of the mRNA degradation rate \(\delta_m\) from the transcription inhibition (actinomycin D) induced decay using the function,

\[ \frac{dM}{dt} = c_M - \delta_m M(t) \]

where \(c_M\) is a small non-negative constant that is close to zero if transcription is fully inhibited.
Refractory Model
Statistical analyses of refractory (off) period modification were performed using a three-state model of transcription (Figure 3B). The parameter values were estimated by analysing the transcription profiles statistically inferred by SSM to real-time luciferase datasets. The maximum likelihood estimates of refractory and Off periods, $T_1$ and $T_2$, were obtained by fitting the sum of two exponential distributions to the durations after a down switch with the assumption of $T_1 < T_2$, while On period, $T_0$, was estimated by fitting an exponential distribution to the durations after an up switch. Both durations in the SSM results were analysed as right-censored ones. These period estimations are detailed below. Low and high rates, $\beta_L$ and $\beta_H$, are the respective transcription rates after a down and up switch.

Maximum Likelihood Estimation of Refractory (T1), and Off (T2) Periods
In the following, capital $T$ denotes a period parameter, while small $t$ its realisation in the SSM-generated Markov chain, which represents the posterior distribution of the parameters. An inter-switch duration ($t$) is called complete if it is flanked by two switches, but right-censored if it is open-ended.

Pre-processing
In the continuous SSM, it is often the happening that a down or up switch is followed by the same-direction switch. To fit the SSM result into the current discrete on-off-primed model, such consecutive durations after same-direction switches are merged to form a single inter-switch duration.

Refractory and Off Periods Estimation
As Poisson processes, refractory and off periods, $t_1$ and $t_2$, follow respective exponential distributions, $t_1 \sim \text{Exp}(-\lambda_1)$ and $t_2 \sim \text{Exp}(-\lambda_2)$, where $\lambda_1$ and $\lambda_2$ denote respectively the inverse of $T_1$ and $T_2$. If $\lambda_1 = \lambda_1$ is assumed, the probability density function ($f_d(t_d)$) of the sum of those two random variables, $t_d = t_1 + t_2$ is given as follows.

$$f_d(t_d) = \frac{\lambda_1 \lambda_2}{\lambda_2 - \lambda_1} \left( \exp(-\lambda_1 t) - \exp(-\lambda_2 t) \right)$$

while its corresponding distribution function is given as follows.

$$F_d(t_d) = \frac{1}{\lambda_2 - \lambda_1} \left( \lambda_2 \left( 1 - \exp(-\lambda_1 t) \right) - \lambda_1 \left( 1 - \exp(-\lambda_2 t) \right) \right)$$

For given sets of complete and censored duration data, $(t^{\text{comp}})$ and $(t^{\text{cens}})$, a logarithmic likelihood is given as follows.

$$l(T_1, T_2) = \sum f_d(t^{\text{comp}}; T_1, T_2) + \sum \left( 1 - F_d(t^{\text{cens}}; T_1, T_2) \right)$$

With the assumption of $T_1 < T_2$, $T_1$ and $T_2$ are simultaneously estimated as argmax $l(T_1, T_2)$.

DATA AND SOFTWARE AVAILABILITY
Raw data of single cell luminescence profiles is available in Table S1.
Refractory Model. Matlab scripts can be freely accessed at Mendeley Data: https://doi.org/10.17632/wjyccvc2zc.2
These Matlab scripts will infer the duration of the refractory period from stochastic switch model-identified transcription periods.