High levels of Dorsal transcription factor downregulate, not promote, *snail* expression by regulating enhancer action

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Abstract

In *Drosophila* embryos, genes expressed along the dorsal-ventral axis are responsive to concentration of the Dorsal (DI) transcription factor, which varies in space; however, levels of this morphogen also build over time. Since expression of high-threshold DI target genes such as *snail* (*sna*) is supported before DI levels peak, it is unclear what role increasing levels have if any. Here we investigated action of two enhancers that control *sna* expression in embryos, demonstrating using genome editing that DI binding sites within one enhancer located promoter proximally, *sna.prox*, can limit the ability of the other distally-located enhancer, *sna.dis*, to increase *sna* levels. In addition, MS2-MCP live imaging was used to study *sna* transcription rate in wildtype, *dl* heterozygote, and a background in which a photo-sensitive degron is fused to DI (*dl-BLID*). The results demonstrate that, when DI levels are high, DI acts through *sna.prox* to limit the activity of *sna.dis* and thereby influence *sna* transcription rate. In contrast, when DI levels are kept low using *dl-BLID*, *sna.prox* positively influences *sna* transcription rate. Collectively, our data support the view that DI's effect on gene expression changes over time, switching from promoting *sna* expression at low concentration to dampening *sna* expression at high concentration by regulating enhancer interactions. We propose this differential action of the DI morphogen is likely supported by occupancy of this factor first to high and then low affinity binding sites over time as DI levels rise to coordinate action of these two co-acting enhancers.

Significance statement

A gradient of the maternal transcription factor Dorsal is important for establishing spatial expression of target genes along the dorsal-ventral axis of *Drosophila* embryos. Dorsal levels are also dynamic as nuclear concentration builds in time. Surprisingly, expression of high-threshold target genes such as *snail* is supported before levels peak, raising the question why levels continue to build. Our data support the view that peak Dorsal levels act to
preferentially support activity of one enhancer over another to effectively decrease \textit{snail} expression. In addition, while the morphogen Dorsal acts early to support gene expression, later it effectively acts as a damper to limit gene expression. Our results suggest other morphogens also have effects on gene expression that change over time.

\textbf{Introduction}

Recent studies have shed light on the importance of the timing of morphogen exposure during development and the duration of morphogen input on target gene expression (rev. in 1, 2–4). Despite accumulating data supporting temporal roles for morphogens during development, it remains unclear in most cases whether morphogens directly support multiple actions over time and/or include indirect responses of the underlying gene regulatory networks. In this study, we investigated the temporal action of Dorsal (DI), an atypical morphogen and transcription factor, which patterns the dorsal-ventral (DV) axis in a concentration-dependent manner. While genes expressed in ventral and lateral regions along the DV axis are responsive to DI levels (5), we found that some target genes such as \textit{snail} (\textit{sna}), once initiated, no longer require DI-input to sustain expression (6). DI becomes dispensable for \textit{sna} expression once the early DI-target gene \textit{twist} (\textit{twi}) is expressed, demonstrating that the gene network functioning downstream of DI, an indirect response, takes over late. However, DI levels also build in time (7) and it remains unclear what role higher levels of DI support if levels present earlier are sufficient to support gene expression of high-threshold targets such as \textit{sna}.

In particular, we hypothesized that DI's effect on gene expression changes over time, depending on its interaction with distinct enhancers driving particular genes. Previous studies have demonstrated that two enhancers regulate \textit{sna} gene expression in the early embryo, one located in a promoter proximal position (\textit{sna.prox}) and the other located distally (\textit{sna.dis}) (8–11). \textit{sna.prox} includes a \textit{\sim}2 k\text{B} region bound by DI, demonstrated by both chromatin
immunoprecipitation (ChIP) and footprinting (9, 12). *sna.dis* also is bound by DI, identified through ChIP (12, 13). Furthermore, previous studies using large reporter constructs to assay the individual roles of *sna.prox* and *sna.dis* demonstrated that these two enhancers differentially influence levels of *sna* gene expression (8, 14). In particular, deletion of the *sna.prox* enhancer in the context of large reporter constructs results in higher levels of *sna* expression when examined during nuclear cycle (nc) 14 (8, 14), supporting the view that this sequence normally acts as a “damper” to decrease *sna* expression levels during this time. In this study, we investigated how DI levels impact *sna* expression and, in particular, examined the role of *sna.prox* enhancer over time.

**Results**

*Deletions of sna embryonic enhancers from the endogenous sna locus lead to changes in sna expression levels*

To examine the roles of *sna.prox* and *sna.dis* in the native context, we used Crispr-Cas9 genomic editing to delete sequences associated with each enhancer from the *sna* gene locus (Fig. 1A). When a 1.8kB region encompassing the distal enhancer is deleted (Fig. 1A, Δdis1.8), *sna* expression levels are lower than wildtype when qPCR is used to measure total transcript levels at late nc14 (i.e. nc14c, Fig. 1C,F, compare with 1B). Conversely, when a 4.4kB segment encompassing the proximal enhancer (Fig. 1A, Δprox4.4) is removed, *sna* expression level is higher than wildtype (Fig. 1D,F). Similar results were obtained with analysis of enhancer deletions in the context of large reporter constructs encompassing ~25 kB surrounding *sna* locus, suggesting that the proximal enhancer drives weak *sna* expression levels, whereas the distal enhancer supports strong *sna* expression levels (8, 14). Previous studies had suggested that coordination of proximal and distal enhancers is required to support correct *sna* expression levels (8), therefore we used genomic editing to create mutant lines with different *sna.prox*
enhancer deletions to identify the minimal portion able to properly control sna levels of expression.

Two mutant lines were generated with different sized deletions of the proximal enhancer region. These deletions were positioned using information obtained from previous reporter analysis that had highlighted important sequences (Fig. 1A). A 2.0kB portion of the proximal enhancer (i.e. A2.0) was shown to support expression similar to endogenous sna expression; whereas, a 1.6kB portion of the proximal enhancer (i.e. A1.6) is compromised, supporting only partial expression (9). Using genome editing to modify the endogenous sna locus, the first deletion, encompassing a 2.6kB distal sequence (Fig. 1A, Δprox2.6) leaves the full, minimally-defined proximal enhancer intact such that full ventral expression remains (i.e. A2.0, Fig. 1A) (9). In Δprox2.6 embryos, while expression is normal in terms of spatial domain (Fig. 1G), sna levels are slightly lower than wildtype (Fig. 1F). In contrast, when a larger 3.0kB region is deleted using genome editing (Fig. 1A, Δprox3.0), with a breakpoint that is similar to the minimally-active A1.6 reporter construct (9), sna levels are variable and higher than wildtype (Fig. 1F). It was unexpected that embryos that contain a partially active proximal enhancer sequence (i.e. Δprox3.0) are associated with higher sna gene expression.

*Partial deletions of the sna.prox enhancer lead to an increase in sna levels due to uncontrolled sna.dis enhancer activity*

Together with the observation that the distal enhancer supports high sna expression (8, 14), we hypothesized that the higher sna expression found to be associated with Δprox3.0 relates to an overactive distal enhancer. To test this idea, the distal enhancer was removed from the Δprox3.0 background, creating double mutants (i.e. Δprox3.0 Δdis1.8, Fig. 2A). In Δprox3.0 Δdis1.8 double mutant embryos, the sna levels are decreased relative to wildtype, and similar to the sna levels detected in Δdis1.8 single mutant embryos (Fig. 2B, C, compared with Fig. 1F).
This result suggests that the high \textit{sna} levels associated with the \textit{Δprox3.0} mutant background relates to a hyper-active distal enhancer and, additionally, that input from the proximal enhancer is pivotal for regulation of \textit{sna} expression levels.

To further characterize the proximal enhancer’s mechanism of action, we mutated DI binding sites in the \textit{Δprox3.0} background that we hypothesized contains a partially active proximal enhancer; since mutating these sites in the context of reporter assays was associated with reduction of proximal enhancer activity (9). When three DI binding sites that match the JASPAR consensus binding site (http://jaspar.genereg.net), likely of high affinity, are mutated in \textit{Δprox3.0} [i.e. \textit{Δprox3.0(mDi)}, Figs. 2A and S1], the resulting \textit{sna} levels are higher than \textit{Δprox3.0} (Fig. 2C; Fig. 2D, compare with 1E). To test whether the increased \textit{sna} levels associated with \textit{Δprox3.0(mDi)} also relate to a hyper-active distal enhancer, the distal enhancer was additionally deleted from this background [i.e. \textit{Δprox3.0(mDi) Δdis1.8}, Fig. 2A]. In embryos containing this double deletion, \textit{sna} levels are lower than wildtype, but similar to levels detected in \textit{Δdis1.8} embryos (Fig. 2C; Fig. 2E, compare with 1E). This result demonstrates that the increased expression associated with mutation of DI sites within the proximal enhancer stems from increased action of the distal enhancer. Taken together, these results and those in the previous section support the view that a functional proximal enhancer limits activity of the distal enhancer, thereby supporting the physiologically appropriate levels of \textit{sna} expression.

\textit{High DI concentration leads to a decrease in \textit{sna} levels by limiting \textit{sna.dis} enhancer activity}

Previous studies have used the MS2-MCP.GFP system to monitor \textit{sna} transcription from large reporter constructs in which the \textit{sna} gene has been replaced by the MS2.yellow dual-reporter sequence and examined the relative roles of \textit{sna.prox} and \textit{sna.dis} in supporting expression over time (14) (Fig. S2A). These studies found that \textit{sna} transcription decreases during the course of nc14 (14), and were independently confirmed using our imaging settings.
(Fig. S2B). On the other hand, DI concentrations continuously increase during nc14 during this same period (7), demonstrating an anti-correlation exists between DI and sna levels during this stage. This is unexpected as DI is thought to be a positive input to sna and that, moreover, high levels of DI are required to support sna. We hypothesized that at nc14, DI’s effect on sna expression changes, possibly negatively impacting sna levels.

To further investigate this idea that DI negatively impacts sna, sna levels were measured by monitoring sna nascent transcription in embryos containing reduced DI levels (i.e. dl⁺ heterozygotes) to determine whether transcription is increased. The sna MS2 reporter construct was introduced into a dl⁻/+ background and the mean spot size (i.e. mean number of pixels per MS2-MCP.GFP spot), which is correlated to the rate of transcription (15), was examined in early embryos. Initially, at early nc14, the mean spot size is lower in dl⁺ heterozygotes than wildtype (Fig. 3B t=4; Fig. 3C and Movie S1). However, as dl⁺ heterozygous embryos age, the mean spot size increases relative to wildtype (Fig. 3B t=44; Fig. 3C and Movie S1). We hypothesized that this trend relates to decreased proximal enhancer activity as a result of limiting DI.

To test this idea, we measured the mean spot size associated with snaMS2.Δprox, in which only the distal enhancer is active, in a wildtype or dl⁺ heterozygous background, and these levels were compared to the levels supported by both enhancers (i.e. snaMS2.wt). In wildtype embryos, in which DI levels are high, the mean spot size associated with snaMS2.Δprox is higher than snaMS2.wt; most clearly demonstrated at late nc14 (Figs. 3D, S4A, and Movie S2). This suggests that input from the proximal enhancer normally dampens sna levels. In contrast, in dl⁺ heterozygotes at this same stage, no difference is observed in the mean spot sizes of snaMS2.wt or snaMS2.Δprox (Figs. 3E, S4B, and Movie S2). These results demonstrate that reducing DI levels is equivalent to greatly reducing sna.prox activity, and also suggest that the DI levels present in a dl⁻/+ background are lower than necessary to support proximal enhancer activity.
*Decreased levels of Di support cooperative action of sna.prox and sna.dis enhancers*

To further investigate the idea that Di and sna levels anti-correlate, we used the MS2-MCP.mCherry system (16) to examine sna levels *in vivo* that result in real-time from controlled modulation of Di levels accomplished using a previously defined optogenetic approach to degrade Di. In brief, this optogenetic approach supports degradation of Di with fine-scale temporal resolution by fusing a sequence encoding a Blue Light-Inducible Degron (BLID) to the C-terminus of Di (Di-BLID) at the endogenous locus (6, 17). In the dark, the degron domain is inaccessible and Di-BLID protein is present and functional; whereas, upon exposure to blue light, the degron is exposed, resulting in Di-BLID protein degradation in less than 5 min (17). In this previous study, we used MCP.GFP to monitor nascent transcripts from sna.MS2 reporters upon blue light application, but we were unable to monitor the ground state (before illumination with blue laser) because excitation of GFP also supported activation of the photosensitive degron. To address this issue, we used MCP.mCherry to monitor sna.MS2 stem loop production. This allows us to monitor sna transcriptional activity in nc14c before blue laser illumination is applied, which serves as a reference/ground state, and also to monitor subsequent effects after blue light illumination, until nc14d (Fig. 4A). As a control, wildtype embryos devoid of any *di-BLID* fusion were illuminated with a blue laser while detecting MCP.mCherry signal associated with snaMS2.wt (Fig. 4A). Only a slight decrease in mean spot size was observed between embryos with 5 min blue laser illumination relative to embryos kept in dark (Figs. 4C, S5A, S5D, and Movie S5).

To confirm that different levels of Di degradation can be achieved with blue laser illumination of embryos for varying lengths of time, flies with a mCherry-BLID domain fused to the C-terminus of *dl* at the endogenous locus (i.e. *dl-mCh-BLID*) were utilized so that Di protein could be tracked (6). This mutant line allows detection of Di levels by monitoring mCherry
fluorescent signal in the presence or absence (dark) of blue laser illumination. To generate low 
or very low DI levels, \textit{dl-mCh-BLID} embryos were illuminated for 3 min and 5 min, respectively. 
As expected, \textit{dl-mCh-BLID} embryos with 3 min blue laser illumination exhibited lower DI levels 
compared to embryos without illumination, whereas \textit{dl-mCh-BLID} embryo with 5 min blue laser 
ilumination exhibited even lower levels of DI (Figs. 4B, S5, and Movies S3, S4). Since 
increasing illumination results in successively lower levels, we refer to the levels reached after 3 
min illumination as intermediate and the levels reached after 5 min illumination as low.

To assay whether levels of DI differentially impact \textit{sna} transcription, \textit{dl-BLID} 
heterozygous embryos were illuminated for 3 min or 5 min to degrade DI to intermediate or low 
levels, respectively, while detecting MCP.mCherry signal associated with \textit{snaMS2.wt} ("prox and 
dis acting"). \textit{dl-BLID} heterozygotes (\textit{dl-BLID/+}) were utilized for \textit{sna.MS2} assays because the 
\textit{dl-BLID} homozygous embryos (\textit{dl-BLID/dl-BLID}) exhibit only partial \textit{sna} expression. When 
\textit{dl-BLID} heterozygous embryos containing the \textit{snaMS2.wt} reporter are illuminated, the mean 
spot size associated with the MS2-MCP.mCherry signal exhibits an incremental decrease during 
the period of illumination (Figs. 4D, S6B, and S6E), suggesting \textit{sna} transcription decreases as 
DI levels decrease. However, following 5 min illumination, the mean spot size recovers to the 
level present before the illumination (Fig. 4D and Movie S7). In contrast, in the case of only a 3 
min illumination, levels rebound to a state higher than that observed with the 5 min illumination 
or with embryos kept in dark (Figs. 4D, S6A, S6D, and Movie S7 compare with Movie S6). To 
test whether this rebound requires action of the \textit{sna.prox} enhancer, we assayed the effect of 
reducing DI on \textit{snaMS2.Δprox} (i.e. "only dis acting") reporter output. In this case, the mean spot 
size continuously decreases after light illumination and no rebound above control (dark) is 
observed (Figs. 4E, S6C, S6F, and Movies S8, S9). This ability of the \textit{snaMS2.wt} reporter to 
achieve a higher level of expression after blue laser illumination is (i) \textit{sna.prox} 
enhancer-dependent (lost in \textit{snaMS2.Δprox}) and (ii) DI-levels dependent (only associated with 3
min not 5 min illumination). These results also suggest that when low levels of DI are present, the sna.prox enhancer promotes sna transcription; in contrast to the enhancer deletion analysis, which supports the view that the proximal enhancer acts to dampen sna levels (e.g. Figs. 3D and S1).

We propose that the role of the sna.prox enhancer can change in a manner dependent on DI levels. In embryos illuminated for 3 min, the mean spot size associated with snaMS2.wt is larger than that associated with snaMS2.Dprox (Fig. 4G). Whereas Illumination of embryos for 5 min (i.e. more DI-degradation resulting in low DI levels) results in no difference in the mean spot size associated with snaMS2.wt or snaMS2.Dprox (Fig. 4H), suggesting that a critical level of DI is necessary to support proximal enhancer action. Taken together, when DI levels are low but not high enough to support the proximal action, the distal enhancer acts alone to drive weak sna levels; when DI levels are a bit higher but not high enough to support “damper” action by the proximal, the distal and proximal enhancers work cooperatively (Fig. 4I). Under wildtype conditions with normal DI levels present, the role of sna.prox enhancer changes to support dampening of sna levels. The observation that DI levels build in time has implications for a changing role of the sna.prox enhancer over time.

Discussion

Collectively, our results suggest that initially the proximal and distal enhancers cooperate under conditions of lower DI but then as DI levels rise, the proximal enhancer becomes more potent, gaining influence to the promoter over the distal enhancer and subsequently leading to dampening of levels (Fig. 4J-L). While sna is considered to be a high-threshold DI target gene, meaning the highest levels of DI are required to support its expression compared to other DI target genes (18), our results suggest that DI may exhibit a changing effect on sna expression due to its action at the sna.prox enhancer. It is possible that DI's changing effect over time
relates to its ability to differentially occupy binding sites of varying affinity within each enhancer (Fig. 4I). Earlier when lower levels of DI are present, high affinity sites associated within one or both enhancers are occupied (cooperative mode, Fig. 4I); subsequently, low affinity binding sites, which happen to be enriched at the proximal enhancer, are occupied later when peak DI levels are achieved. Occupancy of low affinity binding sites within the snaprox enhancer may allow this particular enhancer to compete better later and thus dampen gene expression at the later stage (competitive mode, Fig. 4I).

In this study, our data demonstrate that in addition to regulating the spatial domain of sna expression, the morphogen DI additionally acts to regulate levels of expression of this target gene. It is possible that other morphogens also change their perceived effect on gene expression over time, such as changing from promoting gene expression to dampening gene expression. We propose that morphogens commonly provide feedback to target genes, after specifying spatial domains of expression, to help keep target gene expression robust.

Furthermore, the sna gene may have acquired two regulatory modules that differentially respond to morphogen DI concentration as a mechanism to buffer genetic and environmental fluctuations in DI levels. This idea had been proposed previously as deletion of snaprox in the context of rescue transgenes also was associated with sensitivity to growth at high temperature (8, 11). Here we show that deletion of either enhancer supports viability, but deletion of the snaprox enhancer, specifically, resulted in increased variability in sna levels (Fig. 1F). Our results that DI input to snaprox supports dampening of gene expression, provides a molecular explanation for how sna expression can be kept in check despite variability in transcription factor inputs through action of the snaprox enhancer. Animals may have developed such safety mechanisms, such as multiple co-acting enhancers, to overcome the challenges posed by fluctuating morphogen gradients that can result from genetic and environmental perturbations (11, 19). Dual-input by morphogens may increase precision of target gene outputs.
Materials and methods

Fly stocks/husbandry and crosses

All flies were kept at 25°C, except for flies bearing the \textit{dl-BLID} allele that were kept at 18°C. \textit{yw} is used for wildtype, unless otherwise noted. To test \textit{sna} transcriptional activity in the \textit{dl}\textsuperscript{1} heterozygous background (#7096, Bloomington Drosophila Stock Center), \textit{dl}\textsuperscript{1}/CyO flies were crossed with \textit{MCP.GFP}, \textit{mCherry-PCP}, \textit{His2Av-eBFP2} to generate \textit{dl}\textsuperscript{1}/CyO; \textit{MCP.GFP}, \textit{mCherry-PCP}, \textit{His2Av-eBFP2}. Virgin flies with \textit{MCP.GFP}, \textit{mCherry-PCP}, \textit{His2Av-eBFP2} or \textit{MCP.mCherry} (from Hernan Garcia, UC Berkeley, US; 16) in wildtype, \textit{dl}\textsuperscript{1} heterozygous or \textit{dl-BLID} heterozygous backgrounds were crossed with males having both proximal and distal enhancers (WT, \textit{snaMS2.wt}), a proximal deletion (NoPrimary, \textit{snaMS2.Δprox}), or a distal deletion (NoShadow, \textit{snaMS2.Δdis}) (from Michael Levine, Princeton University, US; 14) for live imaging to detect \textit{sna} MS2 stem loops. The \textit{dl-BLID} line was generated in a previous study by integrating BLID domain at the C-terminus of DI at the endogenous locus using CRISPR/Cas9 (6).

\textbf{CRISPR-Cas9 Mediated Genome Modification}

To target deletions of various sizes to the distal part of the proximal enhancer [i.e. \textit{sna.Δprox4.4}, \textit{sna.Δprox3.0}, \textit{sna.Δprox3.0(mDI)}] or distal enhancer (i.e. \textit{sna.Δdis1.8}), a transgenic line was generated expressing two guide RNAs (gRNAs) targeting the flanking region described as the proximal enhancer or distal enhancer (8). First, the unique PAM recognition sites were identified flanking this region using the flyCRISPR optimal target finder (https://flycrispr.org/target-finder). To delete the full proximal enhancer, two flanking regions that are located outside of the identified proximal enhancer (8) were targeted due to limited
availability of PAM sequences, and subsequently led to the generation of a 4.4kB deletion. These two sites were cloned into the plasmid pCFD4-U6:1_U6:3tandemRNAs (Addgene plasmid #49411). The plasmid was injected into y2cho2v1; P {nos-phiC31\int.NLS}6X; attP2 (III) (NIG-Fly #TBX-0003), resulting in phiC31-mediated site-integration at landing site attP2 (Chr. III) (20). Integration in the genome at this position was confirmed by PCR/sequencing.

To generate the proximal deletion lines [ snaexual\prox3.0, sna.\ prox2.6 sna.\ prox3.0(mDI)] and the full distal deletion (sna.\ dis2.0), Homology-Directed Repair (HDR)/CRISPR-Cas9 was used (21). A donor construct was generated using pHD-DsRed vector (Addgene plasmid #51434). An -1kb 5’ or 3’ homology arm, either upstream or downstream of the region, was cloned with Smal/NheI or Ascl/XhoI, respectively.

y2cho2v1;CyO,P {nos-Cas9,y+,v +} 2A (NIG-Fly #Cas-0004) virgin flies were crossed with gRNA transgenic male flies. Embryos were collected and injected with 300 ng/µl of the donor vector. The deletion lines were screened by DsRed expression in adult fly eyes and sequencing. The DsRed marker was removed by crossing with a Cre expressing fly line (y[1] w[67c23] P{y[+mDint2]=Cre)y1b; D[8]/TM3, Sb[1], BDSC #851). Crispr/Cas9 generated deletions/mutations were confirmed by PCRing the region of interest and subsequently sequencing. The sequences after Cre-mediated DsRed marker excision are provided below; uppercase indicates genomic sequence, and lowercase indicates loxP remnant sequence.

>sna.\prox3.0
CGACAAAGGATGTGA
CTCAGcggcggagacatatgcacacctgcgtagtgcccccaactgggttaacctttgagt
tcctcagttggggcgtagataactctgtataatgtagcagctacgaagttatagaagcactag
ATGACCCACCCAGGTA
GGATG

>sna.\prox2.6
AGTCGACAAAGGGATGTGACTCAGgggccgacatatgcacacctgctgatgtagtggcccaacctggtaaccttt
tgagtctctcacggtgggctagataactctgtataatgtatgctataacgaggtatatagagacatagtTCTCAGCAAAAAT
TGACAAAGAACCAAA

> sna.Δdis2.0
TAAATTCACACATTTTGCTGcatacgacacctgctgataaactctgtataatgtatgctataacgaggtatatagaagagca
catgTAAATCCGTGTTAAATTGTT

To introduce mutations into the predicted DI binding sites of sna.Δprox3.0 (see Figure
2A), first all three DI consensus sequence were replaced with the complementary base (i.e. A>T
or G>C), and 1kb of the proximal enhancer sequence including all of the mutated DI consensus
sequence was synthesized and inserted into pUC57 (GenScript). This plasmid was used as a
DNA template to generate the homology right arm for HDR donor construct.

> sna.Δprox3.0(mDI) (uppercase: genomic sequence, lowercase: mutated DI consensus
sequences)
ATGACCCACCCAGGTAGGTGAGGACATAatctttttgggAGGCCAGTTTTTCCACTCGTGG
CTTGGTTTTGGCTTTACGCCGTGTA ATTGGATAAGATGGGAAATTACTTTAAATCC
TTCGCTGATCCACATCCGGACATTCCGTCAAGGAAGAAATCCATTTGCAGGAAATAACGAAATG
GAAATGCGGCGTGGTTATGGGCGTATTTCCCATTTCCCTCAACGCGATTTGGTGCAAG
ATCGCGGGGAaaccttaagggcGCTGGAATTTTTTGTCAACCTTGGGT GTTGATCACAAACTTTTG
GGTTTGCTATctaaaaaaggtATTTTACC

To generate sna.Δdis1.8 or sna.Δprox4.4, non-homologous end joining (NHEJ)/CRISPR-Cas9
was used. y2cho2v1;sp/Cyo;P {nos-Cas9,y+,y +} 2A (NIG-Fly #Cas-0004) virgin flies were
crossed with gRNA transgenic male flies. The mutant lines with these deletions were confirmed by PCR/sequencing.

> sna.Δdis1.8 (uppercase: genomic sequence, lowercase: sequence added after NHEJ)
TGAACCTGTTGTGAACTCAGtaccGTCTAATGGCCAGAACACCG

> sna.Δprox4.4
TCAAAATCAAAGGCAGGCCagagtgcaagaggttgacgcAGGAGCGCTGCTGTCTCTT
CC

In situ hybridization and imaging

Embryos were collected for 2hrs, aged at 25°C to give 2-4 hr old embryos, fixed and stained by in situ hybridization using standard protocols. Antisense sna RNA probes were labeled with digoxigenin. For fluorescent in situ hybridization, sheep anti-digoxigenin (Life Technology PA185378) was used with Alexa conjugate secondaries (1:400; Thermo Fisher). For alkaline phosphatase (AP) staining samples, a Zeiss Axioxam 506 microscope was used, and images were taken with a 20x objective.

RNA extraction from single embryos and qPCR

Embryos were collected for 1hr and aged for 2hrs at 25°C. They were hand dechorionated and immersed in halocarbon 27 oil to determine the stage of the embryos. When the embryos reached nc14c, in which 75% of cellularization has been completed, the individual embryo was collected in a 1.5ml microcentrifuge tube for RNA extraction using a standard Trizol protocol (ThermoFisher), followed by DNase I treatment (NEB), and cDNA synthesis using SuperScript III First-Strand Synthesis System kit (ThermoFisher). qPCR was performed on
cDNA using SYBR Green I Master Mix (Roche) on a StepOnePlus Real-Time PCR System (Applied Biosciences).

**Confocal blue laser illumination**

*dl-mCherry-BLID* or *dl-BLID* heterozygous embryos were collected for 1hr and aged for 4hr at 18°C. Embryos were hand dechorionated, and placed onto a heptane glued slide. The specimens were immered in water and staged using a bright field microscope. Once an embryo reached mid nc14 (i.e. nc14c) determined by the cellularization front having reached 50% completion (~40 min after entering nc14), the specimen was covered with a cover slip. To degrade DI, the embryo was illuminated with a Zeiss LSM 800 488nm confocal blue laser with 20% laser power using a 40x water immersion objective (see Fig. 4A).

**Live imaging**

To test that *sna* transcriptional activity changes in response to DI levels, the MS2-MCP system was used. In wildtype (high levels of DI) and *dl* heterozygote (*dl*/*+, intermediate level of DI) embryos, nuclei were marked by imaging H2A-BFP [i.e. His2Av.eBFP2; (22)] with 405nm at 1.5% laser power, and MCP.GFP associated with *sna* MS2 were detected with 488nm at 2.4% laser power. For *dl-mCherry-BLID* or *dl-BLID* heterozygotes, mCherry associated with DI-BLID or MCP.mCherry associated with *sna* MS2 constructs (16) were detected with 561nm at 5% laser power before, during, and after blue laser illumination (see Fig. 4A). All images taken contained 20 Z-planes separated by 2.0μm.

**MS2 Foci detection and Quantification**

MS2 foci were detected and quantified similar to previously described methods (6). Briefly, a gaussian filter was used to smooth the image and background was removed using
Top-hat filtering. A threshold was applied to detect MS2 foci. Foci consisting of three or less pixels were removed as often these were not associated with real signal, as confirmed manually, and appeared outside of nuclei or within nuclei already containing a clear focus. For each focus detected, the number of pixels were counted, and the mean intensity of each focus was determined. The number of foci, the mean area (or mean number of pixels per focus), and the average of the mean intensities was computed at every time point for each individual embryo and plotted. Traces were normalized to the first time point before blue laser illumination to account for variability. Error bars were calculated as the standard error of the mean. Codes are deposited at Github under accession number (TBD).

**DI-mCherry-BLID Quantification**

DI-mCherry-BLID was quantified by making a single z- and time-projection and then drawing a rectangular ROI on the image of the embryo. For each time point, the intensity of each pixel was averaged together for the entire ROI. The same sized ROI was used for all time points of every embryo measured. For each embryo, the ROI was positioned centrally to capture the most signal in the embryo. The size of the ROI was determined beforehand to insure it fit within all embryos measured. Traces were normalized to the first time point before blue laser illumination to account for variability in starting amounts of fluorescence, likely due to variability in DI concentration. Individual and unnormalized traces are shown in the supplement (Fig. S5). Codes are deposited at Github under accession number (TBD).

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**Author Contributions:** A.S. and J.I. conceived the project and planned the experimental approach. A.S. directed the project. J.I. performed all experiments, and J.M. performed all quantitative analysis of imaging data. Data were analyzed by J.I., J.M., and A.S. The manuscript was written by J.I. and A.S. with edits provided by J.M.

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Figures

Figure 1. Embryos with various deletions of sna distal or proximal enhancers fail to support endogenous sna expression levels. (A) Schematic of enhancer deletions, proximal (prox, yellow) or distal (dis, green), by CRISPR-Cas9 system. Two regions in the proximal enhancer identified to support incomplete or full sna expression, A1.6 or A2.0, respectively (9). (B-E,G) sna expression in Drosophila nc14d embryos by in situ hybridization using sna riboprobes. Lateral views of embryos are shown, with anterior to the left and dorsal side up. (F) Normalized sna transcript levels in embryos at nc14c for indicated deletion mutations relative to wildtype determined using qPCR. Sample sizes are indicated next to the genotype. Asterisk: p<0.01 by Student’s t-test. For this and all subsequent figures, box plots show median (center line) and 25th or 75th percentiles (box limits). Whiskers represent 1.5 times the interquartile range from the 25th and 75th percentiles.
Figure 2. Mutating DI binding sites in sna.\textit{Δprox3.0} increases sna expression level in a distal enhancer-dependent manner. (A) Schematic of deletions in the proximal enhancer with/without the distal enhancer deletion. Dark pink bar: putative high-affinity DI binding sites in the proximal enhancer identified by JASPAR (http://jaspar.genereg.net). Black bar: mutated putative high-affinity DI binding sites. (B,D,E) sna expression in nc14d embryos by \textit{in situ} hybridization using sna riboprobes. Lateral views of embryos are shown, with anterior to the left and dorsal side up. (C) Normalized sna transcript levels in embryos at nc14c for indicated deletion mutations relative to wildtype determined using qPCR. Sample sizes are indicated next to the genotype. Asterisk: p<0.01 by Student's t-test.
Figure 3. At high DI levels, the proximal enhancer limits the distal enhancer activity, consequently decreasing sna expression levels. (A) Schematic of reporter constructs used to assay sna transcriptional activity by monitoring MS2-MCP.GFP (14). (B) MCP.GFP signal associated with snaMS2.wt (green dots) driven by both enhancers (“prox+dis acting”) in wildtype
(left column) or $dl^{+/}$ (right column) embryos at time point 4 min (t=4) or 44 min (t=44) after entering nc14. Nuclei are visualized by H2A-BFP (blue). Scale bar: 10μm. (C) Quantitative analysis of mean number of pixels per MCP.GFP spot associated with $snaMS2.wt$ in wildtype (yellow, high DI levels) and $dl^{+/}$ (red, intermediate DI levels) embryos. (D,E) Quantitative analysis of mean number of pixels per MCP.GFP spot associated with $snaMS2.Δprox$ (green) in wildtype (D) or $dl^{+/}$ heterozygotes (E). The mean number of pixels per MCP.GFP spot associated with $snaMS2.wt$ (purple) in wildtype or $dl^{+/}$ heterozygotes used in C are replotted for direct comparison to mean number of pixels per MCP.GFP spot associated with $snaMS2.Δprox$. Error bars represent standard error of the mean. For each condition, movies were obtained for three individual embryos and the data averaged. For the individual traces, see Figure S4.
Figure 4. At low DI levels, the distal enhancer activity is dependent on the proximal enhancer to support endogenous sna expression levels. (A) Schematic of sna
MS2-MCP.mCherry system combined with the DI-BLID degradation system at late nc14. MCP.mCherry signal associated with sna MS2 (red dots) driven by both enhancers (snaMS2.wt) in wildtype embryos are shown. Scale bar: 10μm. (B) Quantitative analysis of normalized mean intensity of mCherry fused to DI-BLID proteins [dl-mCh-BLID]; (6). dl-mCh-BLID embryos were kept in dark or illuminated by a confocal blue laser for 3 or 5 min. Mean intensity of mCherry associated with DI-BLID is measured in each time frame, and the values are normalized to the initial value detected in the first frame. For the individual traces, see Figure S5. (C,D) Quantitative analysis of normalized mean number of pixels per MCP.mCherry spot associated with snaMS2.wt in wildtype (C) or dl-BLID/+ embryos (D). (E) Quantitative analysis of normalized mean number of pixels per MCP.mCherry spot associated with snaMS2.Δprox in dl-BLID/+ embryos. Embryos are kept in dark (red) or illuminated with a confocal blue laser for 3 min (blue) or 5 min (yellow). For the individual traces, see Figure S6. (F-H) Normalized mean number of pixels per MCP.mCherry spot associated with snaMS2.wt (purple) or snaMS2.Δprox (green) in dl-BLID/+ embryos kept in dark (F) or with 3 min (G) or 5 min (H) blue laser illumination are replotted with the date used in D and E. Mean spot size of MCP.mCherry associated with sna MS2 are measured in each time frame, and the values are normalized to the initial value detected in the first frame. Error bars represent standard error of the mean. For each condition, movies were obtained for three individual embryos and the data averaged. (I) Schematic showing high (dark pink) or low affinity (light pink) DI binding sites predicted by JASPAR (http://jaspar.genereg.net) at sna locus (top). A conceptual representation of DI (bottom left) from nc11 to nc14 (7). The low threshold (T1: threshold 1) supports cooperative action of enhancers, and the high threshold (T2: threshold 2) supports enhancer competition. Model showing two modes of coordination between the proximal and distal enhancers, dependent on the concentration of DI, to support the correct level of sna expression (bottom right).