

# Concentrating pre-mRNA processing factors in the histone locus body facilitates efficient histone mRNA biogenesis

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The histone locus body (HLB) assembles at replication-dependent histone genes and concentrates factors required for histone messenger RNA (mRNA) biosynthesis. FLASH (Flice-associated huge protein) and U7 small nuclear RNP (snRNP) are HLB components that participate in 3' processing of the nonpolyadenylated histone mRNAs by recruiting the endonuclease CPSF-73 to histone pre-mRNA. Using transgenes to complement a *FLASH* mutant, we show that distinct domains of FLASH involved in U7 snRNP binding, histone pre-mRNA cleavage, and HLB localization are all required for proper FLASH function in vivo. By genetically manipulating HLB composition using mutations in FLASH, mutations in the HLB assembly factor *Mxc*, or depletion of the variant histone H2aV, we find that failure to concentrate FLASH and/or U7 snRNP in the HLB impairs histone pre-mRNA processing. This failure results in accumulation of small amounts of polyadenylated histone mRNA and nascent read-through transcripts at the histone locus. Thus, the HLB concentrates FLASH and U7 snRNP, promoting efficient histone mRNA biosynthesis and coupling 3' end processing with transcription termination.

## Introduction

Structures termed nuclear bodies (NBs) create microenvironments within the nucleus that promote the organized regulation of genome function. Defined microscopically as a concentration and colocalization of a set of specific RNPs and/or proteins, each NB contains a suite of factors associated with particular nuclear processes including RNA biosynthesis, DNA replication, and DNA repair (Gall, 2000; Matera et al., 2009; Nizami et al., 2010; Morimoto and Boerkoel, 2013). NBs assemble through protein-nucleic acid and protein-protein interactions. For example, the nucleolus is seeded by rRNA (Falahati et al., 2016) paraspeckles are seeded by Men  $\epsilon/\beta$  long noncoding RNAs (Mao et al., 2011a), and the protein Coilin organizes the Cajal body (CB; Mao et al., 2011b). NBs are not membrane bound, and their components can freely exchange with the nucleoplasm (Dundr et al., 2004). By concentrating reactants and substrates, NBs are postulated to provide discrete microenvironments that increase rates of nuclear processes (Matera et al., 2009; Dundr, 2011; Mao et al., 2011b). Although there is some evidence for this hypothesis (Klingauf et al., 2006; Stanek et

al., 2008; Strzelecka et al., 2010; Novotný et al., 2011), disruption of NBs does not always obviously impact associated reactions (Deryusheva and Gall, 2009; Liu et al., 2009). Thus, determining the specific contribution that a NB makes to an in vivo reaction has proven challenging. We have been addressing this question by studying the histone locus body (HLB).

Joe Gall et al. identified the HLB in *Drosophila melanogaster* in 2006 (Liu et al., 2006). It then became apparent that an NB initially described as a specialized CB in which U7 small nuclear RNP (snRNP; Frey and Matera, 1995) and the protein NPAT (Ma et al., 2000; Zhao et al., 2000) are concentrated was the mammalian HLB. The HLB is involved in replication-dependent (RD) histone mRNA synthesis (Ye et al., 2003; White et al., 2011), which occurs at high levels only during S phase of the cell cycle to package newly replicated DNA (Marzluff et al., 2008). RD histone genes encode the only known eukaryotic mRNAs that are not polyadenylated, ending instead in a stem loop at the 3' end (Marzluff et al., 2008). Two factors required for formation of the 3' end of histone mRNAs, FLASH and U7 snRNP, are concentrated in and continuously present in the HLB (Yang et al., 2009). In addition to FLASH

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Abbreviations used in this paper: CB, Cajal body; FL, full length; FLASH, Flice-associated huge protein; HCC, histone cleavage complex; HDE, histone downstream element; *mxc*, multi sex combs; NBs, nuclear bodies; pol II, polymerase II; RD, replication dependent; Slbp, stem-loop binding protein; snRNP, small nuclear RNP; WT, wild type.

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and U7 snRNP, formation of the 3' end of histone mRNAs requires a complex of polyadenylation factors termed the histone cleavage complex (HCC; Yang et al., 2013). Association of the HCC with U7 snRNP requires a direct interaction between FLASH and the U7 snRNP component Lsm11 (Sabath et al., 2013; Yang et al., 2013).

The HLB is present throughout interphase of the cell cycle (Ma et al., 2000; Zhao et al., 2000; Liu et al., 2006; White et al., 2007, 2011; Ghule et al., 2008; Tatomer et al., 2014), and expression of the histone genes is activated by cyclin E/Cdk2, which phosphorylates NPAT (Mxc in *Drosophila*; Ma et al., 2000; Zhao et al., 2000; Ye et al., 2003; White et al., 2007, 2011). A specific DNA element in the *Drosophila* histone gene cluster nucleates the HLB, suggesting direct association between the HLB and chromatin at the histone locus (Salzler et al., 2013). The HLB begins to assemble in the *Drosophila* embryo before the onset of zygotic histone mRNA expression (White et al., 2011; Salzler et al., 2013). The recruitment of FLASH and U7snRNP to the HLB does not require the cis elements in the histone pre-mRNA with which they associate (Salzler et al., 2013). These observations indicate that the HLB is not merely a reflection of high, localized gene expression and instead suggest that the HLB has evolved as a feature of the metazoan nucleus that optimizes cell cycle-regulated biosynthesis of histone mRNAs.

An ideal system to understand HLB function requires the ability to manipulate the localization and activity of HLB components without destroying the HLB, as well as the ability to directly assay the molecular and biological effects of such manipulations. Here, we describe such a system for the *Drosophila* HLB. By using both a transgenic complementation system and genetic alteration of HLB composition, we find that when FLASH and/or U7 snRNP are present at normal levels in the nucleus, but not concentrated in the HLB, histone pre-mRNA processing is slowed. This results in transcriptional read-through and the accumulation of unprocessed histone transcripts at the histone locus, as well as small amounts of misprocessed, polyadenylated mRNA. These data indicate that concentrating factors in the HLB increases the rate of histone pre-mRNA processing coupling 3' end formation with transcription termination.

## Results

### FLASH is essential for histone mRNA 3' end formation in *Drosophila*

FLASH plays a critical biochemical role in histone pre-mRNA processing as a bridge between the U7 snRNP and the HCC (Sabath et al., 2013; Yang et al., 2013; Fig. 1 A). To determine the contribution that concentrating FLASH in the HLB makes to RD histone pre-mRNA processing, we first established a transgenic complementation system to identify the protein domains necessary for FLASH function in vivo (Fig. 1 B). A piggyback transposon insertion in the 5' UTR of the *FLASH* gene (Figs. S1 A and 1 B) results in a large reduction of FLASH protein (Fig. 1 C; and see also Fig. 5). This hypomorphic allele (*FLASH<sup>PBac</sup>*) in trans to a deficiency of *FLASH* (*FLASH<sup>Df</sup>*) reduces viability (20% of expected), and the adult flies that eclose are female sterile but male fertile (Fig. 1 D). Expression of C-terminally V5-tagged full-length (FL) FLASH from a single-copy transgene using the endogenous *FLASH* promoter completely rescued the reduced viability and sterility caused by *FLASH<sup>PBac/Df</sup>*

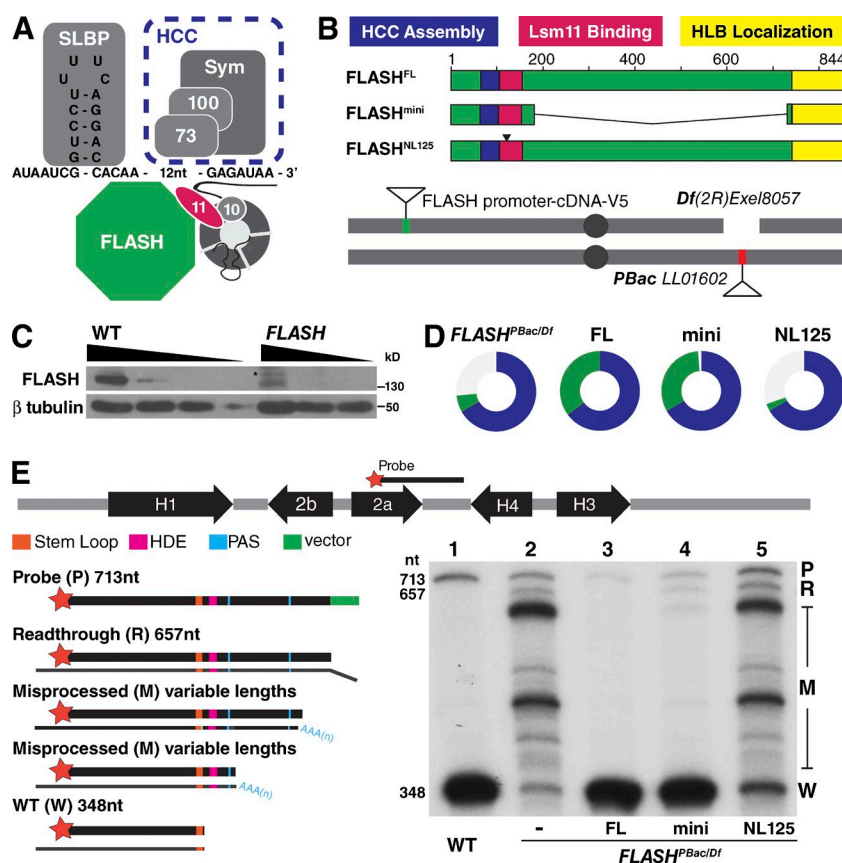
(Fig. 1 D and Table 1). This system provides a means for testing the ability of transgenes expressing different alleles of FLASH to rescue FLASH mutant phenotypes.

We determined whether the developmental phenotypes of the *Drosophila FLASH<sup>PBac/Df</sup>* mutant were caused by histone pre-mRNA processing defects or loss of another FLASH function, as FLASH was originally implicated in Fas-mediated apoptosis in mammalian cells (Imai et al., 1999). The 844-aa FLASH protein has three known biochemical functions (Fig. 1 B). The N-terminal 154 aa of FLASH is essential for histone pre-mRNA processing in vitro, and aa 105–150 in this region bind the N terminus of the U7 snRNP protein Lsm11 (Burch et al., 2011). The Lsm11–FLASH N-terminal interaction creates an interface that recruits the HCC (Sabath et al., 2013; Yang et al., 2013), which contains the endonuclease CPSF-73 (Dominski et al., 2005). The C-terminal 111 aa of FLASH are necessary for localization to the HLB (Burch et al., 2011). To test if these three activities are sufficient for FLASH function in vivo, we expressed a transgene encoding a 547-aa internal deletion of the 844-aa FLASH protein, fusing the HCC assembly and Lsm11 binding domains to the HLB localization domain (FLASH<sup>mini</sup>; Fig. 1 B). This transgene rescued the reduced viability and sterility of the *FLASH<sup>PBac/Df</sup>* mutant (Fig. 1 D and Table 1), suggesting that the essential function of FLASH in *Drosophila* is histone mRNA 3' end formation as a component of the HLB (see Fig. 4).

To understand the molecular basis for the FLASH mutant phenotype, we examined the histone mRNA species produced in various *FLASH* mutant genotypes (Fig. 1 E). When histone pre-mRNA processing is compromised in *Drosophila*, cryptic polyadenylation signals located downstream of the normal processing site are used, resulting in the cytoplasmic accumulation of longer, polyadenylated histone mRNA (Sullivan et al., 2001; Lanzotti et al., 2002; Godfrey et al., 2006). We also detect longer nascent transcripts at the histone locus by in situ hybridization (Lanzotti et al., 2002). To detect misprocessed histone RNA species, we use an S1 nuclease protection assay that provides a quantitative measurement of the amounts of properly processed histone mRNA (Fig. 1 E, “W”) compared with misprocessed, polyadenylated histone mRNA (Fig. 1 E, “M”). The assay also detects “read-through” transcripts that represent the unprocessed nascent pre-mRNA (Fig. 1 E, “R”; Sullivan et al., 2009).

*FLASH<sup>PBac/Df</sup>* mutant third-instar larvae express large amounts of misprocessed, polyadenylated histone mRNA (Fig. 1 E, lane 2; and Fig. S1 C), confirming that a major function of FLASH is histone mRNA 3' end formation. We additionally detected small amounts of properly processed mRNA, consistent with this being a hypomorphic allele, as well as small amounts of unprocessed read-through RNA (Fig. 1 E, lane 2). Expression of FLASH<sup>FL</sup> protein completely rescued this mutant RNA phenotype (Fig. 1 E, lane 3). Expression of FLASH<sup>mini</sup> also rescued the mutant RNA phenotype, although small amounts of both misprocessed and read through histone mRNA were detected in FLASH<sup>mini</sup> larvae (Fig. 1 E, lane 4). FLASH<sup>mini</sup> is almost fully proficient in processing in vivo, consistent with it rescuing viability and fertility of the *FLASH<sup>PBac/Df</sup>* mutant (Fig. 1 D). The developmental phenotypes observed with *FLASH<sup>PBac/Df</sup>* are consistent with our previous in vivo analyses of processing factors stem-loop binding protein (Slbp) and U7 snRNP (Sullivan et al., 2001; Godfrey et al., 2009), supporting the idea that the primary role for FLASH is histone mRNA 3' end formation.

To directly ask if histone mRNA processing is an essential FLASH function, we changed to alanines the asparagine-



**Figure 1. Histone pre-mRNA processing is essential in vivo FLASH function.** (A) The histone pre-mRNA processing machinery. Slbp binds the stem loop on the pre-mRNA. The U7 snRNP interacts with the HDE in histone pre-mRNA. FLASH binds the U7 snRNP protein Lsm11 (11), and the FLASH/U7 snRNP complex recruits the HCC containing the Symplekin scaffold (Sym) and the CPSF100/CPSF73 endonuclease (100/73). 10, Lsm10. (B) Schematic of functional domains of FLASH and the transgene rescue system. Transgenes were inserted at chromosomal location 25C6 and recombined with a deletion of the FLASH locus (*Df(2R)Exel8057*). All *FLASH<sup>PBac/Df</sup>* experimental flies contained a maternal *Df* chromosome in trans to a paternal *PBac LL01602* chromosome (see Fig. S1). NL125 is a 2-aa mutation in the Lsm11 binding region (see Fig. 2 A). (C) Anti-FLASH Western analysis of WT and *FLASH<sup>PBac/Df</sup>* wandering third-instar larvae using threefold serial dilutions. Asterisk, cross-reacting protein. (D) Visual representation of transgenic rescue of *FLASH<sup>PBac/Df</sup>*. Circles indicate the proportion of *FLASH<sup>PBac/Df</sup>* mutant with the indicated transgene (green) and control heterozygous sibling (blue) adult flies obtained. The expected fraction of control siblings is two thirds. Thus, one third green indicates full viability, and the absence of green indicates lethality. Table 1 shows the data for this analysis. (E) S1 nuclease protection assay. The map of a *Drosophila* histone gene repeat is shown. Equal amounts (5 µg) of total RNA from wandering third-instar larvae of WT, the *FLASH<sup>PBac/Df</sup>* mutant, or the mutant complemented with the indicated transgenes were analyzed using a 3'-labeled DNA probe complementary to *H2a* mRNA that extends to the *H4* HDE followed by 56 nt vector sequence (green). The hybrids were incubated with S1 nuclease, and the protected fragments were resolved on a 6% acrylamide-7M urea gel and visualized by autoradiography. Properly processed *H2a* mRNA protects a 348-nt fragment from S1 digestion (W). The *H2a* gene contains multiple downstream cryptic polyadenylation signals (PAS; blue) that protect a heterogeneous set of longer fragments (M). Transcripts that extend beyond the complementary region result in a single protected read-through fragment (R) that is distinct from undigested probe (P). The sizes (in nucleotides) of the processed mRNA, R RNA, and undigested probe are indicated.

leucine residues 125–126 in FLASH that are required for in vitro binding of the N terminus of FLASH with Lsm11 (Burch et al., 2011; Figs. 1 B and 2 A). *FLASH<sup>NL125</sup>* did not rescue *FLASH<sup>PBac/Df</sup>* reduced viability (Fig. 1 D) or the misprocessed histone mRNA phenotype (Fig. 1 E, lane 5). *FLASH<sup>NL125</sup>* protein and U7 snRNP were present in the HLB (see Fig. 4 B), indicating there is an intact HLB in this mutant. We previously showed that a 2-aa mutation in Lsm11 that prevents FLASH binding also results in misprocessed histone mRNA and lethality (Burch et al., 2011). Thus, the interaction between FLASH and Lsm11 is essential for proper histone mRNA 3' end formation and normal development, but not for localization of either component to the HLB.

#### The FLASH N terminus is necessary for histone mRNA 3' end formation in vivo

We have defined a domain N terminal of the Lsm11 binding domain that is essential for histone pre-mRNA processing in vitro (Sabath et al., 2013; Yang et al., 2013). In both mammalian and *Drosophila* nuclear extracts, this domain is necessary for recruitment of the HCC to the FLASH–Lsm11 complex. Deletion of residues 1–77 from *Drosophila* FLASH or mutation to

AAAA of a highly conserved LDIY motif (aa 71–74; Fig. 2 A) inhibited processing in vitro and blocked recruitment of the HCC in nuclear extracts (Sabath et al., 2013).

To determine the in vivo requirement for this region of FLASH, we tested the ability of an N-terminal truncation of FLASH containing aa 71–74 (*FLASH<sup>65–844</sup>*), which functions normally in vitro, as well as *FLASH<sup>78–844</sup>* and *FLASH<sup>LDIY71</sup>* (with aa 71–74 mutated to AAAA), which are defective in processing in vitro, to rescue *FLASH<sup>PBac/Df</sup>* mutant phenotypes (Fig. 2, B–D; and Fig. S2). Expression of *FLASH<sup>65–844</sup>* fully rescued *FLASH<sup>PBac/Df</sup>*-reduced viability and sterility, whereas the mutant lacking 13 additional aa (*FLASH<sup>78–844</sup>*) did not (Figs. 1 D and 2 C and Table 1). Surprisingly *FLASH<sup>LDIY71</sup>*, which has reduced processing activity in vitro (Fig. 2 F, lanes 8 and 9; Sabath et al., 2013), rescued *FLASH<sup>PBac/Df</sup>*-reduced viability and sterility (Fig. 2 C). Consistent with this finding, *FLASH<sup>LDIY71</sup>* mutants contained large amounts of properly processed histone mRNA with a normal 3' end, although we could detect small amounts of misprocessed and read-through histone mRNA species (Fig. 2 D, lane 6; and Fig. S2). We conclude that the *FLASH<sup>LDIY71</sup>* protein supports histone pre-mRNA processing in vivo, in contrast to its low activity in vitro.



There is a second LDIY motif at aa 45–48 in *Drosophila* FLASH that is also conserved in vertebrates (Fig. 2 A). This motif is not essential, because the FLASH<sup>65–844</sup> transgene completely rescues viability and histone pre-mRNA processing. To test whether it might play any role in processing, we mutated these four amino acids to alanine and generated transgenes expressing FLASH proteins with this mutation (FLASH<sup>LDIY45</sup>) or with mutation of both LDIY motifs (FLASH<sup>LDIY45,71</sup>). The FLASH<sup>LDIY45</sup> protein rescued FLASH<sup>PBac/Df</sup> reduced viability and sterility and completely restored histone pre-mRNA processing (Fig. 2, D [lane 7] and E). In contrast, FLASH<sup>LDIY45,71</sup> failed to rescue FLASH<sup>PBac/Df</sup>-reduced viability (Fig. 2 E) and had a more severe histone mRNA processing defect than the FLASH<sup>LDIY71</sup> mutant (Fig. 2 D, lanes 6–8). In addition, fewer FLASH<sup>LDIY45,71</sup> and FLASH<sup>78–844</sup> flies eclosed than FLASH<sup>PBac/Df</sup> flies (Fig. 1 D; Fig. 2, D and E; and Table 1). These mutant FLASH proteins can bind Lsm11 normally, suggesting a dominant-negative effect in the presence of the small amount of wild-type (WT) FLASH present in FLASH<sup>PBac/Df</sup> animals.

We compared the biochemical activity in processing of the FLASH<sup>LDIY45</sup>, FLASH<sup>LDIY45,71</sup>, and FLASH<sup>LDIY71</sup> proteins using an in vitro processing assay (Fig. 2 F, lanes 1–3). Extracts from S2 cells depleted of FLASH by double-stranded RNA are deficient in processing synthetic histone pre-mRNA substrates (Fig. 2 F, lanes 4 and 5), and this deficiency can be biochemically complemented by addition of *Escherichia coli*-derived proteins containing the N-terminal 178 residues of FLASH (Fig. 2 F, lanes 6 and 7; Sabath et al., 2013). In this assay, the FLASH<sup>LDIY45</sup> N-terminal fragment was active (Fig. 2 F, lanes 12 and 13), FLASH<sup>LDIY71</sup> had reduced activity relative to the WT protein (Fig. 2 F, compare lanes 6 and 7 to 8 and 9), and FLASH<sup>LDIY45,71</sup> was inactive (Fig. 2 F, lanes 10 and 11). Together, these genetic

and biochemical data indicate that both the LDIY71 motif and the LDIY45 motif contribute to processing, with the LDIY45 being critical only when the LDIY71 is mutated.

### Concentrating FLASH in the HLB facilitates histone mRNA 3' end formation

The FLASH<sup>LDIY71</sup> mutant functioned in vivo better than expected from its reduced activity in vitro, suggesting that some aspect of the FLASH<sup>LDIY71</sup> processing defect is compensated for in an intact cell. We tested whether concentrating FLASH<sup>LDIY71</sup> in the HLB contributed to histone 3' end formation in vivo. FLASH<sup>1–733</sup> (Fig. 3 A) lacks the C-terminal 111 aa and does not concentrate in the HLB in cultured S2 cells (Burch et al., 2011) or transgenic animals (Figs. 3 D and S3), although it rescued FLASH<sup>PBac/Df</sup>-reduced viability and sterility (Fig. 3 B and Table 1). FLASH<sup>1–733</sup> accumulates to similar levels as FLASH<sup>FL</sup> as assayed by Western blotting (Figs. 3 C and S4), but the low levels of FLASH throughout the nucleoplasm prevent its detection by immunofluorescence. We detected both misprocessed and read through transcripts in FLASH<sup>1–733</sup> animals, suggesting that concentrating FLASH in the HLB may be required for full FLASH function in histone pre-mRNA processing in vivo (Fig. 3 E, lane 4; and Fig. S4). The read-through transcripts in the FLASH<sup>1–733</sup> mutant were present in similar amounts as the misprocessed RNAs (Fig. 3 E, compare R to M in lanes 2 and 4). The histone mRNA species in FLASH<sup>1–733</sup> animals are qualitatively like those in FLASH<sup>LDIY71</sup> animals, which also had similar amounts of read-through transcripts and misprocessed RNAs (Fig. 3 E, lanes 4 and 5). In contrast, when FLASH is severely depleted (e.g., FLASH<sup>PBac/Df</sup>) or inactive in processing (e.g., FLASH<sup>LDIY45,71</sup> and FLASH<sup>NL125</sup>), the misprocessed mRNAs are predominant, as they also are in Slbp and U7

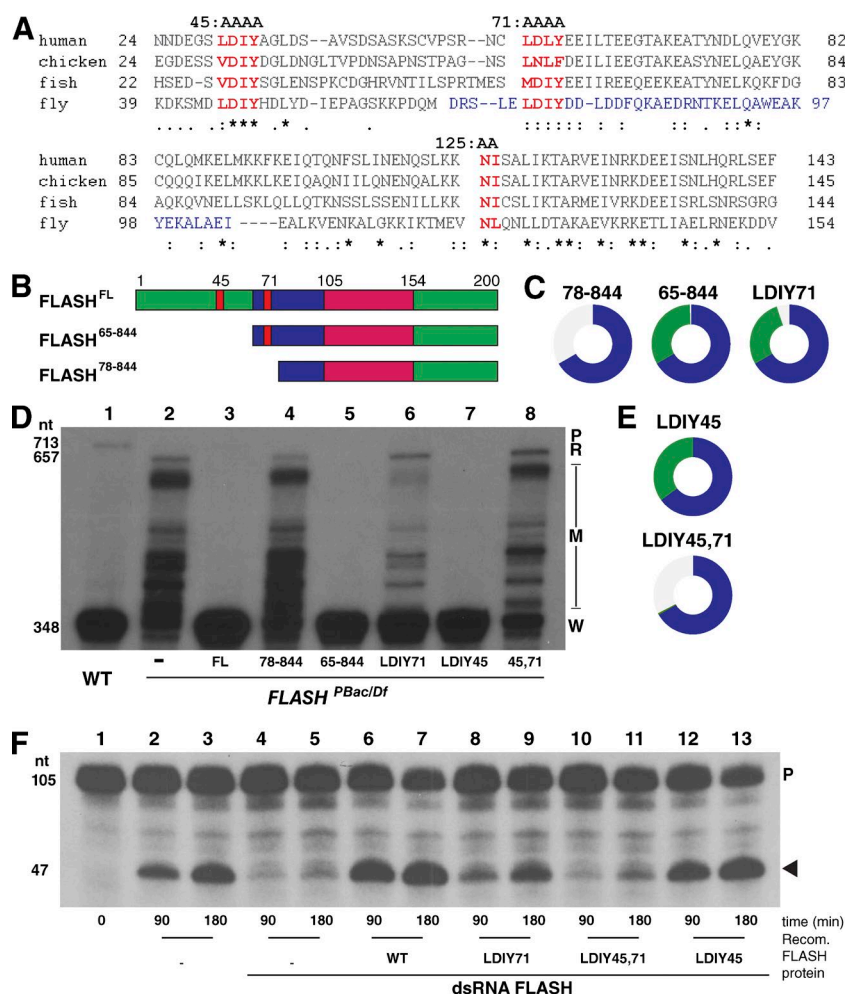
Table 1. Phenotypes of each of the transgenic FLASH constructs in the FLASH<sup>PBac/Df</sup> mutant background

Allele	Flies counted			Mendelian? ( $\chi^2$ P = 0.05)	Student's <i>t</i> test Significantly different from <i>Df<sup>mat</sup>/pBac<sup>pat</sup></i> mutant?	Fertility		HLB
	Control sibling	Experimental	Total			Female	Male	
Df(maternal) × pBac(paternal)	610	62	672	N		N	Y	
pBac(maternal) × Df(paternal)	509	213	722	N	Y, P < 0.001 <sup>a</sup>	N	Y	
pBac × pBac	450	84	534	N	Y, P < 0.05	N	Y	
FL	606	329	935	Y	Y, P < 0.001 <sup>a</sup>	Y	Y	Y
Mini	443	213	656	Y	Y, P < 0.001 <sup>a</sup>	Y	Y	Y
NL 125	607	27	634	N	Y, P < 0.05 <sup>b</sup>	N	n/d	Y
78–844	608	0	608	N	Y, P < 0.001 <sup>b</sup>	n/a	n/a	Y
65–844	596	294	890	Y	Y, P < 0.001 <sup>a</sup>	Y	Y	Y
LDIY 71	387	164	551	Y	Y, P < 0.001 <sup>a</sup>	Y	Y	Y
LDIY 45	381	206	587	Y	Y, P < 0.001 <sup>a</sup>	Y	Y	Y
LDIY 45,71	621	7	628	N	Y, P < 0.001 <sup>b</sup>	N	n/d	Y
1–733	338	189	527	Y	Y, P < 0.001 <sup>a</sup>	Y	Y	N
LDIY 71, 1–733	589	255	844	Y	Y, P < 0.001 <sup>a</sup>	N	Y	N
1–733, 101	467	221	688	Y	Y, P < 0.001 <sup>a</sup>	N	n/d	Y
LDIY71, 1–733, 101	479	195	674	Y	Y, P < 0.001 <sup>a</sup>	N	n/d	Y

The data for each of the genetic rescue experiments presented as circle charts in Figs. 1, 2, and 3 is shown. For the transgenic rescue crosses, a recombinant chromosome containing FLASH<sup>Df</sup> and a FLASH transgene was provided maternally. For each experiment, a  $\chi^2$  analysis was performed under the null hypothesis that the phenotypic classes of adult flies were present in expected Mendelian ratios (two thirds CyO/mutant control siblings and one third homozygous mutant genotype). A Student's *t* test was used to determine whether the number of flies in the experiment class was statistically different from when no transgene was present in the FLASH<sup>PBac/Df</sup> background. Note that three of the transgenes, FLASH<sup>NL125</sup>, FLASH<sup>78–844</sup>, and FLASH<sup>LDIY45,71</sup>, behaved as dominant negatives, consistent with them being incorporated into the HLB and competing with the small amount of endogenous FLASH. The FLASH<sup>PBac/PBac</sup> phenotype suggests the presence of a second site mutation on the PBac chromosome. Assessment of fertility for each genotype is also noted. The last column indicates transgenic FLASH proteins capable of localizing to the HLB. N, no; n/a, not applicable; n/d, not determined; Y, yes.

<sup>a</sup>A difference caused by transgene rescue of partial lethality.

<sup>b</sup>A difference caused by a more severe mutant phenotype resulting from transgene expression.



**Figure 2. Function of FLASH N-terminal mutants in histone pre-mRNA processing.** (A) Alignment of FLASH N-terminal sequences indicating the region required for HCC binding (blue), the conserved LDIY motifs (red; mutated to AAAA in FLASH<sup>LDIY45</sup> and FLASH<sup>LDIY71</sup>), and the NL residues (red) of the Lsm11 binding region mutated to AA in FLASH<sup>NL125</sup>. (B) Diagram of the first 200 aa of the N-terminal FLASH deletion mutants. (C) Visualization of rescue of FLASH<sup>Pac/Df</sup> with transgenic FLASH N-terminal mutants as in Fig. 1 D. (D) Equal amounts total RNA from WT and mutant larvae complemented with the indicated FLASH transgenes were analyzed by S1 nuclease protection as in Fig. 1 E. A PhosphorImager analysis of the same gel is in Fig. S1. (E) Visualization of rescue of FLASH<sup>Pac/Df</sup> with transgenic FLASH LDIY mutants. (F) Biochemical activity of FLASH LDIY mutant proteins was assessed using an in vitro histone pre-mRNA processing assay (Sabath et al., 2013). FLASH-depleted nuclear extracts were complemented with 100 ng of the indicated recombinant (Recomb.) N-terminal FLASH-GST fusion protein (aa 1–178). <sup>32</sup>P-labeled synthetic H3 pre-mRNA was incubated with the extract for 90 or 180 min, and the resulting undigested probe (P) and cleaved probe (arrowhead) were resolved on an 8% polyacrylamide-7M urea gel. See also Fig. S1.

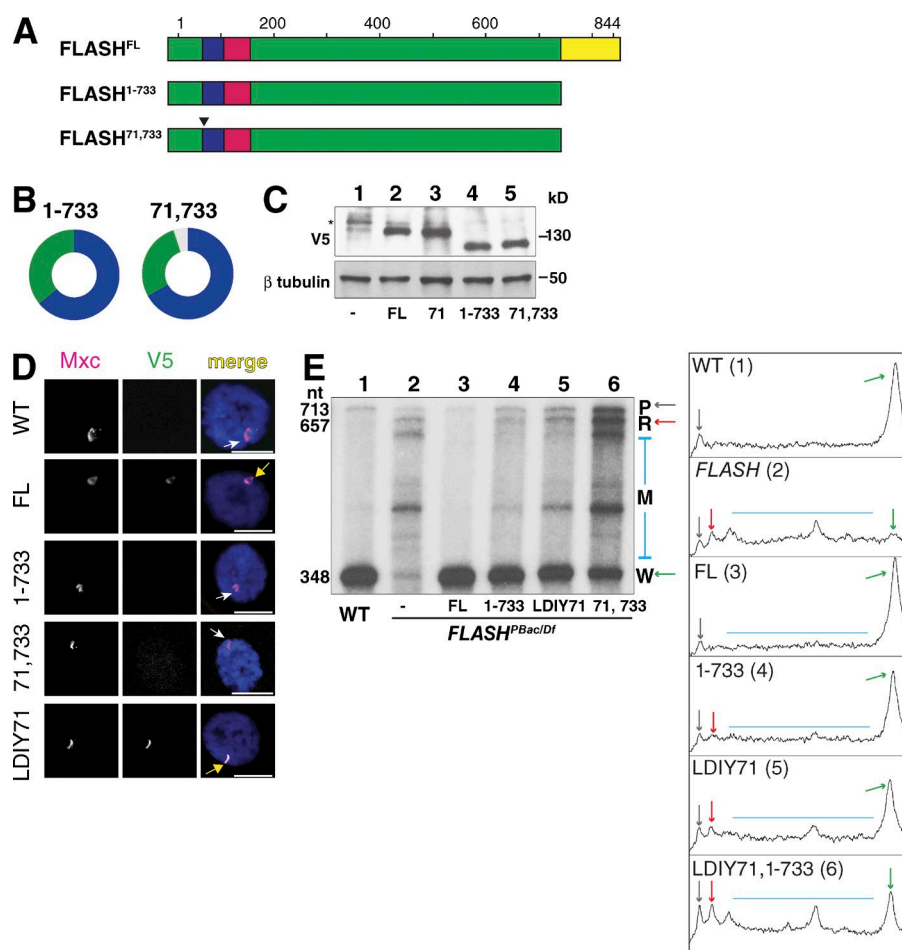
mutants (Lanzotti et al., 2002; Godfrey et al., 2006). Thus, these two FLASH mutants (FLASH<sup>I-733</sup> and FLASH<sup>LDIY71</sup>) with very different biochemical defects resulted in a similar alteration in histone mRNA biosynthesis in vivo.

These results suggest that both concentrating FLASH in the HLB and FLASH's intrinsic processing activity may independently contribute to efficient pre-mRNA processing in vivo. If this is true, mislocalizing the partially functional FLASH<sup>LDIY71</sup> protein by deleting the C-terminal 111 aa (Fig. 3 A, FLASH<sup>71,1-733</sup>) should exacerbate its mutant phenotype. FLASH<sup>71,1-733</sup> did not concentrate in the HLB (Fig. 3 D) but accumulated to similar levels as FLASH<sup>FL</sup>, FLASH<sup>LDIY71</sup>, and FLASH<sup>I-733</sup> (Fig. 3 C). Although FLASH<sup>71,1-733</sup> rescued FLASH<sup>Pac/Df</sup>-reduced viability (Fig. 3 B), it did not restore fertility (Table 1). Consistent with this more severe developmental phenotype, the histone pre-mRNA processing defect is exacerbated in FLASH<sup>LDIY71,1-733</sup> relative to either single mutant (Fig. 3 E, compare lanes 4 and 5 with lane 6). Thus, the FLASH<sup>LDIY71</sup> protein is more effective in promoting histone pre-mRNA processing when it is concentrated in the HLB. The synergistic effect of disrupting both processing activity and localization is consistent with independent contributions from each domain in 3' end formation in vivo.

A second possibility to explain the FLASH<sup>I-733</sup> RNA phenotype is that a region in the C terminus of FLASH also contributes to processing, independent of its role in localizing FLASH to the HLB. This does not seem likely, because aa 1–178 of

FLASH are fully active in vitro, and we have not detected any interaction of the C terminus of *Drosophila* FLASH with Lsm11 using in vitro binding assays (unpublished data). However, to address this possibility, we examined a situation in which FL FLASH is mislocalized by using a mutant allele of the *multi sex combs* (*mxc*) gene. *mxc* encodes a 1,837-aa protein orthologous to human NPAT that likely acts as a scaffold for HLB assembly (White et al., 2011; Terzo et al., 2015). *mxc*<sup>G46</sup> encodes a C-terminal truncation of Mxc protein lacking aa 1643–1837, and *mxc*<sup>G46</sup> mutants are viable but female sterile. In the *mxc*<sup>G46</sup> mutant, FLASH is also mislocalized (Fig. 4 C). Importantly, the total amount of FLASH protein in *mxc*<sup>G46</sup> mutant ovaries was similar to the normal levels of WT FLASH (Fig. 5 D). Thus the C-terminal region of Mxc is required for localization of FLASH to the HLB (Fig. 4 C). Because our C-terminal anti-Mxc antibody does not detect Mxc<sup>G46</sup> protein (White et al., 2011; Fig. 4 C), we stained salivary glands of transgenic GFP-Mxc<sup>G46</sup>-rescued animals with anti-GFP antibodies to confirm that Mxc<sup>G46</sup> protein supports HLB assembly (Fig. 4 C; Terzo et al., 2015).

In *mxc*<sup>G46</sup> mutants, we detected small amounts of both misprocessed H2a RNA and read-through transcripts, although the majority of the H2a mRNA was properly processed (Fig. 5 B). These results are similar to the FLASH<sup>I-733</sup> and FLASH<sup>LDIY71</sup> mutants (Fig. 2 D). We also observed read-through transcripts by in situ hybridization in both the FLASH<sup>I-733</sup> and Mxc<sup>G46</sup> nurse cells (Fig. 6, yellow arrows). Thus, mislocalizing WT, FL



**Figure 3. Concentrating FLASH in the HLB promotes histone 3' end formation.** (A) Diagram of C-terminal FLASH deletion mutants. (B) Visualization of rescue of *FLASH<sup>PBac/Df</sup>* with transgenic FLASH C-terminal mutants as in Fig. 1 D. (C) Anti-V5 Western analysis of total protein from larvae expressing *FLASH<sup>FL</sup>*, *FLASH<sup>LDIY71</sup>*, *FLASH<sup>1-733</sup>*, and *FLASH<sup>LDIY71, 1-733</sup>* rescue transgenes. Asterisk, cross-reacting protein. (D) Localization of FLASH transgenic proteins (V5) in larval polyploid salivary gland nuclei stained for an HLB marker (Mxc). WT transgenic FLASH protein localizes to the HLB (yellow arrow), but the C-terminal truncation mutants do not (white arrows). Bars, 10  $\mu$ m. (E) Total RNA from WT third-instar larvae, the *FLASH<sup>PBac/Df</sup>* mutant and the mutant rescued with the indicated FLASH transgenes was analyzed by S1 nuclease mapping as in Fig. 1 E. An autoradiograph is shown and the band intensities of a PhosphorImage of each lane were plotted using the ImageJ gel-analysis function.

FLASH by truncating Mxc resulted in essentially identical effects on histone mRNA as removal of the FLASH C terminus. Although we cannot rigorously exclude another contribution to processing from the FLASH C terminus, these results are consistent with the effect being caused by mislocalizing FLASH.

#### High concentrations of FLASH are necessary for U7 snRNP accumulation in the HLB

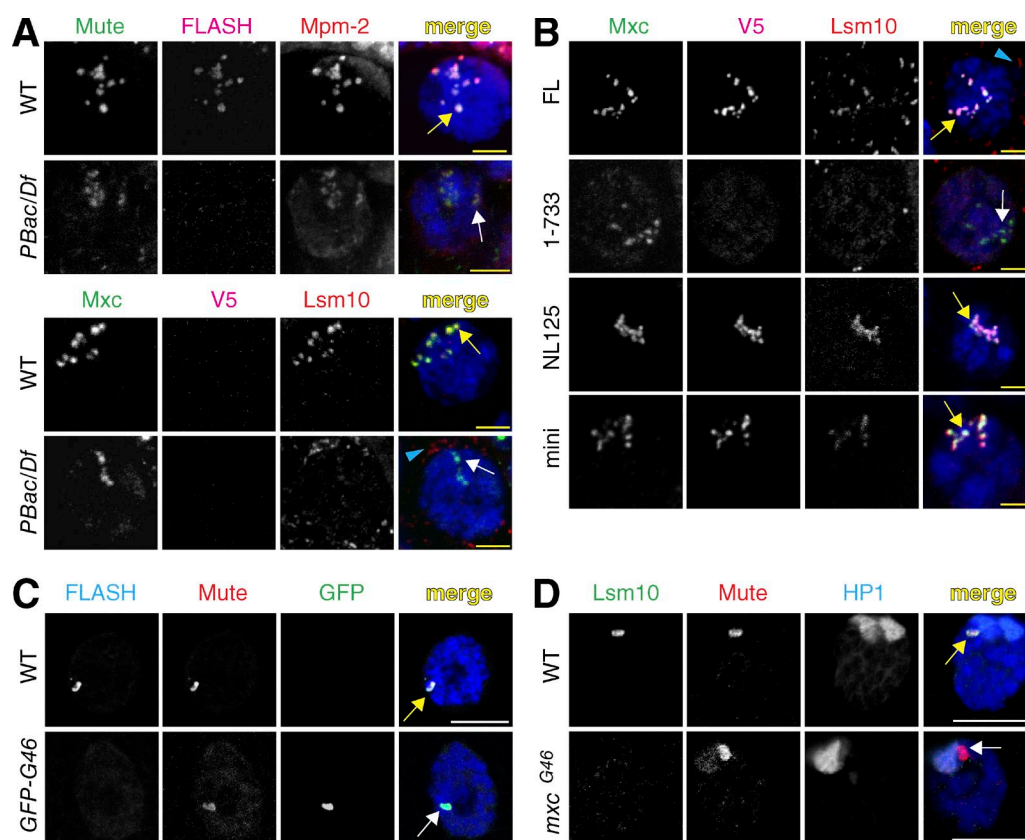
To determine whether some of the effects of mutating FLASH might be mediated through changes in HLB composition, we visualized HLBs in different FLASH mutant animals with antibodies against a panel of HLB markers (Figs. 4 and S3). We analyzed ovaries of the small number of *FLASH<sup>PBac/Df</sup>* mutant females that survive to adulthood. Although these females are sterile and we do not observe fully developed eggs, oogenesis consistently proceeded to stage 8, allowing us to examine stage 5 polyploid nurse cells, which provide excellent cytology for HLBs (Liu et al., 2006). We did not detect FLASH in the HLBs of *FLASH<sup>PBac/Df</sup>* mutant ovaries, although Mxc and Mute were present (Fig. 4 A). We also did not detect the U7 snRNP-specific protein Lsm10 in these HLBs (Fig. 4 A). Expressing *FLASH<sup>FL</sup>* protein in the *FLASH<sup>PBac/Df</sup>* mutant restored Lsm10 localization to the HLB, as did expressing *FLASH<sup>mini</sup>* and *FLASH<sup>NL125</sup>* (Fig. 4 B). In contrast, expressing the *FLASH<sup>1-733</sup>* mutant protein resulted in accumulation of low levels of Lsm10 in the HLB (Fig. 4 B), which was only detectable in some experiments and is possibly because of a small amount of FL FLASH produced by the *FLASH<sup>PBac/Df</sup>* genotype. These data suggest that high

concentrations of FLASH in the HLB are needed for U7 snRNP accumulation in the HLB. In addition, the *FLASH<sup>NL125</sup>* results demonstrate that the interaction of the N terminus of FLASH and Lsm11 is not necessary for concentration of U7 snRNP in the HLB, even though this interaction is required for pre-mRNA processing. Lsm10 also does not accumulate in the HLBs of the *mxc<sup>G46</sup>* mutant (Fig. 4 D), consistent with the results with the mislocalized *FLASH<sup>1-733</sup>* protein. Collectively, our data suggest that two distinct regions of FLASH function within the HLB to (a) localize FLASH and U7 snRNP to the HLB (C terminus of FLASH) and (b) bind the Lsm11 protein of U7 snRNP and recruit the HCC to the U7 snRNP to activate histone pre-mRNA processing.

#### Factors in addition to FLASH are required to concentrate U7 snRNP in the HLB

The *Drosophila* H2aV gene produces a polyadenylated mRNA that encodes a variant H2a histone protein with the combined roles of mammalian H2AX and H2AZ, functioning in transcriptional regulation and the DNA damage response (Redon et al., 2002). The proper ratio of H2aV to canonical H2a histone protein is critical for viability (Li et al., 2014). We have previously shown that U7 snRNP was not localized to the HLB in *H2aV* mutant animals or H2aV-depleted cells (Wagner et al., 2007). Because these studies were done before our discovery of FLASH, we revisited *H2aV* mutants to determine whether loss of H2aV also affects FLASH localization. FLASH and two other HLB markers, Mxc and Mute, colocalize in a prominent focus in *H2aV<sup>810</sup>* mutant salivary glands (Fig. 5 A). Lsm11 was absent from these HLBs (Fig. 5 A).





**Figure 4. Analysis of the HLB in FLASH mutants.** (A and B) The presence of four HLB components was assessed in nurse cells of the indicated genotypes at stage 5 of oogenesis with antibodies against FLASH; Lsm10; Mute, a constitutive HLB factor of unknown function (Bulchand et al., 2010); Mxc; and MPM-2 monoclonal antibodies, which detect phosphorylated Mxc (White et al., 2011). Note that at these stages of oogenesis, the chromatids of the polyploid nurse cells are dispersed, resulting in many HLB foci in close proximity. Yellow arrows indicate proper localization, and white arrows indicate mislocalization. Blue arrowheads indicate signal from the Lsm10 antibody that is not present in the nucleus; the source of this signal is unknown. Bars, 5  $\mu$ m. (C and D) HLB assembly in polyploid salivary gland nuclei of *mxc*<sup>G46</sup> mutants (D) or *mxc*<sup>G46</sup> null mutants expressing transgenic GFP-Mxc<sup>G46</sup> protein (GFP-G46; C) detected with anti-GFP antibodies. Note that salivary gland chromosomes are polytene, resulting in a single, large HLB. HP1 staining visualizes the chromosome center located near the histone locus. Bars, 15  $\mu$ m.

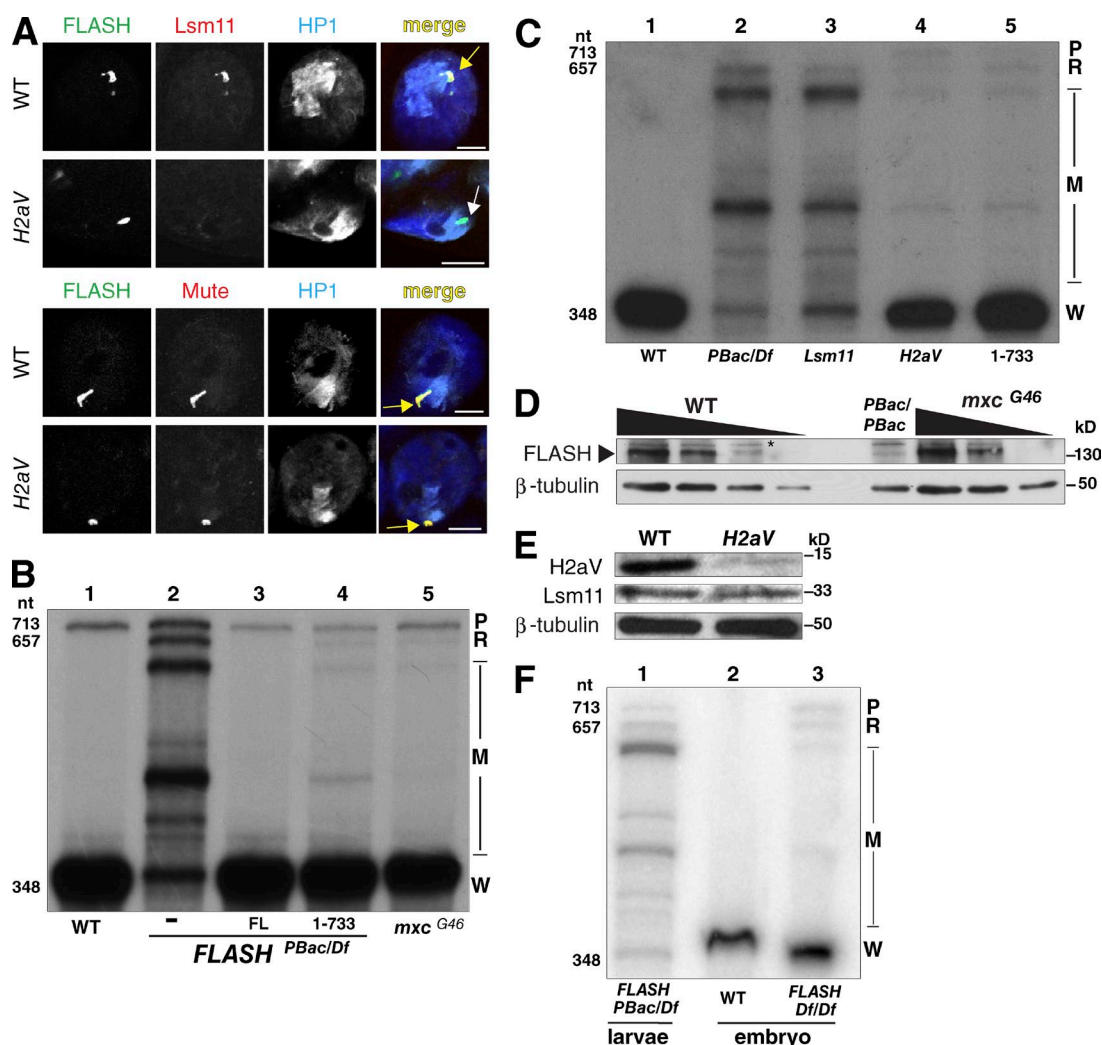
In *H2a*<sup>V810</sup> mutants we detected both misprocessed H2a RNA and read-through transcripts, although the majority of the H2a mRNA was properly processed (Fig. 5, B and C). These results are essentially indistinguishable from the RNA phenotype of the *mxc*<sup>G46</sup> and *FLASH*<sup>1-733</sup> mutants (Fig. 5, B and C), supporting the idea that localization of FLASH and U7 snRNP is necessary for efficient processing of histone mRNA. Importantly, the levels of Lsm11 were similar to WT in the H2aV mutant larvae (Fig. 5 E), and we previously showed that nuclear extracts prepared from H2aV depleted cells are as active as extracts from control cells in histone pre-mRNA processing (Wagner et al., 2007). As with FLASH, these results indicate that high local concentrations of processing-competent U7 snRNP in the HLB, and not its overall availability in the nucleus, are necessary for efficient histone pre-mRNA processing in vivo.

#### The HLB promotes rapid cotranscriptional histone pre-mRNA processing

Our transgenic analysis found that mutants with reduced local FLASH activity, either as a result of mislocalizing FLASH (*FLASH*<sup>1-733</sup>, *mxc*<sup>G46</sup>) or U7 snRNP (*H2a*<sup>V810</sup>), or reducing FLASH activity in the HLB (*FLASH*<sup>LDLY71</sup>), produce read-through and misprocessed RNA in similar amounts (as well as large amounts of WT mRNA). The production of read-through and misprocessed histone mRNAs indicates that RNA

polymerase II (pol II) must have transcribed well past the normal cleavage site. Analysis of pol II occupancy by ChIP sequencing in WT embryos and cultured Kc cells from the modENCODE project revealed that there is little or no RNA pol II present in the intergenic regions between histone genes (Fig. 6 A). This profile suggests that normally there is rapid cleavage followed by efficient transcription termination.

Failure to properly process histone pre-mRNA and terminate transcription results in longer transcripts that are detected at the site of transcription using a probe derived from the region downstream of the H3 gene (H3-ds; Fig. 6 A; Lanzotti et al., 2002; Godfrey et al., 2006, 2009). We tested whether similar transcripts can be detected in mutants with mislocalized FLASH. The fluorescent H3-ds probe does not detect any transcripts in WT cells but detected abundant misprocessed, polyadenylated histone mRNA in the nurse cell cytoplasm of the *FLASH*<sup>Pbac/Df</sup> mutant (Fig. 6 B). This phenotype was rescued by *FLASH*<sup>FL</sup> (Fig. 6 B). *FLASH*<sup>1-733</sup> and *mxc*<sup>G46</sup> nurse cells contained predominantly cytoplasmic WT H3 mRNA that was detected with a coding probe, but not the H3-ds probe (Fig. 6 B). With the H3-ds probe, we detected foci of nascent transcripts in *FLASH*<sup>1-733</sup> and *mxc*<sup>G46</sup> nurse cells, but no cytoplasmic misprocessed mRNA (Fig. 6 B). Because the steady-state level of normally processed histone mRNAs is similar in the *FLASH*<sup>1-733</sup>, *mxc*<sup>G46</sup> and WT animals, many of these longer,



**Figure 5. Failure to concentrate FLASH and/or U7snRNP in the HLB results in inefficient histone pre-mRNA processing.** (A) Larval salivary glands from *H2aV*<sup>B10</sup> mutants were stained for HLB markers and HP1. Arrows as in Figs. 2 and 3. Bars, 10  $\mu$ m. (B) Equal amounts of RNA from ovaries of WT (lane 1) or *FLASH*<sup>Pb<sup>Df</sup> (lane 2), *FLASH*<sup>FL</sup> (lane 3), *FLASH*<sup>1-733</sup> (lane 4), and *mx<sup>c</sup>G46* mutants were analyzed by S1 nuclease protection assay as in Fig. 1 E. (C) Equal amounts of RNA from WT (lane 1), *FLASH*<sup>Pb<sup>Df</sup> (lane 2), *Lsm11*-null mutant (Godfrey et al., 2009; lane 3), *H2aV* mutant (lane 4), and *FLASH*<sup>1-733</sup> (lane 5) third-instar larvae were analyzed by S1 nuclease protection assay as in Fig. 1 E. (D) Western analysis of threefold serial dilutions of extracts from WT, *FLASH*<sup>Pb<sup>Df</sup>, and *mx<sup>c</sup>G46* ovaries using anti-FLASH antibody and tubulin as a loading control. Asterisk, cross-reacting protein. (E) Anti-*H2aV* or anti-*Lsm11* Western analysis of proteins from *H2aV*<sup>B10</sup> mutant larvae, with tubulin as a loading control. See also Fig. S5. (F) Equal amounts of RNA from 14–18 h WT embryos (lane 2) or *FLASH*<sup>Df/Df</sup> embryos (lane 3) were analyzed by S1 nuclease mapping as in E. Lane 1 is analysis of RNA from *FLASH*<sup>Pb<sup>Df</sup> larvae.</sup></sup></sup></sup>

nascent transcripts are likely ultimately cleaved at the normal site. The increase in steady-state amount of unprocessed, nascent histone H3 pre-mRNA suggests that when FLASH or U7 snRNP is not concentrated in the HLB, the rate of histone pre-mRNA processing slows.

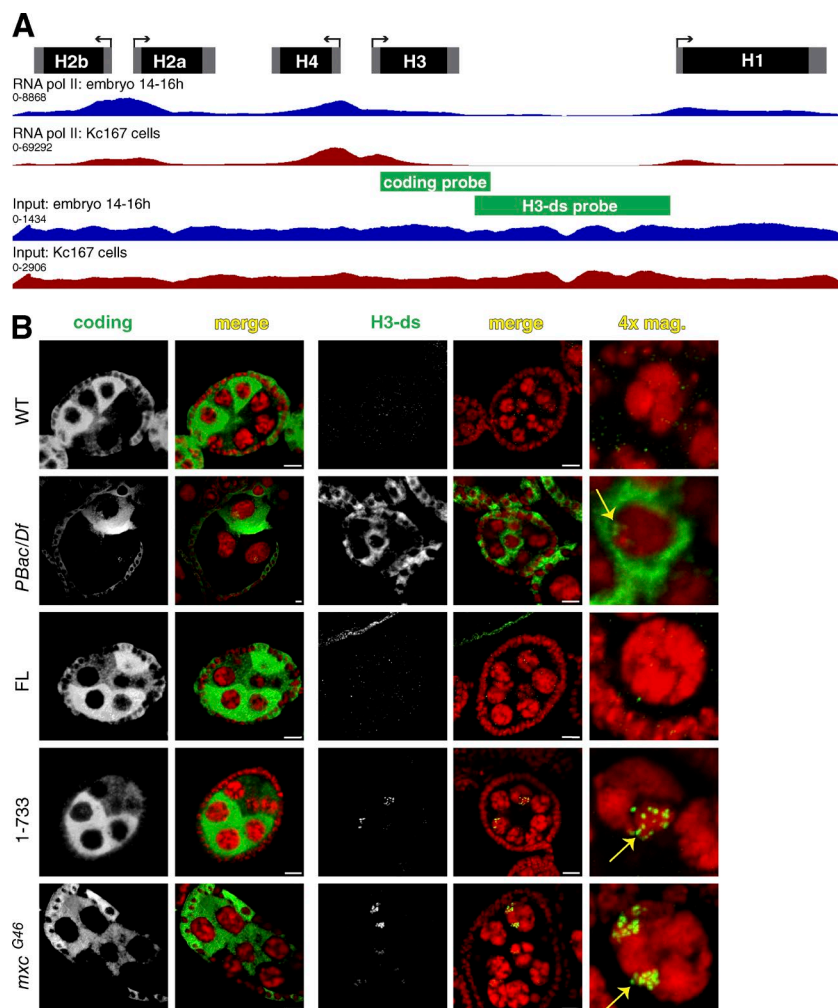
To further support the possibility that the efficiency of 3' end formation is sensitive to the amount of FLASH activity in the HLB, we examined the expression of histone mRNA in a FLASH-null mutant during embryogenesis. Maternally deposited WT FLASH protein accumulates in the HLB before activation of zygotic histone transcription (White et al., 2011). Therefore, zygotic *FLASH* mutant phenotypes will not be apparent until multiple rounds of cell division deplete the maternal FLASH protein. This “maternal run out” situation provides an opportunity to examine embryos with reduced concentrations of FLASH in the HLB but before FLASH is completely depleted. Stage 17 (14–18 h) *FLASH*<sup>Df/Df</sup> mutant embryos had both misprocessed and read-through histone mRNAs in similar

amounts (Fig. 5 F, lane 3), a phenotype qualitatively different from the effect of severely depleting FLASH that occurs by larvae stages. Thus, the initial phenotype as maternal FLASH is depleted in the *FLASH* mutant is similar to the effect of mislocalizing normal amounts of FLASH, as seen in the *mx<sup>c</sup>G46* and *FLASH*<sup>1-733</sup> mutants. This effect is likely a result of reducing the local concentration of FLASH in the HLB, supporting the conclusion that concentrating factors in the HLB is important for efficient histone mRNA 3' end formation.

## Discussion

Our systematic analysis of *Drosophila* FLASH mutants and mutants in other HLB components has allowed us to attribute a function in histone mRNA metabolism to the HLB: promoting rapid histone pre-mRNA processing, resulting in efficient coupling of transcription termination with 3' end formation.





**Figure 6. Histone mRNA expression analyzed by in situ hybridization.** (A) RNA pol II occupancy data for 14–16 h embryos and Kc167 cells (Kharchenko et al., 2011) visualized on a single histone repeat. The location of in situ hybridization probes used in B is indicated. (B) Fluorescent in situ hybridization of ovaries of the indicated genotypes using an H3 coding probe or a probe (H3-ds) that only detects misprocessed or read-through transcripts. Note that histone H3 mRNA accumulates only during S phase and that the nurse cell endocycles are asynchronous. The cytoplasmic H3-ds signal in *FLASH<sup>PBac/Df</sup>* is from misprocessed H3 transcripts that were polyadenylated and exported. The small amounts of misprocessed RNA in *FLASH<sup>1-733</sup>* and *mxc<sup>G46</sup>* (see Fig. 5 B) mRNAs are below our threshold of detection. The nuclear foci in *FLASH<sup>1-733</sup>* and *mxc<sup>G46</sup>* mutants detected with the H3-ds probe (yellow arrows) are nascent transcripts at the histone locus. Bars, 10  $\mu$ m.

### The HLB contributes to multiple steps in histone mRNA biosynthesis

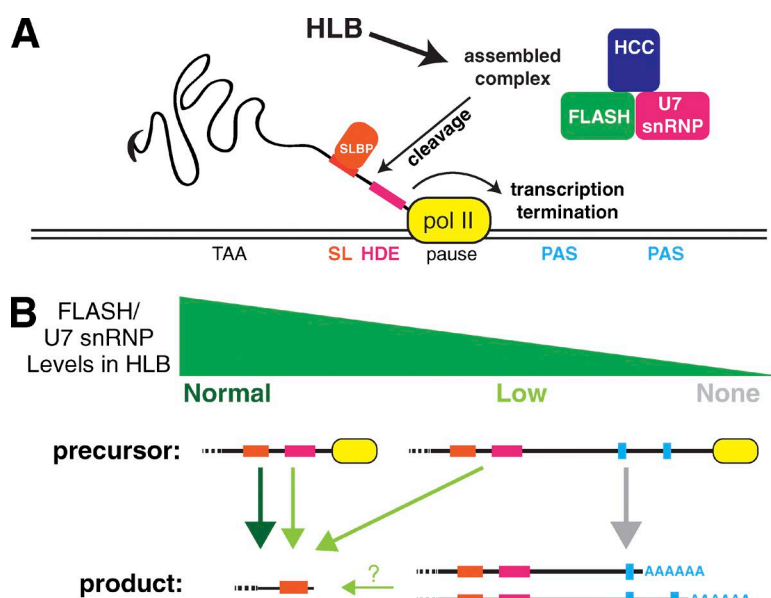
HLB assembly initiates with formation of a complex containing FLASH and Mxc, followed by incorporation of other components including U7 snRNP (White et al., 2011). How Mxc and FLASH are specifically targeted to the histone locus is not known, but this process does not require the 3' processing signals in the histone pre-mRNA and is coordinated by the *Drosophila* H3-H4 promoter (Salzler et al., 2013). An emerging property of HLB components is the presence of independent biochemical and HLB targeting domains (Yang et al., 2014; Terzo et al., 2015). For instance, the FLASH C terminus is required for localization to the HLB, whereas the N-terminal region required for histone pre-mRNA processing interacts with U7 snRNP. Similarly, U7 snRNP recruitment to the HLB requires the C termini of FLASH and Mxc but does not require the interaction between the N termini of FLASH and Lsm11 that is required for histone pre-mRNA processing (Burch et al., 2011).

Once the HLB assembles in early development, Mxc, FLASH, and U7 snRNP are concentrated in the HLB throughout interphase, independent of whether the histone genes are being expressed (White et al., 2011). Thus, activation of histone mRNA biosynthesis is distinct from recruitment of HLB components to the histone locus. The constitutive presence of FLASH and U7 snRNP contrasts with cell cycle-dependent enrichment of the HCC component Symplekin in the HLB (Tatomer et al., 2014).

The uncoupled process for HLB localization of and interaction between U7 snRNP and FLASH suggests that changes within the HLB during S phase may facilitate HCC recruitment to and subsequent cotranscriptional cleavage of histone pre-mRNA.

### Concentrating factors within the HLB ensures efficient histone mRNA synthesis

Here, we have analyzed three different situations to determine that high concentrations of FLASH and U7 snRNP in the HLB are necessary for efficient histone mRNA synthesis: (1) FLASH mutants with impaired localization, (2) mutants in Mxc and H2aV that result in mislocalization of WT FLASH and/or U7 snRNP, and (3) diminished WT FLASH levels as the maternal supply of FLASH is reduced during development of FLASH mutant embryos. A direct way to do this would be to retarget *FLASH<sup>1-733</sup>* to the HLB by replacing the C terminus of FLASH with another HLB localization signal. We attempted to directly test the possibility that localization of FLASH is critical for histone pre-mRNA processing by retargeting *FLASH<sup>1-733</sup>* to the HLB with the N-terminal 101 aa of Mxc, which can target GFP to the HLB and is essential to target Mxc to the HLB (Terzo et al., 2015). Although this signal targeted *FLASH<sup>1-733</sup>* to the HLB in cultured cells and partially rescued the FLASH mutant phenotype (Fig. S4), only a small amount of the fusion protein accumulates in transgenic animals in some tissues (Fig. S5), precluding us from making any firm conclusions regarding additional functions of the FLASH C terminus.



**Figure 7. Model of HLB participation in histone pre-mRNA processing.** (A) RNA pol II (yellow) pauses 3' of the HDE, allowing SLBP (orange) to bind the stem loop (SL) in the pre-mRNA downstream of the stop codon and U7 snRNP (pink), FLASH (green), and the HCC (blue) to assemble into a complex that associates with the pre-mRNA at the HDE. Processing occurs followed by transcription termination. (B) High levels of active FLASH and U7 snRNP (dark green arrow) in the HLB result in rapid cleavage while pol II is paused leading to transcription termination, preventing RNA pol II from encountering downstream polyadenylation signals (PASs). Low levels of FLASH (light green arrows) slow normal processing resulting in release of some pol II from the pause site. This leads to production of longer pre-mRNAs, which are detected at the site of histone gene transcription by in situ hybridization. Most of these transcripts are ultimately processed at the normal site, and a small proportion of these are polyadenylated (thin, green arrow). In this situation, there are multiple possible precursor RNAs that could give rise to properly processed histone mRNA. Some will be processed while the RNA pol II is paused, others will be processed after the pol II releases from the pause site, and it is possible (question marks) that some of the polyadenylated transcripts could be "reprocessed" as described in the text. Absence or severe reduction in any histone pre-mRNA processing factor, whether or not it is an HLB component, results in production of a long precursor pre-mRNA that is processed by the polyadenylation machinery (gray arrow).

The processing of histone pre-mRNAs in *Drosophila* is normally very efficient. There is a strong pause site just downstream of the U7 binding site (the histone downstream element [HDE]; Adamson and Price, 2003), and no transcripts are detected extending 3' of the processed RNA in WT animals (Lanzotti et al., 2002). In addition, RNA pol II is not present downstream of the HDE in both *Drosophila* (Fig. 6 A) and mammalian cells (Anamika et al., 2012; Cheng et al., 2012). Thus, termination of transcripts from the histone genes is normally tightly coupled with histone pre-mRNA processing. Mutations in *Drosophila* FLASH, SLBP, or U7 snRNP that prevent histone pre-mRNA processing allow transcription to proceed 3' of the histone genes, resulting in formation of polyadenylated histone mRNAs via cryptic, downstream signals (Lanzotti et al., 2002; Godfrey et al., 2006). Here, we showed that subtler perturbations that affect the localization of FLASH and/or U7 snRNP to the HLB, or that partially impair FLASH activity within the HLB, also result in read-through transcripts at the histone locus, although large amounts of properly processed histone mRNA still accumulate. The appearance of these longer, nascent transcripts suggests that normal processing is delayed, resulting in loss of coupling between processing and transcription termination.

What is the basis for the mutant RNA phenotype in cases where the activity of processing factors is reduced rather than abolished? Mechanistic differences between processing 3' ends of polyadenylated mRNAs and histone mRNAs suggest an answer. The interaction of polyadenylation factors with the C-terminal domain of RNA pol II couples polyadenylation with transcription (McCracken et al., 1997; Adamson et al., 2005) and the C-terminal domain may participate directly in promoting cleavage or polyadenylation (Hirose and Manley, 1998). In contrast, there are no known interactions between RNA pol II and components specific to the histone-processing complex. Unlike polyadenylation, cotranscriptional processing does not increase the rate of histone mRNA 3' end formation in vitro (Adamson and Price, 2003). These results suggest that process-

ing histone pre-mRNA at the normal location can occur after the polymerase has transcribed well past the HDE (Fig. 7).

A second possible way to accumulate properly processed histone mRNA from longer transcripts is to "reprocess" a transcript that has already been cleaved and polyadenylated. Our previous study suggested in mammalian cells that utilization of a processing site downstream of the stem loop or HDE might stabilize a longer transcript, providing time for partially defective histone processing machinery to generate a normal histone mRNA 3' end (Liu et al., 1989). Indeed, in *Caenorhabditis elegans*, which lacks FLASH and U7 snRNP but contains SLBP, there is evidence that the normal pathway of histone mRNA processing is polyadenylation followed by cleavage after the stem loop by an siRNA-like mechanism (Mangone et al., 2010; Avgousti et al., 2012).

### Cellular microenvironments that enhance biological processes

Our results support the idea that NBs facilitate reactions by concentrating factors (Matera et al., 2009; Mao et al., 2011b). The exchange of NB components with the adjacent environment is slower than expected by diffusion, promoting concentration in NBs (Dundr et al., 2004). Such concentration of components creates discrete microdomains with different physical chemical properties than the surrounding nucleoplasm (Brangwynne et al., 2011; Han et al., 2012; Kato et al., 2012). A role for CBs in facilitating snRNP assembly by concentrating factors has been previously suggested (Klingauf et al., 2006; Stanek et al., 2008; Novotný et al., 2011). In addition, a reduced rate of tri-snRNP assembly was postulated to cause the splicing defects observed in zebrafish lacking CBs because of Coilin depletion (Strzelecka et al., 2010). In the case of the HLB, our data provide evidence for two roles: enhancing the rate of pre-mRNA processing and promoting coupling of processing with transcription termination. Identification and characterization of additional HLB components in the future will facilitate our ability to further test this model of NB function.

## Materials and methods

### Drosophila strains

PBac[PB]FLASH<sup>LL01602</sup> was a gift from G. Matera (University of North Carolina, Chapel Hill, NC), who obtained it from the *Drosophila* Genomics Resources Center. The site of insertion was determined by PCR and sequencing (Fig. S1). FLASH constructs previously generated in pIZ/V5 (Thermo Fisher Scientific) were inserted into pATTB (Bischof et al., 2007) with KpnI and XhoI for  $\phi$ C31-mediated transgenesis (Best Gene) and integration into the attP40 landing site. Transgenic GFP-Mxc<sup>G46</sup> integrated at VK00033 was expressed with the ubiquitin promoter of pUGW in the *mxc*<sup>G48</sup> null background (White et al., 2011). Additional strains are summarized in Table 2. The features of the FLASH transgenes are summarized in Table 3.

### Genomic PCR

Genomic DNA was prepared by incubating a squashed fly in 50  $\mu$ l of squash buffer (10 mM Tris, pH 8.2, 25 mM NaCl, 1 mM EDTA, and 200  $\mu$ g/ml Proteinase K; New England Biolabs, Inc.) at 37°C for 30 min. After 10 min at 85°C, 1  $\mu$ l per reaction was used to amplify each product with TAQ DNA polymerase (Thermo Fisher Scientific). PCR products were purified (Thermo Fisher Scientific) and subjected to Sanger sequencing. The primers used in each PCR reaction are summarized in Table 4.

### Histone mRNA analysis

A total of 5  $\mu$ g total cellular RNA extracted with Trizol reagent (Invitrogen) was used for each S1 nuclease protection reaction. The probe was created by 5' end labeling BspEII cut H2a DNA with  $\alpha$ -<sup>32</sup>P-dCTP using the Klenow fragment of DNA polymerase I (New England Biolabs, Inc.). The probe was gel purified and hybridized to either total larval RNA or control yeast tRNA followed by digestion with S1 nuclease (Lanzotti et al., 2002). Protected fragments were resolved on a 6% polyacrylamide-7M urea gel and visualized by autoradiography. The band intensities for each lane were plotted by the ImageJ gel-analysis function.

### Immunoblotting

Ovaries of the indicated genotypes were dissected in Grace's media (Gibco) and lysed in RIPA buffer (150 mM NaCl, 1% Triton X-100, 50 mM Tris, pH 7.5, 0.1% SDS, 0.5% sodium-deoxycholate, and protease inhibitors; Thermo Fisher Scientific). Brains and imaginal discs were dissected from wandering third-instar larvae and lysed by boiling in sample buffer. All samples were passed through a 25-gauge needle 100 times before resolving through a 10% gel by SDS-PAGE and detection with ECL Prime (GE Healthcare) using the antibodies summarized in Table 6.

### Immunofluorescence

For immunofluorescence, salivary glands were dissected in PBS + 0.1% Triton X-100 and ovaries were dissected in Grace's medium

Table 2. *Drosophila* strains

Strain	Source	Reference
PBac LL01602	Drosophila Genomics Resource Center: 140418	—
Df (2R) 8057	Bloomington: 7871	—
PBac c02047 ( <i>Lsm11</i> )	Exelixis Harvard: c02047	Godfrey et al., 2009
<i>His2a</i> <sup>V810</sup>	Bloomington: 9264	van Daal and Elgin, 1992
<i>Mxc</i> <sup>G46</sup>	Bloomington: 32114	White et al., 2011
<i>Mxc</i> <sup>G48</sup>	Bloomington: 7141	White et al., 2011

Table 3. Construct features

FLASH transgene feature	Sequence
attB-KpnI-promoter	5'-GATCTCTAGAGGTACCcgagattattgttattgtatgtgttttttaacgaatttaa-3'
V5-XhoI-attB	5'-GGTAAGCCTATCCCTAACCCTCTCCTCGGTCTCGATTCTACGTAACTCGAGCCGCGGCCGAGAT-3'
5' UTR-EcoRI-coding	5'-GAAGAAACGTAAGCGATTGGTGAATTCGAGCCGCGGCCGAGAT-3'
EcoRI-FL	5'-GAATTCGAGCCGCGGCCGAGAT-3'
EcoRI-aa 65	5'-GAATTCGAGCCGCGGCCGAGAT-3'
EcoRI-aa 78	5'-GAATTCGAGCCGCGGCCGAGAT-3'
aa 844-SacII-V5	5'-CTGCTGCgGACAAACCAATTCCGCGGTTTCGAA-3' GGTAAGCCTATCCCTAACCCTCTCCTCGGTCTCGATTCTACG-3'
aa 733-SacII-V5	5'-CTCACCAGACTCCAAACAGGCTCCGCGGTTTCGAA-3' GGTAAGCCTATCCCTAACCCTCTCCTCGGTCTCGATTCTACG-3'
FLASH <sup>mini</sup> junction	5'-CGAAAAGGACGATGT/ACCCAGACTCCAAAA-3'

Engineered restriction sites are underlined, start and stop codons are bold, the FLASH promoter sequence is lowercase, the V5 tag is italicized, and "/" indicates the unique junction in the FLASH<sup>mini</sup> open reading frame.

Table 4. Sequences for primers used to identify the PBac LL01602

Primer name	Sequence
FLASH sequencing F	5'-GAACGGTACCCGAGATTATTGTTTGTATTGTGATGT-3'
FLASH sequencing R	5'-GTCTTCCAAATCCATCAAGTCCTC-3'
5' Pbac	5'-CGCGATAAATCTTTCTCTCTCG-3'
3' Pbac	5'-CCGATAAAACACATGCGTCA-3'



(Gibco). Fixation and permeabilization conditions are summarized in Table 5. Confocal images were taken at room temperature at a zoom of 1.0–3.0 with a 40 $\times$  (numerical aperture 1.30) Plan Neo-fluor objective on a 510 laser scanning confocal microscope (ZEISS) using the LSM data acquisition software (ZEISS). Image false coloring and contrast was adjusted, and images were digitally zoomed using Photoshop (Adobe). Staining conditions and antibodies are presented in Table 6.

### Fluorescence in situ hybridization

Ovaries were dissected and fixed in 4% formaldehyde in PBS for 20 min and postfixed in 4% formaldehyde in PBS + 0.1% Tween-80 (PBTw). Samples were washed three times in PBTw for 2 min and then incubated in 3  $\mu$ g/ml proteinase K in PBTw for 10 min at room temperature followed by 30 min on ice. Digestion was quenched by incubation in 2  $\mu$ g/ml glycine/PBTw for 2 min twice. After two PBTw rinses, ovaries were postfixed for 20 min in 4% formaldehyde. Ovaries were washed five times in PBTw (Lécuyer et al., 2008). Samples were then hybridized to the indicated probe as previously described (Tomancak et al., 2002). Detection was performed with 1:200 peroxidase-conjugated anti-DIG (Roche) and 1:100 Cy5 tyramide reagent (PerkinElmer).

Fluorescent in situ probes were synthesized by in vitro transcription from vector template DNA that was linearized by digestion with ApaI for H3-coding and XhoI for H3-ds as previously described (Lanzotti et al., 2002). The H3 coding probe was transcribed with T3 RNA polymerase (Ambion), and the H3-ds probe was transcribed with T7 RNA polymerase (Ambion) using the DIG RNA labeling mix (Roche). RNA synthesis was verified by agarose gel electrophoresis. The sequence boundaries for each probe are summarized in Table 7.

### In vitro processing assay

Depletion of FLASH from S2 cells, preparation of nuclear extracts, and in vitro processing assays were performed as previously described (Sabath et al., 2013). The details for nuclear extract preparation are as follows. Cells were harvested by centrifugation (5 min, 3,000 rpm, 4°C) and washed once in PBS. The cell pellet was resuspended in three times the estimated pellet volume in buffer A (10 mM Hepes/KOH, pH 7.9, 1.5 mM MgCl<sub>2</sub>, 10mM KCl, and 0.5 mM DTT [add fresh]). Cells were transferred and lysed with 10–12 strokes in a 2-ml glass Dounce. Nuclei were pelleted by centrifugation (5 min, 5,000 rpm, 4°C). The volume of the nuclear pellet was estimated and resuspended in 1.5 vol of low-salt buffer C (20 mM Hepes/KOH, pH 7.9, 25% glycerol, 1.5 mM MgCl<sub>2</sub>, 0.2 mM EDTA, 0.5 mM DTT [add fresh], and 0.02 M NaCl). The resuspended nuclei were transferred to a fresh tube with a micro-stir bar. High-salt buffer C (20 mM Hepes/KOH, pH 7.9, 25% glycerol, 1.5 mM MgCl<sub>2</sub>, 0.2 mM EDTA, 0.5 mM DTT [add fresh], and 1.2 M NaCl) was added drop by drop to the nuclei. The total amount of high-salt buffer added brought the final concentration of Na<sup>+</sup> to between 0.26 and 0.28 M. The amount was calculated with the following equation: high-salt buffer C = (volume resuspended nuclei  $\times$  0.25/0.95 + volume resuspended nuclei  $\times$  0.28/0.98)/2. After 1 h mixing on ice, the lysate was transferred to a new Eppendorf and cleared by centrifugation (10 min, 10,000 rpm, 4°C). The supernatant was transferred to a Slide-A Lyzer Dialysis cassette (3,500 molecular weight cut-off and 0.1–0.5 ml capacity; Thermo Fisher Scientific) with a syringe and 18-gauge needle. The nuclear extract was dialyzed buffer D (20 mM Hepes/KOH, pH 7.9, 20% glycerol, 0.1 M KCl, 0.2 mM EDTA, and 0.5 mM DTT [add fresh]) for 2 h at 4°C, and this step was repeated with fresh buffer D. The extract was removed with a syringe and cleared by centrifugation (10 min, 10,000 rpm, 4°C). The supernatant was aliquoted, frozen on dry ice/EtOH, and stored at –80°C.

Table 5. Tissue preparation for antibody staining for immunofluorescence

Tissue	Fixation	Permeabilization
Salivary gland (except in Figs. 4 C and 5 A)	7% formaldehyde, 20 min	0.2% Tween-80, 15 min
Salivary gland (Fig. 4 C)	3.7% formaldehyde, 20 min	Not performed
Salivary gland (Fig. 5 A)	3.7% formaldehyde, 15 min	1% Triton X-100, 15 min
Ovary	7% formaldehyde, 20 min	0.2% Tween-80, 15 min

Table 6. Summary of antibodies used for both immunofluorescence and Western blot analysis

Antibody	Raised in or recognizes	Source	Concentration	Incubation
Primary				
FLASH	Rabbit	Yang et al., 2009	1:2,000	4°C, overnight
V5	Mouse	Invitrogen	1:1,000	4°C, overnight
Mxc	Guinea pig	White et al., 2011	1:2,000	4°C, overnight
Mute	Guinea pig	Bulchand et al., 2010	1:5,000	4°C, overnight
Lsm10	Rabbit	Liu et al., 2006	1:2,000	4°C, overnight
Lsm11	Rabbit	Liu et al., 2006	1:2,000	4°C, overnight
HP1	Mouse	Developmental Studies Hybridoma Bank	1:1,000	4°C, overnight
H2aV	Rabbit	Leach et al., 2000	1:1,000	4°C, overnight
GFP	Chicken	EMD Millipore	1:1,000	4°C, overnight
$\beta$ -Tubulin	Rabbit	Abcam	1:5,000	Room temperature, 2 h
Secondary				
Alexa Fluor 488	Rabbit	Invitrogen	1:2,000	Room temperature, 2 h
Alexa Fluor 488	Mouse IgG2a	Invitrogen	1:1,000	Room temperature, 2 h
Cy3	guinea pig	Jackson ImmunoResearch Laboratories, Inc.	1:1,000	Room temperature, 2 h
Alexa Fluor 555	Mouse IgG1	Invitrogen	1:1,000	Room temperature, 2 h
Cy5	rabbit	Abcam	1:2,000	Room temperature, 2 h
Cy5	rabbit	Jackson ImmunoResearch Laboratories, Inc.	1:1,000	Room temperature, 2 h
ECL donkey HRP	rabbit	GE Healthcare	1:10,000	Room temperature, 1 h
ECL prime donkey HRP	mouse	GE Healthcare	1:10,000	Room temperature, 1 h

Table 7. Boundaries for fluorescent in situ hybridization probe sequences

Probe	Sequence
H3-coding 5' end of probe	5'-ATGGCTCGTACCAAGCAAAC-3'
H3-coding 3' end of probe	5'-GCTCAACATTTATTAATTA-3'
H3-ds 5' end of probe	5'-CGGTACTGGGTCTTAATCA-3'
H3-ds 3' end of probe	5'-AGTGCTCTCTCTCGATTC-5'

## RNA pol II localization

High-throughput sequencing reads were processed as previously described (McKay and Lieb, 2013). The following exceptions were made to unambiguously map reads to the histone locus. The exact number of histone gene repeats in the *Drosophila* genome is unknown, and the reference genome sequence is thus incomplete. To circumvent this problem, a custom reference genome was created by removing all canonical histone gene-repeat sequences and adding back a single 5-kb histone gene-repeat unit. An unlimited number of reads were then mapped to this custom genome with bowtie (Langmead et al., 2009) using the options “–nomaqround” and “–best.” Coverage values were then calculated for each base in the genome, and the data were visualized using Integrative Genomics Viewer (Robinson et al., 2011). Datasets modENCODE\_5122 and modENCODE\_5569 were downloaded from the modENCODE ftp site (Kharchenko et al., 2011).

## Online supplemental material

Fig. S1 is a characterization of *FLASH*<sup>LL01602</sup> PBac. Fig. S2 shows expression and localization of transgenic FLASH proteins. Fig. S3 shows RNA phenotypes in animals where FLASH concentration and/or biochemical activity of FLASH in the HLB is altered. Fig. S4 shows attempts to target *FLASH*<sup>1–733</sup> to the HLB using the HLB localization site from Mxc. Fig. S5 shows HLB localization analysis of transgenic FLASH-mxc fusion constructs. Online supplemental material is available at <http://www.jcb.org/cgi/content/full/jcb.201504043/DC1>.

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