Architecture of Pol II(G) and molecular mechanism of transcription regulation by Gdown1

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Abstract

Tight binding of Gdown1 represses RNA polymerase II (Pol II) function in a manner that is reversed by Mediator, but the structural basis of these processes is unclear. Although Gdown1 is intrinsically disordered, its Pol II interacting domains were localized and shown to occlude TFIIF and TFIIB binding by perfect positioning on their Pol II interaction sites. Robust binding of Gdown1 to Pol II is established by cooperative interactions of a strong Pol II binding region and two weaker binding modulatory regions, thus providing a mechanism both for tight Pol II binding

and transcription inhibition and for its reversal. In support of a physiological function for Gdown1 in transcription repression, Gdown1 co-localizes with Pol II in transcriptionally silent nuclei of early Drosophila embryos but re-localizes to the cytoplasm during zygotic genome activation. Our study reveals a self-inactivation through Gdown1 binding as a unique mode of repression in Pol II function.

Introduction

In eukaryotes, Pol II is the central machine responsible for transcription of genes that specify mRNAs, snRNAs, and microRNAs. In response to signals that result in expression of specific sets of genes, Pol II is recruited to cognate gene promoters through the concerted functions of transcription activators, co-activators that include the Mediator, and the general transcription factors (GTFs)[1]. The GTFs (TFIIB, -D, -E, -F, and –H) play essential roles in transcription through interactions with Pol II at core promoters during preinitiation complex (PIC) formation and function (initiation)[2]. The large (25–30 subunit) Mediator complex acts principally in the regulation of transcription through joint enhancer-bound activator and Pol II interactions that facilitate both Pol II recruitment and enhancer-promoter interactions[3–5]. However, the Mediator has multiple functions in transcription that are not fully understood[3,4].

While conventional Pol II is composed of 12 subunits (POLR2A to 2L), Pol II(G) is a newly discovered form of Pol II containing a stoichiometric, tightly associated Gdown1 subunit (POLR2M)[7]. Although Pol II and Mediator are highly conserved from yeast to human[6], Gdown1 is metazoan-specific[7]; and its ubiquitous expression[8,9] suggests a global role in transcriptional regulation. However, the biological roles of Gdown1 remain largely unknown. In vitro studies revealed that in the absence of Mediator, Pol II(G), unlike Pol II, inhibits transcription, suggesting a primary function of Gdown1 in transcription repression. This effect is attenuated in the presence of Mediator[7], indicating that Mediator somehow reverses the inhibitory effect of Gdown1. Thus, the formation of Pol II(G) elicits an enhanced dependence upon Mediator for transcription. Although the underlying mechanism is unclear, the Mediator-dependent Pol II(G)-mediated transcription appears to differ from Pol II-mediated transcription.

In vitro studies also revealed that Gdown1 prevents TFIIF from associating with Pol II, which in turns leads to inhibition of PIC assembly[10]. TFIIF plays a critical role in transcription initiation[2,6] by stabilizing the PIC through interactions with other GTFs (including TFIIB and TFIIE) as well as promoter DNA[11]. However, the fact that Pol II interactions with Gdown1 and TFIIF are mutually exclusive indicates that for successful PIC formation, either Gdown1 must be dissociated from Pol II or, alternatively, the inhibitory effect of Gdown1 on TFIIF binding to Pol II must somehow be neutralized (for example, by structural changes). Our previous studies were unable to deduce the fate of Gdown1 when Pol II(G)-mediated repression was reversed due to the extremely strong binding to Pol II[10].

Here, we examine in detail both physical and functional interactions between Pol II and Gdown1. We map Gdown1 locations on Pol II by cryo-EM, CX-MS and integrative modeling approaches and further establish Gdown1 Pol II interaction regions by

biochemical analyses. We find that Gdown1 locations on Pol II perfectly overlap contact sites for TFIIF and TFIIB. Moreover, we identify mutations in Gdown1 that bypass the Mediator requirement for reversal of the repression. Analyses of these mutants show that the primary Pol II binding region(s) of Gdown1 are regulated by two essential regulatory regions that further stabilize Gdown1-Pol II interactions. An integrative modeling approach uniquely reveals the location of these regulatory regions, providing insights into the molecular mechanisms underlying both the robust binding of Gdown1 to Pol II and the Mediator-facilitated dissociation of Gdown1 from the Pol II binding site. This model is consistent with the further demonstration of an essential role of Gdown1 in Drosophila melanogaster early embryonic development, where Gdown1 co-localization with Pol II in nuclei is inversely correlated with active gene transcription.

Results

Molecular architecture of Pol II(G) based on cryo-EM and CX-MS analyses

Gdown1 and TFIIF associate with Pol II in a mutually exclusive manner, which prevents efficient formation of PIC and leads ultimately to inhibition of transcription[10]. To further elucidate the molecular mechanism of the inhibition, we first set out to visualize Gdown1 location on Pol II by single-particle cryo-electron microscopy. Human Pol II(G) was purified from nuclear extracts of cells expressing flag-tagged Gdown1[10], and Pol II(G) cryo-EM images were computationally screened and clustered to obtain a cryo-EM map of Pol II(G) at ~4 Å resolution (Methods, Supplementary Fig. 1). All major Pol II domains were well resolved in the Pol II(G) cryo-EM map, with the exception of the clamp and RPB4 and RPB7 subunits (Fig.1a and Supplementary Fig. 1d, e) that appeared blurred (possibly due to high mobility). An atomic model of human Pol II was obtained by real-space refinement of a model derived from relevant portions of the published model of bovine Pol II[12]. To identify Gdown1 density in the Pol II(G) map, we calculated a difference map between the Pol II(G) cryo-EM map and a 4 Å molecular map derived from corresponding portions of the bovine Pol II atomic model. Three major densities were observed: (1) around RPB3 and RPB10; (2) at the RPB2 protrusion and wall domains; and (3) at the RPB1 dock domain (Fig. 1a).

To identify which Gdown1 residues might correspond to Gdown1 densities in the Pol II(G) cryo-EM map, we performed CX-MS analyses[13,14] with purified Pol II(G) (Methods, Supplementary Fig. 1a), and obtained 40 cross-links (Methods, Supplementary Table 1) that were used to build maps of Gdown1-Pol II inter-molecular interactions (Fig. 1b) and Gdown1 intra-molecular interactions (Fig. 1c). Inter-molecular cross-links were detected in Pol II subunits RPB1, RPB2, RPB3, and RPB10. With the exception of RPB1 dock contacts, these crosslinks were in accord with cryo-EM data.

Gdown1 interaction sites overlap TFIIF and TFIIB contact sites on Pol II

Relative to effects of Gdown1 on its function, TFIIF is composed of Rap74 and Rap30 subunits that form a hetero-dimer whose functional domains, dimerization domain and WH domain are connected by the Rap30 linker domain[6] (Fig. 2a). Strikingly, Gdown1 densities on Pol II overlapped perfectly with the interaction site of the Rap30 linker domain on RPB2[11] (Fig. 2b) and also with the interaction site of the TFIIF dimerization domain on the

RPB2 lobe[15–17] (Supplementary Fig. 2a). In addition, four lysines in the Gdown1 C-terminus cross-linked to the RPB2 protrusion near the binding site of the Rap30 WH domain[18] (Fig. 1b, 2c). This perfect overlap unequivocally confirms, and further details, the molecular basis of Gdown1 inhibition of TFIIF binding to Pol II[10].

Our cryo-EM analyses also revealed Gdown1 density overlaps with the Pol II interaction sites (RPB1 dock and RPB2 wall domains) of the TFIIB B-core (Fig. 2a, Supplementary Fig. 2b) and B-ribbon domains (Fig. 2d)[19,20]. Consistent with these results, we observed crosslinking of the Gdown1 C-terminal region to residues (K820, K821) in the RPB2 wall domain (Fig. 1b, Fig. 2d); and although a corresponding density was not detected, the Gdown1 C-terminal region was also crosslinked to residues (K213, K331) in the RPB1 clamp domain (Fig. 1b and Supplementary Fig. 2c) that is recognized by the TFIIB B-linker[19,20]. These striking coincidences of Gdown1 C-terminal region and TFIIB interaction sites on Pol II prompted us to examine whether Gdown1 could also inhibit TFIIB binding to Pol II. To this end, we performed a Pol II binding assay with ^{35}S-labeled TFIIB and an in vitro transcription assay with a pre-melted promoter template that retains the TFIIB requirement for transcription but partially bypasses the normal TFIIF requirement[21] and, correspondingly, decreases Gdown1 transcriptional inhibition through TFIIF[22]. Notably, the results show that Gdown1 indeed interferes with the interaction of TFIIB with Pol II and inhibits transcription (Supplementary Fig. 2d, e). Since the Pol II clamp loop rudder extends into the active site, directly interacts with DNA and facilitates separation of the RNA transcript from DNA[23], these results also suggest a hindrance to the Gdown1 C-terminal interaction with this domain once RNA synthesis has begun.

Identification of Gdown1 domains that stabilize the interaction of Pol II binding region I with Pol II

We next determined Pol II binding and functional domains of Gdown1 by biochemical studies. GST-pulldown assays showed robust binding of the Gdown1 C-terminal fragment (181–368) to Pol II, but only weak binding of the N-terminal fragment (and only when fused C-terminally to GST) (Supplementary Fig. 3a, b). In complementary in vitro transcription assays (Supplementary Fig. 3c), the C-terminal fragment, but not the N-terminal fragment, showed a dose-dependent inhibition that was significantly less than that observed for full-length, suggesting that while the N-terminal region (analyzed further below) cannot act alone it somehow contributes to the intrinsic inhibitory activity of the C-terminal region. In a further analysis of the C-terminus based in part on conserved amino acids, we generated various C-terminal deletion mutants and tested their effects on basal transcription and Pol II-binding (Supplementary Fig. 3a, d, e, f). Fragments containing C-terminal deletions to residue 314 maintained basal transcription inhibitory activity (Supplementary Fig. 3d), whereas fragments with further deletions either lost inhibitory activity while maintaining significant (but not full) Pol II binding (fragments 1–298 and 1–269) or lost both Pol II binding and inhibitory activity (fragment 1–226) (Supplementary Fig. 3e, f). Based on these results, high sequence conservation (Fig. 3c), and further results in Fig. 3d (discussed below), we identify residues 299–314 as a C-terminal transcription inhibitory region (C-TIR) important for Gdown1 inhibitory activity but not for general Pol II binding. Based on several Gdown1 deletion and binding studies (Supplementary Fig. 3f, g, h, i, j), we define

Gdown1 (227–298) as Pol II binding region I. Gdown1 (315–340) is defined as Pol II binding region II based on crosslinking results (Fig. 1b). Although its Pol II binding activity is significantly weaker than that of Pol II binding region I (Supplementary Fig. 3g, also see below), the C-terminal region (315–368) that includes Pol II binding region II is clearly involved in the inhibitory activity (Supplementary Fig. 3d, e).

In a further analysis of the C-TIR region by mutation of conserved residues, full length Gdown1 with L303A and L304A mutations showed a near-complete loss of transcriptional inhibitory activity (Fig. 3d), while retaining full Pol II binding activity (Fig. 3e). These results indicate that the two hydrophobic residues, and thus C-TIR, are critical for Gdown1 inhibitory activity and, further, that the loss of Gdown1 activity is not due to an overall loss of Pol II binding. Therefore, we postulated that C-TIR might have qualitative and/or quantitative effects on one or both of the flanking Pol II binding regions. In this regard, the C-TIR L303/4A mutation significantly reduced the binding to Pol II of the Gdown1 (227–368) and (194–368) fragments containing Pol II binding regions I and II, (Supplementary Fig. 3h, j and Fig. 3g, lane 9) -- consistent with its effect on Gdown1 function and a role in altering specific Pol II interactions or functional properties of Pol II binding region I or II [Note that the severe effect of the L303/304 mutation on binding of the Gdown1(194–368) fragment (Fig. 3g, lane 9) relative to the full length Gdown1 (Fig. 3e, lane 5) reflects the loss of N-terminal sequences that, like C-TIR, also stabilize binding of the C-terminal domains]. Although the exact role of C-TIR in modulating or stabilizing the Pol II interactions of the Gdown1 C-terminal domains is unclear, the extensive cross-linking of these domains (Fig. 1b) suggests that it may facilitate their direct cooperative binding to Pol II.

In further consideration of Pol II binding region I, Gdown1 residues K228 and K240 were found to cross-link to K95 in the RPB2 protrusion (Fig. 1b and Fig. 3b). The protrusion domain surrounding Gdown1 density, where Rap30 linker also binds (Fig. 2b), is enriched in hydrophobic amino acids (shown in orange in Fig. 3b) and, reciprocally, there are several highly conserved hydrophobic amino acids in the crosslinked Gdown1 region (227–248) (Fig. 3a), suggesting the potential involvement of hydrophobic interactions in the Gdown1-Pol II interaction at this site. In support of this idea for Pol II binding domain I, Gdown1(194–298) fragments in which selected hydrophobic residues were changed to alanine (Fig. 3a) lost Pol II binding (except for the P244A mutant) (Fig. 3f). Surprisingly, the same mutations failed to elicit any obvious deficit in Pol II binding when analyzed in the context of the large C-terminal Gdown1(194–368) fragment (Fig. 3g), although this may simply reflect compensation by Pol II binding region II and/or the C-TIR for general Pol II binding and does not eliminate a role for the indicated binding region I hydrophobic interactions in the inhibitory functions of Gdown1. The detrimental effect of the L303/4A mutation on binding of the Gdown1(194–368) fragment to Pol II (Fig. 3g, lane 9) is consistent with the results discussed earlier and again indicative of a stabilizing effect of C-TIR on Pol II binding region I.

C-TIR acts cooperatively with N-terminal TIR to stabilize the interaction of Pol II binding region I with Pol II

In a further analysis of the large 180-residue Gdown1 N-terminal fragment that alone does not repress transcription but nonetheless contributes significantly to Gdown1 inhibitory activity (Supplementary Fig. 3c), a small N-terminal deletion significantly reduced transcription inhibition by Gdown1 (Fig. 4b) without affecting Gdown1 binding to Pol II (Supplementary Fig. 4a). However, an electrophoretic mobility-shift assay (EMSA) demonstrated that the normal Gdown1 inhibition of PIC assembly was significantly reduced (Supplementary Fig. 4b), suggesting that the Gdown1 N-terminus somehow affects Gdown1-Pol II interactions.

To investigate the mechanism, we sought to identify N-terminal amino acids involved in Gdown1- Pol II interactions. Notably, the Gdown1 cryo-EM density near RPB3 (Fig. 4c), where Gdown1 K39 and K43 were cross-linked to K17 in RPB10 (Fig. 1b and Fig. 4e), overlapped with an acidic amino acid-enriched region in RPB3 (Fig. 4c, d). The presence of highly conserved arginine or lysine residues in the Gdown1 N-terminal region (Fig. 4e) thus suggested a salt-bridge interaction. In support of this prediction, the binding to Pol II of a GST-fused Gdown1(1–96) fragment in which these basic amino acids were mutated to aspartic acid was almost completely lost, whereas mutations of highly conserved acidic amino acids to arginine stabilized the Pol II interaction (Fig. 4f), indicating that these amino acids are important for Pol II interactions. Therefore, we defined this region (1–67) as an N-terminal transcription inhibition region (N-TIR). Although N-TIR is located more than 160 amino acids away from Pol II binding region I (Fig. 4a), cross-links between the two regions (Fig. 1c) indicate their close apposition in Pol II-bound Gdown1. A further analysis showed that K29/30D mutations in a Gdown1(1–244) fragment eliminated its binding to Pol II (Fig. 4g), thereby explaining the loss of transcription inhibition upon the N-terminal deletion (Fig. 4a).

Although roles for N-TIR and C-TIR in stabilizing the Pol II binding region I interaction in Gdown1 repression became evident (Fig. 4a), localizations of these regions on Pol II (especially C-TIR, which did not show any significant binding to Pol II), remained unknown. To determine the locations, we performed cross-linking-based integrative structure modeling[24,25] entailing a four-step procedure (Methods, Supplementary Fig. 5, Fig. 6). This analysis indicated an N-TIR localization near RPB10 and RPB3 where Gdown1 density was detected (Fig. 4h), consistent with the biochemical results (Fig. 4f). Although corresponding cryo-EM density was not observed, the N-terminal region (65–94) was localized to RPB10 and RPB3 and mapped adjacent to Pol II binding region I. Thus, N-TIR interacts with Pol II binding region I through the (65–94) region (Fig. 4h), explaining how N-TIR could stabilize the Gdown1-Pol II interaction. Pol II binding region I covers the RPB2 protrusion surface where two distinct densities overlap well, and Pol II binding region II also localizes to the RPB2 protrusion. The location of C-TIR was identified as an overlapping region between (216–314) and (300–335) (Supplementary Fig. 4c), whereas about two-thirds of the region does not overlap with the density (Fig. 4h). This may explain why C-TIR does not interact with Pol II. Overall, the modeling study revealed that both TIRs are located near Pol II binding regions, thus furnishing evidence for stabilization of Pol II binding region I by TIRs.

Gdown1 nuclear co-localization with Pol II is inversely co-related with transcriptional activation in early fly embryos

Whereas in vitro studies clearly show that Pol II(G) has unique properties relative to Pol II, little is known about the biological roles of Pol II(G) and why higher metazoans need two distinct forms of Pol II to regulate gene transcription. Beyond earlier indications of Gdown1 functions in mammalian cells[10], we sought to establish in vivo functions of Gdown1 in the context of global gene regulation in Drosophila embryogenesis. Initial studies established that Pol II(G), evidenced by reciprocal co-immunoprecipitation of Pol II and Gdown1, is present both in Drosophila embryo nuclear extracts (Fig. 5a) and in derived chromatographic fractions that are distinct from those containing Pol II (data not shown). Gdown1 is present at all life cycle stages and was detected in both nuclear (more abundant) and cytoplasmic (less abundant) fractions at the embryonic stage (Fig. 5b) but appeared predominantly cytoplasmic at the adult stage (Fig.5b, c). Strikingly, Gdown1 co-localizes with Pol II in nuclei at the transcriptionally silent syncytial blastoderm stage (Fig. 5d), but is detected only in the cytoplasm and not in the nuclei that retain Pol II at the later cellular blastoderm stage at which global transcription is initiated (Fig. 5e). Moreover, the pole cells that are transcriptionally silent retain nuclear Gdown1 at stage 5 (Fig. 5f). These data suggest a role for Gdown1, through formation of Pol II(G), in transcriptional repression.

We also demonstrated that whereas homozygous fly embryos carrying *Gdown1* alleles with P element insertions grow to 1st-instar larva, adult flies are not obtained. This requirement for Gdown1 was further confirmed by siRNA-mediated Gdown1 knockdown in the embryo. In related genetic studies, maternal Gdown1 knockout embryos were not obtained, further indicating that Gdown1 plays a critical role in early embryonic development in the fly.

Although we have not yet been able determine which embryonic gene expression events are influenced by Gdown1 knockout, these studies have provided strong evidence that Gdown1 has essential functions *in vivo*.

Discussion

The present study has identified key Pol II binding domains (I and II) and modulatory domains (N-TIR and C-TIR) in metazoan-specific Gdown1 (Fig. 4a), which effects a unique mode of repression of Pol II function. Although Pol II interactions of individual Gdown1 functional domains are not sufficient to inhibit transcription, their inter-connected (cooperative) interactions with Pol II establish robust binding. At the same time, weak interactions of individual domains could potentially facilitate Gdown1 dissociation by Mediator and subsequent transcription.

The C-terminal Pol II binding regions (I and II) are the primary cause of the inhibitory activity of Gdown1 by preventing both TFIIF and TFIIB from binding to Pol II. Among the functional domains in TFIIF, the Rap30 linker has been shown to be essential both for growth in yeast[15] and for transcription initiation in vitro[26]. Moreover, recent cryo-EM studies of the PIC[18] and an initially transcribing complex[27] showed that the Rap30 linker makes contacts with multiple sites, including the RPB2 external domain, the RPB2 protrusion, TFIIB, TBP, and a downstream promoter region. Since these sites are essential

for PIC formation, it was proposed that the linker interactions in the PIC might position other essential TFIIF domains[18]. Hence, the Gdown1 C-terminal interaction with the RPB2 protrusion domain could explain the mutually exclusive interactions of Pol II with TFIIF and Gdown1.

Notably, whereas the Gdown1(227–298) region interaction with the Pol II protrusion domain is stable, the Gdown1(315–368) region interaction appears quite fluid as evidenced by crosslinking to multiple domains in RPB2 and RPB1. Although deletion of this region did not result in a complete loss of Gdown1 inhibitory activity in the in vitro transcription assay, its interactions with the Pol II wall and rudder domain were found to inhibit TFIIB binding to Pol II and could be critical in a cellular context. Since TFIIB plays a pivotal role in PIC formation by direct binding both to Pol II and promoter DNA[6], the Gdown1 interaction could thus further impede Pol II incorporation into the PIC. Also, the Gdown1 interactions with the Pol II clamp loop may interfere with DNA loading to the cleft. Taken together, a role of the Gdown1 C-terminal Pol II binding region in transcription repression is to prevent TFIIF and TFIIB from binding to Pol II, which contributes to the maintenance of Pol II an inactive state in the absence of Mediator.

Whereas Pol II binding region I is the major site for Gdown1 binding to Pol II, the interaction itself does not suffice for the Gdown1 transcription inhibition activity. As mutations in both TIRs can bypass the Mediator requirement for transcriptional activation, TIRs play essential roles in the inhibitory activity. Notably, whereas C-TIR, which consists of only 15 amino acids, does not directly bind to Pol II, point mutations in C-TIR cause a complete loss of repression. Although it remains unclear how C-TIR stabilizes the Gdown1 C-terminus interaction with Pol II, the modeling study predicts that C-TIR (300–314) is localized on the RPB2 protrusion parallel to the Gdown1(216–299) region. Since C-TIR likely possesses substantial secondary structure[28], it may directly associate with the Gdown1(216–299) region to stabilize the C-terminal Pol II interaction. The study also indicates that half of the Gdown1(300–314) region does not overlap with the density on the RBP2 protrusion, which may explain the absence of an independent C-TIR binding activity.

While C-TIR is located near Pol II binding region I, N-TIR, which is positioned at RPB3 and RPB10, is far from this region. However, the Gdown1(65–94) region, which is connected to N-TIR, directly contacts to the Pol II binding region I. This may explain how N-TIR deletion leads to the loss of transcription inhibition. Although N-TIR and C-TIR both show little independent binding to Pol II, their joint stabilization of the Gdown1 C-terminal interaction may account for the robust binding of Gdown1 to Pol II that is resistant to dissociation even by 2 M urea[7].

Since the mutations in N-TIR and C-TIR can relieve the normal Mediator requirement for transcription by Pol II(G), it is possible that Mediator interactions with Pol II disrupt N-TIR and/or C-TIR interactions with Pol II. In this regard, our cryo-EM study revealed at least two Mediator interacting sites on RPB3 and the RPB1 dock domain. While it remains unknown which region of Gdown1 interacts with the RPB1 dock domain, which is a contact site for MED18 in the Mediator head module[27], N-TIR is located around the RPB3 site where the Mediator tail module is likely to be located[29]. Interestingly, the location of Mediator tail

domain subunit Med2 in the yeast PIC complex has been mapped near RPB3[29], where, more intriguingly, N-TIR also interacts (Supplementary Fig. 4d). This result raises the possibility that a Mediator tail domain interaction with RPB3 may lead to a concomitant N-TIR dissociation and de-stabilization of the Gdown1-Pol II interaction, which could be at least part of the mechanism by which Mediator reverses Gdown1-mediated transcription inhibition in vitro. In relation to biological relevance, we observed a joint Gdown1 and Pol II nuclear localization in transcriptionally silent embryos and selective re-localization of Gdown1 to the cytoplasm during zygotic genome activation. These results, the Mediator-dependent reversal of Gdown1 inhibition of Pol II[7], and Pol II(G) localization upstream of transcription start sites[10] suggest a biological role for Gdown1 that maintains Pol II in an inactive, potentially poised, state until an appropriate activation signal is generated.

METHODS

Cell line

HeLa S cell line that stably expresses FLAG-tagged human Gdown1[10] was used for Pol II(G) purification. The cell line was examined and tested negative for mycoplasma contamination.

Purification of Pol II and Pol II(G).

Nuclear extracts were prepared from FLAG-tagged Gdown1 stable cell line[10] and dialyzed in TGE buffer (20 mM Tris-HCl pH 7.9 at 4 °C, 25% Glycerol, 0.1 mM EDTA, 2 mM DTT, 0.5 mM PMSF) containing 0.1 M Ammonium sulfate. The dialyzed nuclear extracts were fractionated on a DE52 column. The 0.3 M ammonium sulfate fraction was subjected to anion exchange chromatography (Hi-trapQ) and eluted with a linear gradient from 0.1 M to 1 M KCl in buffer B. The elution was dialyzed in buffer C containing 0.3 M KCl and was subjected to affinity chromatography using M2-agarose and eluted with $3 \times$ FLAG peptide.

Preparation of recombinant Gdown1 proteins.

Human Gdown1 was cloned into pET21b (C-terminal His $\times$ 6 tag). After transformation, *Escherichia coli* BL21(DE3) RIL cells were grown at 37 °C to *A* 600 nm of 0.6 before protein expression with 1 mM IPTG for 4 h at 25 °C. Subsequent steps were completed at 4 °C unless otherwise noted. Cells were lysed by sonication in Lysis buffer (50 mM HEPES pH 7.6, 500 mM NaCl, 10 mM imidazole, 1 mM PMSF, 10% glycerol,1 mM DTT). Cleared lysate was subjected to affinity chromatography using Ni-NTA agarose (Qiagen), and excess chaperone was removed by washing the resin with a 5 mM ATP and denatured *E. coli* protein in Lysis buffer at room temperature. Protein was eluted with Lysis buffer supplemented with 250 mM imidazole. Elution was exchanged into buffer B (20 mM HEPES pH 7.9, 100 mM KCl, 0.2 mM EDTA, 20% glycerol, 1 mM DTT). Diluted protein was subjected to anion exchange chromatography (Hi-trapQ) and eluted with a linear gradient from 0.1 M to 1 M KCl in buffer B. Gdown1-containing fractions were pooled, and dialyzed in buffer B.

Antibodies.

Antibodies against Gdown1 was raised in rabbits and have been described elsewhere[10]. Antibodies against RPB3, was purchased from Bethyl Laboratories. Antibodies against full-length drosophila Gdown1 was validated by western blot, immunoprecipitation, and immunostaining with purified recombinant drosophila Gdown1 protein, purified drosophila Pol II(G) from drosophila embryo nuclear extracts, and Gdown1 null drosophila embryos.

In vitro transcription and electrophoretic mobility-shift assays.

Pol II, and GTFs were purified as described previously[30]. Bovine Pol II and Pol II(G) were purified as described previously[7]. In vitro transcription with purified components was performed as described[30]. Electrophoretic mobility-shift assays were performed as described[31].

Protein interaction assays.

For the GST pulldown assays, approximately 5 μg of each GST protein was immobilized on glutathione-Sepharose beads and incubated for 2 hr at 4°C with HeLa nuclear extract or the ^{35}S-labeled proteins that were expressed in the TNT expression system (Promega). After washing with Buffer C containing 0.1 M or 0.3M KCl and 0.1% NP40, bound proteins were analyzed by SDS-PAGE and were subjected to immunoblot or autoradiography.

Chemical cross-linking and mass spectrometry.

Five μg of the natively isolated Pol II(G) was chemically cross-linked with 0.5 mM disuccinimidyl suberate (Creative Molecules) for 30 min at 25 °C with constant agitation. The reaction was then quenched with 50 mM ammonia bicarbonate. After disulfide reduction and cysteine alkylation, the cross-linked Pol II(G) complexes were separated on a SDS PAGE gradient gel (4–12%) which was briefly stained by GelCode Blue Stain Reagent (Thermo Fisher) to enable the visualization of the cross-linked complexes. Efficiently cross-linked materials that correspond to gel regions of > 250 kDa were excised and digested in-gel using trypsin [14,32–35]. After peptide extractions, the cross-linked peptides were separated into six different fractions by a self-packed pH10 C18 reverse resin, acidified and analyzed by LC/MS. For cross-link identifications, the purified peptides were dissolved in the sample loading buffer (5% MeOH, 0.2% FA) and analyzed by an Orbitrap Q Exactive Plus mass spectrometer (Thermo Fisher). The instrument was coupled online to an EASY-nLC 1000 System (Thermo Fisher) for chromatography separation of peptides. Peptide mixtures were loaded onto an Easy-Spray column (C18, 3mm particle size, 200 Å pore size, and 50 μm × 15cm, Thermo fisher). The top 8 most abundant ions (with charge stage of 4–7) were selected for fragmentation by HCD. The raw data were searched by pLink. All spectra were manually verified as previously described [14,32,34–36]. The cross-link data was visualized and analyzed by the CX-Circos software.

Cryo-EM analysis.

For grid preparation and data acquisition, 3.5 μL of 0.2 mg/mL purified Pol II(G) complex was applied to the glow-discharged C-flat R2/2 grid coated with a home-made continuous thin layer of carbon. In order to overcome an orientation bias, the grid was treated with 5 μl

of 0.1% (w/v) poly-L-lysine hydrobromide (Polysciences) prior applying the sample. Excess sample was manually blotted and vitrified in liquid ethane. The entire procedure was carried out at 4 °C and 98% relative humidity. Cryo grids were loaded into a Titan Krios transmission electron microscope operating at 300 keV. Images were automatically acquired with Leginon[37] at a nominal magnification of 22,500X (1.31 Å/pixel at the specimen level) using a total dose of ~40 electrons per $Å^2$ and a nominal defocus range of 1.5 – 2.8 μm. A total of 2,710 images were acquired using a Gatan K2 Summit direct electron detector, operated in electron counting mode. Each image was acquired as a 35 frame dose fractionated movie over a 7 s exposure time.

Electron microscopy data processing.

Dose-fractioned movies were aligned using dosefgpu_driftcorr[38] with a frame offset of 7 and a B factor of 1000 $pixels^2$. The Pol II(G) particles were automatically picked from the motion-corrected images using FindEM in the Appion pipeline[39], via the templates derived from 5000 picked particles using DoGpicker. Particles were then extracted using Relion 1.4[40] with a box size of 224 pixels. CTF parameters were estimated using the programs CTFFIND4[41]. A total of 201,527 particles were extracted using a box size of 224 $pixels^2$. The 2D, 3D classification and refinement were performed with Relion 1.4. Two rounds of 2D classification and one round of 3D classification were performed to select the homogenous particles. After 3D classification, one set of 141,619 particles were then submitted to particle-based motion correction and radiation damage weighting, and followed by 3D auto-refinement. All 3D classification and 3D refinements were started from a 60 Å low-pass filtered version of x-ray crystal structure of bovine RNA polymerase II (PDB: 5FLM)[12]. The refinement resulted in an overall structure at a resolution of 4.0 Å based on the gold-standard FSC = 0.143 criteria[42]. Prior to visualization, all density maps were corrected for the modulation transfer function (MTF) of the detector, and then sharpened by applying a negative B-factor that was estimated using automated procedures[43].

In order to improve the quality of the reconstructed map at the N-terminal region of Gdown1, we used the signal-subtracted focus classification and refinement in RELION[44]. We subtracted projections from the Pol II of the reconstruction in the experimental particle images using the relative orientation of each experimental image from the last 3D auto-refinement run of the overall polished particles. The subtracted experimental images were then subjected to the 3D classification with a soft binary mask around the N-terminal region of Gdown1 and no alignments. 35,026 signal subtracted particles from the 3D class with the strongest Gdown1 signals were selected, and the corresponding experimental polished particles were subjected to 3D auto-refinement by continuing with the last angular and translational search parameters from the 3D auto-refinement step of the overall Pol II(G). This approach produced a better quality map around the N-terminal region of Gdown1.The same strategy was also applied for the C-terminal region of Gdown1, but didn't further improve the map which indicates a dynamic interaction between the C-terminal region and Pol II.

To build the Pol II atomic model, the cryo-EM structure of bovine Pol II (PDB: 5FLM) served as the reference. The starting model was placed in the density by rigid-body fitting in

UCSF Chimera[45]. Further model adjustments were done manually using Coot[46]. Consistent with previous observations, the apo Pol II preserves a flexible clamp. We were not able to model the clamp part of the Pol II(G). Focus refinement using a soft mask around the C-terminal region of Gdown1 improved the local density, and we were able to trace the main chain. The map is not good enough to assign the sequence. Refinement of the Pol II and Gdown1 models against the Pol II(G)1 cryo-EM map was done by using the real space refinement function implemented in Phenix[47]. Final models were validated using Molprobity[48] and the FSC of the final model versus the Pol II(G)1 map (Supplementary Fig. 1).

Preparation of drosophila nuclear extracts.

Embryos (0–12h old) were collected from a mass population of Drosophila melanogaster. The nuclear extracts were made as previously described[49].

Immunostaining and confocal imaging.

Embryos were washed, dechorionated, and fixed as previously described[50]. After devitellinization in methanol, embryos were rinsed with PBST (PBS containing 0.2% Tween 20) three times, and incubated with blocking solution (Roche) for 1 h at room temperature. Embryos were then incubated with primary antibody overnight at 4 °C. After washing with PBST three times, embryos were incubated with secondary antibody for 1 h at room temperature, washed with PBST three times, and mounted in Vectashield (Vector Laboratories) for microscopy. Primary antibody was diluted in blocking solution as follows: anti-dGdown1 (Roeder lab) 1:200, anti-CTD phosphor-ser5 (Clone H14, BioLegend) 1:100. Confocal images of immuno-stained embryos were obtained using a LSM 780 laser scanning confocal microscope (Zeiss).

Data availability

The cryo-EM density maps of Pol II(G) has been deposited in the Electron Microscopy Data Bank under accession number EMD-7997. Additional source data are available upon reasonable request.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgements

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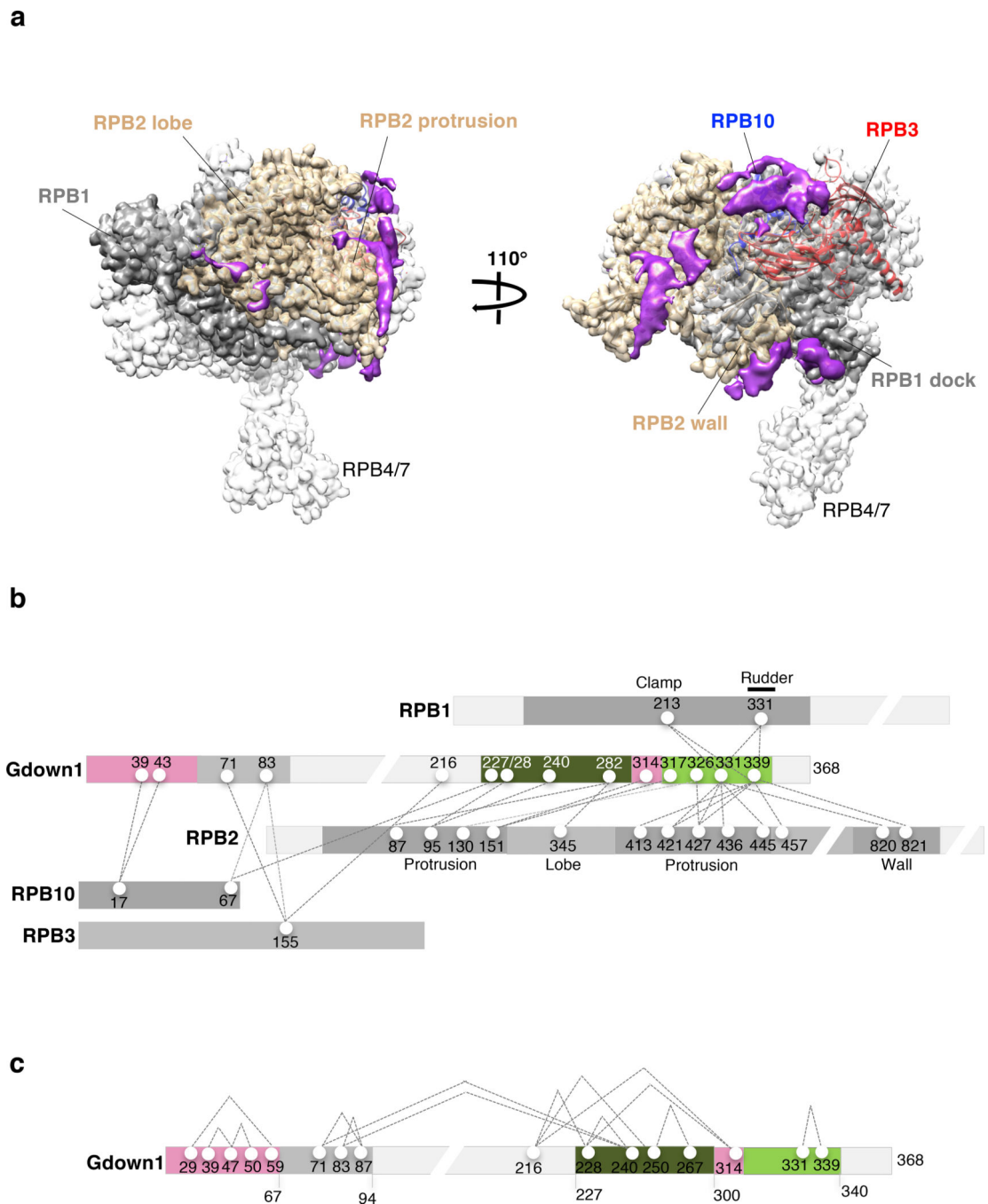
Figure 1. Molecular architecture of Pol II(G) by cryo-EM and CX-MS analyses.
a, Cryo-EM structure of Pol II(G). Non-Pol II (Gdown1) density is shown in purple. An atomic model of human Pol II was obtained by real-space refinement of a model derived from relevant portions of the published model of bovine Pol II (PDB 5FLM). To identify Gdown1 density in the Pol II(G) map, the difference between all stable portions of the Pol II(G) cryo-EM map was calculated. **b,** Diagram of cross-links between Pol II subunits and Gdown1. Purified Pol II(G) was subjected to crosslinking with the amine-specific cross-
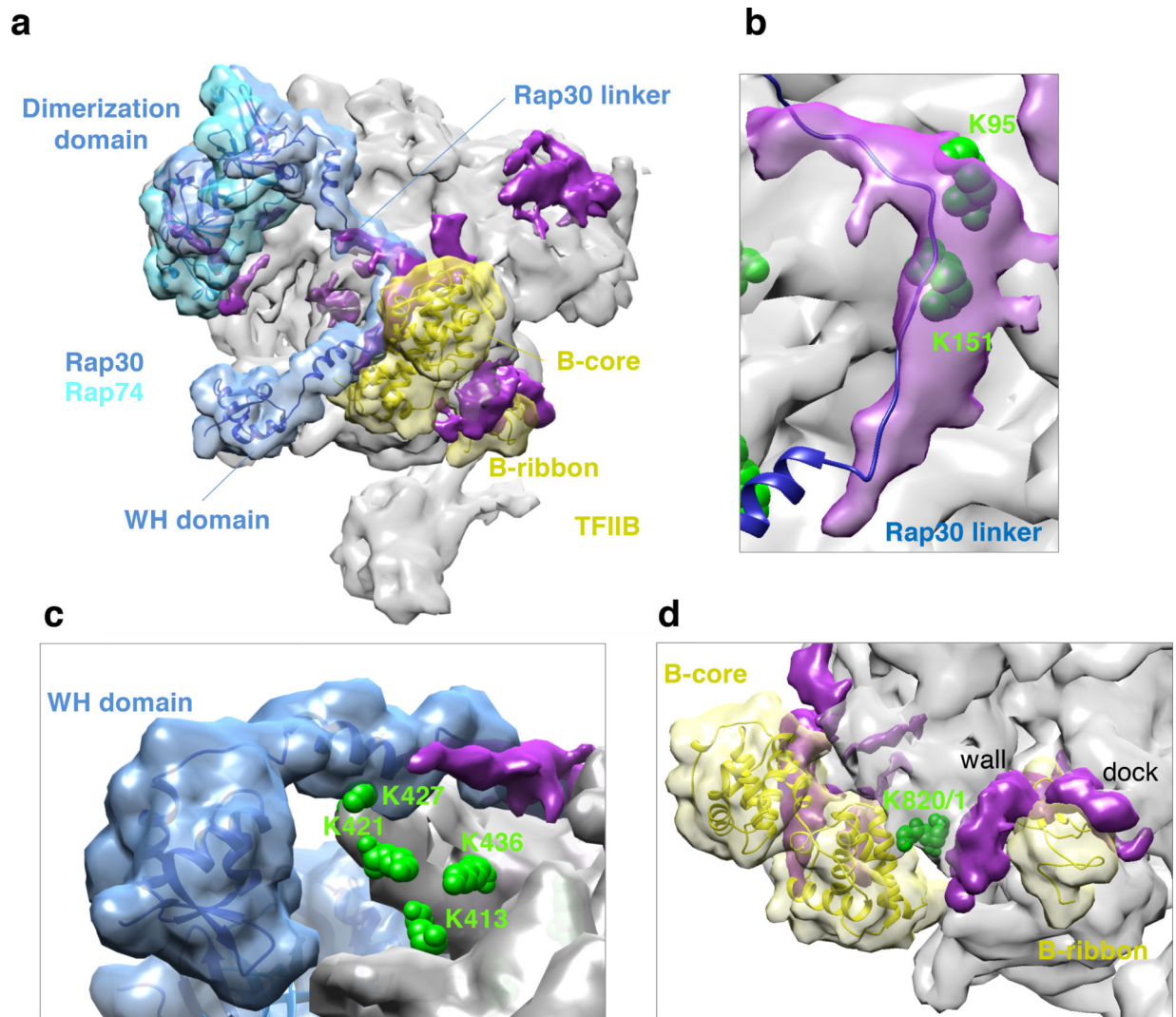
linker, disuccinimidyl suberate (DSS), followed by high-resolution MS. **c,** Intra-molecular cross-links map of Gdown1.

Figure 2. Gdown1 interaction sites overlap TFIIF and TFIIB contact sites on Pol II.
**a,** Positions of TFIIF (Rap74 and Rap30) and TFIIB relative to Gdown1 density (shown in purple) on Pol II. Rap74, Rap30, and TFIIB are shown in light blue, dark blue and yellow, respectively. **b,** Overlap of Gdown1 density on the RPB2 protrusion domain with the Rap 30 linker domain. Two RBP2 lysines (K95 and K151) that crosslink to the C-terminal region of Gdown1 are shown in green. The Rap30 linker domain is shown in dark blue. **c,** Position of the Rap 30 WH domain (shown in dark blue) relative to Gdown1 density (shown in purple). Four RPB2 lysines that cross-link to the C-terminal region of Gdown1 are shown in green. **d,** Positions of TFIIB B-core and B-ribbon domains (shown in yellow) relative to Gdown1 density. Two RPB2 lysines (K820, K821) that cross-link to the C-terminal region of Gdown1 are shown in green.
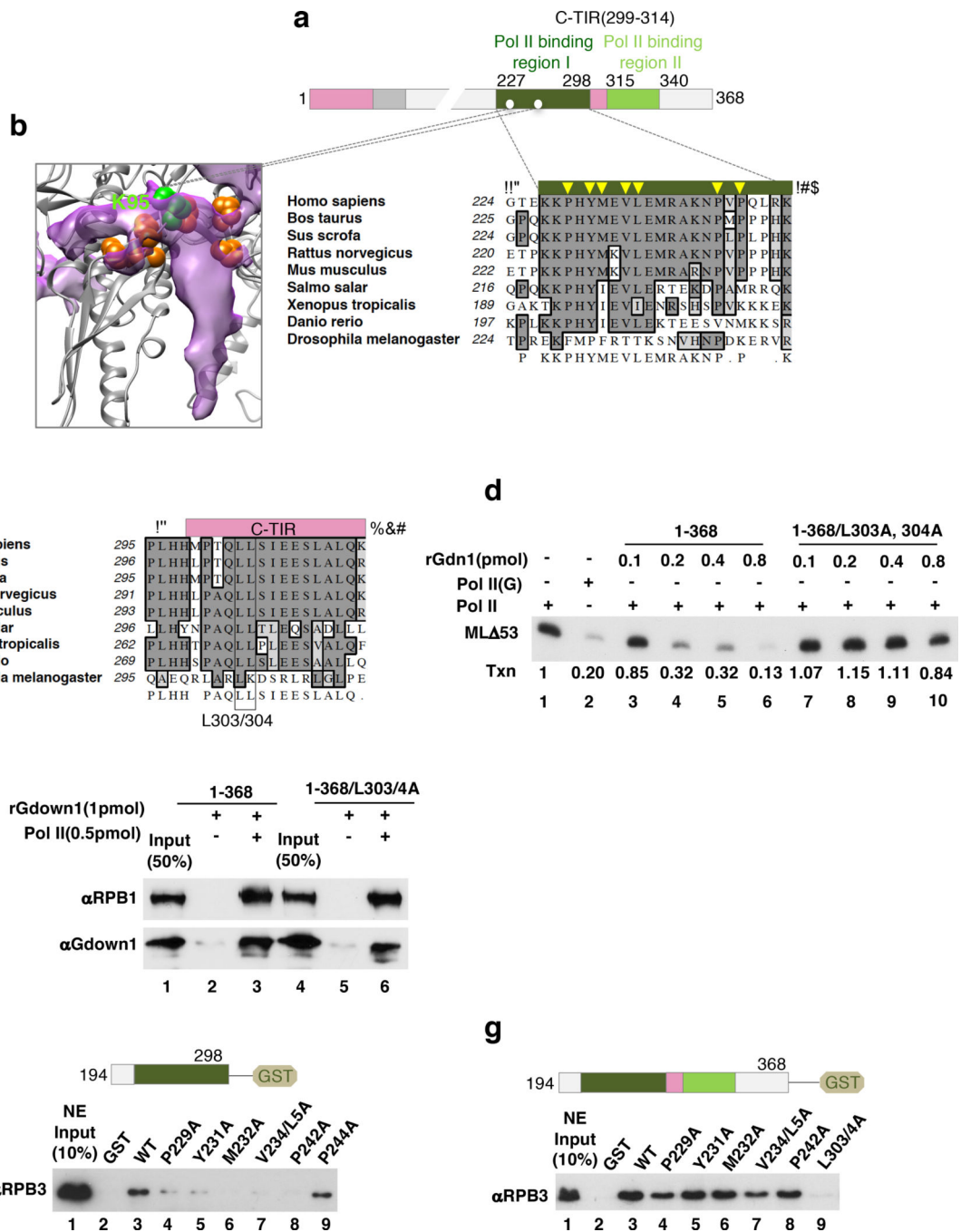
Figure 3. C-TIR stabilizes the interaction of Pol II binding region I with Pol II.
a, Schematic of Gdown1 functional domains. Pol II binding region I (227–298), C-TIR (299–314), and Pol II binding region II (315–340) are shown in dark green, pink, and light green, respectively. Cross-linked residues (K228 and K240) in Pol II binding region I are shown in white circles. Alignment of binding region I sequences from various species. Amino acid residues that were changed to alanine by site-directed mutagenesis are shown in yellow triangles. **b,** RPB2 protrusion domain surrounding Gdown1 density (shown in purple). K95 (shown in green) in the RPB2 protrusion domain (shown in gray ribbon) cross-

links to K228 and K240 in the Gdown1 Pol II binding region I. Hydrophobic amino acids in the protrusion domain are shown in orange. **c,** Alignment of TIR sequences from various species. Amino acid residues that were changed to alanine by site-directed mutagenesis are boxed in black. **d,** In vitro transcription assay with Gdown1 WT (1–368) and L303/4A mutant (1–368/L). Reactions contained 10 ng of TBP, 10 ng of TFIIB, 25 ng of TFIIF, 10 ng of TFIIEα, 5 ng of TFIIEβ, 50 ng of Pol II or Pol II(G) and indicated amounts of His-tagged full-length Gdown1(1–368) or L303/4A mutant. Reactions were incubated at 30°C for 1 h and purified RNA products resolved by PAGE. The experiment was repeated at least three times, with the same results each time. Uncropped image is shown in Supplementary Dataset 1. **e**, Pol II interaction assays. Full-length Gdown1 or L303/4A mutant fragments were incubated with Pol II and subjected to immunoprecipitation with anti-CTD antibody (8W16). Bound proteins were analyzed by immunoblot. The experiment was repeated at least three times, with the same results each time. Uncropped western blot image is shown in Supplementary Dataset 1. **f, g**, Pol II binding assay. Immobilized GST or indicated C-terminally fused GST-Gdown1 proteins with or without indicated mutations were incubated with HeLa nuclear extract in Buffer C containing 0.3 M KCl and 0.1% NP40. After washing, bound proteins were analyzed for bound Pol II by immunoblot. The experiment was repeated at least three times, with the same results each time. Uncropped western blot image is shown in Supplementary Dataset 1.
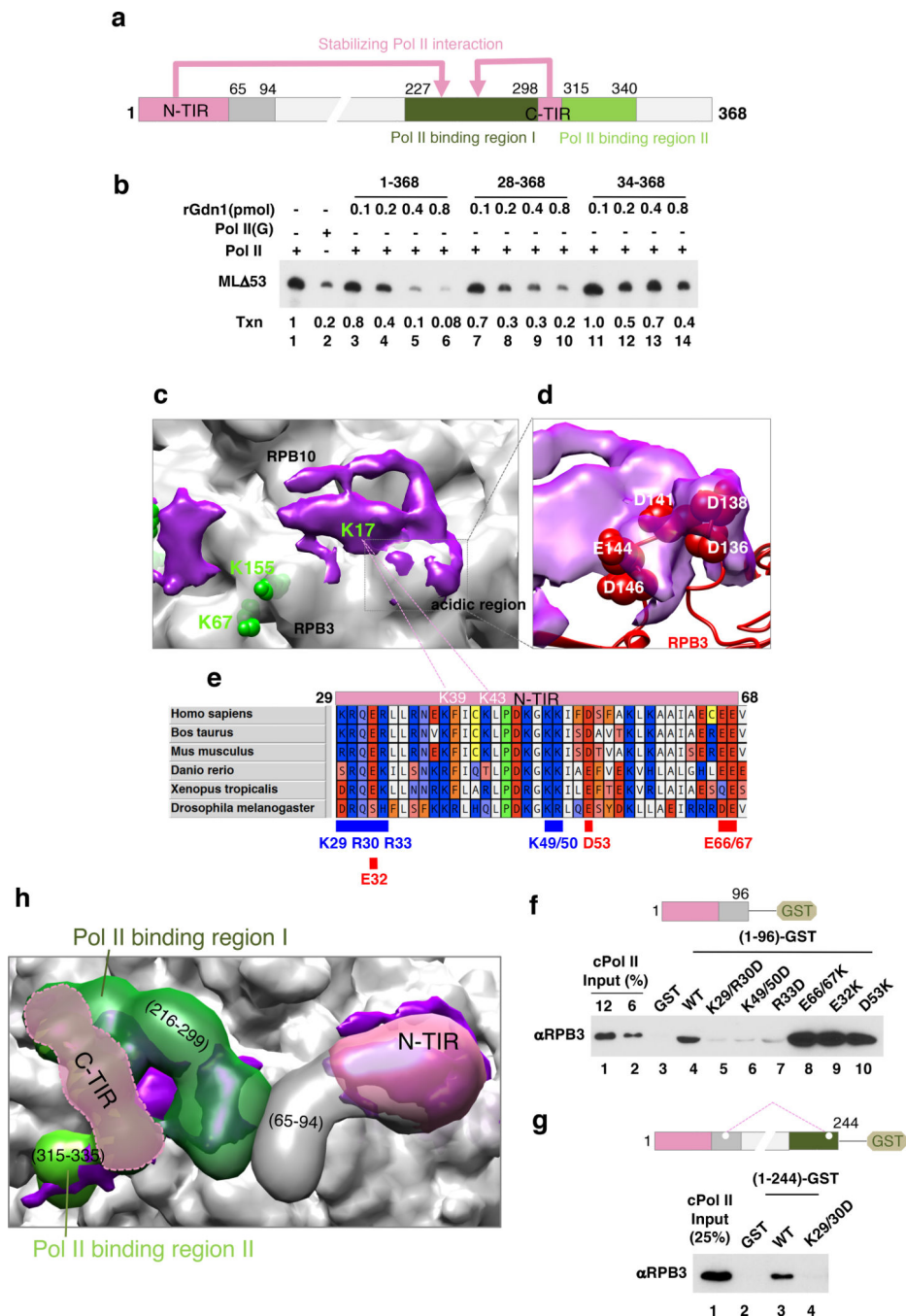
Figure 4. N-TIR and C-TIR stabilize the interaction of Pol II binding region I with Pol II.
a, Schematic of Gdown1 functional domains. **b,** In vitro transcription assay with Gdown1 WT (1–368) and N-terminal deletion mutants. Assay as described in Figure 3d. The experiment was repeated at least three times, with the same results each time. Uncropped image is shown in Supplementary Dataset 1. **c,** Position of Gdown1 density on RPB10 and RPB3. Cross-linked RPB3/10 lysines (Fig. 1b) are shown in green. **d,** Overlap of Gdown1 density (purple) with the acidic region residues (red) in RPB3. **e,** Schematic of cross-linked N-TIR residues and alignment of N-TIR sequences (28–68) from various species. Residues

changed by site-directed mutagenesis from basic to acidic or from acidic to basic are shown in red or blue, respectively. **f, g,** Binding of WT and mutant Gdown1 N-terminal fragments to Pol II. Immobilized GST fusion protein were incubated with purified Pol II in Buffer C containing 0.1 M KCl and 0.1% NP40. After washing, bound Pol II was monitored by immunoblot. The experiment was repeated at least three times, with the same results each time. Uncropped western blot image is shown in Supplementary Dataset 1. **h**, Location of Gdown1 functional domains determined by integrative structure modeling.
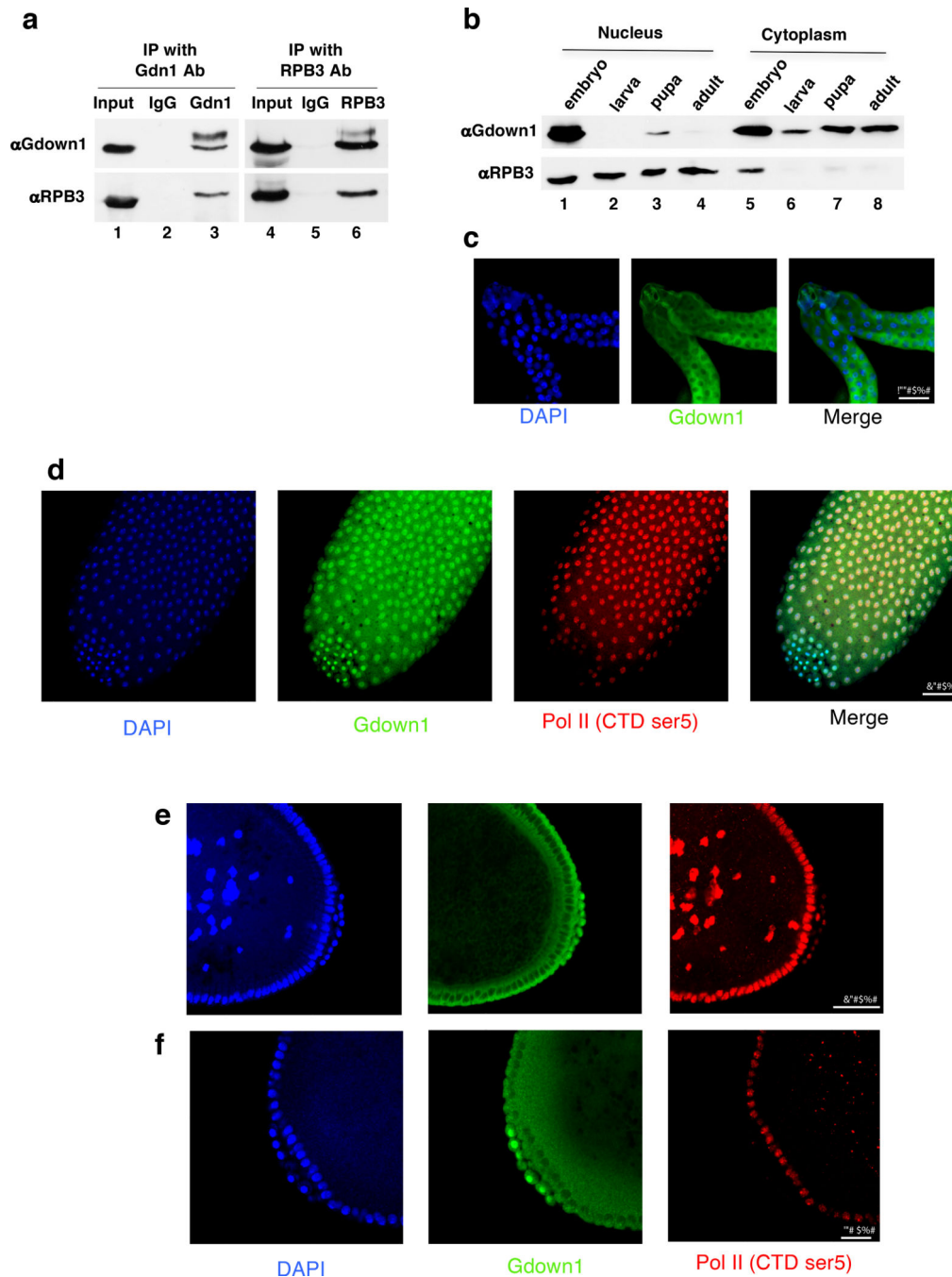
Figure 5. Gdown1 nuclear co-localization with Pol II is inversely co-related to transcriptional activation in early drosophila embryos.

a, Immunoprecipitation with drosophila embryo nuclear extracts. Anti-Gdown1 (lane 3) or anti-RPB3 (lane 6) or IgG (lane 2 and 5) was incubated with fly embryo nuclear extract in Buffer C containing 0.3 M KCl and 0.1% NP40. After washing, bound proteins (Gdown1 and RPB3) was detected by immunoblot. The experiment was repeated at least three times, with the same results each time. Uncropped western blot image is shown in Supplementary Dataset 1. **b**, Gdown1 cellular localization during drosophila development. Nuclear fractions or cytoplasm fractions were prepared from each development stages. Gdown1 and RPB3

were detected by immunoblot. The experiment was repeated at least three times, with the same results each time. Uncropped western blot image is shown in Supplementary Dataset 1. **c,** Gdown1 Immunostaining in salivary gland. **d, e, f**, Immunostaining of Gdown1 and Pol II (CTD ser5) at a syncytial blastoderm stage (d) and a cellular blastoderm stage (e and f).