SUMOylation down-regulates rDNA transcription by repressing expression of upstream-binding factor and proto-oncogene c-Myc

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ABSTRACT

Ribosome biogenesis is critical for proliferating cells and requires the coordinated activities of three eukaryotic RNA polymerases. We recently showed that the small ubiquitin-like modifier (SUMO) system controls the global level of RNA polymerase II (Pol II)-controlled transcription in mammalian cells by regulating cyclin-dependent kinase 9 (CDK9) activity. Here, we present evidence that the SUMO system also plays a critical role in the control of Pol I transcription. Using an siRNA-based knockdown approach, we found that multiple SUMO E3 ligases of the protein inhibitor of activated STAT (PIAS) family are involved in SUMO-mediated repression of rDNA gene transcription. We demonstrate that endogenous SUMO represses rDNA transcription primarily by repressing upstream-binding factor (UBF) and proto-oncogene c-Myc (c-Myc) expression and that ectopic overexpression of SUMO-associated enzymes additionally represses rDNA transcription via c-Myc SUMOylation and its subsequent degradation. The results of our study reveal a critical role of SUMOylation in the control of rDNA transcription, uncover the underlying mechanisms involved, and indicate that the SUMO system coordinates Pol I– and Pol II–mediated transcription in mammalian cells.
Ribosome biogenesis is a major cellular process that occurs in specific nuclear compartments, the nucleoli. A rate-limiting step in this process is ribosomal DNA (rDNA) transcription by RNA polymerase I (Pol I), which accounts for up to 60% of cellular RNA synthesis in proliferating cells (1-3). Pol I transcription is initiated by binding of UBF and selectivity factor 1 (SL1) complex to the rDNA promoter. The UBF/SL1 complex in turn promotes the recruitment of a subpopulation of RRN3 (also known as TIF-1A)-associated Pol I to form a Pol I pre-initiation complex (PIC) at the rDNA promoter (4,5). In mouse and human cells, ~200 rDNA gene copies per haploid genome are distributed in 5 clusters on different chromosomes. Despite the need for a high level of rDNA transcription, typically only a fraction of the rDNA genes is transcriptionally active and the remaining is epigenetically silenced (6,7).

While much progress has been made in study of regulation of Pol I transcription by signaling pathways and epigenetic mechanisms (7-12), the less well understood is the molecular mechanism(s) that coordinates the transcription by Pol I and transcription by Pol II. In this regard, transcription factor c-Myc has been shown to enhance Pol I transcription directly by binding to the rDNA promoter region (13,14) and in a model of granulocyte differentiation c-Myc also stimulates Pol II-dependent transcription of a cohort of factors associated with Pol I transcription (termed “Pol I regulon”) (15,16).

Reversible posttranslational modification with the small ubiquitin-related modifier SUMO (SUMOylation) is involved in essentially all fundamental cellular processes in eukaryotic cells (17,18). SUMOylation is governed by a conserved cascade consisting of an E1-activating enzyme, an E2-conjugating enzyme UBC9, and multiple E3 ligases. By modification of a large number of transcription factors and regulatory proteins, SUMO has a broad role in regulation of transcription by Pol II (19-22). Furthermore, we recently reported that SUMO has a novel role in regulating the global level of Pol II transcription, via inhibiting transcription elongation through CDK9 SUMOylation (23). SUMO has also been shown to regulate the stability of rDNA repeats in S. cerevisiae (24) and rRNA processing and ribosome maturation in metazoan (25-28). However, it is surprisingly not known if SUMO also regulates Pol I transcription.

In this study we investigated if and how SUMO regulates Pol I transcription in mammalian cells. We show that, similar to its role in repressing global Pol II transcription, SUMO also represses Pol I transcription. Furthermore, we present evidence that SUMO represses Pol I transcription indirectly, primarily through control of UBF and c-Myc expression.

Results

SUMO represses Pol I transcription

We recently showed that SUMO represses the global Pol II transcription by SUMOylation of CDK9 and consequently prevents the formation of P-TEFb complex (23) that is required for efficient transcriptional elongation of Pol II-transcribed genes. To determine if SUMO also represses Pol I transcription, we knocked down UBC9, the sole SUMO E2-conjugating enzyme in HEK293T and HeLa cells by two distinct
Repression of rDNA transcription by SUMO

shRNAs (Fig.1A and Fig.1B, left panel). Subsequent analysis of newly transcribed rDNA products, the 47S rRNA precursor, by quantitative RT-PCR (RT-qPCR) as reported (11,12) showed that knockdown of UBC9 led to substantially increased levels of the ribosomal RNA precursor in both cell lines (Fig.1A and Fig.1B, right panel). This observation was further confirmed by knockdown of UBC9 in these cells using a synthetic small interference RNA (siRNA) (Fig.1C and Fig.1D). Thus, knockdown of UBC9 elevated the level of rDNA transcription, suggesting that the Pol I-mediated rDNA transcription is repressed by the endogenous SUMO system. In support of a repressive role of SUMO on Pol I transcription, we found that ectopic overexpression of Flag tagged SUMO E2 enzyme UBC9 inhibited rDNA transcription in HeLa cells and this inhibition is further enhanced by co-expression of a GFP-tagged SUMO1 (Fig.1E). However, no inhibition of rDNA transcription was observed when a SUMO E2-inactive UBC9 mutant (C93S) was expressed alone or together with GFP-SUMO1. Thus, UBC9 suppresses rDNA transcription in a SUMOylation-dependent manner.

To further evaluate the effect of SUMO on rDNA transcription, we used an assay based on incorporation of the uridine analog 5-ethynyluridine (EU) into newly synthesized RNAs (29). As rRNA synthesis accounts for ~60% of total transcription and is restricted in the nucleolus, imaging detection of incorporated EU after a 30 min pulse labeling reaction using a click chemistry (29) revealed unique bright EU spots that showed a perfect co-localization with a nucleolus resident protein nucleolin (NCL), thus reflecting the levels of rDNA transcription (Fig. 1F). Using this assay, we found that knockdown of UBC9 led to increased size and intensity of EU foci (Fig.1G). In contrast, ectopic overexpression of UBC9 markedly diminished the size and intensity of EU foci and this effect was further enhanced by co-expression of GFP-SUMO1 (Fig.1H). Although increased Pol II transcription activity may also partially contribute to increased EU incorporation in UBC9-knockdown cells, marked reduction of EU foci in Flag-UBC9- or Flag-UBC9 plus GFP-SUMO1-expressing cells (Fig. 1H) provided clear evidence that SUMO plays a role in repression of rDNA transcription.

Multiple PIAS family E3 ligases are involved in repression of rDNA transcription

To further confirm a role of SUMO in repression of rDNA transcription, we tested if the PIAS family proteins, the major SUMO E3 ligases (30), are involved in this repression. We used smart pools of siRNAs against various PIAS mRNAs to treat HeLa cells for two days and measured the efficacy of knockdown by RT-qPCR analysis (Fig. S1A). Subsequent RT-qPCR analysis of 47S rRNA precursor showed that knockdown of PIAS1, PIAS2 and PIAS4 resulted in variously increased levels of rDNA transcription (Fig.2A). To substantiate the siRNA-based observation, we designed two different shRNAs targeting PIAS1 and PIAS4, respectively. We validated by RT-qPCR analysis that transfection of HeLa cells with these shRNAs but not shVector (shVec) were able to down-regulated the levels of corresponding PIAS1 and
PIAS4 mRNAs (Fig. S1B). Importantly, we found that knockdown of PIAS1 or PIAS4 by their specific shRNAs led to elevated levels of rDNA transcription in HeLa cells (Fig. 2B). Furthermore, by EU incorporation assay, we observed elevated levels of rDNA transcription in shPIAS1- or shPIAS4 transfected but not control shVect transfected cells (Fig. 2C). Together these loss of function assays suggest that multiple PIAS E3 ligases are likely involved in repression of rDNA genes by SUMO.

In agreement with the loss-of-function results by siRNA/shRNA knockdown, we found that ectopic overexpression of PIAS1, PIAS2β and PIAS4 repressed rDNA transcription as revealed by EU incorporation assay, whereas ectopic overexpression of PIAS2α and PIAS3 failed to do so (Fig. 2D). Repression of rDNA transcription by PIAS1 was dependent on its SUMO E3 ligase activity, as a mutant PIAS1 defective in E3 ligase activity failed to repress rDNA transcription in EU incorporation assay (Fig. 2E). Furthermore, we confirmed by RT-qPCR that ectopic overexpression of PIAS1, but not its E3 ligase mutant, repressed rDNA transcription in HeLa cells (Fig. 2F) and HEK293T cells (Fig. S1C) and this repression was further enhanced by co-expression of GFP-SUMO1. Taken together, both loss and gain of function assays provide evidence that multiple PIAS family E3 ligases are involved in repression of rDNA transcription and experiments with PIAS1 indicate that PIASs are likely to repress rDNA transcription in a SUMO E3 activity-dependent manner.

Repression of rDNA by SUMO correlates with reduced levels of UBF and C-Myc proteins

Thus far, we have demonstrated that the SUMO system represses rDNA transcription. To define the mechanisms underlying SUMO-mediated rDNA transcriptional repression, we first examined if key transcription factors required for Pol I transcription are sumoylated. Intriguingly, our extensive effort failed to detect significant SUMOylation of endogenous NCL, UBF, c-Myc, RRN3, and TAF1B, a subunit of the SL1 complex, even under the conditions with ectopically overexpressed UBC9 plus SUMO1 (Fig. S2A) or PIAS1 plus SUMO1 (Fig. S2B), with which rDNA transcription was markedly suppressed (Fig. 1E and Fig. 2F and Fig. S1C). In addition, we also failed to detect SUMOylation of RPA194, the large subunit of RNA Pol I (Fig. S2A and S2B). The failure in detecting SUMOylation on these proteins was unlikely due to technical problems, because an increase in SUMOylation was detected for endogenous HDAC1 and HDAC2 under the same condition (Fig. S2A).

Although we could not detect SUMOylation on aforementioned Pol I regulatory proteins, we consistently observed a reduced level of UBF and c-Myc proteins upon overexpression of PIAS1 plus SUMO1 (Fig. 3A). Taken together, both loss and gain of function assays provide evidence that multiple PIAS family E3 ligases are involved in repression of rDNA transcription and experiments with PIAS1 indicate that PIASs are likely to repress rDNA transcription in a SUMO E3 activity-dependent manner.

Repression of rDNA by SUMO
analysis revealed that ectopic overexpression of PIAS1 and SUMO1 also led to a substantial reduction of UBF and c-Myc mRNAs, but not that of NCL and TAF1B (Fig. 3B), suggesting that the reduction of UBF and c-Myc proteins upon overexpression of PIAS1 plus SUMO1 is likely due to repression of their transcription. Similarly, we found that ectopic overexpression of PIAS1 plus SUMO1 also reduced the levels of endogenous UBF and c-Myc proteins (Fig. S3A) and the levels of UBF and c-Myc mRNAs (Fig. S3B) in an E3 activity-dependent manner in HEK293T cells. Thus, repression of rDNA transcription by ectopically expressed UBC9 or PIAS1 plus SUMO1 correlates with repression of UBF and c-Myc expression. In contrast, we found that knockdown of UBC9 in HeLa cells led to an elevated level of UBF and c-Myc proteins (Fig. 3C) and elevated UBF and c-Myc mRNAs (Fig. 3D). Similarly, we found that knockdown of UBC9 in HEK293T cells by either shRNA or siRNA also led to an elevated level of UBF and c-Myc proteins (Fig. S3C). Thus, these data reveal a role of SUMO in repression of UBF and c-Myc expression.

We also used immunofluorescence assay to examine if SUMO represses UBF and c-Myc expression. First, we observed that ectopic co-expression of PIAS1 and SUMO1, but not PIAS1m and SUMO1, substantially reduced the levels of c-Myc (Fig. 3E) and UBF proteins (Fig. S4A), but not NCL and RPA194 proteins in HeLa cells (Fig. S4B). In contrast, knockdown of UBC9 by shRNAs resulted in elevated levels of UBF and c-Myc proteins, but not NCL in shUBC9-transfected cells (Fig. 3F). Together, these data suggest that SUMO may repress rDNA transcription through its ability to repress UBF and c-Myc expression. In support of this, we found that UBF and c-Myc are both important for rDNA transcription, as knockdown of either UBF (Fig. S5A) or c-Myc (Fig. S5C) led to reduced rDNA transcription in HeLa cells (Fig. S5B and S5D), consistent with previous reports that UBF and c-Myc are essential for rDNA transcription (13,14,31,32).

**Endogenous SUMO represses rDNA transcription indirectly and mainly through repression of UBF and c-Myc expression**

We recently showed that SUMO globally represses Pol II transcription through SUMOylation of CDK9 (23). SUMOylation of CDK9 blocks its interaction with Cyclin T1/T2 and formation of a functionally active P-TEFb complex, thus controlling global Pol II transcriptional elongation and gene expression (23). As repression of rDNA transcription correlates with reduced expression of UBF and c-Myc, we wished to test if SUMO represses rDNA transcription through repression of UBF and c-Myc expression. To this end, we made use of two engineered HEK293T cell lines that were deleted of endogenous CDK9 genes by CRISPR-Cas9 and stably expressed either a Myc-tagged wild-type CDK9 (CDK9WT) or a SUMO-deficient CDK9 mutant (CDK9K/R) with all lysine residues mutated to arginine (Fig. 4A and reference (23)). We showed previously that the CDK9K/R cell line was viable, whereas straight knockout of CDK9 in HEK293 cells was lethal (23). We found that, while knockdown of UBC9 by
either shRNA or siRNA in control CDK9WT cells led to elevated transcription of UBF and c-Myc, this was not observed in the CDK9K/R cell line (Fig. 4B and 4C). We found by western blot analysis that, while knockdown of UBC9 by either shRNA or siRNA led to elevated levels of UBF and c-Myc proteins, it failed to alter the levels of UBF and c-Myc proteins in the CDK9K/R cells (Fig. 4D and 4E). We further confirmed by immunofluorescent staining assay that knockdown of UBC9 resulted in elevated levels of UBF and c-Myc in CDK9WT but not CDK9K/R cells (Fig. S6). Together these data indicate that in the CDK9K/R cells the transcription of UBF and c-Myc genes and consequently their levels of proteins are not repressed by endogenous SUMO system.

We next examined if rDNA transcription is repressed by SUMO in CDK9K/R cells. Importantly, we found that while knockdown of UBC9 by either shRNA or siRNA led to an elevated level of rDNA transcription in the CDK9WT cells, it failed to elevate rDNA transcription in the CDK9K/R cells (Fig. 4F and 4G). By EU incorporation assay, we confirmed that knockdown of UBC9 led to an elevated level of rDNA transcription in CDK9WT but not in CDK9K/R cells (Fig. 4H). Together, these data indicate that the rDNA transcription in CDK9K/R cells is not repressed by endogenous SUMO, indicating that endogenous SUMO represses rDNA transcription primarily through its ability to repress transcription of UBF and c-Myc via CDK9 SUMOylation.

Ectopic overexpression of SUMO system components represses rDNA transcription through SUMOylation-induced c-Myc degradation

The above results suggest that endogenous SUMO represses rDNA transcription primarily in a CDK9 SUMOylation-dependent manner. To explore additional mechanisms that SUMO may repress rDNA transcription, we tested if ectopic overexpression of the SUMO system is able to repress rDNA transcription in the CDK9K/R cells. RT-qPCR analysis revealed that overexpression of PIAS1 and SUMO1 was able to repress rDNA transcription in the CDK9K/R cells, albeit to a lesser extent compared to that in the CDK9WT cells (Fig. 5A, upper panel). We confirmed by EU incorporation assay that overexpression of PIAS1 and SUMO1 was able to repress rDNA transcription in both CDK9WT and CDK9K/R cells (Fig. S7A). However, overexpression of PIAS1 and SUMO1 did not significantly affect the transcription of c-Myc (Fig. 5A, lower panel) and UBF in the CDK9K/R cells (Fig. S7B), in agreement with their repression by SUMO being dependent on CDK9 SUMOylation.

To understand how overexpression of PIAS1 and SUMO1 repressed rDNA transcription in the CDK9K/R cells, we performed western blot analysis of Pol I regulatory proteins. We found that, similar to the CDK9WT cells, ectopic overexpression of PIAS1 plus SUMO1 in CDK9K/R cells resulted in downregulation of c-Myc, whereas UBF was downregulated only in the CDK9WT but not CDK9K/R cells (Fig.5B). Consistent with the previous results, no change in NCL, RRN3, TAF1B and RPA194 was observed (Fig.5B). We
confirmed by immunostaining assay that ectopic overexpression of PIAS1 and SUMO led to a reduced level of c-Myc proteins not only in the CDK9WT but also CDK9K/R cells (Fig. 5C), whereas downregulation of UBF was only observed in the CDK9WT cells (Fig. S6C). Thus, ectopic overexpression of SUMO system components is able to repress rDNA transcription in the CDK9 SUMOylation-resistant cells, possibly through downregulation of c-Myc proteins.

It was reported that SUMOylation of c-Myc can lead to subsequent ubiquitination and proteasome degradation (33,34). We detected c-Myc SUMOylation when HeLa cells were overexpressed with PIAS1 and SUMO1 and ubiquitin-dependent protein degradation was blocked by MG132 (Fig. 5D). Thus, consistent with previous studies (33,34), ectopically expressed SUMO is likely to reduce the level of c-Myc proteins through SUMOylation-induced ubiquitin-dependent degradation. In support of this, we found that ectopic overexpression of PIAS1 and SUMO1 reduced the half-life of c-Myc proteins in CDK9K/R cells (Fig. S8).

To test if reduced c-Myc proteins were responsible for reduced rDNA transcription in the CDK9K/R cells, we tested if ectopic overexpression of c-Myc could relieve PIAS1 and SUMO1-induced rDNA repression. We observed that ectopic overexpression of Myc-tagged c-Myc proteins (Fig. 5E, left panel) was able to abrogate rDNA repression instigated by ectopically overexpressed PIAS1 and SUMO1 in the CDK9K/R cells (Fig. 5E, right panel). Thus, ectopic overexpression of the SUMO system components can additionally repress rDNA transcription through a SUMOylation-mediated degradation of c-Myc proteins.

**Discussion**

In this study we unravel a critical role for SUMO in regulation of rDNA transcription. Knockdown of SUMO E2 enzyme UBC9 or E3 PIAS proteins in various cells significantly elevated rDNA transcription, whereas ectopic overexpression of SUMO system components can additionally repress rDNA transcription. We present evidence that the endogenous SUMO system represses rDNA transcription primarily through regulating the expression of transcription factors UBF and c-Myc, whereas ectopic overexpression of SUMO system components can additionally repress rDNA transcription via SUMOylation-induced c-Myc degradation.

In principle, SUMO is a dynamic modification and regulates substrate proteins through its effect on protein-protein interaction, subcellular localization, activity and stability (17,18). As transcription factors and coregulators are among the most frequently identified and characterized SUMO substrates, it is not surprising that SUMO has been shown to play a broad role in regulation of transcription by Pol II (20,21,35,36). Interestingly, SUMO in general is involved in transcriptional repression and multiple mechanisms have been proposed to explain the roles of SUMOylation in transcriptional repression (21,37). We recently uncovered a novel role of SUMO in repression of the global level of transcription via control of P-TEFb complex formation by SUMOylation of CDK9 (23). This finding provides a
novel mechanism for transcriptional repression by SUMO. In contrast to the well-recognized role of SUMO in regulation of Pol II transcription and to the best of our knowledge, it is surprisingly not known prior to this study if SUMO regulates rDNA transcription by Pol I. As knockdown of UBC9 or multiple PIAS proteins all led to increased transcription of rDNA genes (Fig. 1 and Fig. 2), SUMO has a role in repression of Pol I transcription. In this regard, the EU incorporation assay, although not entirely quantitative, provides a sensitive measurement for the levels of newly synthesized rRNAs in nucleoli. As rDNA transcription accounts for up to 60% RNA synthesis and is restricted to nucleoli compartments, our findings that knockdown or overexpression of SUMO system components led to elevated or diminished EU foci staining in nucleoli (Fig.1 and Fig.2) provide compelling evidence for repression of rDNA transcription by SUMO.

In search of the potential mechanisms by which SUMO represses rDNA transcription, we initially focused our attention on SUMOylation of key transcription factors required for rDNA transcription. However, no significant SUMOylation of endogenous UBF, NCL, RRN3, TAF1B and c-Myc was observed under the regular culture condition, despite the previous reports on SUMOylation of nucleolin (38) and c-Myc proteins (33), both required for efficient rDNA transcription (13,14,39). Even with overexpression of UBC9 or PIAS1 and SUMO1, we could not observe significant SUMOylation on these proteins (Fig. S1). Instead, we observed elevated or reduced levels of UBF and c-Myc proteins and mRNAs upon UBC9 knockdown or overexpression of the SUMO system components, respectively (Fig. 1, 2 and 3). These observations point to an alternative mechanism that SUMO represses rDNA transcription indirectly through regulation of UBF and c-Myc transcription. In support of this idea, we find that rDNA transcription is no longer repressed by the endogenous SUMO system in the CDK9K/R cell line in which SUMO does not repress UBF and c-Myc transcription (Fig. 4). Thus, endogenous SUMO most likely represses rDNA transcription through repression of UBF and c-Myc transcription, although we could not exclude the possibility that SUMOylation of the Pol I regulatory proteins may also partially contribute to repression of rDNA transcription.

While our study suggests that endogenous SUMO represses rDNA transcription via repression of UBF and c-Myc expression, we show that ectopic overexpression of the SUMO system components can repress rDNA transcription through SUMOylation-induced degradation of c-Myc proteins (Fig. 5), in agreement with previous studies (33,34). Indeed, c-Myc SUMOylation can be detected upon overexpression of PIAS1 and SUMO1 and in the presence of the proteasome inhibitor MG132 (Fig. 5D).

In sum, our study demonstrates that, similar to transcription by Pol II, rDNA transcription by Pol I is also repressed by SUMO. We uncover two mechanisms for SUMO-mediated transcriptional repression of rDNA genes, an indirect one by repression of UBF and c-Myc transcription via CDK9 SUMOylation that appears to be the major repression
force by the endogenous SUMO system and the other one by SUMO-mediated c-Myc degradation that may dominate under an overexpression condition. However, these two mechanisms are not mutually exclusive and may act together to control rDNA transcription and cell proliferation. Furthermore, both mechanisms converge on the control of c-Myc protein levels, in well agreement with c-Myc as a master regulator of rDNA transcription and cell proliferation. Given the critical role of c-Myc in regulating global as well as specific programs of Pol II transcription (15,40,41), our study suggests that SUMO can coordinate Pol I and Pol II transcription by controlling c-Myc expression. In light of reported regulation of Pol III transcription by SUMO (42,43), it is tempting to propose that SUMO has a role in coordinating transcription by all three RNA polymerases.

**Experimental procedures**

**Cell lines, plasmids and antibodies**

HeLa and HEK293T cells were cultured in Dulbecco’s Modified Eagle’s medium (GIBCO, Thermo Fisher, Waltham, MA, USA) with 10% fetal bovine serum (GIBCO) in a humidified incubator at 37 °C with 5% CO2. Plasmids encoding UBC9, UBC9m (C93S), SUMO1, CDK9 and CDK9K/R mutant were constructed as previously described (23). The CDK9 knockout cell lines with expression of Myc-CDK9 or Myc-CDK9K/R mutant were generated as described (23). The plasmids for pCDNA3.0-HA-PIAS1m(C346SC351SC356S) was derived from pCDNA3.0-HA-PIAS1 by site-directed mutagenesis. Various PIAS expression constructs were generated by sub-cloning the corresponding coding sequences into the pCDNA3.0-HA vector. The antibodies used in this study were listed as follows: rabbit anti-UBC9 (CST#4918, dilution 1:5000 for WB), rabbit anti-c-Myc (abcam#ab32072, dilution 1:5000 for WB and 1:1000 for IF, 0.5 μg for IP), mouse anti-UBF (santa cruz#sc-13125, dilution 1:1000 for WB and 1:100 for IF), rabbit anti-NCL (homemade Polyclonal antibody, dilution 1:10000 for WB and 1:2000 for IF), anti-RRN3 (abcam#ab112052, dilution 1:5000 for WB, rabbit anti-TAF1B (absci#AB44736, dilution 1:1000 for WB), mouse anti-RPA194 (santa cruz#sc-48385, dilution 1:1000 for WB and 1:100 for IF), rabbit anti-CDK9 (Abclonal#A1564 dilution 1:1000 for WB), rabbit anti-HDAC1 (Abclonal#A0238, dilution 1:5000 for WB), rabbit anti-HDAC2 (Abclonal#A2084, dilution 1:5000 for WB), mouse anti-HA (Abmart#M2003, dilution 1:5000 for WB and 1:1000 for IF), rabbit anti-HA (santa cruz#sc-805 dilution 1:1000 for WB and 1:200 for IF), mouse anti-GFP (Abmart#M2004L dilution 1:5000 for WB and 1:1000 for IF), mouse anti-Flag (Sigma#F7425 dilution 1:5000 for WB and 1:1000 for IF) and mouse anti-β-Actin (Sigma#A5316, dilution 1:5000 for WB).

**Transfections, western blotting analysis, immunoprecipitation and immunofluorescence staining**

DNA and siRNA transient transfection were performed using lipofectamine 2000 (Invitrogen) according to manufacturer's instructions. Western blot, immunoprecipitation and immunofluorescence staining were performed essentially as described (23), using the antibodies as indicated. Images
were obtained at 40X magnification using Olympus IX73P2F microscope and processed using Image J. Images in Figure 1F, Figure 2B and Figure 2C, were taken with a Leica DM4000BLED microscope and processed using Image J.

**Knockdown with siRNAs or shRNAs and quantitative RT-PCR**

The sequence information for siRNAs targeting human UBC9, PIAS1, PIAS2, PIAS3 and PIAS4 were provided in Supplementary Information, Table S1. All siRNAs were synthesized by Genepharma. The sequences for shRNA targeting human UBC9, c-Myc and UBF are also listed in Supplementary Information, Table S1. The vector for shRNAs was pLKO.1. For RT-qPCR analysis, an equal number of cells for each sample was collected and preparation of total RNA was carried out using the RNAiso Plus kit (Takara#D9108A). All cDNAs were prepared using the TransScript One-Step gDNA Removal and cDNA Synthesis SuperMix (TransGen Biotech#AT311) according to manufacturer’s instructions. Quantitative PCR analysis was performed using CFX96 Touch Real-Time PCR Detection System (Bio-RAD). The following cycle conditions were applied for PCR analysis: initial denaturation at 94 °C for 10 min, followed by 40 cycles at 94°C for 20s, 60°C for 20s, and 72°C for 20s. The delta Ct (ΔCt) obtained was used to find the relative expression of genes according to the formula: relative expression n = 2-ΔΔCt, where ΔΔCt= ΔCt of respective genes in experimental groups – ΔCt of the same genes in control group. GAPDH was used as an internal control. GraphPad Prism 8 software was used for plotting. The primer sequences used in the RT-qPCR are listed in Supplementary Information, Table S1.

**EU labeling assay**

The EU incorporation assay was carried out essentially as described (29). Briefly, 48 hours after initial transfection, EU was added to the complete culture medium at a final concentration of 1 mM and cells were incubated at 37 °C for 30 min. After EU labeling, cells were washed with PBS and fixed in 125 mM Pipes, pH 6.8, 10 mM EGTA, 1 mM magnesium chloride, 0.2% Triton X-100 and 3.7% formaldehyde for 30 min at room temperature. The cells were then washed with TBS and stained with 10 μM Alexa594-azide.

**Data analysis**

Statistical analyses were performed using IBM SPSS statistics 23. Data are presented as mean of 2 or 3 independent experiments. Statistical relevance was determined using the unpaired Student’s test for RT-qPCR. The error bars represent ± SD. Differences of P < 0.05 were considered significant. *0.01 < P < 0.05; ** P < 0.01.

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Author contributions: YP, ZW, ZW, FY, JL, and J.W. designed the study. YP, ZW, ZW, and FY carried out the experiments and analyzed the data. JL, and JW supervised experiments. YP, and J.W. wrote the manuscript. All authors read and approved the final version of the manuscript.

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FIGURE LEGENDS
Figure 1. rDNA transcription is repressed by endogenous SUMO system. (A and B) HEK293T cells (A) or HeLa cells (B) were transfected with two different shRNAs targeting UBC9, and western blot and RT-qPCR analyses were performed to detect the levels of UBC9 proteins (left panel) and the relative levels of 47S rRNA precursor (right panel), respectively. (C and D) HEK293T cells (C) and HeLa cells (D) were treated with siRNA targeting UBC9, and western blot and RT-qPCR analyses were performed
to detect the levels of UBC9 proteins (left panel) and the relative levels of 47S rRNA precursor (right panel), respectively. Note the level of pre-rRNA in scramble control siRNA (NC)-transfected cells was set as 1. (E) Ectopic overexpression of UBC9 and SUMO1 repressed rDNA transcription. HeLa cells were transfected with or without Flag-UBC9 and/or GFP-SUMO1 as indicated. Expression of UBC9 and SUMO1 was confirmed by western blot analysis (upper panel). The levels of rDNA transcription were determined by RT-qPCR. Note that western blot with anti-GFP antibody revealed that co-transfection of Flag-UBC9 but not Flag-UBC9m(C93S) promoted global SUMOylation. (F) Fluorescent imaging showing complete co-localization of EU staining with nucleolin. HeLa cells were pulse labeled with ethyneluridine for 30 min and processed for detection of RNA-incorporated EU by click chemistry and nucleolin by immunostaining. Scale bar, 20 μm. (G) The EU incorporation assay showing a significantly increased EU incorporation in shUBC9-transfected cells (marked by arrows) in comparison to untransfected control cells. HeLa cells were transfected with two different shRNAs against UBC9 as indicated. Two days after transfection, the cells were pulse labeled with EU for 30 min and then processed for imaging of EU. Scale bar, 20 μm. (H) EU incorporation assay showing the effect of ectopic overexpression of UBC9 and/or SUMO1 on rDNA transcription. Note that the EU incorporation was drastically repressed in UBC9 and UBC9 plus SUMO1-transfected HeLa cells (marked by arrows). Scale bar, 20 μm. All western blots were performed at least three times and data were highly reproducible. The immunofluorescent staining experiments were also repeated at least three times, with the rate of transfected cells with increased or reduced levels of EU incorporation counted from all 3+ representative experiment (mean ± SD).

Figure 2. Multiple PIAS family proteins are involved in repression of rDNA transcription. (A) Effect of knockdown of PIAS family proteins on rDNA transcription. Hela cells were transfected with siRNA against each of the PIAS family proteins and two days after transfection the relative levels of 47S rRNA precursor were measured by RT-qPCR. (B) Effect of knockdown of PIAS1 and PIAS4 by specific shRNAs on rDNA transcription. Hela cells were transfected with either control shVec or two different PIAS1 or PIAS4-specific shRNAs. Two days after transfection the relative levels of 47S rRNA precursor were measured by RT-qPCR. (C) The EU incorporation assay showing that knockdown of PIAS1 or PIAS4 in HeLa cells all led to increased rRNA synthesis. The shRNA-transfected cells were marked by arrows. Also shown on the right are the rates of transfected cells with increased levels of EU incorporation counted from 3+ representative experiment (mean ± SD). (D) The EU incorporation assay showing the effect of ectopic overexpression of various PIAS proteins with or without SUMO1 on rDNA transcription in HeLa cells. Transfected cells were marked by arrows. Note that PIAS1, PIAS2b and PIAS4 were able to repress rDNA transcription when they were ectopically overexpressed. Also shown on the right are the rates of transfected cells with decreased levels of EU incorporation counted from 3+ representative experiment (mean ± SD). Scale bar, 20 μm. (E) The EU incorporation assay showing that ectopically expressed PIAS1 repressed rDNA transcription in HeLa
cells in an E3 ligase activity-dependent manner. Transfected cells were marked by arrows. Also shown on the right are the rates of transfected cells with decreased levels of EU incorporation counted from 3+ representative experiment (mean ± SD). Scale bar, 20 μm. (F) RT-qPCR analyses showing that ectopically expressed PIAS1 repressed rDNA transcription in HeLa cells in an E3 activity-dependent manner and the repression was augmented by co-expressed SUMO1.

Figure 3. Repression of rDNA transcription by SUMO correlates with transcriptional repression of UBF and c-Myc. (A) Western blot analysis showing the effect of ectopically overexpressed PIAS1 and/or SUMO on a panel of Pol I core transcription factors, regulators and subunit RPA194. (B) RT-qPCR analyses showing the effect of ectopically overexpressed PIAS1 and/or SUMO on the relative levels of mRNAs encoding Pol I core transcription factors and regulators. (C) Western blot analyses showing the effect of knockdown of UBC9 in HeLa cells by using either shRNA or siRNA on a panel of Pol I core transcription factors, regulators and subunit RPA194. (D) RT-qPCR analyses showing the effect of knockdown of UBC9 on the relative levels of mRNAs encoding Pol I core transcription factors and regulators. The level of each mRNA in shVec-transfected cells was set as 1. (E) Immunofluorescent staining assay showing ectopically expressed PIAS1 proteins downregulated the levels of c-Myc proteins in an E3 activity-dependent manner in HeLa cells. Also shown on the right are the rates of transfected cells with decreased levels of c-Myc proteins based on 3+ representative experiment (mean ± SD). Scale bar, 20 μm. (F) Immunofluorescent staining assay showing elevated levels of UBF and c-Myc proteins, but not NCL, in shUBC9-transfected HeLa cells. The rates of shUBC9-transfected cells with increased levels of UBF, NCL or c-Myc were shown on the right, based on 3 representative experiments. Scale bar, 20 μm. All western blots were performed at least three times and data were highly reproducible.

Figure 4. rDNA transcription is not repressed by endogenous SUMO in cells deficient of CDK9 SUMOylation. (A) Characterization of CDK9WT and CDK9K/R mutant cell lines. These HEK293T cell lines, with both endogenous CDK9 genes disrupted by CRISPR-Cas9, expressed either Myc-tagged wild-type CDK9 (CDK9WT) or a mutant CDK9 with all lysine residues converted to arginines (CDK9K/R). (B) RT-qPCR analysis showing the effect of knockdown of UBC9 by shRNAs on the relative levels of UBF, c-Myc and NCL mRNAs in both CDK9WT and CDK9K/R cells. Note the level of mRNA for each protein in shVec-transfected cells was set as 1. (C) RT-qPCR analysis showing the effect of knockdown of UBC9 by siRNA on the relative levels of UBF, c-Myc and NCL mRNAs in both CDK9WT and CDK9K/R cells. Note the level of mRNA for each protein in scramble control siRNA-transfected cells was set as 1. (D) Western blot analysis showing the effect of knockdown of UBC9 by shRNAs on the levels of c-Myc, UBF and NCL proteins in both CDK9WT and CDK9K/R cells. (E) Western blot analyses showing the effect of knockdown of UBC9 by siRNA on the levels of c-Myc, UBF and NCL in both CDK9WT and CDK9K/R cells.
cells. (F) Western blot and RT-qPCR analyses showing that knockdown of UBC9 by shRNAs led to elevated Pol I transcription in CDK9WT but not in CDK9K/R cells. Note the level of pre-RNA in shVec-transfected cells was set as 1. (G) Western blot and RT-qPCR analyses showing that knockdown of UBC9 by siRNAs led to elevated Pol I transcription in CDK9WT but not in CDK9K/R cells. Note the level of pre-RNA in control scramble siRNA-transfected cells was set as 1. (H) The EU incorporation assay showing that knockdown of UBC9 by transfected shRNAs elevated the level of rDNA transcription in CDK9WT, but not in CDK9K/R cells. Transfected cells were marked by arrows. The rates of cells with increased EU incorporation vs counted transfected cells were shown on the right, based on 3 representative experiments. Scale bar, 20 μm.

Figure 5. Ectopic overexpression of SUMO system components can repress rDNA transcription by downregulation of c-Myc through SUMO-induced c-Myc degradation. (A) RT-qPCR analysis showing that ectopic overexpression of PIAS1 and SUMO1 was able to repress rDNA transcription in both CDK9WT and CDK9K/R cells. Note that ectopic overexpression of PIAS1 and SUMO1 repressed c-Myc expression in CDK9WT but not in CDK9K/R cells. (B) Western blot analysis showing that ectopic overexpression of PIAS1 and SUMO1 downregulated the levels of c-Myc proteins in both CDK9WT and CDK9K/R cells. Note the levels of UBF proteins were downregulated in CDK9WT but not in CDK9K/R cells. (C) Immunofluorescent staining assay showing that ectopic overexpression of PIAS1 and SUMO1 downregulated the levels of c-Myc proteins in both CDK9WT and CDK9K/R cells. The rates of cells with reduced c-Myc vs counted transfected cells were shown on the right, based on analysis of 3 representative results. Scale bar, 20 μm. (D) Western blot analysis showing addition of MG132 was able to block c-Myc downregulation induced by ectopic co-expression of PIAS1 and SUMO1 and allowed the detection of sumoylated/ubiquitinated c-Myc proteins. (E) Ectopic overexpression of c-Myc abrogated PIAS1 and SUMO1-induced rDNA repression in CDK9K/R cells. The CDK9K/R cells were transfected with PIAS1/SUMO1, together with or without Myc-tagged c-Myc, as indicated. Two days after transfection the cells were collected for western blot analysis (left panel) or RT-qPCR analysis of the 47S rRNA precursor (right panel).
Figure 1

A) HEK293T

B) HeLa

C) HEK293T

D) HeLa

E) HeLa WCL

F) NCL, EU, Merge, DAPI

G) shRNA, EU, Merge, DAPI

H) GFP, EU, Merge, DAPI

Figure 1
Figure 2
Figure 3
Figure 4

(A) HEK293 CDK9WT CDK9K/R CDK9K/R = no SUMO
WB: CDK9
Myc-CDK9
Endo-CDK9
Actin

(B) UBF

(C) UBF

(C-Myc

(NCL

(D) CDK9WT CDK9K/R

+ -

- + shVector

- + shUBC9#1+2

UBC9

UBF

NCL

Actin

(F) CDK9WT CDK9K/R

(shVector shUBC9#1+2 shVector shUBC9#1+2

UBC9

UBC9

Actin

(G) CDK9WT CDK9K/R

NC siUBC9 NC siUBC9

UBC9

UBF

NCL

Actin

(H) CDK9WT

shRNA EU Merge DAPI

shU BC9 #1

0.71 ± 0.06

0.83 ± 0.15

0.03 ± 0.03

0.16 ± 0.01

CDK9K/R

shRNA EU Merge DAPI

shU BC9 #1

0.71 ± 0.06

0.83 ± 0.15

0.03 ± 0.03

0.16 ± 0.01

Figure 4
Figure 5
SUMOylation down-regulates rDNA transcription by repressing expression of upstream-binding factor and proto-oncogene c-Myc
Yu Peng, Zhenxing Wang, Zhiqiang Wang, Fang Yu, Jiwen Li and Jiemin Wong

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