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DNA Topoisomerase 1α Promotes Transcriptional Silencing of Transposable Elements through DNA Methylation and Histone Lysine 9 Dimethylation in Arabidopsis

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Abstract

RNA-directed DNA methylation (RdDM) and histone H3 lysine 9 dimethylation (H3K9me2) are related transcriptional silencing mechanisms that target transposable elements (TEs) and repeats to maintain genome stability in plants. RdDM is mediated by small and long noncoding RNAs produced by the plant-specific RNA polymerases Pol IV and Pol V, respectively. Through a chemical genetics screen with a luciferase-based DNA methylation reporter, LUCL, we found that camptothecin, a compound with anti-cancer properties that targets DNA topoisomerase 1α (TOP1α) was able to de-repress LUCL by reducing its DNA methylation and H3K9me2 levels. Further studies with Arabidopsis top1α mutants showed that TOP1α silences endogenous RdDM loci by facilitating the production of Pol V-dependent long non-coding RNAs, AGONAUTE4 recruitment and H3K9me2 deposition at TEs and repeats. This study assigned a new role in epigenetic silencing to an enzyme that affects DNA topology.

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Introduction

DNA methylation and histone H3 lysine 9 (H3K9) methylation are two chromatin modifications widely employed by eukaryotes to maintain genome stability [1,2]. H3K9 methylation and DNA methylation are targeted via small interfering RNAs (siRNAs) to repeats and transposable elements (TEs) and are required for their transcriptional silencing [1,2].

In plants, cytosine methylation is established through a process known as RNA-directed DNA methylation (RdDM), which involves small and long noncoding RNAs produced by plant-specific RNA polymerases, Pol IV and Pol V, respectively [2]. Pol IV is thought to transcribe RdDM target loci and generate long precursor RNAs. These are eventually processed into 24-nucleotide (nt) siRNAs that are loaded into the Argonaute protein AGO4 [3,4,5,6,7]. In parallel, Pol V generates long non-coding RNA transcripts from RdDM target loci, and these transcripts recruit siRNA-AGO4 to chromatin [8,9]. Through the concerted action of these two polymerases, siRNA-AGO4 becomes localized to target loci, and this ultimately recruits the methyltransferase DRM2, which effects de novo DNA methylation. In plants, DNA methylation occurs in three sequence contexts, CG, CHG, and CHH. In contrast to CG and CHG methylation, which can be maintained through the DNA methyltransferases MET1 and CMT3, respectively, CHH methylation is propagated by constant de novo methylation through RdDM [2,10].
Author Summary

DNA topoisomerase is an enzyme that releases the torsional stress in DNA generated during DNA replication or transcription. Here, we uncovered an unexpected role of DNA topoisomerase 1α (TOP1α) in the maintenance of genome stability. Eukaryotic genomes are usually littered with transposable elements (TEs) and repeats, which pose threats to genome stability due to their tendency to move or recombine. Mechanisms are in place to silence these elements, such as DNA-directed DNA methylation (RdDM) and histone H3 lysine 9 dimethylation (H3K9me2) in plants. Two plant-specific RNA polymerases, Pol IV and Pol V, generate small and long noncoding RNAs, respectively, from TEs and repeats. These RNAs then recruit protein factors to deposit DNA methylation or H3K9me2 to silence the loci. In this study, we found that treatment of plants with camptothecin, a TOP1α inhibitor, or loss of function in TOP1α, led to the de-repression of RdDM target loci, which was accompanied by loss of H3K9me2 or DNA methylation. The role of TOP1α in RdDM could be attributed to its promotion of Pol V, but not Pol IV, transcription to generate long noncoding RNAs.

In plants, H3K9 dimethylation (H3K9me2) is another repressive chromatin mark associated with TE and repeat silencing [11,12,13]. H3K9me2 and CHG methylation act in a self-reinforcing loop to promote the maintenance of these marks by histone methyltransferases KRYPTONITE (KYP or SUVH4), SUIVH5 and SUIVH6 and the DNA methyltransferase CMT3 [14]. How H3K9me2 is initially deposited is less well understood, but the RdDM pathway plays a role, as mutations in RdDM pathway genes cause marked reductions in H3K9me2 levels at RdDM target loci [7,8,15]. In fact, a recent study revealed a strong genome-wide inter-dependence between non-CG (CHG and CHH) DNA methylation and H3K9 dimethylation [16].

DNA topoisomerases are enzymes that maintain proper DNA topology [17]. During replication or transcription, the DNA helical structure opens to promote the replication or transcription fork, and the DNA in front of the fork becomes positively supercoiled, while the DNA behind the fork becomes negatively supercoiled. Topoisomerases bind these regions, nick the DNA to relieve the torsional stress, and re-ligate the DNA. Topoisomerases are divided into two major types, I and II, and further subtypes depending on their mode of action and structure [17,18].

In Arabidopsis, there are two genes encoding type IB topoisomerases, TOP1α and TOP1β, which are tandemly arrayed in the genome. top1α mutants exhibit gross morphological defects, while top1β mutants are phenotypically normal [19]. DNA-mediated knockdown of TOP1β in a top1α background is lethal [19]; thus these two genes are functionally redundant.

Here, we uncover a role of TOP1α in transcriptional silencing of TEs. We exploited a luciferase-based reporter (LUCL) that undergoes transcriptional silencing by DNA methylation [20] to perform a chemical genetics screen. We found that camptothecin (CPT) released the DNA methylation of LUCL and de-repressed its expression. CPT is a well-studied natural quinoline alkaloid that targets type IB topoisomerases [21,22]. Both the addition of CPT and loss-of-function in TOP1α led to the de-repression of RdDM target loci accompanied by a release of DNA methylation and/or a decrease in H3K9me2 levels. TOP1α is dispensable for Pol IV-mediated siRNA biogenesis but is required for the production of Pol V-dependent, long non-coding RNA transcripts. Consistent with the current model that these transcripts recruit siRNA-AGO4 to chromatin, inactivation of TOP1α resulted in reduced AGO4 occupancy at these loci. Taken together, through the identification of TOP1α as a player in RdDM, we have assigned new roles to a protein affecting DNA topology.

Results

Camptothecin releases the silencing of LUCL

To identify genes involved in DNA methylation, we performed a chemical genetics screen with LUCL, a transcriptionally-silenced luciferase (LUCL)-based reporter line [20]. In LUCL, LUCL is driven by a dual 35S promoter and both the 35S promoter and the LUCL coding region harbor DNA methylation [20]. The DNA methylation at LUCL, and consequently its transcriptional silencing, is controlled by MET1, and to a lesser extent, by the RdDM pathway [20].

Over 3,000 compounds were screened against LUCL seedlings for their effects on LUCL expression. A hit compound, camptothecin (CPT) (Figure 1A), was found to release LUCL silencing in a concentration- and time-dependent manner (Figure 1B and C). Interestingly, CPT released LUCL silencing in a bi-phasic manner, with optimal levels at 10 μM. Further, the release of LUCL activity was not observed until one day of chemical addition in a time course assay (Figure 1B). Consistently, continuous live imaging revealed that an increase in LUCL activity occurred at about 15 hr after the addition of the chemical (Figure 1C). The slow kinetics suggested that cell division is likely necessary for the de-repression of the reporter. The effects of CPT on LUCL protein activity reflected a release of LUCL silencing, as the addition of CPT led to an increase in LUCL transcript levels (Figure 1D). Consistent with the dose-dependent effects of CPT on LUCL activity, LUCL transcript levels were most de-repressed at 10 μM of CPT (Figure 1D).

Previous experiments with LUCL ruled out that it reports miRNA activity, even though it contains the miR172 binding sequence [20]. Consistently, we found that the addition of CPT did not release the LUCL activity of a miRNA reporter line, Pro35S::LUCL Pro35S::miR-LUC (Figure 1G; [23]). Thus, CPT released the LUCL activity of LUCL through a miRNA-independent mechanism.

To determine whether CPT increased LUCL transcript levels by reducing DNA methylation, we performed McrBC-PCR to examine the methylation status of LUCL. After digestion of genomic DNA with McrBC, an enzyme that only cuts methylated DNA [24], 35S promoter sequences were amplified by PCR. In the DMSO-treated control sample, little product was observed, indicating that this region was highly methylated in LUCL. However, after CPT treatment, the amount of PCR products increased (Figure 2A), suggesting that CPT treatment led to a reduction in 35S promoter methylation. In addition, the DNA methylation status of the 35S promoter and the LUCL coding region was examined by bisulfite sequencing (Figure 2B). The addition of 10 μM CPT resulted in a drastic reduction of CHH methylation, and to some extent CHG methylation, in region #1 (Figure 2C). CG methylation was largely unaffected upon CPT treatment, with the exception of region #4 (Figure 2C).

The CPT target, TOP1α, promotes CG methylation at 5S repeats

Due to their potent anti-cancer properties, CPT and its analogs have been intensely studied. The cellular target of CPT is topoisomerase I and the mechanism by which CPT inhibits topoisomerase I is well understood [25]. Given this knowledge, our finding that CPT de-represses LUCL implicated TOP1α in transcriptional gene silencing. A top1α mutant allele, top1α-2, had been found in an unrelated project (Xigang Liu and Xuemei Chen, unpublished...
defects than mine the status of enzyme that cuts unmethylated DNA in a CG context, to deter-
methylated DNA, as indicated by the increase in intensity of the (also known as
reports miRNA activity [23], RdDM regulation [5]. We digested genomic DNA with
in terms of hours:minutes after addition of 10 μM CPT or DMSO. The y-axis indicates the relative fold change of LUC activity upon
reduction of CPT as compared to DMSO. CPT affects the activity of LUC but not Pro35S::miR LUC, a reporter line in which the LUC transgene reports miRNA activity [23], Pro35S::LUC, [23], or LUC, a reporter line in which the LUC transgene is silenced specifically by CHH methylation [36], (D) qRT-PCR measuring LUC transcript levels upon addition of different concentrations of CPT. Three biological replicates, each with three technical replicates, were performed. Error bars represent standard deviation from the three biological replicates.
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results). The top1α-2 mutant carried a C→T point mutation in the second exon, which generates a premature stop codon (Figure S1A). top1α-2, which had been isolated in the Landsberg erecta background, was introgressed into Col-0 through five backcrosses to derive top1α-2Col. top1α-2Col was then crossed to LUC in the Col-0 background. Unlike CPT, which released LUC activity, the top1α-2Col mutation was not able to release LUC activity (Figure S1B), probably due to activity of the partially redundant TOP1β gene.

We next asked whether TOP1α inactivation or CPT treatment affected DNA methylation of endogenous RdDM loci. 5S rDNA is present with thousands of copies in the genome and is under RdDM regulation [5]. We digested genomic DNA with HpaII, an enzyme that cuts unmethylated DNA in a CG context, to determine the status of 5S rDNA methylation. We found that, like mgo1-7, a Pol V mutant, top1α-2 and CPT-treated seedlings had less methylated DNA, as indicated by the increase in intensity of the lower molecular weight restriction fragments (Figure 2D). top1α-7 (also known as mgo1-7 [26]; Figure S1A) has weaker developmental defects than top1α-2Col. The top1β-1 loss-of-function mutant in the Col-0 background (Figure S1A) has no obvious morphological defects (Xigang Liu and Xuemei Chen, unpublished results). CG methylation at 5S repeats was only weakly reduced in top1α-7 mutants and unaffected in top1β-1 mutants. Similarly, DNA blot analyses were conducted to examine CHG methylation at MEA-ISR and 180 bp repeats (Figure S1C and D), and CHH methylation at 5S rDNA repeats. Only a slight reduction in CHG methylation at the 180 bp repeats was detected in top1α-2 (Figure S1D), top1α-2 was indistinguishable from the isogenic Ler parental line in terms of CHG methylation at MEA-ISR (Figure S1C) or CHH methylation at 5S repeats (Figure S1D).

TOP1α has a limited role in DNA methylation in the genome

The studies above on a small number of loci revealed a limited role of TOP1α in DNA methylation. In order to obtain a global view of the function of TOP1α in DNA methylation, we performed whole genome bisulfite sequencing (MethylC-seq) on Ler, top1α-2, Col-0, top1α-7, mgo1-3 (a Pol IV mutant) and mgo1-11 seedlings. A total of 10 libraries representing one to three biological replicates of the genotypes (Table S1) were sequenced. Acceptable bisulfite conversion efficiency (Table S1) and read coverage (Table S2) were achieved for each library.

We identified differentially methylated regions (DMRs) using established procedures in the literature (see Material and Methods and Text S1). We compared each mutant to its wild-type control in the same biological replicate. We also called DMRs among the three Col-0 replicates to establish the background of spontaneous DMRs in wild type. Despite the high degree of reproducibility of the biological replicates (Table S3), when the three Col-0
Figure 2. Addition of CPT or loss of TOP1α releases DNA methylation. (A) McrBC-PCR-based methylation analysis of the 35S promoter in LUCL seedlings treated with CPT. At2g19920 is unmethylated and serves as an internal loading control. The number listed above indicates the concentration (in μM) of CPT added. D = DMSO. (B) A schematic diagram of the LUCL transgene. The numbered black lines indicate the regions for which bisulfite sequencing was performed (Figure 2C). (C) Levels of DNA methylation of LUCL in DMSO- and CPT-treated seedlings as determined by bisulfite sequencing. The regions correspond to the numbered lines in Figure 2B. (D) Loss of TOP1α results in reduced 5S rDNA methylation. Genomic DNA was digested with HpaII followed by Southern blotting. Less methylated DNA is expected to yield a higher intensity of bands lower down the gel as in nrpe1-11 (a Pol V mutant). In the CPT-treated sample, 25 μM of CPT was used. top1α-7, top1β-1, and nrpe1-11 are to be compared to Col-0 (wild type), and top1α-2 is to be compared to Ler (wild type).

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TOP1α Promotes Transcriptional Silencing

TOP1α silences transposons through DNA methylation and H3K9 dimethylation

Since the methylation-sensitive DNA blot analyses only revealed an effect of top1α alleles on DNA methylation at the SS and 100 bp repeats and the methylome profiling studies did not support a global role of TOP1α in DNA methylation, we sought to evaluate whether TOP1α is required for the transcriptional silencing of endogenous RdDM loci. qRT-PCR was performed to determine transcript levels from seven well-known RdDM loci. In both wild-type seedlings treated with CPT as well as top1α (both top1α-2 and top1α-7) seedlings, these endogenous siRNA target loci were derepressed (Figure 4A). This confirmed a role of TOP1α in silencing the RdDM target loci.

We asked whether the release of transcriptional silencing of endogenous RdDM target loci (Figure 4A) in top1α or CPT-treated seedlings was accompanied by a loss of DNA methylation. We performed McrBC-qPCR assays to quantify the levels of DNA methylation amongst different genotypes/treatments at six endogenous RdDM loci. At most of the loci, DNA methylation was reduced in the two top1α mutants, but the reductions were small in top1α-7 (Figure 4B). Treatment of wild-type [Ler] plants with CPT resulted in reductions in DNA methylation at four of the six tested loci (Figure S1E). Although the overall trend of reduced DNA methylation in the two top1α mutants and CPT treated plants agreed with the observed de-repression of these loci, there were also inconsistencies whereby de-repression was not accompanied by reductions in DNA methylation, such as at sir02 in top1α-2 and CPT-treated plants.

This incomplete correlation between TE de-repression and a reduction in DNA methylation prompted us to ask whether TOP1α silences TEs through another mechanism. Previous studies have shown that H3K9me2 is a major repressive mark for transposon silencing and that H3K9me2-dependent silencing acts in concert or in parallel with RdDM [31,32,33]. Like DNA methylation, H3K9me2 is targeted to specific TEs through siRNA-AGO4 [7]. Thus, we investigated whether loss of TOP1α function or CPT treatment altered H3K9me2 levels at TEs. Chromatin immunoprecipitation (ChIP)-qPCR showed that H3K9me2 levels at 24nt, sir02, cluster4, and At4G1 showed in both top1α-7 and top1α-11 (Figure 4C). We also performed ChIP-qPCR on LUC transgenic seedlings treated with DMSO or CPT. CPT treatment was found to cause a strong reduction in H3K9me2 levels at four TE loci (Figure 4D). As CPT was initially isolated through a chemical genetics screen with LUC, we asked whether the LUC transgene in LUC also harbored H3K9me2 and, if so, whether CPT treatment reduced its H3K9me2 levels. Indeed, ChIP-qPCR showed that the d35S of the LUC transgene (region #1 in Figure 2B) harbored H3K9me2 with CPT treatment reducing H3K9me2 levels (Figure 4D).

As H3K9me2, which is introduced by KYP and its paralogs, and H3K9 dimethylation, which is deposited by CMT3, acts in a self-reinforcing loop, and both H3K9me2 and CMT3 contribute to CHH methylation [14,16], we asked whether the role of TOP1α in DNA methylation depends on KYP or CMT3. To address this question, we treated Ler (wild-type), kyp-2 and cm5-7 plants with CPT to inhibit topoisomerase I activity and then assayed DNA methylation at six TE loci. CPT treatment of wild-type plants resulted in reduced DNA methylation at four of the six loci (Figure S1E). The reduction in DNA methylation caused by CPT treatment was minimal at these four loci in either cm5-7 or kyp-2 (Figure S1E). This suggested that the effects of TOP1α in DNA methylation require CMT3- and KYP-mediated H3K9 dimethylation.

TOP1α does not affect small RNA levels

The promotion of DNA methylation and/or H3K9me2 deposition at TEs implicates a role of TOP1α in RdDM, a process that involves Pol IV and Pol V. As topoisomerases are required to release DNA topological tension generated by transcription [17], it would be reasonable to expect that TOP1α is required for the activities of either Pol IV or Pol V. We first tested whether TOP1α is required for the activities of Pol IV, the output of which is the accumulation of 24-nt siRNAs from RdDM target loci. RNA blot analysis showed that siRNA accumulation at several loci was similar in Ler and top1α-2 (Figure S3A). To gain a global view on the potential relationship between TOP1α and Pol IV, we compared deep sequencing profiles of small RNAs from Ler, top1α-2, Col-0, top1-3, and top1α-11. The size distributions of all small RNA reads in Ler and top1α-2 were almost identical (Figure S3B). To determine whether TOP1α affects siRNA accumulation at specific regions of the genome, we identified differential small RNA regions (DSRs). While large numbers of DSRs were found in top1-3 or top1α-11 relative to the wild-type control, consistent with the essential role of Pol IV and the auxiliary role of Pol V in siRNA biogenesis [3,5,6], very few were found in top1α-2 (Table S7). Furthermore, analysis of small RNA abundance throughout the genome did not support a global role of TOP1α in small RNA accumulation (Figure S3C). Therefore, Pol IV activity does not appear to require TOP1α.

Given that we had found 71 WT-top1α DSRs (Table S7), we asked whether the reduced CHH methylation at the 97 WT-top1α DMRs was associated with reduced siRNA levels. We found that only 11 of the 97 DMRs overlapped with WT-top1α DSRs (Figure 3D). A representative of such a locus is shown in Figure S2A. Most of the 97 DMRs did not overlap with the 71 WT-top1α DSRs; two such loci are shown in Figure S2B and C. Therefore, the reduced CHH methylation in top1α could not be explained by...
reduced siRNA levels. On the other hand, more than 60% of the 97 WT-top1a DMRs overlapped with WT-nrpd1 DSRs (Figure 3D; Figure S2A and B), suggesting that these regions, which require TOP1a for CHH methylation, undergo Pol IV-dependent siRNA production. Therefore, TOP1a must promote CHH methylation at these RdDM loci independently of siRNA biogenesis.

TOP1α promotes the production of Pol V-dependent transcripts and AGO4 occupancy at TEs

We next tested whether TOP1α promotes the production of Pol V-dependent transcripts. We performed qRT-PCR and RT-PCR to detect Pol V-dependent transcripts from eight loci, MEA-ISR, AtSN1, and six IGN loci that produce such transcripts [9,30]. At all eight loci, the levels of the Pol V-dependent transcripts were reduced in top1a-2 as compared to Ler (Figure 5A and B). We previously showed that Pol II generates long noncoding transcripts at the soloLTR locus [34]. The accumulation of these transcripts at soloLTR was also reduced in top1a-2 (Figure 5B). Therefore, TOP1α contributes to the production of Pol V-dependent or Pol II-dependent long noncoding transcripts.

As the Pol V- or Pol II-dependent long noncoding transcripts facilitate the recruitment of siRNA-AGO4 to chromatin to ultimately result in RdDM or H3K9me2 deposition, we asked whether TOP1α promotes AGO4 occupancy at these RdDM target loci. ChIP-qPCR was conducted with anti-Myc antibodies in Myc-AGO4 [35] and Myc-AGO4 top1a-2 plants. At four well-known RdDM target loci, AGO4 occupancy was reduced in top1a-2 (Figure 5C).

To determine whether TOP1α might act directly at these RdDM loci, we examined TOP1α occupancy at these loci. We first generated a TOP1α-HA fusion driven by the TOP1α promoter (TOP1α-HA) and introduced it into top1a-2. The morphological phenotypes of top1a-2 plants were completely rescued by TOP1α-HA, indicating that the transgene was functional. We then performed ChIP-qPCR using anti-HA antibodies. TOP1α was found at all six loci examined (Figure 5D).

Discussion

Beginning with a forward chemical genetics screen with a transcriptionally silenced reporter, LUXL, we have discovered that the well-studied anti-cancer compound CPT can de-repress loci undergoing transcriptional silencing by releasing H3K9 methylation and/or DNA methylation. As topoisomerase I is the cellular target of CPT, this implicates topoisomerase I in transcriptional silencing. Indeed, two top1a alleles, top1a-2 and top1a-7, mimic CPT treatment in de-repressing the expression of endogenous RdDM target loci and reducing H3K9me2 or DNA methylation levels at these loci.

Figure 3. TOP1α does not globally impact DNA methylation but promotes CHH methylation at a small number of loci. (A) Pie charts showing that the great majority of WT-top1a DMRs show reduced DNA methylation in top1a. Each circle represents total WT-top1a DMRs in a methylation context (CG or CHH). The red and blue areas represent DMRs with reduced and increased DNA methylation in top1a, respectively. The numbers indicate the numbers of DMRs in each category. The numbers in the parentheses represent the percentage of the DMR category in total DMRs. (B) The majority of WT-top1a CHH DMRs showing reduced DNA methylation in top1a overlap with TEs (74%, blue). Those that overlap with genes and intergenic regions are shown in red (7%) and green (19%), respectively. (C) The majority of WT-top1a CHH DMRs overlap with CHH DMRs between WT and nrpd1 or WT and nrpe1 (91%, red), suggesting that these regions require Pol IV or Pol V for CHH methylation. The portion of WT-top1a CHH DMRs not overlapping with WT-nrpd1 or WT-nrpe1 CHH DMRs are shown in blue (9%). (D) Overlap between WT-top1a CHH DMRs with DSRs (differential small RNA regions) between WT and nrpd1 or WT and top1a. There is little overlap between WT-top1a CHH DMRs and WT-top1a DSRs.
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Here, we first consider whether TOP1α acts through RdDM or independently of RdDM to silence TEs. RdDM requires Pol IV and Pol V, which generate siRNAs and long noncoding RNAs, respectively. We show that TOP1α is dispensable for siRNA accumulation, but is required for the production of Pol V-dependent long noncoding RNAs, which are known to recruit siRNA-AGO4 to chromatin. Consistently, TOP1α promotes the recruitment of AGO4 to RdDM target loci. Moreover, 88 out of 97 WT-top1α CHH DMRs with reduced methylation in top1α also require Pol IV or Pol V for CHH methylation (Figure 3C). 58 rDNA loci lose CG methylation in top1α-2 and nrpe1-11 mutants, and provide an example of a genomic region where CG methylation requires TOP1α, Pol IV, and Pol V. These data suggest that TOP1α acts at least in part through RdDM to silence TEs and repeats. However, MethC-seq analyses revealed that TOP1α has a limited role in DNA methylation. We envision two possibilities for the limited role in DNA methylation observed for TOP1α. First, TOP1α may have a much broader role in DNA methylation in the genome, and the limited effects of top1α mutants on DNA methylation could be due to the redundant functions of TOP1β. So far, our efforts to knock down TOP1β in the top1α-2 background have been unsuccessful. Second, TOP1α’s primary functions may lie in the promotion of H3K9 dimethylation, with DNA methylation being a secondary effect of H3K9 dimethylation. From our studies of a limited number of RdDM loci, we found that reduced H3K9me2 levels, but not necessarily reduced DNA methylation, always accompany the de-repression of these loci by CPT treatment or by mutations in TOP1α. Therefore, it is likely that the primary function of TOP1α lies in facilitating H3K9me2 deposition. Consistent with this model, the observed effects of CPT treatment on DNA methylation at four loci require CMT3 and KTP, both of which promote H3K9 dimethylation. Another observation consistent with this hypothesis is that CPT treatment had no effect on LUCH (Figure 1C), a reporter gene that is strictly repressed by CHH methylation and is insensitive to loss of function in CMT3 [36]. As CMT3-mediated DNA methylation requires H3K9me2 [14], we presume that LUCH is not repressed by H3K9me2. The lack of an effect of CPT treatment on LUCH would be consistent with TOP1α acting in TGS through H3K9me2 deposition.

Our finding that TOP1α promotes the production of Pol V-dependent transcripts is consistent with what is known about the function of topoisomerases in bacteria and yeast. Topoisomerases are thought to facilitate transcription elongation by relaxing supercoils [37]. Consistent with this model, loss of Top1 in Schizosaccharomyces pombe results in the accumulation of Pol II in gene bodies [38,39]. The parallels of Pol V- and Pol II-mediated transcription have recently been highlighted [40], and we propose that TOP1α promotes transcription elongation by Pol V as it does for Pol II.

Although we prefer a model in which TOP1α acts in RdDM by facilitating the production of long noncoding RNAs by Pol V or Pol II, an alternative model cannot be overlooked. Studies in other systems have shown that topoisomerases interact with SMC-containing proteins acting in chromosome compaction [41,42]. DMS3, a player of the RdDM machinery, contains an SMC domain [43]; therefore, there is a possibility that TOP1α may facilitate RdDM through DMS3.

In summary, we have discovered a role for DNA topoisomerase I in H3K9 methylation and DNA methylation in Arabidopsis. Another study showed that chemical inhibitors of topoisomerases I and II release the epigenetic silencing of an imprinted gene in mouse [44]. Together, these studies point to a role of topoisomerases in epigenetic silencing. Given that CPT is a canonical anti-cancer compound and several of its derivatives are presently used in cancer treatment, this raises the possibility that pharmacologically induced DNA topoisomerase inhibition could also have epigenetic consequences.

Figure 4. TOP1α promotes transposon silencing at endogenous RdDM target loci through H3K9me2 deposition. In (A) to (D), error bars represent standard deviation calculated from three biological replicates. In (A) and (B), top1α-7, top1α-2 and CPT-treated LUC were compared to Col-0, Ler, and DMSO-treated LUC respectively. The relative levels to these controls (set to 1.0) are shown. The loci tested are labeled on the x axis. (A) Loss of TOP1α or addition of CPT results in RdDM target loci de-repression. (B) Loss of TOP1α or addition of CPT results in a release of DNA methylation at some loci. MrcBC-qPCR analysis was performed to quantify DNA methylation levels in top1α or CPT-treated plants. Higher DNA levels in this assay correlate with lower levels of DNA methylation. At1g40129 served as an internal unmethylated control. (C) Loss of TOP1α results in reduced H3K9me2 levels at endogenous RdDM loci. ChiP-qPCR was performed to measure H3K9me2 levels at five RdDM target loci. elf4A1, which does not harbor H3K9me2, was used as an internal control. – = samples processed without antibody. + = samples processed with anti-H3K9me2 antibodies. (D) CPT treatment results in reduced H3K9me2 levels at the LUC transgene and four endogenous RdDM loci. LUC plants were treated with either DMSO or 10 μM CPT and subjected to ChiP. qPCR was performed with the immunoprecipitated DNA for the LUC transgene and four endogenous RdDM loci. elf4A1 was used as an internal negative control. – = samples processed without antibody. + = samples processed with anti-H3K9me2 antibodies.
and “-” signs represent “no antibody” or “anti-Myc antibodies”, respectively. Error bars represent standard deviation calculated from three technical replicates. (D) TOP1A-HA top1a-2 seedlings were subjected to ChIP with anti-HA antibodies. qPCR was then performed on the immunoprecipitated DNA for six endogenous RdDM loci. “-” and “+” signs represent “no antibody” or “anti-HA antibodies”, respectively. Error bars represent standard deviation calculated from three technical replicates. Two biological replicates were performed and showed the same trend. One biological replicate is shown. doi:10.1371/journal.pgen.1004446.g005

MethylC-seq analysis: Identification of Differentially Methylated Regions (DMRs)

Raw data from Illumina sequencing were filtered to remove reads that failed to pass the Illumina quality control and to condense multi-copy reads to a single copy. Hereafter, the reads were mapped to TAIR 10 Arabidopsis genome as well as a C-to-T converted genome using BS SEEKER [45] with default settings. Only perfectly and uniquely mapped reads were retained. For Ler and top1a-2, which are in the Landsberg ecotype, the reads were mapped to a pseudo-Ler genome generated by incorporating the Ler polymorphisms into the TaIR10 Columbia genome (ftp://ftp.arabidopsis.org/Polymorphisms/Ecker_ler.homozygous_sup.txt). This enables the direct comparison of DMR regions between the Columbia and Landsberg samples.

DMRs were identified following a published method [27] with some modifications. In brief, the genome was split into continuous 100 bp windows. The Cs or Ts were counted in each window in the three different contexts (CG, CHG or CHH) separately. Only windows with less than 4 Cs each sequenced at least 4 times in the wild-type sample were kept for the DMR analysis. The methylation level for a window was determined as:

$$\text{methylation level} = \frac{\sum a_i}{\sum (a_i+b_i)}$$

in which $a_i$ denotes the number of read “C”s and $b_i$ denotes the number of read “T”s mapping to the $i$th cytosine site. The methylation level in each window in wild type is then compared to the corresponding window in a mutant. A methylation difference of 0.4, 0.2, and 0.1 for CG, CHG, and CHH, and an adjusted p-value (FDR)<0.01 (Fisher’s exact test) were used as the cutoff for defining DMRs.

Additional measures were taken to reduce experimental noise. First, two or three biological replicates/alleles were examined. In deriving initial DMRs, we compared each wild type/mutant pair from the same biological replicate (Table S5). Then, DMRs located within 200 bp of each other were merged. Next, the overlap in DMRs from the two biological replicates/alleles was identified (Table S5). Finally, we removed the DMRs that overlapped with the hypervariability (HV) regions found to be prone to changes in DNA methylation [28,29] (Table S5).

See Text S1 for Supplemental Methods and Table S8 for oligonucleotides used in this study.

Accession numbers and data deposition

The gene accession numbers used in this study are At5g55310 (TOP1a), At5g55300 (TOP1b), At1g05460 (NRD1), and At2g40030 (NRD2).
Supporting Information

**Figure S1** The nature of top1x and top1β alleles and the effects of top1x-2 on DNA methylation at several loci. (A) Schematic representation of TOP1x and TOP1β and several mutant alleles. The white triangles represent T-DNA insertions, top1x-2 is a point mutation that causes an early stop codon (star). (B) top1x-2 <sup>2clo</sup> does not de-repress LUCI, top1x-2 <sup>2clo</sup> is top1x-2 introgressed into Col-0 through five backcrosses. LUCI, ago4-6 and LUCI drm2-6 were included as positive controls, as ago4-6 and drm2-6 weakly de-repress LUCI [20]. (C) DNA blot analysis of the MEA-ISR locus. Genomic DNA from ten-day old seedlings was digested with MspI and hybridized with a probe corresponding to the MEA-ISR locus. MspI cuts unmethylated DNA in a CHG context. The upper and lower bands represent methylated and unmethylated DNA. *nrpe1-11* is a Pol V mutant in the Col-0 background. No change was observed between top1x-2 and Ler (the wild-type control for top1x-2). (D) DNA blot analysis of 180 bp and 5S repeats. Left panel: Genomic DNA from ten-day old seedlings was digested with MspI and hybridized with a probe corresponding to the 180 bp centromeric repeats. *cm3-7* is a control with reduced CHG methylation. *top1x-2* has a slight reduction in CHG methylation at the 180 bp repeats as compared to Ler. Right panel: Genomic DNA from ten-day old seedlings digested with *HaeIII* and hybridized with a probe corresponding to the 5S loci. *HaeIII* recognizes the GGC sequence, but cannot cut when the last C is methylated, thus it is sensitive to CHH methylation. *nrpd1-3* is a Pol IV mutant and *nrpd1-11* is a Pol V mutant. Both serve as controls with reduced CHH methylation and are to be compared to Col-0 as wild type. No change was observed between top1x-2 and Ler (the wild-type control for top1x-2). (E) CPT treatment results in reductions in DNA methylation at several RdDM loci in a CMT3- <sup>2</sup>- and KLP-dependent manner. McrBC-qPCR analysis was performed to quantify DNA methylation levels in CPT-treated Ler (wild type), *cm3-7* or *kyp-2* ten-week-old seedlings. Relative DNA ratios between CPT treatment and DMSO treatment are shown. Higher DNA levels in this assay correlate with lower levels of DNA methylation. *At1g40129* served as an internal unmethylated control. Error bars represent standard deviations calculated from three biological replicates.

**Figure S2** Representative screen shots of an overlay of MethylC-seq and small RNA-seq at three genomic regions in various genotypes. The three different loci (A-C) are indicated by their genomic coordinates above the tracks. The top five tracks depict CHH methylation (blue vertical lines). The y-axis indicates the methylation level from 0 (0%) to 1 (100%). The next four tracks represent small RNAs (purple vertical lines). The y-axis indicates small RNA abundance normalized by read depth. The positions of genes or transposable elements (TEs) are indicated below the small RNA tracks, with the green and brown rectangles representing genes and TEs, respectively. The position of these boxes (above or below the line) indicates which DNA strand those features are transcribed from. The black, orange, and purple rectangles at the bottom indicate the positions of WT-top1x DMRs, Col-ntpd1-3 DSRs, and Ler-top1x-2 DSRs, respectively. DMRs, differentially methylated regions; DSRs, differential small RNA regions.

**Figure S3** TOP1x does not globally contribute to small RNA accumulation. (A) Loss of TOP1x did not significantly change siRNA (cluster4, soloLTR, siR1003) and miRNA (miR173) levels. RNA blots were performed for Ler (wild type) and *top1x-2*. U6 was used as an internal loading control. The numbers indicate the relative abundance of the small RNAs in the mutant (with that in the wild type set to 1). (B) The size distribution of total small RNA reads in Ler and *top1x-2* is largely similar. (C) Box-and-whisker plots of global small RNA abundance in various genotypes. The whiskers extend to the most extreme data points that are no more than 1.5 times the interquartile range from the box. Significant reduction is indicated by *''* (P<10<sup>-10</sup> Mann-Whitney U test). *nrpe1-11* and *nrpd1-11* have mutations in Pol IV and Pol V, respectively, and are to be compared to Col-0 (wild type). Small RNAs were mapped to the genome, which is divided into 500 bp static windows. Only windows in which read abundance was at least 10 RPM in Col-0 or Ler are considered. The x-axis represents the genotypes as indicated. The y-axis shows normalized read abundance (in RPM, reads per million) in 500 bp windows. Small RNA levels were unaffected in the *top1x-2* mutant, whereas they were reduced in *ntpd1-3* and *nrpd1-11* as compared to Col-0.

**Table S1** Summary of bisulfite conversion efficiency for each genotype.

**Table S2** Read coverage of whole genome bisulfite sequencing libraries.

**Table S3** Correlation coefficient values for the different biological replicates of each genotype in MethylC-seq.

**Table S4** DMRs between wild-type samples.

**Table S5** Derivation of DMRs between wild type and *top1x*, *ntpd1-3*, or *nrpd1-11*.

**Table S6** Final WT-top1x DMRs.

**Table S7** Only a small number of differential small RNA regions (DSRs) were found between wild type and *top1x-2*. Whole genome high throughput sequencing was performed for small RNAs in wild type (Col-0 and Ler), *ntpd1-3*, a Pol IV mutant, *nrpd1-11*, a Pol V mutant, and *top1x-2*. *ntpd1-3* and *nrpd1-11* are to be compared to Col-0 and *top1x-2* to be compared to Ler. The genome was divided into 500 bp static windows and small RNA reads in each window were counted and compared between each mutant and its corresponding wild type. Thousands of DSRs were found in *ntpd1-3* or *nrpd1-11* as compared to Col-0, but only 71 were found in *top1x-2* relative to Ler (see Experimental Procedures for the derivation of DSRs). The numbers of DSRs mapping to different genomic features (TE, gene, and inergenic region) are listed. TE = transposable element. “Reduced” and “increased” refer to DSRs with reduced and increased small RNA read counts in the mutants, respectively.

**Table S8** Oligonucleotides used in this study.

**Text S1** Supplemental methods.

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**Author Contributions**

Conceived and designed the experiments: TTD LG XL SheL YZ RJS PM. Performed the experiments: TTD LG XL SheL YZ. MO BL RJS PM ShaL OP. Analyzed the data: TTD LG XL SheL YZ. MO BL RJS PM ShaL OP XC. Wrote the paper: TTD LG XL BL DW XC.

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