G9a selectively represses a class of late-replicating genes at the nuclear periphery

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We have investigated the role of the histone methyltransferase G9a in the establishment of silent nuclear compartments. Following conditional knockout of the G9a methyltransferase in mouse ESCs, 167 genes were significantly up-regulated, and no genes were strongly down-regulated. A partially overlapping set of 119 genes were up-regulated after differentiation of G9a-depleted cells to neural precursors. Promoters of these G9a-repressed genes were AT rich and H3K9me2 enriched but H3K4me3 depleted and were not highly DNA methylated. Representative genes were found to be close to the H3K9me2 enriched but H3K4me3 depleted and were not highly DNA precursors. Promoters of these G9a-repressed genes were AT rich and H3K9me2 enriched but H3K4me3 depleted and were not highly DNA methylated. Representative genes were found to be close to the H3K9me2 enriched but H3K4me3 depleted and were not highly DNA precursors. Promoters of these G9a-repressed genes were AT rich and H3K9me2 enriched but H3K4me3 depleted and were not highly DNA methylated. Representative genes were found to be close to the H3K9me2 enriched but H3K4me3 depleted and were not highly DNA methylated. Representative genes were found to be close to the H3K9me2 enriched but H3K4me3 depleted and were not highly DNA methylated.

Results

Changes in Histone Methylation Following G9a Knockout. Stable and irreversible genetic knockout often leads to compensatory genetic and epigenetic changes that can confuse interpretation of the resulting phenotypes (15). For example, Suv39h1,2-knockout cells lose H3K9me3 but also gain H3K27me3 in pericentric heterochromatin, where it is not seen in wild-type cells (4). This kind of adaptation can be largely circumvented through the use of a conditional-knockout, in which cellular responses to the acute loss of the gene product can be monitored. We have constructed a conditional knockout of G9a in mouse ESC line T2 (Fig. 1). This cell line has a single copy of the G9a gene flanked by loxP recombinase sites and expresses 4-hydroxytamoxifen (OHT)-inducible Cre fusion protein (16). Addition of OHT results in the rapid deletion of G9a (Fig. 1B), a partial reduction in total H3K9me1 and H3K9me3, and a substantial reduction in total

Edited by Mark T. Groudine, Fred Hutchinson Cancer Research Center, Seattle, WA, and approved September 25, 2009 (received for review June 4, 2009)
H3K9me2 (Fig. 1C), but has no effect on mono-, di- or trimethylated H3K27 or H4K20me3. The effects on the different methylated forms of H3K9 were similar with either commonly used polyclonal antibodies or more recently reported monoclonal antibodies (17) and were similar in a stable G9a-null ESC line. Hence, this conditional-knockout cell line provides the opportunity to observe the effects of acute G9a loss in the absence of complicating adaptive changes.

We previously reported that H3K9me2 is enriched at the nuclear periphery in human, mouse, and hamster fibroblasts, as well as in mouse ESCs, and that stable G9a-knockout ESCs preferentially lose peripheral H3K9me2 (13). To determine whether acute G9a loss also preferentially affects peripheral H3K9me2, we performed immunofluorescence detection of mono-, di- and trimethyl H3K9 before and after the addition of OHT (Fig. 2). H3K9me1 was distributed throughout the interior of the nucleus, excluding the periphery and clusters of pericentric heterochromatin (chromocenters), whereas H3K9me2 was localized primarily to the chromocenters; both these modifications show some reduction in signal after G9a loss but no change in localization. In contrast, H3K9me2
was most prominent at the periphery but also localized to some interior chromatin. G9a knockout led to complete loss of the peripheral signal and reduction of the internal signal but also a detectable increase at chromocenters. This increase also was detected in a stable G9a-knockout cell line (13). Hence, after G9a knockout, H3K9me2 remains that is difficult to detect by Western blotting but is detected by immunolocalization, mostly at chromocenters. Taking these findings together, we conclude that G9a is the primary methyltransferase responsible for H3K9me2 at the nuclear periphery but that other mechanisms maintain low levels of this modification in other nuclear compartments.

**G9a Loss Leads to Up-Regulation of a Select Group of Genes.** To identify genes affected by G9a loss, we examined changes in transcript levels 7 days after OHT addition using microarray analysis. Comparison with mock-treated G9a^{flox/flox} cells identified 167 genes as more than 4-fold up-regulated and 10 genes as slightly more than 3-fold down-regulated (Fig. 3A). Analysis of a selection of the best-annotated (RefSeq) of these genes by RT-PCR verified 7/7 up-regulated genes, whereas down-regulated genes were affected less than 2-fold in the presence and absence of G9a (supporting information (SI) Fig. S1A). ChIP analysis revealed significant enrichment of H3K9me2-containing promoters relative to an unaffected gene (Fig. 3B), consistent with these genes being targets of the G9a HMTase activity. Next, 4 days after the addition of OHT, we differentiated ESCs to NPCs for 9 days using a defined differentiation medium in adherent culture (18). These results revealed a set of 119 genes that were more than 4-fold up-regulated in OHT-treated vs. mock-treated cells (Fig. S1B), approximately one-third of which overlapped with the genes up-regulated in undifferentiated cells (Fig. 3C). Interestingly, genes up-regulated uniquely after differentiation to NPCs were developmentally regulated genes, mostly involved in vascular development and/or expected to be silent in NPC-derived lineages. A complete list of these genes and their properties can be found in Table S1. Inspection of results from a recent ChIP-chip analysis of H3K9me2 in mouse ESCs (11) revealed that the promoter regions of most up-regulated genes were significantly enriched for H3K9me2 (Fig. 3D), extending our individual gene results (Fig. 3B) and suggesting that the promoters of most of these genes are direct targets of the G9a HMTase activity. In fact, the set of genes that were found to be up-regulated in both ESCs and NPCs were significantly more enriched for H3K9me2 than those promoters of either cell type alone (Table S1). Although we are not certain whether G9a represses all of these genes directly or indirectly, for the purposes of discussion we refer to them collectively as “G9a-repressed” genes.

Interestingly, G9a-repressed promoters are only a small subset of genes highly enriched for H3K9me2. Because the majority of H3K9me2 is lost after G9a deletion (Fig. 1), the repression of most of these H3K9me2-containing promoters must be maintained by other mechanisms. Most G9a-repressed genes do not fall within the previously identified LOCKs (11), but they are located within >50-kb regions of generally high H3K9me2 (Fig. S2) just below the threshold used to identify LOCKs. Although G9a-repressed genes were found on all chromosomes, they were highly enriched on the single X-chromosome in these male cells, particularly in differentiated cells (Fig. S3). The promoters of G9a-repressed genes also were unusually low in GC content (Fig. 3E), with 74.2% having low to intermediate CpG density (more than twice the fraction found in total genes as defined in ref. 19), mostly low in H3K4me3, and not enriched for either RNA pol II or DNA methylation relative to total genes. Interestingly, these genes are distinct from 126 genes whose promoters have high levels of G9a-dependent DNA methylation after retinoic acid-induced differentiation (9). Together these results identify a specific set of genes that are dependent upon G9a for their repression, many of which are H3K9 methylated but not highly DNA methylated despite the role of G9a in recruiting DNA methyltransferase.

**Most G9a-Repressed Genes Are Induced Within Late-Replicating Chromatin.** Because the nuclear periphery is a late-replicating compartment (13), G9a-dependent enrichment of H3K9me2 at the nuclear periphery (Fig. 2) suggested that the HMTase activity of G9a is preferentially targeted to late-replicating chromatin. In fact, that H3K9me2 was strongly enriched within mid- to late-replicating chromosome domains (Fig. 4A). Furthermore, G9a-repressed

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**Fig. 3.** Transcriptional changes after G9a knockout. (A) Following transcription microarray analysis, data normalization, and consolidation of probe values into 1 value per gene, gene expression values for all 24,210 genes identified on the NimbleGen microarray were plotted on the y-axis for mock-treated cells vs. the fold change for each gene after OHT treatment relative to mock treatment. A 4-fold increase in transcription was considered a cut-off (red line) for up-regulated genes; a 3-fold decrease in transcription was considered as a cut-off (blue line) for down-regulated genes. (B) ChIP analysis for enrichment of H3K9me2 at the promoters of 8 genes that were either repressed (Magea1, Magea2, Rhox2, Dub1, Aszl1, Akr1c12, Gm397) or unaffected (Oct4) by G9a knockout in ESCs. Immunoprecipitated DNA as well as DNA from the input was quantified by real-time PCR. (C) At 5 days after the addition of OHT, cells were differentiated to NPCs for 9 days. Microarray analysis was performed as in (A) (Fig. S1). Shown is a Venn diagram of the number of genes up-regulated after G9a deletion in ESCs (blue) and in NPCs (green) and those genes common to both ESCs and NPCs (red). (D) The total amount of H3K9me2 integrated over a region from 2 kb upstream to 0.5 kb downstream of the transcription start site of each gene was determined using previously published ChIP-chip data in mouse ESCs (11), and gene number was plotted as a function of H3K9me2 level. (Top) The number of total genes with each H3K9me2 value. (Middle) The number of up-regulated genes from (A) with each H3K9me2 value. (Bottom) The degree of up-regulation following G9a deletion vs. H3K9me2 for each gene. (E) Top) The GC content surrounding each promoter ± 1 kb from the transcription start site (total genes). (Middle) The GC content for the up-regulated promoter as in (D). (Bottom) A plot of the fold transcriptional change following G9a deletion vs. promoter GC%.
and Magea1 replication, potentially as part of the mechanism by which it these results raised the possibility that G9a may mediate late-replicating genes. Hence, G9a is the gatekeeper for the transcription of a set of 

Shown are 5-Mb regions surrounding 3 highly up-regulated genes regulated by more than 5-fold were late replicating (Table S1). 

\[ \text{[Log}_2(\text{Early/Late}) \] of 2 replicate samples (dye-swap) are plotted vs. the map position of each probe. (A–F) Scatter plots of probe values between the indicated conditions. (G) LOESS curves of replication timing in ESCs (blue) vs. NPCs (green) for either mock-treated (darker shade) or OHT-treated (lighter shade) cells. Shown are 5-Mb regions surrounding 3 highly up-regulated genes Asz1, Dub1a, and Magea1. The red dotted line indicates the map position of the gene.

genes are located within AT/long interspersed nuclear element-rich regions (LINE) (Table S1) that can switch replication timing during differentiation (14, 20). Finally, although ∼75% of genes are replicated in the first half of S-phase, nearly 75% of G9a-repressed genes were late replicating (Fig. 4B), and almost all genes up-regulated by more than 5-fold were late replicating (Table S1). Hence, G9a is the gatekeeper for the transcription of a set of late-replicating genes. 

Because late replication is associated with gene silencing (21), these results raised the possibility that G9a may mediate late replication, potentially as part of the mechanism by which it represses these genes. To investigate the influence of G9a on replication timing, TT2 G9a−/−floxed cells were treated with or without OHT for 2 days, followed by 5 additional days of culture. Spectral karyotyping (SKY) was performed to verify genomic integrity after OHT treatment (Fig. S4). Replication timing was analyzed genome-wide using a previously described protocol (14). Briefly, cells were labeled with BrdU, sorted by flow cytometry into early- and late-S-phase populations, and BrdU-substituted DNA was immunoprecipitated, differentially labeled, and co-hybridized to a high-density whole-genome oligonucleotide microarray. This process generates a “replication-timing ratio” [Log2(Early/Late)] for each of the tiled probes, which are positioned every 5.8 kb throughout the mouse genome. Replicates (dye-swap) showed high correlation [R = 0.74–0.83 for raw data and R = 0.95–0.96 for locally weighted scatter plot-smoothed (LOESS) data], and smoothed values were averaged. Fig. 4C shows a comparison of such averaged values for each probe across a 60-Mb segment of chromosome 19. Visual inspection of many such segments revealed no detectable changes in replication timing. Plotting all data points from mock- vs. OHT-treated cells relative to each other demonstrated a very high correlation between these data sets across the genome (Fig. 4D). We did detect an advance in the replication timing of major (but not minor) satellite DNA contained within the pericentric heterochromatin in G9a-knockout ESCs (22), consistent with a prior report (23). This finding could be related to the redistribution of H3K9me2 to pericentric heterochromatin (Fig. 2).

To investigate the possibility that G9a might play a role in replication timing changes that occur during differentiation, we differentiated TT2 G9a−/−floxed to NPCs using defined medium conditions (18) following mock- or OHT-treatment as in Fig. 3. Neural differentiation proceeded similarly with or without G9a, as verified by both transcription microarray and individual gene RT-PCR analyses. Replication timing was profiled genome wide after differentiation. As shown in Fig. 4E, differentiation elicited many changes in replication timing encompassing nearly 20% of the genome, consistent with previous results (14). However, comparison of mock- vs. OHT-treated NPCs showed high correlation (Fig. 4F), indicating that differentiation-induced changes in replication timing were unaffected by the loss of G9a. To determine whether localized changes in replication timing took place at G9a-repressed genes, LOESS tracings of the average replication timing values for 5-Mb regions surrounding 3 highly up-regulated genes were overlaid (Fig. 4G), revealing no detectable changes in replication timing as a result of G9a loss in either differentiated or undifferentiated cells. We conclude that G9a depletion leads to the induction of a class of late-replicating genes without affecting their replication timing or programmed changes in replication timing during differentiation. All replication timing and transcription microarray data are available for public viewing and downloading at ref. 24.

**G9a-Repressed Genes Localize to the Nuclear Periphery.** The above results demonstrate that H3K9me2 is enriched at the nuclear periphery and G9a is required to maintain H3K9me2 in the peripheral compartment (Fig. 2). Moreover, the nuclear periphery is a late-replicating (13) and generally repressive nuclear compartment (12, 25–27). These findings raised the question as to whether G9a-repressed genes are localized to the periphery. To address this question, we determined the subnuclear locations of 6 genes that were up-regulated in both ESCs and NPCs after G9a knockout by FISH and scored their radial distances from the periphery in mock- and OHT-treated NPCs relative to an unaffected gene. Results (Fig. 5) revealed that all 6 G9a-repressed genes were located near the nuclear periphery and remained near the periphery after G9a knockout, suggesting that G9a-repressed genes are enriched in the late-replicating peripheral compartment of the nucleus.

**Discussion** We demonstrate that the acute loss of HMTase G9a in mouse ESCs results in rapid depletion of H3K9me2 from the nuclear periphery and de-repression of 167 genes. G9a-deficient cells are capable of differentiation to NPCs, accompanied by the failure to repress a partially overlapping, set of 119 genes. Surprisingly, these genes are mostly late replicating, but their activation is not associated with a switch to early replication as usually is observed.
for late-replicating genes that become activated during differentiation (14, 28, 29). In fact, almost complete depletion of H3K9me2 and partial depletion of H3K9me1 and H3K9me3 genome-wide resulting from G9a loss had no detectable effect on replication timing throughout the genome, except for a small effect on pericentric heterochromatin. We show that H3K9me2 is depleted selectively at the nuclear periphery upon G9a loss, and consistently we find G9a-repressed genes to be near the nuclear periphery. Our results demonstrate that, although G9a may act at many promoters and is known to affect the DNA and histone methylation of chromatin throughout the genome, it is directly responsible for the repression of a small set of late-replicating genes that are localized within peripheral heterochromatin.

Our results demonstrate that G9a is responsible for the majority of H3K9me2, as well as a fraction of H3K9me1 and H3K9me3, confirming and extending prior observations (2, 3, 5, 13). We also detect some remaining residual H3K9me2, including an increase in H3K9me2 in the pericentric heterochromatin, where G9a loss has been shown to cause a reduction in DNA methylation (7) and an increase in HP1-gamma (30). These changes in chromatin structure may account for a moderate karyotypic instability we observed following G9a loss and a slight advance in replication timing of major-satellite DNA (22, 23). The most dramatic qualitative effect of G9a loss is the selective loss of detectable H3K9me2 at the nuclear periphery (13).

Although some have reported results with specific genes derived from transcription microarray analysis of G9a-knockout mouse embryonic fibroblasts (30) or ESCs (31), we report the full set of genes affected by G9a. We find that, both in ESCs and after differentiation to NPCs, a surprisingly small number of genes are significantly de-repressed when G9a is lost. Most G9a-repressed genes harbor H3K9me2 at their promoters, suggesting that the repression by G9a is direct; however, few of these genes are DNA methylated. The lack of DNA methylation at G9a-repressed promoters may be related simply to the fact that many of them have low CpG content and are poor substrates for DNA methyltransferase. However, G9a-repressed genes in NPCs did not overlap with genes whose promoter DNA methylation during retinoic acid-induced NPC differentiation was G9a dependent (9). Our differentiation system did not employ retinoic acid, but both differentiation systems enrich for NPCs, so these results suggest that many promoters that require G9a for DNA methylation during differentiation must be repressed by additional redundant mechanisms (e.g., the factors to express these genes may be missing). At these promoters, DNA methylation may act as a fail-safe means to prevent their activation rather than the primary cause of their silent state (9).

Most interestingly, we find that the set of G9a-repressed genes is highly enriched in the late-replicating peripheral compartment of the nuclear that is selectively depleted of H3K9me2 after G9a loss. In fact, we find that H3K9me2, which has been shown to correlate with nuclear lamina-associated regions (11), is highly correlated with nuclear periphery. (A) FISH of mock- vs. OHT-treated cells using BAC probes for the indicated gene loci. DNA is counterstained in blue. White arrowheads point to the FISH signals (green). (B) At least 100 FISH signals for each experimental condition were scored, and the distance of each FISH signal to the nuclear periphery was normalized to the radius of the nucleus. Data are plotted as cumulative frequency graphs; the x-axis represents the relative radial distance to the nuclear periphery, where 0 represents the center and 1 the periphery of the nucleus. Each graph shows 1 G9a-repressed gene (indicated above each graph) in mock-treated (solid black line) and OHT-treated (dashed line) cells, plotted alongside an internally localized gene (Oct4; solid gray line) that is unaffected by G9a knockout. P-values were calculated using a 2-sample Kolmogorov-Smirnov test.
with late replication. Moreover, of the 126 previously identified genes with G9a-dependent promoter DNA methylation (9), more than 90% are early replicating, and at least 1 of these promoter-genes with G9a-dependent promoter DNA methylation (9), more with late replication. Moreover, of the 126 previously identified promoters strong enough to become easily accessible even within the nuclear periphery may render these genes held in check by G9a are distinct from those whose promoters with late replication. Moreover, of the 126 previously identified promoters strong enough to become easily accessible even within the context of a generally silent compartment. Our results also demonstrate that, consistent with observations in several other chromatin modifications (reviewed in ref. 21), replication timing is remarkably resilient to even dramatic changes in the amount of specific histone modifications.

**Methods**

**Construction, Culture, and Differentiation of G9a–/– ESCs.** Construction of conditional G9a–/– ESCs from TT2 parental ESCs was described in Figure S2 of ref. 33. ESCs were cultured as described (14). All experiments were set up as follows: 106 cells were treated with 0.78 μM tamoxifen (4-OHT) or vehicle (ethanol) for 48 h and harvested 5 days later. SKY analysis was performed as a fee-for-service by the Roswell Park Cancer Institute SKY facility. Cells were differentiated 2 days after OHT or mock treatment, as described (18), the medium was changed every 2 days for 9 days.

**Fluorescence in Situ Hybridization.** 2D FISH was performed as previously described (14), except that fixation was in 3:1 methanol:acetic acid, using digoxigenin-labeled commercially available bacterial artificial chromosome (BAC) probes prepared using the PrepEase BAC purification kit (USB #78722, USB Corporation), which were collected with a DeltaVision imaging system. Measurements of the distance of FISH probes to the nuclear periphery and cumulative distance plots were made with the computer program FIISH Finder (36).

**ACKNOWLEDGMENTS.** We thank S. Ty, J. Shirley, and X. Liu for development of the FISH Finder program used to analyze subnuclear position, H. Kimura and T. Jenuwein for supplying antibodies, X. Li for help with transcriptional analysis, R. Didier for flow cytometry, and S. Matsui for SKY analysis. This work was supported by National Institutes of Health Grant GM083377 to D.M.G. and The Leukemia and Lymphoma Society Special fellowship to Y.T.