Histone H3 lysine 9 di-methylation as an epigenetic signature of the interferon response

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Type I IFN (IFN-α/β) is a potent proinflammatory cytokine that facilitates innate and adaptive immune responses against viruses and bacterial pathogens (Stetson and Medzhitov, 2006). The proinflammatory potential of IFN-α/β (hereafter defined as IFN) implies the necessity for tight control of this cytokine expression. In the absence of a clinically manifested infection, IFN levels in the organism are largely supported by plasma-cytoid DCs (Barchet et al., 2005), which are characterized by exceptionally high spontaneous or signal-induced IFN production. Heightened IFN production by plasmacytoid DCs reflects an overall potent capacity of the innate immune cells, such as macrophages, DCs, and monocytes, to transcribe IFN and IFN-stimulated genes (ISGs) in response to pathogens or pathogen-derived ligands (Diebold et al., 2003; Pietras et al., 2006; Kumagai et al., 2007). In contrast to cells of the innate immune system, parenchymal cells, as well as neurons and cardiac myocytes, must exert tight control over IFN expression to avoid tissue injury (Trinchieri, 2010).

Effective antiviral immunity depends on the ability of infected cells or cells triggered with virus–derived nucleic acids to produce type I interferon (IFN), which activates transcription of numerous antiviral genes. However, disproportionately strong or chronic IFN expression is a common cause of inflammatory and autoimmune diseases. We describe an epigenetic mechanism that determines cell type–specific differences in IFN and IFN-stimulated gene (ISG) expression in response to exogenous signals. We identify di-methylation of histone H3 at lysine 9 (H3K9me2) as a suppressor of IFN and IFN-inducible antiviral gene expression. We show that levels of H3K9me2 at IFN and ISG correlate inversely with the scope and amplitude of IFN and ISG expression in fibroblasts and dendritic cells. Accordingly, genetic ablation or pharmacological inactivation of lysine methyltransferase G9a, which is essential for the generation of H3K9me2, resulted in phenotypic conversion of fibroblasts into highly potent IFN–producing cells and rendered these cells resistant to pathogenic RNA viruses. In summary, our studies implicate H3K9me2 and enzymes controlling its abundance as key regulators of innate antiviral immunity.

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requires that lysine residues be free from other forms of posttranslational modification, in particular lysine methylation (Wang et al., 2008). Di- or tri-methylation of H3K9 is capable of preventing activation of gene expression not only passively, by prohibiting acetylation, but also actively, by recruiting transcriptional repressors of the HP1 family (Nakayama et al., 2001; Fischle et al., 2003). Therefore, it is plausible that the levels of H3K9me2 at IFN and ISG promoters may contribute to the cell type–specific differences in IFN and ISG expression.

Here, we present data that reveal H3K9 di-methylation as an epigenetic determinant of IFN and ISG expression by cells of various types. We describe a reverse correlation between H3K9me2 occupancy at IFN and ISG promoters and the ability of cells to express these genes. In mice and men, di-methylation of H3K9 is catalyzed by the G9a/GLP enzymatic complex, where G9a is essential for the catalytic function and stability of the complex (Tachibana et al., 2005). We demonstrate that ablation or pharmacological suppression of G9a and ensued deficiency in euchromatic H3K9me2 converts nonprofessional IFN-producing cells into potent IFN and ISG expressers and renders them resistant to viral infection.
RESULTS AND DISCUSSION

"Natural" H3K9me2 deficiency at Type I IFN and ISGs in DCs

Di-methylation of lysine 9 of histone H3 (H3K9me2) is widely linked to transcriptional repression of euchromatic genes and heterochromatic gene silencing (Peters et al., 2003). Although gene transcription is largely governed by upstream signals that recruit transcription factors, the abundance of H3K9me2 at individual gene promoters, including those of IFN and ISGs, could be a limiting factor in what ultimately determines the likelihood of transcription. We speculated that developmentally predetermined and/or signal-controlled H3K9me2 at Ifnα/β promoters could contribute to cell type–specific differences in IFN and possibly ISG expression. In agreement with this model, we found significantly lower levels of H3K9me2 at IFN and ISG promoters in splenic DCs as compared with mouse embryonic fibroblasts (MEFs), which have been used as an example of “nonprofessional” IFN–expressing cells. In MEFs, H3K9me2 occupies the entirety of Ifnα as well as the Mx1 ISG (Fig. 1 A). The pattern of H3K9me2 occupancy of the Ifnα and Mx1 loci is similar in DCs. However, the gene-specific levels of H3K9me2 in these cells are significantly lower than in MEFs (Fig. 1 A). Similar to the Ifnα and Mx1 loci, Ifnα4 and ISG Ifit1 were nearly depleted of H3K9me2 in DCs (unpublished data). Collectively, H3K9me2 enrichment around the promoters of >300 ISGs (Schoggins et al., 2011) was much lower in DCs than in MEFs, highlighting this difference of epigenetic patterning of antiviral genes in two cell types (Fig. 1 B). Overall, H3K9me2 levels were also low at the promoters of many known proinflammatory genes, including LPS-inducible genes (Fig. 1 C; Ramirez-Carrozzi et al., 2009; Nicodeme et al., 2010). This finding points to “natural” H3K9me2 depletion as an epigenetic hallmark of proinflammatory genes. However, in our studies we choose to focus on poly I:C–induced ISGs as a representative group of genes involved in inflammatory responses. The low abundance of H3K9me2 at IFN and ISGs in DCs is not a consequence of auto- and/or paracrine DC stimulation by IFN as supported by similarly low levels of H3K9me2 at IFN and ISGs promoters in WT and IFNαR1−/− DCs (Fig. 1 A). Reduced levels of H3K9me2 at IFN and ISG promoters in DCs are also associated with a relative enrichment of H3K4me3 at these loci (Fig. 1 B). The latter histone mark is indicative of transcriptionally poised and/or actively transcribed genes (Bernstein et al., 2002). In contrast to IFN and ISGs, neuron- or liver-specific Nog1 or Alb genes, which are silent in both DCs and MEFs, display equally high levels of H3K9me2 (Fig. 1 D, top). Accordingly, both cell types lack H3K9me2 at highly expressed housekeeping genes such as Hprt or Gapdh (Fig. 1 D, bottom).

Discordances in H3K9me2 levels at Ifnα/β and ISG promoters could also be extended to other cell types. In mice, BM-derived macrophages and DCs, as well as ex vivo isolated splenic plasmacytoid and conventional DCs, showed lower H3K9me2 levels at the Ifnβ promoter as compared with MEFs, cardiac myocytes, or neuroblastoma cells. (Fig. 1 E, left). Moreover, the disparity in H3K9me2 levels between DCs and MEFs was conserved between mice and humans, as indicated by high and low levels of H3K9me2 at the Ifnβ promoter in human fibroblasts and monocyte-derived DCs, respectively (Fig. 1 E, right).

The mechanism of the locus–specific differences in H3K9me2 levels between DCs and MEFs is not known. DCs express G9a and GLP at slightly lower levels compared with MEFs (Fig. 1 F). However, similar global levels of H3K9me2 in DCs and MEFs (Fig. 1 F), as well as equally high H3K9me2 levels at silent gene promoters in both cell types (Fig. 1 D), suggest equal capacity of DCs and MEFs to catalyze H3K9me2. Therefore, the difference in H3K9me2 levels at IFN and ISG loci in DCs and MEFs might reflect differences in cell type–specific targeting of IFN and ISG loci by G9a/GLP and/or by H3K9me2 demethylases.

G9a deficiency converts MEFs into potent IFN- and ISG-expressing cells

The low levels of H3K9me2 in DCs pointed to a potential role of natural loss of this modification in regulation of IFN and ISG expression. According to this model, removal of H3K9me2 from the Ifnα locus could result in phenotypic conversion of MEFs into DC-like IFN producers. In lieu of naturally low H3K9me2 levels at Ifnα, we expected that...
Figure 3. G9a silences Ifnβ and IFN-inducible gene expression in fibroblasts. (A) G9a WT and G9a−/− MEFs were stimulated with poly I:C and Ifnβ mRNA levels were determined by qPCR (top). The number and levels of genes in control and G9a-deficient MEFs treated with poly I:C were determined by microarray analysis (bottom). Venn diagrams show the differences in the number of genes that were up-regulated more than twofold in G9a WT and G9a−/− MEFs. Red, G9a WT; blue, G9a−/−. The hierarchical clustered heat maps show expression of the greater than twofold poly I:C-induced genes in G9a−/− as compared with G9a WT MEFs. Color bar represents log2 scale of raw signal values for each time point. Data represent triplicate samples. (B) Abundance of the chromatin marks associated with active gene transcription at Ifnb1 (left) and ISG (right) loci. The distribution and levels of epigenetic marks representing active transcription at Ifnb1 were measured by ChIP-Seq in G9a WT and G9a−/− MEFs before and after poly I:C stimulation. Scatter plots display differential abundance of transcription-associated histone marks at poly I:C-induced genes 6 h after stimulation. X- and y-axes are FPKM values for epigenetic marks in G9a WT and G9a−/− MEFs 6 h after stimulation. (C) G9a WT and G9a−/− splenic DCs were stimulated with poly I:C and Ifnβ stimulation.
H3K9me2 deficiency in DCs would not have a large impact on gene expression. To study the impact of G9a deficiency on IFN and ISG expression in MEFs, the G9a gene was conditionally inactivated by tamoxifen treatment of ERT2-cre;G9a<sup>fl/fl</sup> MEFs. G9a-deficient DCs were isolated from Vav-cre;G9a<sup>fl/fl</sup> mice, where expression of the Vav1 promoter-driven Cre resulted in G9a inactivation within the hematopoietic lineage.

Deficiency of the G9a gene in MEFs or DCs resulted in loss of G9a protein and global reduction of euchromatic H3K9me2 levels in MEFs and DCs (Fig. 2 A, left). Genome-wide analysis of H3K9me2 distribution, exemplified by the pattern of H3K9me2 abundance along chromosome 4, shows nearly complete loss of H3K9me2 at most genes in DCs and MEFs, except loci that are likely to reside within heterochromatin (Fig. 2 A, right). These results show the essential role of G9a in catalyzing H3K9me2 in DCs in a fashion that has been reported previously for G9a-deficient cells of other lineages (Tachibana et al., 2002; Peters et al., 2003; Rice et al., 2003). However, the consequences of G9a deficiency on IFN and ISGs differed significantly between these cell types. In MEFs, deficiency in G9a drastically reduced the amount of H3K9me2 at <i>I<sub>ifn<sub>β</sub></i> and ISG promoters to DC-like levels (Fig. 2 B). Because DCs already had naturally low levels of H3K9me2 at <i>I<sub>ifn<sub>β</sub></i> and ISGs, lack of G9a had little additional impact on this histone modification at these gene promoters. (Fig. 2 B).

Deficiency in G9a and associated loss of H3K9me2 had little impact on basal gene expression in either MEFs or DCs (unpublished data). However, there was a significant and selective impact of G9a deficiency on IFN and ISG expression in MEFs upon poly I:C stimulation. Triggering of G9a-deficient MEFs with poly I:C resulted in significantly stronger and faster up-regulation of <i>I<sub>ifn<sub>β</sub></i> as compared with WT MEFs (Fig. 3 A, top). In addition, more poly I:C–induced genes were expressed at higher levels in G9a-deficient MEFs as compared with poly I:C–treated control MEFs (Fig. 3 A, bottom; and Table S1). This impact of G9a deficiency was highly specific, as mutant MEFs did not display any significant changes in the pattern and levels of the genes that are not involved in the poly I:C response (Table S1). With the exception of six genes, including <i>Il15</i> and <i>Il6</i>, the expression levels of non-ISG proinflammatory genes were similar in WT and G9a-deficient MEFs (unpublished data). Earlier studies by Ramirez-Carrozzi et al. (2009) revealed several genetic and epigenetic determinants of transcriptional dynamics of proinflammatory genes in macrophages. It is plausible that the differential H3K9me2 contribution to poly I:C–induced ISGs or other proinflammatory gene activation reflects a unique feature of the transcriptional network involved in ISG activation.

The augmented up-regulation of <i>I<sub>ifn<sub>β</sub></i> and ISGs in G9a-deficient MEFs was associated with a signal-dependent enrichment of transcription-associate marks such as H3K9Ac, H3K4me3, H3K36me3, and Pol II at <i>I<sub>ifn<sub>β</sub></i> and ISG promoters (Fig. 3 B). Contrary to MEFs, deficiency of G9a in DCs had little additional impact on already robust poly I:C–induced <i>I<sub>ifn<sub>β</sub></i> and ISG expression (Fig. 3 C).

The positive impact of G9a deficiency on IFN and ISG expression was reproduced by suppression of G9a enzymatic activity (Kubiczek et al., 2007; Vedadi et al., 2011). Treatment of MEFs with either BIX1294 or UNC0638 G9a inhibitors lead to reduction of H3K9me2 at the <i>I<sub>ifn<sub>β</sub></i> locus, as well as an increase in poly I:C–induced <i>I<sub>ifn<sub>β</sub></i> expression in a fashion similar to G9a-deficient cells (Fig. 3, D and E). Furthermore, inhibition of G9a resulted in enhanced <i>I<sub>ifn<sub>β</sub></i> expression not only in MEFs but also in poly I:C–stimulated primary mouse keratinocytes (Fig. 3 E).

Enhanced poly I:C–induced IFN and ISG expression in the absence of G9a suggested a key role of G9a-mediated H3K9me2 in the regulation of IFN and ISG promoter accessibility to signal-induced transcription factors. Several lines of evidence support a chromatin-dependent role of G9a in regulation of <i>I<sub>ifn<sub>β</sub></i> and ISG expression. Deficiency of G9a did not affect poly I:C–induced phosphorylation of IRF3 as evident from appearance of the phosphorylated IRF3 species at 6 h after stimulation (unpublished data). Additionally, a nonchromatinized plasmid-based <i>I<sub>ifn<sub>β</sub></i> promoter displayed equally strong activity in both mutant and control cells (unpublished data). The levels of STAT1 phosphorylation and activation of a reporter ISG, which carries a STAT–dependent ISRE sequence, were similar in IFN-treated G9a-deficient and control cells (unpublished data). These data ruled out any significant contribution of G9a to signaling processes upstream of IFN-inducing and IFN-induced transcriptional networks. Instead, we found that G9a determines the outcome of the IFN expression even under conditions of excessive transcription factor function. Expression of a constitutively active IRF3 in MEFs resulted in higher <i>I<sub>ifn<sub>β</sub></i> expression in G9a-deficient as compared with control MEFs (unpublished data).

Previous findings had implicated G9a in the regulation of <i>I<sub>ifn<sub>β</sub></i> expression by the transcriptional suppressor Blimp1 (Keller and Maniatis, 1991; Gyory et al., 2004). According to the data, binding of G9a to Blimp1 was important for termination of <i>I<sub>ifn<sub>β</sub></i> mRNA levels were determined by qPCR. The number and levels of genes in control and G9a-deficient splenic DCs treated with poly I:C were determined by microarray analysis (bottom). Venn diagrams show the number of genes that were up-regulated more than twofold in G9a<sup>−/−</sup> and G9a<sup>+/+</sup> DCs. Red, G9a<sup>−/−</sup>; blue, G9a<sup>+/+</sup>. The hierarchical clustered heat maps show expression of the greater than twofold poly I:C–induced genes in G9a<sup>−/−</sup> as compared with G9a<sup>+/+</sup> MEFs. Color bar represents log<sub>2</sub> scale of raw signal values for each time point. Data represent duplicate samples. (D) WT MEFs were treated with pharmacological inhibitors of G9a, UNC0638 (1 μM), or BIX1294 (1 μM) for 7 d. The abundance and distribution of H3K9me2 was determined by ChIP-Seq. Y-axes represent the number of reads per million mapped reads. (E) MEFs or primary mouse keratinocytes were treated with DMSO, or with G9a inhibitors BIX1294 and UNC0638 for 7 d. Control or inhibitor-treated cells were stimulated with poly I:C and <i>I<sub>ifn<sub>β</sub></i></i> mRNA levels were determined by qPCR. Data are representative of three independent experiments. Error bars are SD.
resulted in higher ISG expression in G9a-deficient MEFs (Fig. 4 C). Enhancement of antiviral gene expression rendered G9a<sup>−/−</sup> MEFs resistant to viral infection. Sindbis or VSV viruses, engineered to express GFP to monitor the frequency of infected cells and relative viral replication levels per cell, could not replicate in G9a-deficient cells as efficiently as in control cells (Fig. 4 D).

We provide evidence for a key role of G9a-mediated H3K9me2 in the regulation of IFN expression and antiviral transcriptional program. We established a reverse correlation between the abundance of H3K9me2 at IFN or ISG promoters with the scope and amplitude of activation of these genes. We also show that H3K9me2 deficiency, which follows G9a genetic ablation or pharmacological suppression, boosts IFN and ISG expression in MEFs to high levels and renders MEFs highly resistant to viral infection. Importantly, stimulation of MEFs or other cell types with poly I:C or viruses does not dramatically change the abundance of H3K9me2 at the IFN or ISG promoters (unpublished data). Therefore, the amount of H3K9me2 at IFN or ISG could be viewed as a new developmentally installed lineage-specific epigenetic determinant in higher ISG expression in G9a-deficient MEFs (Fig. 4 C). Enhancement of antiviral gene expression rendered G9a<sup>−/−</sup> MEFs resistant to viral infection. Sindbis or VSV viruses, engineered to express GFP to monitor the frequency of infected cells and relative viral replication levels per cell, could not replicate in G9a-deficient cells as efficiently as in control cells (Fig. 4 D).

Enhanced resistance to viral infection in the absence of G9a
Enhanced poly I:C–stimulated gene expression in G9a<sup>−/−</sup> MEFs indicated a potential regulatory role of G9a and H3K9me2 in cell susceptibility to viral infection. In G9a-deficient MEFs, Ifnβ and ISG expression were greatly enhanced in response to vesicular stomatitis virus (VSV) or influenza A (Fig. 4, A and B). The heightened antiviral gene response reflects both increased production of Ifnβ and augmented responsiveness of ISGs to Ifnβ stimulation. Incubation of mutant and control MEFs with equal amounts of recombinant IFN-β resulted in higher ISG expression in G9a-deficient MEFs (Fig. 4 C). Enhancement of antiviral gene expression rendered G9a<sup>−/−</sup> MEFs resistant to viral infection. Sindbis or VSV viruses, engineered to express GFP to monitor the frequency of infected cells and relative viral replication levels per cell, could not replicate in G9a-deficient cells as efficiently as in control cells (Fig. 4 D).
of the IFN-driven innate response. Mechanistically, abundance of H3K9me2 within a nucleosome that carries two histone H3 molecules defines the number of nucleosomes and, hence, gene alleles that could be driven into active transcription by signal-induced histone H3 acetylation. Therefore, our data suggest that the ability of DCs to express high levels of IFN, as compared with MEFs, is based on a developmentally determined relatively higher number of IFN and ISG alleles accessible for transcriptional activation.

The key role of H3K9me2 in the IFN response raises a question of the role of H3K9me2 methylating and demethylating enzymes in inflammatory and autoimmune diseases that are rooted in increased IFN production. In this context, it would be important to determine the abundance of H3K9me2 at IFN and ISG promoters in monocytes from patients with systemic lupus erythematosus who display heightened levels of IFN expression (IFN signature) in the absence of stimulation (Pascual et al., 2006).

MATERIALS AND METHODS

Mice. C57BL/6 mice were obtained from Charles River. G9a−/− and G9a+/− mice were described previously (Sampath et al., 2007; Schaefer et al., 2009). Rosa26-ER73-Cre mice were provided by T. Ludwig (Columbia University, New York, NY). Vav-cre mice were obtained from The Jackson Laboratory. IFNAR1−/− mice were obtained from B&K Universal Ltd. All mice were bred onto a C57BL/6 background. Mice were housed under specific pathogen-free conditions and experimental protocols were approved by the Rockefeller University Institutional Animal Care and Use Committee.

Cells. MEFs were generated from day 12.5 or 13.5 embryos from G9a−/−; Rosa26-ER73-Cre and immortalized with SV40-LTA before deletion of G9a was induced with 4-OH Tamoxifen. MEFs were cultured in DME supplemented with 10% FBS, L-Glutamine, penicillin-streptomycin, and 2-mercaptoethanol. Cardiac myocytes were isolated from day 6 embryonic stem cells expressing α-MHC and grown in puromycin as described previously (Kolossov et al., 2006). N2A neuroblastoma cells were cultured in DME supplemented with 10% FBS, L-Glutamine, penicillin-streptomycin, sodium pyruvate, Hepes, 2-mercaptoethanol, and 100 ng/ml G418.

ChIP. Cells were fixed with 1% formaldehyde and fixation was stopped by addition of glycine. ChIP and preparation of samples for ChIP-Seq was performed as previously described (Lee et al., 2006; Nicodeme et al., 2010). Quantitative PCR was performed with a Light Cycler 480 (Roche) using SYBR green for detection of real-time transcripts. Standard curves were generated using input DNA to normalize for differences in primer efficiencies. Data are represented as the amount of ChIP DNA/Input DNA or ChIP DNA associated with Histone H3.

Analysis of ChIP-Seq. Samples were amplified onto flow cells using a standard cluster station (Illumina) per manufacturer protocol. Cluster kits were used version 4 single read cluster amplification kits. Samples were sequenced on the GAIIx platform (Illumina) for 36 cycles using version 4 sequencing kits, and raw sequencing data were processed using the onboard SCS/RTA software version 2.6 yielding 36 bp reads. Sequencing reads were aligned to the mouse genome (NCBI37/mm9) using Bowtie (Langmead et al., 2009). Reads were kept if they aligned with two errors or less over the 36 bp read length and did not align to more than one location in the genome.

Normalization of G9a−/− and G9a+/− MEF and DC H3K9me2 ChIP-Seq. As the distribution of reads between the WT and G9a−/− cells will reflect markedly different regions and levels of enrichment of H3K9me2, the typical method of normalization based purely on the number of sequencing reads obtained would not provide a useful means of equilibrating enrichment signals, and a method based on Major Satellite repeat enrichment was used. Quantitative PCR (qPCR) data indicated increases in signal for H3K9me2-associated Major Satellite enrichment in G9a−/− cells with respect to WT cells in both MEFs and DCs. Repeat analysis was performed by aligning reads to the rodep.ref repeat file from RepBase15.01, allowing two errors over 36 bp. The number of reads aligning to Major Satellite sequence was tabulated for WT and G9a−/− cells. Using the qPCR data, a target value was obtained that reflected the number of Major Satellite-associated reads expected for G9a−/− (WT multiplied by qPCR—observed increase). The ratio of observed-to-expected counts was calculated for MEF and DC datasets and used to adjust enrichment values and browser visualizations.

ChIP-Seq enrichment (FPKM). To determine the enrichment of H3K9me2, H3K4me3, H3K9Ac, and H3K36me3, the total number of reads aligning to the specified region of interest was calculated and reported as fragments per kilobase of region per million mapped reads (FPKM). This normalizes the enrichment values based on read number and length of region of interest. Regions of interest were: ±1 kb from the TSS for H3K9me2; ±3 kb for H3K4me3 and H3K9Ac; and full transcript length for H3K36me3. RefSeq data downloaded from the University of California, Santa Cruz table browser was used.

Per-gene profiles. Browser tracks were displayed using the Integrative Genome Viewer, and Made using the igtools context function set for 100 bp extensions in the 3′ direction and either 5 bp or 25 bp windows. ChIP-Seq data are accessible through Gene Expression Omnibus, accession no. GSE22102.

RNA isolation, reverse transcription, and qPCR. RNA was isolated with the RNeasy mini kit (QIAGEN). RNA was reverse transcribed using SuperScript III (Invitrogen). qPCR was performed using SYBR green and products were quantified with a standard curve. Gene expression was displayed relative to Tbp.

Microarray analysis. 1 μg of total RNA from two (DC) or three (MEF) samples per group was used to prepare biotin-labeled RNA using Ambion Illumina TotalPrep RNA Amplification kit (Applied Biosystems) and hybridized to Illumina MouseRef-8 v2.0 expression BeadChip kits. The chips were scanned using a BeadArray Reader (Illumina) followed by analysis using Genespring (Agilent Technologies). The raw expression data were quality assessed
and subjected to background adjustment and normalization. The individual gene expression levels were compared by using an unpaired Student’s t test (P < 0.05) and by pairwise comparison. The microarray datasets are available in the Gene Expression Omnibus under the accession no. GSE24826.

Stimulations and transfections. MEFs were transfected with 2 µg/ml poly I:C with Lipofectamine 2000 (Invitrogen) according to the manufacturer’s instructions. Reporter constructs were transfected with CaPO4. The IFN-β reporter was provided by T. Taniguchi (University of Tokyo, Tokyo, Japan). The ISRE reporter was obtained from Invitrogen. DCs were directly stimulated with 10 µg/ml poly I:C.

Viruses and viral infection. Stocks of GFP-expressing VSV (designated VSV-GFP M51R) were kindly provided by GlaxoSmithKline. C.M. Rice were supported by the Greenberg Medical Research Institute, the Starr Health grant R21AI077018 and the Starr Cancer Consortium. M.R. MacDonald and G9a/GLP inhibitors.

Antibodies. H3 (ab1791), H3K9me2 (ab1220), H3K9Ac (ab4441), H3K36me3 (ab9080), total Pol II (ab5408), and Pol II S5 (ab5153) antibodies were all obtained from Abcam. H3K4me3 was from Millipore (17614). Phospho-IRF3 (4947) and phospho-STAT1 (9171) were from Cell Signaling Technology. GLP (BO422) was from Perseus Proteomics. IRF3 (S1–3290) was from Invitrogen, and STAT1 (610185S) was from BD. The G9a antibody was described previously (Sampath et al., 2007).

G9a/GLP inhibitors. BIX1294 and UNC0638 were obtained from Sigma-Aldrich. Both inhibitors were used at 1 µM. Cells were incubated with inhibitor for 7 d and the media was replenished with fresh inhibitor every 2 d.

Online supplemental material. Table S1 shows genes that were upregulated more than twofold in G9a−/− and G9a+/+ MEFs stimulated with poly I:C at 2, 6, and 12 h. Online supplemental material is available at http://www.jemb.org/cgi/content/full/jem.20112343/DC1.

We are grateful to Tada Taniguchi, Adolfo Garcia-Sastre, Dan Stetson, and David Levy for many valuable reagents. We thank Shihari Sampath for the initial discussions about the role of G9a in antiviral responses; Angela Santana for technical assistance; the Genomics Resource Center for performing microarray and sequencing; the Flow Cytometry Resource Center for cell sorting; and Rebecca Rizzo for editorial assistance.

T.C. Fang was supported by the Irvington Institute Fellowship Program of the Cancer Research Institute. A. Tarakhovsky was supported by National Institutes of Health grant 2R1AI077018 and the Starr Cancer Consortium. M.R. MacDonald and C.M. Rice were supported by the Greenberg Medical Research Institute, the Starr Foundation, and National Institutes of Health grant R01 AI091707. A. Stienen was supported by the Medical School RWTH Aachen University.

I. Rioja, V. Paravcini, R.K. Piriija, and K. Lee are employees of GlaxoSmithKline. Research support, including ChIP-Seq analysis, supplies, and other expenses, was provided by GlaxoSmithKline.

Submitted: 4 November 2011
Accepted: 14 February 2012

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