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Interferon Stimulation Creates Chromatin Marks And Establishes Transcriptional Memory

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Abstract

Epigenetic memory for signal dependent transcription has remained elusive. So far, the concept of epigenetic memory has been largely limited to cell-autonomous, preprogramed processes such as development and metabolism. Here we show that interferon β (IFN β) stimulation creates transcriptional memory in fibroblasts, conferring faster and greater transcription upon restimulation. The memory was inherited through multiple cell divisions and led to improved anti-viral protection. Of $\sim 2,000$ IFN β stimulated genes (ISGs), about half exhibited memory, which we define as memory ISGs. The rest, designated non-memory ISGs, did not show memory. Surprisingly, mechanistic analysis showed that IFN memory was not due to enhanced interferon signaling, or retention of transcription factors on the ISGs. We demonstrated that this memory was attributed to accelerated recruitment of RNA polymerase II and transcription/chromatin factors, which coincided with acquisition of the histone H3.3 and H3K36me3 chromatin marks on memory ISGs. Similar memory was observed in bone-marrow macrophages after IFN γ stimulation, suggesting that IFN stimulation modifies the shape of innate immune response. Together, external signals can establish epigenetic memory in mammalian cells that imparts lasting adaptive performance upon various somatic cells.

Significance Statement

Epigenetic memory for experience-based gene expression has not been well studied in higher organisms. Here we demonstrate that cells previously exposed to interferons exhibit a memory response, and mount faster and higher transcription upon restimulation in fibroblasts and macrophages. Genome-wide analysis showed that memory was ascribed to accelerated recruitment of transcription factors to the genes. This process rested upon a distinct chromatin state involving the histone H3.3 and H3K36 modification. Our findings provide mechanistic framework for the previously proposed idea of "trained innate immunity" representing memory, independent of adaptive immunity. Together, this study highlights learning as a fundamental faculty of mammalian somatic cells.

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Lineage specific gene expression in differentiated cells is somatically inherited through epigenetic mechanisms, conveyed in part by chromatin(1-3). In particular, enhancer DNAs and associated chromatin modifications, such as H3K4me1 and H3K27ac have a large impact on lineage specification and extension(4-7). In addition, the replacement histone, H3.3 contributes to epigenetic memory during transition to embryonic pluripotency(8-10). H3.3 is deposited in the genome by coupling with transcription and distributed across actively expressed genes(11-15).

Epigenetic memory for signal dependent transcription has not been well studied in multi-cellular organisms, although transcriptional memory has been reported in yeast following environmental cues (16-19). Nutrient response genes in yeast, such as GAL1 and INO1, are induced more rapidly and at higher levels when cells had previously experienced nutrient signaling. This trait is transmitted across cell generations and may involve histone H2A.Z and other pathways(20-22).

Given the conservation of transcriptional processes in the eukaryotes, one can assume that the mechanism of signal dependent memory has also withstood evolutionary selection. An impetus for studying transcriptional memory for interferon (IFN) stimulated genes (ISGs) came from our previous observation that ISGs accumulate histone H3.3 as they are transcribed after IFN treatment(10, 23, 24). IFNs, both type I IFN (α/β) and type II IFN (IFN γ) stimulate many ISGs in various cell types and enhance anti-microbial activity (25-27). Here we show that IFN stimulation confers transcriptional memory that permits faster and greater ISG transcription in mouse embryonic fibroblasts (MEFs) and bone-marrow (BM) derived macrophages. The memory was attributed to faster and greater recruitment of phospho-STAT1 and RNA polymerase II (Pol II), but not Pol II pausing. Further, memory establishment coincided with acquisition of chromatin marks by H3.3 and H3K36 trimethylation. This study highlights a previously ill-defined epigenetic process through which external signals give rise to transcriptional memory that endows adaptive behavior in response to changing external milieu.

IFN β Stimulation Generates Heritable Transcriptional Memory

To investigate whether IFN affords epigenetic memory, MEFs were treated with IFN β (hereafter IFN) for 3 h or 24 h, washed, and left without IFN for 24 h. Pretreated cells were then restimulated with IFN, and ISG induction was compared with naïve cells that were not treated with IFN before (See experimental diagram in Fig. 1a, top). Time course analysis in Fig. 1a and SI Appendix, Fig. S1a showed that typical ISGs, such as Mx1, Ifit1, Oas1a, and others were induced faster and at higher levels in pretreated cells than naïve cells. Another ISG, Irf1 and constitutive Gtf2b did not (see below also). Thus, pretreated cells exhibited a memory response similar to that in yeast. To assess the duration of memory, pretreated cells were left without IFN for up to 96 h and then restimulated to examine memory. Mx1, Ifit1, Oas1a and other ISGs maintained memory over this period (Fig. 1b and SI Appendix, Fig. S1b). Moreover, Oas1a and Mx1 retained memory farther, for up to eight days (Fig. 1c). IFN memory was also detected with nascent ISG transcripts, showing that this memory response represents *de novo* transcription, rather than mRNA carry-over (SI Appendix, Fig. S1c). Some ISGs, such as Irf1, did not exhibit memory, in that induction levels were similar in naïve and pretreated cells, indicating variability in memory formation (Fig. 1a,b). Additionally, the memory response was

not detected in constitutively expressed genes that did not respond to IFNs (e.g., Gtf2b in Fig. 1a,b). Flow cytometry analysis of CFSE stained cells showed that both naïve and pretreated cells divided multiple times equally up to 8 days, demonstrating that IFN memory was transmitted across cell divisions (Fig. 1d). Other MEF cell lines and NIH3T3 fibroblasts displayed similar memory responses after IFN stimulation (SI Appendix Fig. S1d). Thus, IFN stimulation generates epigenetic memory that confers some ISGs faster and greater transcription upon restimulation.

To assess the functional significance of the memory, we examined IFN's anti-viral activity. Naïve and memory cells were infected with encephalomyocarditis virus (EMCV) with or without brief IFN treatment. Naïve cells underwent extensive cytopathic death after infection, irrespective of IFN treatment, as detected by sparse crystal violet staining (Fig. 1e and SI Appendix, Fig. S1e). In contrast, most of pretreated cells remained viable upon viral infection. Accordingly, the EMCV viral titers in supernatants were >10 fold higher in naïve cells compared to pretreated cells. Thus, IFN memory provides improved antiviral activity, suggesting the ability to create adaptive response.

IFN β activates JAK/STAT signaling pathway to elicit its biological activity(25). To assess whether the memory is ascribed to changes in IFN signaling, we tested expression of IFNAR1 receptor and phosphorylated (p) STAT1, critically required for ISG transcription. Flow cytometry and Western data (Fig. 1f,g) showed that expression of IFNAR1 and pSTAT1 was virtually identical in naïve and pretreated cells. These data support the view that IFN memory is not attributed to substantial changes in IFN signaling, but to a subsequent event(s).

Memory Response Is Selective: RNA-seq Analysis

To globally identify ISGs that gain memory after IFN treatment, RNA-seq analysis was performed for naïve and pretreated MEFs that were restimulated by IFN up to 6 h (Fig. 2a). About 2,000 ISGs were upregulated or downregulated by IFN both in naïve and pretreated cells (SI Appendix, Fig. S2a and Fig. S2b for GO analysis)(28). Of upregulated ISGs, approximately half (1,057 genes) displayed a typical memory response, showing faster and/or higher expression in pretreated cells at some point during 6 h of IFN stimulation (Fig. 2b). Of these, 546 ISGs showed strong memory (RPKM levels >2 fold higher in pretreated cells than naïve cells). The rest of 511 ISGs showed somewhat weaker memory where mRNA levels in pretreated cells were higher than naïve cells, but by less than two-fold. On the other hand, about 600 ISGs did not show memory, and were classified as non-memory ISGs, since their RPKM levels were similar in naïve and pretreated cells. This group included Irf1 that did not show memory in the initial qRT-PCR analysis (see Fig. 1a,b). Moreover, additional 275 genes were designated refractory ISGs, as they were induced in naïve cells but not in pretreated cells. qRT-PCR data in SI Appendix, Fig. S2c confirmed the loss of induction in pretreated cells for these ISGs. This acquired unresponsiveness is analogous to "endotoxin tolerance" caused by bacterial lipopolysaccharide (LPS)(29-31). GO analysis showed that memory and non-memory ISGs share overlapping and related categories, such as innate immune- and defense-responses (SI Appendix, Fig. S2d). In contrast, GO categories enriched in refractory ISGs were unrelated to those in memory and non-memory ISGs, pointing to negative transcriptional regulation, indicating distinct functional traits for this group of ISGs. Kinetic profiles of memory

ISG expression in Fig. 2c highlighted faster transcript induction in pretreated cells compared to naïve cells. A fraction of ISGs showed somewhat elevated expression levels at 0 h in pretreated cells, suggestive of carry-over transcripts. Nevertheless, their expression greatly exceeded that in naïve upon subsequent restimulation, verifying authentic memory response. Thus, IFN stimulation imparts transcriptional memory to more than 1,000 ISGs. Further, this memory response varied in different ISGs, revealing selectivity in memory formation. These data indicate that the second IFN stimulation does not produce a carbon copy of the first response, but it likely imposes reprogramming of gene expression patterns.

Pol II Is Recruited to Memory ISGs Faster and at Higher Levels Upon Restimulation

RNA polymerase II (Pol II) is paused near the transcription start sites (TSS) of various genes(32, 33). Some genes with paused Pol II, such as heat shock genes and LPS responsive genes are induced rapidly after stimulation(34-36). However, we previously observed that many ISGs do not have paused Pol II prior to IFN, rather Pol II is recruited after stimulation along with binding of phosphorylated STAT1 and BRD4(24, 37). Here we asked whether IFN memory is attributed to a change in the Pol II binding status, from non-paused to paused state. qChIP analysis of memory ISGs (Mx1, Ifit1 Oas1) showed virtually no Pol II on the TSS prior to restimulation both in naïve and pretreated cells (Fig. 3a). However, upon restimulation, Pol II was recruited to the ISGs faster in pretreated cells than naïve cells. In contrast, Irf1, a non-memory ISG had sizable amounts of prebound Pol II in both naïve and pretreated cells where Pol II was recruited similarly after restimulation. As expected, Pol II binding on Gtf2b was not affected by IFN treatment. Likewise, memory ISGs had neither prebound pSTAT1, nor prebound BRD4. But these factors were recruited faster in pretreated cells upon restimulation (Fig. 3b,c). Mirroring the Pol II data, pSTAT1 and BRD4 binding was very similar for the non-memory ISG, Irf1 in naïve and pretreated cells. These data support the possibility that accelerated Pol II recruitment, rather than Pol II pausing accounts for IFN memory.

To further evaluate this possibility, ChIP-seq was performed for global Pol II occupancy on memory and non-memory ISGs (Fig. 3d and SI Appendix, Fig. S3a,b). Consistent with the above qChIP data, little to no prebound Pol II was detected on memory ISGs prior to IFN restimulation, both in naïve and pretreated cells. After 30 min of IFN restimulation, however, Pol II was recruited at higher levels in pretreated cells than in naïve cells. Thus, memory ISGs lose Pol II after IFN washout and are without it until restimulation. In contrast, non-memory ISGs were already bound by Pol II prior to IFN stimulation in both naïve and pretreated cells (Fig. 3d and SI Appendix, Fig. S3b). Interestingly, refractory genes also had prebound pol II (SI Appendix, Fig. S3c). Constitutively expressed genes had constitutive Pol II binding, whereas silent genes were without Pol II, as expected (Fig. 3d). Together, memory and non-memory ISGs showed contrasting Pol II binding patterns. These results show that memory and non-memory ISGs differ in the Pol II binding status and that acquisition of IFN memory is not due to conversion of Pol II binding status, but to enhanced factor accessibility.

Memory ISGs Are Marked by the Histone H3.3 and H3K36me3

Given that Pol II and other transcription factors do not remain on memory ISGs after IFN washout, but gain improved accessibility, IFN memory information may be stored elsewhere,

presumably in the chromatin milieu. To identify chromatin marks that coincide with IFN memory, we investigated distribution of replacement histones, H3.3 and H2A.Z, since they are reported to play a role in epigenetic memory(8, 9, 20). The H3K36me3 mark was also tested, since we previously found that IFN stimulation triggers H3K36me3 marking on ISGs along with H3.3 deposition(23, 24). ChIP-seq analysis was performed for these marks in naïve and pretreated cells without restimulation (Fig. 4a-c). For H3.3, we analyzed MEFs derived from mice in which the endogenous H3f3b was replaced by the HA-tagged H3.3 cDNA. The HA-tag on H3.3 allowed us to detect H3.3 distribution with higher reliability than using anti-H3.3 antibodies. Memory and non-memory ISGs exhibited disparate H3.3 distribution patterns. Memory ISGs did not have H3.3 in naïve cells, but gained this mark in pretreated cells. Conversely, non-memory ISGs were already marked with H3.3 in naïve and pretreated cells. To delineate whether the gain of H3.3 mark is important for memory response, we knocked down H3.3 in pretreated cells by siRNA during 48 h of IFN washout (SI Appendix, Fig. S4a, b). H3.3 specific siRNA, but not control siRNA significantly reduced IFN memory response, as evidenced by reduced ISG mRNA expression and reduced Pol II recruitment. However, H3.3 siRNA, when added to naïve cells, did not inhibit ISG induction as much under these conditions. These data are in line with the idea that H3.3 deposition contributes to the acquisition of functional IFN memory. We also observed reduced ISG induction in cells stably expressing H3.3 shRNA relative to control shRNA (SI Appendix, Fig. S4c). The H3K36me3 mark displayed the same feature, showing a clear dichotomy between memory and non-memory ISGs. Thus, memory ISGs acquired H3.3 and H3K36me3 marks only after initial IFN stimulation, while non-memory ISGs possessed these marks without IFN stimulation. Like Pol II binding, refractory ISGs also possessed both H3.3 and H3K36me3 (SI Appendix, Fig. S4h). As expected, both marks were present in constitutively expressed genes, irrespective of IFN, whereas they were absent in silent genes, (Fig. 4a,b). In contrast, H2AZ did not show an obvious correlation with ISG memory: H2AZ showed a sharp peak at the TSS of all ISGs and constitutively expressed genes (Fig. 4c-e, and SI Appendix, Fig. S4d-h) (12). Other histone marks representing expressed genes such as H3K4me3 and H4ac did not show a clear correlation with memory, although H3K4me1, an enhancer mark showed a modest correlation (SI Appendix, Fig. S4i-p).

IFN γ Stimulation Generates Transcriptional Memory in Macrophages

To test if memory is generated in other cells, we next examined IFN γ stimulated BM derived macrophages. IFN γ activates another JAK/STAT pathway distinct from IFN β to establish anti-microbial activity(25, 26). Macrophages are a major cell type responsible for innate immunity in the body and unlike MEFs, they are post-mitotic under normal conditions. Naïve and pretreated macrophages were stimulated with IFN γ , and tested for ISG expression (experimental diagram in Fig. 5a). Initial qRT-PCR analysis found that some ISGs were expressed higher in prestimulated macrophages, indicative of IFN memory (SI Appendix, Fig. S5a). Microarray analysis found that naïve macrophages expressed about 425 ISGs (SI Appendix, Fig. S5b and SI Appendix, Fig. S5c.d for GO analysis). Of these, 66 ISGs were deemed memory ISGs, as their expression was at least >1.5-fold higher in prestimulated macrophages than naïve ones (Fig. 5b, e.g. Gbp), while 251 ISGs were judged non-memory

ISGs, as their levels were similar in naïve and prestimulated macrophages (e.g., *Irf1* in SI Appendix, Fig. S5a). In addition, as many as 108 ISGs were found refractory genes, in that their expression was abrogated or markedly lower upon restimulation (e.g., *Il12d* in SI Appendix, Fig. S5a). Clustering profiles in Fig. 5c illustrate that memory ISGs were expressed earlier and/or higher in prestimulated macrophages than naïve cells, whereas, refractory ISGs were not induced or reduced in prestimulated macrophages. Some memory and non-memory ISGs were found both in MEFs and macrophages (e. g., *Mx1*, *Irf7*, *Ifi44* for memory, *Irf1* for non-memory ISGs), suggesting a common feature, while others were unique to IFN γ and/or macrophages (*Nos2*, *Il12br*, *Ciita*, *Tlr11*). GO analysis revealed that memory and non-memory ISGs have similar processes and pathways related to immunity and defense (SI Appendix, Fig. S5d). In contrast, refractory ISGs showed distinct categories such as biosynthesis and transcription, a feature similar to MEFs. These results indicate that IFN γ pretreatment alters the overall character of macrophage host defense programs.

To assess if IFN memory generated in macrophages and fibroblasts share a common mechanism, we tested the status of Pol II recruitment. ChIP-Seq analyses were performed for global distribution of Pol II along with H3.3 and K3K36me3 in naïve and pretreated macrophages, without restimulation. For Pol II, binding was tested 1 h after IFN γ stimulation as well. IGV examples of memory ISGs (*Gbp* cluster), and non-memory ISG (*Il4ra*) (Fig. 5d, upper and lower panels) showed a remarkable similarity with those in fibroblasts. For *Gbp* genes, Pol II was absent before stimulation, but vigorous binding ensured after stimulation.

Global Pol II distribution shown Fig. 5e largely reproduced the pattern in fibroblasts, in that Pol II was already bound to non-memory and refractory ISGs even in naïve cells, while memory ISGs were without prebound Pol II. We noted that a fraction of memory ISGs exhibit prebound Pol II, suggesting contribution of paused Pol II to memory or to other aspects (38).

The H3.3 and H3K36me3 marking pattern found in macrophages were also similar to those in fibroblasts MEFs: memory ISGs gained these marks after IFN γ stimulation, whereas these marks were present in non-memory and refractory ISGs even in naïve cells before IFN γ stimulation (Fig. 5f,g, and SI Appendix, Fig. S5e-g). Other histone mark, H3K4me1 showed a modest correlation (SI Appendix, Fig. S5e-h). Together, transcriptional memory, generated by different IFNs in different cell types, exhibited similar molecular features, underscoring a shared mechanism and biological significance.

Discussion

IFN stimulation generated lasting transcriptional memory in MEFs and macrophages, conferring faster and higher ISG induction upon restimulation. This memory was inherited through mitoses in MEFs, while sustained in a post-mitotic state in macrophages. The memory led to improved resistance to viral infection, despite that not all ISGs acquired memory. The memory was largely attributed to accelerated Pol II recruitment along with accelerated binding of phosphor-(p) STAT1, rather than Pol II pausing, which were associated with acquisition of H3.3 and H3K36me3 marks.

A salient feature of IFN memory was faster ISG induction both in MEFs and macrophages. Faster induction is also a main characteristic of transcriptional memory in

yeasts(16-19, 39). IFN memory we observed here also share common features with that reported for HeLa cells and mouse macrophages upon IFN γ stimulation(6, 38).

Genome-wide analyses revealed three types of ISGs, memory, non-memory and refractory ISGs in both cell types, highlighting non-uniformity among ISGs. Memory and non-memory ISGs shared a number of GO categories that were mostly related to innate immune responses in both cell types. In contrast, GO categories for refractory ISGs were unrelated to innate immunity. This indicates that the memory response is not a simple repeat of ISG expression, but represents readjustment of gene expression programs in the cell to accommodate changing environments.

Pol II binding status in memory ISGs differed from that in non-memory and refractory ISGs in both cell types. Pol II occupancy was promptly lost from memory ISGs after IFN washout, while some prebound Pol II was present on non-memory and refractory genes even without IFN. Likewise, p-STAT1 was recruited to memory ISGs only after IFN stimulation in both naïve and pretreated cells. Thus, Pol II pausing and STAT1 signaling are not likely to be a major mechanism of memory. However, paused Pol II was shown to be associated with memory in some genes in yeast and IFN γ treated HeLa cells(38). This report and our results are not mutually exclusive, since multiple mechanisms likely contribute to transcriptional memory. Indeed, prebound Pol II was detected in a small fraction of memory ISGs in macrophages in this study as well. Given that Pol II was prebound on non-memory ISGs and refractory ISGs, it is possible that these ISGs have already been epigenetically engaged before IFN stimulation.

Acquisition of IFN memory correlated with the histone H3.3 and H3K36me3 chromatin marks in both MEFs and macrophages. On memory ISGs, these marks were absent in naïve cells but accorded after IFN stimulation. Our study identifies H3.3 as a chromatin signature that denotes IFN memory. Our data that H3.3 knockdown during IFN washout decreased memory response may support the possibility that this mark has a functional importance. With an additional correlation found between FN memory and H3K4me1, albeit weaker, IFN memory may be associated with the recently proposed “latent enhancers”(6, 7). While constitutive and poised enhancers, which already carry Pol II and certain chromatin marks, latent enhancers do not have prebound Pol II, but is associated with cytokine stimulation(6). H3.3 deposition was another memory feature found in our study, suggesting that memory acquisition is linked to histone replacement. The role of H3.3 histone replacement in transcriptional memory was reported during iPS transition in frog embryos(9, 10).

There is growing evidence that innate immune responses have a memory aspect, termed “trained” innate immunity, that is independent of classical immunological memory, which depends on lymphocyte-mediated adaptive immunity (40, 41). For example, earlier non-lethal infection with *Candida albicans* provides resistance to the subsequent lethal infection in mice (42). This memory depends on cells of the monocyte/macrophage lineage and is associated with changes in chromatin. Trained Innate immunity to candida is shown to require intact STAT1, but not STAT3(30). Given that ISG induction depends on STAT1, we surmise that trained innate immunity may partly include IFN memory described here. Also, BCG that stimulates IFN γ response is shown to provide protection against heterologous, viral or bacterial infection(43). In contrast to IFNs which increase innate protection, LPS exposure, after a brief

burst of inflammatory responses, leads to a period of profound tolerance and immunosuppression(30). While detailed mechanisms are still elusive, LPS tolerance is shown to depend on the transcription factor ATF7(44). LPS tolerance likely affects the course of various infectious diseases, particularly sepsis, a prevalent, but hard-to-treat disease. It is interesting to note that since ISGs are induced after LPS through IFN β dependent feedback, LPS tolerance may well be modulated by aspects of IFN memory.

In conclusion, this study offers new insight into signal-induced epigenetic memory that conveys enhanced adaptive performance upon ordinary cells.

Materials and Methods

Cells and treatment.

MEFs and BM derived macrophages were prepared from wild type and mutant mice in which the H3f3b locus was replaced by a H3.3-HA cDNA. Details of the derivation of the mouse strain will be presented elsewhere. The procedures used in this work were approved by the NICHD animal care and use committee with the ASP # 14-044. MEFs were prepared from day 13.5 embryos and maintained in Dulbecco's minimal essential medium (DMEM) with 10% fetal bovine serum (FBS) and antibiotics, and treated with 100 U/ml of murine recombinant IFN β (PBL Interferon Source) for indicated periods. BM derived macrophages were cultured in DMEM/F12 medium containing 10% FBS, 20% L929 conditioned medium as a source of M-CSF, 10% glutamine, and antibiotics treated with 100 U/ml of murine recombinant IFN γ (Invitrogen) for indicated periods (45).

For EMCV infection, MEFs treated with or without IFN β for 3 h were infected with EMCV (Moi =10) in serum-free and antibiotics-free medium for 1h at 37°C, washed and allowed to proceed in culture for 24h. Cells were fixed with methanol and incubated with crystal violet solution (1% crystal violet, 10% ethanol) for 20 min at room temperature. For viral titration, plaque assays were performed with L929 cells incubated with serial dilutions of culture supernatants (10^{-2} ~ 10^5) followed by incubation with agar overlay for 24h at 37°C. The plaques were visualized by staining with a 0.02% neutral red. The viral titers were expressed as plaque forming units (pfu).

Cell proliferation assay was carried out with 1.5 μ M CellTrace CFSE (Molecular Probes) according to the manufacturer's instructions and fluorescent signals were detected by flow cytometry in FACSCalibur (BD) with the Flow Jo Software.

ISG mRNA, ISG nascent transcripts detected by qRT-PCR with appropriate primers (24, 37) listed in SI Appendix, Dataset S3. pSTAT1, total STAT1 and surface IFNAR1 were detected by immunoblot assay and flow cytometry, respectively. Antibody for murine pSTAT1 was from Santa Cruz Biotechnology (sc-417), STAT1 Cell Signaling Technology (#9172), and PE anti-mouse IFNAR1 from Biolegend (#127311).

For siRNA transfection to knockdown H3.3, negative control (si-Ctrl) siRNA and siRNA for H3.3a and H3.3b were transfected into MEF cells with Lipofectamine 2000 (Invitrogen) for 48 h prior to IFN stimulation. H3f3a and H3f3b siRNA oligomers were purchased from Life Technology. siRNA sequences tested here are for H3f3a : Sense GAUUGCGAGUGGAAACAUAAtt Antisense UAUGUUUCCACUCGCAAUCat (5'-4'). Si RNA ID#: s234173M. And for H3f3b: Sense CAGCAUCAUCUAUUACUAAtt Antisense UUAGUAAUAGAUGAUGCUGgt. si RNA ID#: s67338 and): Sense CCCAGGAUUUCAAACCGAtt Antisense UCGGUUUUGAAAUCUGGGcg, si RNA ID#: s67336.

Knockdown of H3.3 were performed by lentiviral vectors and transduction or siRNA transfection. The shRNA expressing lentiviral vector to knockdown both H3f3a and H3f3b genes was constructed with pLKO.1-puro vector and same shRNA sequence as described previously (23). pLKO.1-puro control shRNA vector (CAACAAGATGAAGAGCACCAA) was used as a control. The shRNA lentiviruses were produced by transient transfection of HEK293T cells with a pLKO.1-puro vector and the packaging plasmids using

Polyethyleneimine following the standard procedure. Lentiviral supernatants were collected at 24 and 48 h and MEF cells were transduced with lentiviruses with 8 µg/ml Polybrene (Sigma).

RNA-seq.

Total RNA was obtained from TRIzol (Invitrogen) and purified by the RNeasy mini kit (Qiagen). Oligo(dT) selection was performed by using Dynal magnetic beads (Invitrogen) according to the manufacturer's protocol. mRNAs were fragmented by heating at 94 °C for 3 min in the fragmentation buffer (40 mM Tris-acetate (pH 8.2), 100 mM potassium acetate and 30 mM magnesium acetate). RNA fragments were precipitated with GlycoBlue (Ambion) as a carrier and reverse transcribed by SuperScript II reverse transcriptase (Invitrogen) with random primer and RNasin (Promega). Second-strand cDNAs were synthesized using *E. coli* DNA polymerase I and RNaseH (Invitrogen). Purification of second-strand cDNAs was performed with ZYMO DNA clean & concentrator-5 kit. Library preparation, including ligating barcode was conducted using a Mondrian SP (NuGEN Technologies Inc.) and the Ovation SP Ultralow Library system (NuGEN). Fragments ranging from 250 to 450 b.p. were selected and subjected to paired-end sequencing on a HiSeq2000 (Illumina). For RNA-Seq analysis, paired-end reads were aligned to the *Mus musculus* reference genome mm10 using Bowtie/Tophat allowing up to two mismatches. Transcript abundance was quantified using Cufflinks 1.2.1. Genes showing an RPKM of more than 2-fold increase in treated samples at least one time point during IFN treatment (1, 2, 4, 6 h) over untreated samples (0 h) were considered upregulated genes (ISGs) and were analyzed further. Raw data files are available at the NCBI Gene Expression Omnibus (GEO) server under the accession number 89440.

qChIP and ChIP-seq.

qChIP was performed essentially as described (24, 37). Chromatin (4×10^5 cells) was incubated with antibodies bound to Dynabeads protein G (Invitrogen) for 3 h, followed by immunoprecipitation, de-crosslinking, DNA purification and qPCR. The following antibodies were used: Antibody for RNA polymerase II (8WG16) obtained from Covance, antibodies for HA (ab9110), H3 (ab1791), H3K36me3 (ab9050), H3K4me1 (ab8895), H3H4me3 (ab8580) were from abcam, antibody for H4Ac antibody (06-866) was from Millipore. BRD4 was described (37) Primer sequences are listed in SI Appendix, Dataset S3.

For ChIP-seq, cells were crosslinked with 1% formaldehyde for 10 min and nuclei were pelleted after sequential wash and resuspended in sonication buffer (10 mM Tris-HCl, 1 mM EDTA, 0.1% SDS, protease inhibitor cocktail) and sonicated in a Bioruptor (Qsonica) to shear the chromatin into 200-300 b.p. fragments. Chromatin from $2-10 \times 10^6$ cells was used for each ChIP-seq experiment. Chromatin was incubated with antibody and Protein G beads (Dynabeads) at 4 °C for overnight, washed sequentially with RIPA buffers. Immunoprecipitated chromatin was de-crosslinked and DNA purified. ChIP-seq libraries were constructed using a Mondrian SP (NuGEN Technologies Inc.) and the Ovation SP Ultralow Library system (NuGEN). Fragments ranging from 250 to 450 b.p. were selected for single-read sequencing on a HiSeq2000 (Illumina). Resulting reads were aligned to the mouse reference genome (mm10) using BWA(Li H. and Durbin R. (2009)) with default parameters. Uniquely mapped

reads were retained for downstream analysis. Peak calling was performed using SICER (Bioinformatics 25:1952-1958). SICER was used with window size setting of 200 b.p., a gap setting of 600 b.p., and FDR setting of 0.001. Data from immunoprecipitated samples were compared to those with input DNA. Raw data files are available at the NCBI Gene Expression Omnibus (GEO) server under the accession number 89440.

Microarray analysis of BM macrophages.

Macrophages were stimulated with 100 U/ml IFN γ and total RNA purified with the RNeasy mini kit (Invitrogen), and then subjected to microarray analysis using the Mouse Exon 1.0 ST Array (Affymetrix) essentially as described (46). Quality analysis of total RNA, cDNA synthesis, hybridization, and data extraction were performed at Expression Analysis, Inc. Data were analyzed using the GeneSpring software (Silicon Genetics). Values with $p \leq 0.05$ and two-fold cut-off were considered significant. ANOVA was used to identify differentially expressed genes. Genes showing more than twofold increase or decrease with $p < 0.05$ either at 3 or 6 h after IFN γ treatment were considered up- or downregulated genes. Among upregulated genes (ISGs), those showing >1.5-fold higher expression ($p < 0.05$) in pretreated macrophages over naïve macrophages were selected as memory ISGs. Genes whose expression levels did not differ (<1.5-fold difference) in pretreated and naïve cells non-memory ISGs. ISGs that did not increase the expression (<1.5 over 0 h) in prestimulated macrophages after 3 or 6 h of IFN γ were classified as refractory ISGs. Raw data files are available at the NCBI Gene Expression Omnibus (GEO) server under the accession number 89440.

Author Contributions

K.O. conceived the project, steered the work throughout. K.O. and R.K. wrote the manuscript with the help of other authors. J.Z. directed genome-wide analyses and gave critical advice. R.K., W.Y., and Y.Z., planned and executed experiments, and performed bioinformatics analysis. R.K. prepared the main and supplemental Figures. R.O., M.P., and A.D. performed additional experiments. Y.Y. and Y.W. instructed on sequencing and data analysis. K.S., T.F., and T.T. gave technical and conceptual guidance to this project.

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Figure Legends:

Fig. 1. IFN β Stimulation Generates Transcriptional Memory in MEFs

(a) Top: experimental design. Naïve and pretreated MEFs were first treated with vehicle alone or 100 U/ml of IFN β (IFN) or for 24 h, respectively, washed and incubated without IFN for 24 h. These MEFs were then stimulated with IFN (100 U/ml) for indicated times (h).

Bottom: ISG mRNAs were measured by qRT-PCR, normalized by Gapdh and expressed as fold induction. Mx1, Ifit1 and Irf1 are examples of memory response and non-memory response, respectively. Constitutively expressed Gtf2b was run as a control. The data represent the mean of three independent experiments $\pm$ S.D. Statistically significant differences are indicated (Student's t test, * p <0.05, ** p <0.01).

(b) Naïve and pretreated cells were left without IFN for indicated times (h) and stimulated with IFN for 3 h. ISG mRNAs were measured as above. See SI Appendix, Fig. S1c for nascent ISG mRNA.

(c) Naïve and pretreated cells were left without IFN for 196 h (8 days), stimulated with IFN for 1 h or 3 h, and ISG mRNAs were measured as above. The data represent the average of three independent experiments $\pm$ S.D. Statistically significant differences are indicated (Student's t test, * p <0.05, ** p <0.01).

(d) Naïve and pretreated cells were labeled with CFSE, treated with or without IFN for 24 h, washed and then incubated without IFN for indicated days. CFSE staining was detected by flow cytometry. Repeated cell divisions were detected for in three separate 96 h washout experiments.

(e) Naïve and pretreated cells were treated with IFN for 2 h and infected with EMCV for 24h (moi=10). Cell survival was assessed by crystal violet staining (top), and EMCV viral titers in the culture supernatants were determined by plaque assay (bottom). Similar data were obtained in three independent assays. This applies to Figure 1f and 1g below.

(f) Naïve and pretreated cells were stimulated with IFN for 6 h and IFNAR1 surface expression was detected by flow cytometry using anti-mouse IFNAR1 antibody.

(g) Naïve and pretreated cells were stimulated with IFN for indicated times (h) and expression of pSTAT1 and STAT1 was detected by immunoblot analysis of whole cell extracts using corresponding antibodies. β -actin was used as a loading control. Uncropped western blots are presented in SI Appendix, Fig. S1f.

Fig. 2. Memory Response is Selective RNA-seq Analysis

(a) Experimental design for RNA-seq. Naïve and pretreated cells were stimulated with IFN for 0,1, 2, 4 and 6 h. ISGs were defined by those genes showing >2-fold higher transcript expression (RPKM) at any time point during 6 h of IFN treatment in naïve cells.

(b) Total memory ISGs (1057) were subdivided into two groups. *Strong memory*: ISG mRNA levels were more than 2-fold higher in pretreated cells than naïve cells (>2 in RPKM). *Weak memory*: ISG mRNA levels were more than 1.5-fold higher (>1.5 in RPKM), but less than 2-fold (<2 in RPKM). Non-memory ISGs were those with similar expression levels in naïve and pretreated cells (difference less than 1.5-fold (<1.5 in RPKM). Refractory ISGs showed more than 1.5-fold lower levels of expression in pretreated cells than naïve cells (>1.5 in RPKM).

See qRT-PCR examples in SI Appendix, Fig. S2c, GO analysis in SI Appendix, Fig. S2d. Expression data for individual ISGs are available on the website. Strong memory ISGs and Non-memory ISGs were subjected to further analysis. List of Memory, Non-memory, and Refractory ISGs is in SI Appendix, Dataset S1.

(c) Hierarchical clustering analysis for memory ISGs (strong) was performed by the Ward's minimum variance method using JMP software (JMP®, SAS Institute). ISG expression levels are depicted by differential coloring. The clustering analysis was done by using Ward's minimum variance method for unsupervised hierarchical clustering.

Fig. 3 Pol II Is Recruited to Memory ISGs Faster and at Higher Levels Upon Restimulation

(a,b,c) Naïve and pretreated cells (24 h pretreated and 6 h interval time) were stimulated with IFN for indicated times (0, 1, 3h) and qChIP assays were performed to detect binding of Pol II (a), pSTAT1 (b), and BRD4 (c) at the TSS/promoter region of memory ISGs (Mx1, Ifit1), non-memory ISG (irf1) and Gtf2b (control). Values represent the average of three independent experiments $\pm$ S.D.

(d) ChIP-seq analysis for global distribution of Pol II on the memory ISGs, non-memory ISGs, constitutively expressed genes or silent genes. See SI Appendix, Fig. S3a-c for IGV examples.

Fig. 4. Memory ISGs Are Marked by the Histone H3.3 and H3K36me3

(a,b,c) ChIP-seq analysis was performed for genome-wide distribution of H3.3 (a), H3K36me3 (b) and H2AZ (c) over memory ISGs, non-memory ISGs (left panels), and constitutively expressed or silent genes (right panels) in naïve or pretreated MEFs without restimulation. Pretreated cells were left without IFN for 48 h. See SI Appendix, Fig. 4Sd-h for IGV examples.

(d) IGV images of H3.3, H3K36me3 and H2AZ peaks over memory ISGs (Mx1, Mx2) in naïve and pretreated cells.

(e) IGV images of H3.3, H3K36me3 and H2AZ peaks over non-memory ISG (Irf1) in naïve and pretreated cells.

Fig. 5. IFN γ Stimulation Generates Transcriptional Memory in Macrophages

(a) Experimental design for microarray analysis.

(b) ISGs were defined as those genes expressed at least 2 fold higher in naïve macrophages treated with IFN γ for 3 or 6 h than untreated macrophages ($p < 0.05$). ISGs were classified into three groups. Memory ISGs: showing at least 1.5-fold higher in expression (>1.5) in prestimulated macrophages than naïve cells. Non-memory ISGs: showing less than 1.2-fold (<1.2) difference between naïve and prestimulated macrophages. Refractory ISGs: showing at least 1.5-fold less (<1.5) expression in prestimulated macrophages with IFN γ for 3 or 6 h. List of Memory, Non-memory, and Refractory ISGs is in SI Appendix, Dataset S2.

- (c)** Hierarchical clustering of memory (left) and refractory ISGs (right). Memory ISGs exhibited faster and/or greater ISG induction in prestimulated macrophages than naïve cells. Refractory ISGs failed to express or showed reduced ISG expression in prestimulated macrophages.
- (d)** IGV images of Pol II, H3.3 and H3K36me3 distribution over the Gbp gene cluster (memory ISGs) and Il4ra (non-memory ISG).
- (e)** Global Pol II distribution was analyzed for naïve and prestimulated macrophages at 0 h or 1 h after IFN γ stimulation. Distribution of H3.3 and H3K36me3 was tested for untreated naïve and memory Macrophages (24 h after IFN washout).
- (f,g)** Global distribution of H3.3 (f) and H3K36me3 (g) over memory ISGs, non-memory ISGs or refractory ISGs in naïve and prestimulated macrophages.









