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The Small Ubiquitin-like Modifier-Deconjugating Enzyme Senstrin-Specific Peptidase 1 Switches IFN Regulatory Factor 8 from a Repressor to an Activator during Macrophage Activation

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Macrophages, when activated by IFN-γ and TLR signaling, elicit innate immune responses. IFN regulatory factor 8 (IRF8) is a transcription factor that facilitates macrophage activation and innate immunity. We show that, in resting macrophages, some IRF8 is conjugated to small ubiquitin-like modifiers (SUMO) 2/3 through the lysine residue 310. SUMO3-conjugated IRF8 failed to induce IL12p40 and other IRF8 target genes, consistent with SUMO-mediated transcriptional repression reported for other transcription factors. SUMO3-conjugated IRF8 showed reduced mobility in live nuclei and bound poorly to the IL12p40 gene. However, macrophage activation caused a sharp reduction in the amount of SUMOylated IRF8. This reduction coincided with the induction of a deSUMOylating enzyme, senstrin-specific peptidase 1 (SENP1), in activated macrophages. In transfection analysis, SENP1 removed SUMO3 from IRF8 and enhanced expression of IL12p40 and other target genes. Conversely, SENP1 knockdown repressed IRF8 target gene expression. In parallel with IRF8 deSUMOylation, macrophage activation led to the induction of proteins active in the SUMO pathway and caused a global shift in nuclear protein SUMOylation patterns. Together, the IRF8 SUMO conjugation/deconjugation switch is part of a larger transition in SUMO modifications that takes place upon macrophage activation, serving as a mechanism to trigger innate immune responses. The Journal of Immunology, 2012, 189: 3548–3556.
ation of myeloid progenitor cells to functional macrophages (21). Similarly, IRF8 stimulates development of DC subsets and controls production of IL12p40 and type I IFNs (20, 22). IRF8 activates other factors, including chemokines, transcription factors, and those involved in many antipathogenic activities (23–26). IRF8 also enhances CIITA transcription, thereby increasing MHCII expression and Ag presentation (27, 28). Supporting the critical importance of IRF8 in host defense, point mutations in IRF8 are associated with severe immunodeficiency and increased susceptibility to infectious pathogens in humans and mice (22, 29, 30). Various posttranslational modifications influence the activity of IRF8. For example, phosphorylation and dephosphorylation alter IRF8’s functions, affecting innate immune responses and leukemia pathogenesis (26, 31). IRF8 is conjugated to ubiquitin through an E3 ligase TRIM21 that alters IRF8’s ability to regulate IL12p40 transcription positively and negatively (32, 33).

This study began with our observation that IRF8 was SUMOylated in resting macrophages. We show in this study that the lysine (K) at 310 is a major SUMOylation site in IRF8, and SUMO conjugation abrogates IRF8’s ability to stimulate transcription of its target genes, including IL12p40 and IFN-β. A defining observation was that the levels of SUMOylated IRF8 decreased upon macrophage activation triggered by IFN-γ/TLR stimulation. The change in IRF8 SUMOylation coincided with a global shift in the nuclear protein SUMOylation and the induction of enzymes that regulate SUMO pathways. Among multiple SUMO-deconjugating enzymes, SENP1 was induced most robustly in macrophages after stimulation and removed SUMO from IRF8, leading to the reversal of SUMO-mediated repression of target gene transcription. Our results are consistent with the paradigm in which deSUMOylation causes derepression of key genes to allow progression of cellular maturation, examples of which have been reported for monocytes, erythroid cells, and keratinocytes (34–36).

**Materials and Methods**

**Plasmids and vectors**

pcDNA3.1 vectors (Invitrogen) containing mouse IRF8 cDNA or IRF8 tagged to GFP (GFP-IRF8), CFP-IRF8, or Flag-IRF8 and those for T7- or V5-tagged SUMO1, SUMO2, SUMO3, and SUMO3GA were described (19, 32, 33). pcDNA3.1 vector for mutant Flag-IRF8 was constructed by site-directed mutagenesis. SUMO3 fusions of IRF8 are described in Supplemental Fig. 2. pcDNA 3.1 vector for a murine full-length SENP1 with or without hemagglutinin (HA) tag was constructed using pCMV-SPOSPORT vector mSENP1 (Invitrogen). The catalytic mutant C599S was constructed by standard cloning procedures for a murine full-length Senp1 with or without Flag epitope (wt) IRF8 and GFP-IRF8 were described, and those for GFP-SUMO3, YFP-SUMO3, and YFP-SUMO3GA were constructed from pEGFP-C1 or pEYFP-C1 vector by inserting PCR-amplified SUMO3 constructs of GFP-SUMO1, SUMO2, SUMO3, and SUMO3GA were described (19, 32, 33). pcDNA3.1 vectors (Invitrogen) containing mouse IRF8 cDNA or IRF8 tagged to GFP (GFP-IRF8), CFP-IRF8, or Flag-IRF8 were constructed by standard cloning procedures for a murine full-length Senp1 with or without Flag epitope (wt) IRF8 and GFP-IRF8 were described, and those for Flag-IRF8, SENP1, and the C599S mutant were constructed by standard cloning procedures (22, 38) (Clontech).

**Cells, communoprecipitation, and SUMOylation assays**

RAW 264.7 (RAW), NIH3T3, and 293T cells (American Type Culture Collection) were maintained in DMEM with 10% FBS or 10% donor serum (for NIH3T3). IRF8−/− macrophage cell line, CL2 cells were cultured in RPMI 1640 and 10% FBS plus rM-CSF (5 ng/ml) (32). Bone marrow (BM)-derived macrophages were obtained by culturing BM mononuclear cells in the presence of 20 ng/ml M-CSF (Invitrogen) for 5–6 d. BM-derived IRF8−/− DCs were cultured in the presence of Flt3L for ~4 wk. These cells exhibited a DC progenitor-like property and differentiated into DCs after IRF8 transduction. Detailed procedures and the properties of these DCs will be presented elsewhere (P. Tailor, unpublished observations). Procedures for quantitative RT-PCR (qRT-PCR), in vivo SUMOylation, and communoprecipitation assays were performed, as described (19).

**Fluorescence resonance energy transfer and fluorescence recovery after photobleaching**

Fluorescence resonance energy transfer (FRET) analyses were performed with NIH3T3 cells transfected with CFP-IRF8 and/or YFP-SUMO3 or YFP-GA for 30 h on a Zeiss microscope Axiovert 200 with the Axiovision FRET software (39). Fluorescence recovery after photobleaching (FRAP) assays were performed with NIH3T3 cells transfected with pMSCV vector for GFP-IRF8 or GFP-IRF8 fused to SUMO1 or SUMO3, as described (38). Detailed procedures are in Supplemental Fig. 3.

**Chromatin immunoprecipitation analysis**

Chromatin immunoprecipitation (ChIP) analysis was performed essentially as described (19). Briefly, CL2 cells expressing GFP-SUMO3, GFP-IRF8, or GFP-IRF8-SUMO3 from the pMSCV vector were cross-linked with 1% formaldehyde for 10 min and sonicated in Misonix Sonicator 3000. Chromatin preparations corresponding to 0.2 × 10^6 cells were incubated with rabbit anti-GFP Ab (Abcam) or control rabbit IgG bound to Dynabeads protein G (Invitrogen) overnight at 4°C. Immunoprecipitated chromatin was decross-linked, and purified DNA was analyzed by quantitative PCR using appropriate primers. The percentage input was calculated as 2^[(cycle threshold (CT) IP sample – CT input)].

**Results**

**IRF8 is conjugated to SUMO2/3 in resting macrophages**

To determine whether IRF8 is SUMOylated in macrophages, immunoprecipitation analysis was performed with nuclear extracts from unstimulated RAW264.7 (hereafter RAW) cells or RAW cells stimulated with IFN-γ or IFN-γ plus CpG (IFN-γ/CpG). Materials precipitated with anti-IRF8 Ab were blotted with anti-SUMO2/3 Ab. As shown in Fig. 1A (left panels), in unstimulated cells, a fraction of IRF8 consistently exhibited SUMO2/3 reactivity, which migrated more slowly than the rest of IRF8 (~62 kDa versus ~51 kDa), indicating conjugation of SUMO2/3 to IRF8. However, the levels of SUMO 2/3-linked IRF8 fell noticeably when cells were stimulated with IFN-γ or IFN-γ/CpG. The total IRF8 levels, however, increased after stimulation, as expected (32, 33).

The detailed time course analysis in Fig. 1A (right panels) showed very similar results, in that IRF8 was SUMOylated before stimulation, which was attenuated upon macrophage activation. Those data indicate that IRF8 is modified by SUMO2/3 in resting macrophages, but macrophage activation prompts a reduction in SUMOylated IRF8. We did not detect conjugation of SUMO1 to IRF8, possibly suggesting that IRF8 may be predominantly modified by SUMO2 or SUMO3 in RAW cells under these conditions.

To confirm that IRF8 is a SUMO substrate, immunoprecipitation experiments were carried out in 293T cells transfected with Flag-tagged IRF8 (Flag-IRF8) and V5-tagged SUMO1, SUMO2, or SUMO3. Data in Fig. 1B showed that all three SUMO proteins are conjugated to IRF8 (upper panel). A single, SUMO-conjugated IRF8 band was predominant after a short exposure (upper middle panel). However, additional, multiple SUMO-conjugated bands were visible upon a longer exposure (lower middle panel), suggesting that polymeric SUMO chains were formed on IRF8, although a covalent linkage with other proteins may also have occurred (40). In these assays, SUMO3 consistently gave more robust IRF8 modification than SUMO1 and SUMO2, the basis of which has not been studied further. As expected, immunoblot of whole-cell extracts (WCE) revealed that proteins other than IRF8 were also conjugated to V5-SUMOs (lower panel). Based on these data and the structural similarity between SUMO2 and SUMO3, we tested mostly the activity of SUMO3 in the current study.

FRET analysis detects transfer of fluorescent energy from a donor to an acceptor that takes place in two proteins of close physical proximity. This assay has been used to study SUMO...
conjugation events in vitro and in vivo (41). To further solidify IRF8 SUMO conjugation in vivo, NIH3T3 cells were transfected with CFP-IRF8 and YFP-SUMO3, or YFP-mutant SUMO3 (SUMO3GA), in which the C-terminal amino acids GG were mutated to AA, eliminating the SUMOylation activity (42). In microscopic inspection in Fig. 1C (left panel), we detected FRET signals when CFP-IRF8 was coexpressed with YFP-SUMO3, but not YFP-SUMO3GA. We quantified the microscopic observations by ranking FRET signals into four levels, as recommended (39). More than 200 CFP- and YFP-positive cells were ranked (Fig. 1C, right panel). About 50% of cells showed FRET signals with varying scores. In contrast, nearly 90% cells expressing YFP-SUMO3GA were FRET negative. These results further support a close physical link between IRF8 and SUMO3.

Identification of K310 as a major IRF8 SUMO conjugation site

IRF8 has several amino acid motifs that conform to or resemble the consensus SUMO site, C[KXE. To identify a functional SUMO conjugation site in IRF8, the K residues in the motifs were mutated to arginine (R), and the resultant mutants were tested for SUMO1 or SUMO3 conjugation. Results in Fig. 2 showed that mutation of K310 abolished conjugation of both SUMO1 and SUMO3, whereas mutation of K72 did not. Immunostaining analysis showed that both wt IRF8 and the IRF8 K310R mutant localized to the nucleus along with SUMO3, confirming that the absence of SUMO conjugation was not due to dysregulated localization of the mutant (Supplemental Fig. 1A). We also tested additional four K residues that may serve as SUMO conjugation sites as assessed by a National Center for Biotechnology Information database. Mutation of these sites had no effect on IRF8 SUMO conjugation (Supplemental Fig. 1B, summary in Supplemental Fig. 1C). These data show that K310 is a single, main SUMO conjugation site in IRF8.

**FIGURE 1.** IRF8 SUMOylation status in resting and activated macrophages. (A) Left panels, Nuclear extracts (500 µg) from unstimulated RAW cells or stimulated with IFN-γ (150 U/ml) for 8 h or IFN-γ overnight, followed by CpG (150 ng/ml) for 6 h, were immunoprecipitated with Ab for IRF8 and blotted against SUMO2/3 or IRF8. Extracts were blotted for TFIIB for loading control. Right panels, Unstimulated RAW cells (0) or stimulated with IFN-γ overnight, followed by CpG for indicated times and nuclear extracts, were tested, as above. TFIIB was tested as a loading control. (B) 293T cells (1 x 10⁶) were transfected with Flag-IRF8 (1 µg), V5-SUMO1, SUMO2, or SUMO3 for 30 h, and extracts were precipitated with anti-Flag Ab and blotted with anti-V5 or anti-Flag Ab. The lower middle panel depicts results of a longer exposure revealing multiple SUMOylated IRF8 bands. The bottom panel indicates immunoblot of WCE with anti-V5 Ab. (C) NIH3T3 cells (1 x 10⁶) were transfected with 1 µg CFP-IRF8 and YFP-SUMO3 or SUMO3GA for 30 h and tested for a real-time interaction of IRF8 with SUMO3 by FRET analysis. Only SUMO3, not SUMO3GA, gave FRET signals (upper panel). In the lower panel, FRET signals were quantified by scoring fluorescent intensity of ~200 cells, according to the modified Youvan method.

**FIGURE 2.** Identification of a SUMOylation site in IRF8. The 293T cells transfected with Flag-IRF8 mutants (K72R or K310R) and T7-SUMO1 or V5-SUMO3 were immunoprecipitated with anti-Flag Ab and blotted with anti-T7 and anti-V5 Ab (upper middle panel) or anti-Flag Ab (lower middle panel). Vector alone and wt Flag-IRF8 (wt) were tested as controls. The bottom panel indicates immunoblot analysis of WCE.
SUMO3 conjugation inhibits IRF8 target gene transactivation

To assess the functional consequence of IRF8 SUMOylation, we tested wt IRF8, the K310R mutant, and two additional IRF8-SUMO conjugate constructs for the ability to stimulate Il12b (IL12p40) and Ifnb (IFN-β) promoters. For the IRF8-SUMO conjugates, SUMO3 was covalently conjugated to either the N terminus or C terminus of IRF8 (SUMO3-IRF8 or IRF8-SUMO3, respectively; see diagram in Supplemental Fig. 1D). When transfected in 293T cells, IRF8-SUMO3 retained the SUMO moiety and migrated more slowly than IRF8, but SUMO3-IRF8 was reverted to the size of unconjugated IRF8 within 30 h, presumably due to the cleavage of SUMO from IRF8, indicating that the former was more stable than the latter (see immunoblot data in Supplemental Fig. 1E). Molecularly constructed SUMO conjugates such as these have been widely used for studying the functional significance of SUMOylation, mostly because of improved stability (43). In luciferase assays performed in RAW cells, wt IRF8 enhanced activities of both reporters by 2- to 5-fold, with a further increase after IFN-γ/CpG stimulation (Fig. 3A). K310R gave modest, but consistently higher luciferase activities (25–50% increase) in both reporters. The SUMO3-IRF8 conjugate, from which SUMO3 was removed during culture, gave reporter activity comparable to wt IRF8. In contrast, the stable SUMO conjugate, IRF8-SUMO3, did not stimulate either reporter. Next, these IRF8 constructs were stably expressed in IRF8−/− macrophages (named CL2) and tested for mRNA expression of Il12b and Ccl9, IRF8 targets, that are important for innate immune responses (Fig. 3B) (19, 32, 33). Wt IRF8 enhanced mRNA expression of both genes before and after IFN-γ/CpG stimulation. The K310R mutant further increased the expression, although moderately, suggesting that SUMOylation represses IRF8 transactivation. Supporting this view, K310R increased expression of additional IRF8 targets, Pml and Ciita, over wt IRF8 by 40–60% (Supplemental Fig. 1F). IRF8-SUMO3, in contrast, failed to enhance expression of these genes. IRF8-SUMO3 repression of gene expression was not due to a non-specific effect, because this construct did not repress induction of TNF-α by IFN-γ/TLR (Supplemental Fig. 1G). Similarly, when stably expressed in IRF8−/− DCs, wt IRF8 enhanced expression of Il12b and Ifnb mRNA after CpG stimulation, but IRF8-SUMO3 did not (Fig. 3C) (22). Furthermore, wt IRF8, but not IRF8-SUMO3, stimulated surface expression of MHC II on DCs upon CpG stimulation (Fig. 3D). When tested in RAW cells that express endogenous IRF8, IRF8-SUMO3 repressed IFN-γ/CpG induction of Il12b and Ifnb, indicating that IRF8-SUMO3 repressed endogenous IRF8 as well (Supplemental Fig. 1H). To study whether IRF8-SUMO3 negatively affects activity of endogenous IRF8, these results indicate that SUMO3 conjugation abrogates IRF8’s capacity to stimulate target gene expression.

FIGURE 3. SUMO3 conjugation represses transactivation of IRF8 target genes. (A) RAW cells (1 × 10⁶) were transfected with 600 ng indicated luciferase reporters, 800 ng pcDNA-IRF8 vectors, and 60 ng pRL-TK vector for 30 h and stimulated with IFN-γ overnight and CpG for 6 h. Reporter activity was normalized by Renilla luciferase activity. Values represent the average of three assays ± SD. Statistical significance was tested by Student t test; *p < 0.01, **p < 0.005. (B) IRF8−/− macrophages (CL2) were transduced with pMSCV vectors for wt IRF8, K310R mutant or SUMO3-IRF8, or IRF8-SUMO3 for 5 d and stimulated with (filled bars) or without (open bars) IFN-γ and CpG. Levels of Il12b and Ccl9 transcripts were measured by qRT-PCR. Values represent the average of three determinations ± SD. Statistical significance was tested by Student t test; *p < 0.01, **p < 0.005. (C) IRF8−/− DCs were transduced with the above vectors for 5 d and stimulated with (filled bars) or without (open bars) CpG for 7 h and Il12b and Ifnb transcript levels were measured, as above. Statistical significance was tested by Student t test; **p < 0.005. (D) IRF8−/− DCs transduced with indicated vectors were stimulated with or without CpG for 36 h, and expression of MHCII (I-Ab) was detected by flow cytometry. The gray lines indicate signals by control IgG.
SUMO conjugation impairs genome-wide mobility of IRF8

SUMO conjugation alters intranuclear localization of some transcription factors (44, 45). We checked localization of GFP-IRF8 and GFP-IRF8-SUMO3 in the nucleus of living cells, both un-conjugated and SUMO3-conjugated IRF8 distributed evenly in the nucleus, indicating that SUMO3 conjugation did not alter the uniform distribution pattern reported for IRF8 (32) (Fig. 4A). GFP-SUMO3, not conjugated to IRF8, tested as a control also localized mostly in the nucleus, whereas free GFP distributed both in the cytoplasm and the nucleus. We then asked whether SUMOylation affects an interaction of IRF8 with chromatin by FRAP analysis. Most transcription factors, including IRF8, are highly mobile within the nucleus and bind to chromatin with a rapid, on-and-off mode, which reflects a global, genome-wide chromatin-scanning activity of the transcription factor, which is critical for their function (38, 46). NIH3T3 cells expressing GFP-IRF8 or GFP-IRF8-SUMO3 were briefly photobleached within a small area in the nucleus, and fluorescence recovery was measured for the subsequent 28 s. Cells expressing free GFP or GFP-SUMO3 were also tested as controls. Fig. 4B depicts the recovery profile for each construct obtained from 15 separate cells. Free GFP recovered almost instantly following photobleach (blue). Although more slowly than free GFP, GFP-IRF8 also recovered rapidly, showing a t1/2 of ∼2 s (red) (38). GFP-IRF8 regained original fluorescence intensity fully by 16 s. In contrast, GFP-IRF8-SUMO3 showed a dramatically slower recovery, reclaiming only ∼75% of the original fluorescence even at the end of measurement (black). GFP-SUMO3, in contrast, showed the slowest mobility (green) particularly evident in an early stage, suggesting its tendency to bind to chromatin. In line with our results, SUMO1 was reported to move very slowly in a similar FRAP assay (44). These results indicate that SUMO3 conjugation markedly reduces IRF8’s genome-wide mobility. Consistent with these data, we found that conjugation of SUMO1 also dramatically reduced IRF8’s real-time mobility (Supplemental Fig. 2B).

SUMOylated IRF8 binds poorly to the Il12b promoter

Having observed inhibition of global IRF8 mobility by SUMO conjugation, it was of interest to assess whether SUMO conjugation affects binding of IRF8 to specific target sites. ChIP analysis was performed for the Il12b gene in CL2 macrophages stably expressing GFP-IRF8 or GFP-IRF8-SUMO3. Anti-GFP Ab was used to detect binding of IRF8 to four regions within the Il12b gene, that is, 1.8 kb upstream promoter, the transcription start site (TSS), and two coding regions (see Fig. 4C for the Il12b gene map and the location of the primers used for ChIP in the bottom). After IFN-γ stimulation, GFP-IRF8-SUMO3 bound less efficiently to another target, Cst3, than wt IRF8 (data not shown).

IFN-γ and TLR stimulation triggers a global shift in SUMOylated nuclear proteins in macrophages

We next asked whether macrophage activation changes SUMOylation of proteins other than IRF8. This question was asked, because we previously observed that IFN-γ/TLR stimulation of macrophages causes a large increase in ubiquitinated nuclear proteins (47). To detect SUMO2/3-conjugated nuclear proteins, extracts from unstimulated or stimulated RAW cells were blotted with anti-SUMO2/3 Ab (Fig. 5A). SUMO2/3-conjugated proteins were markedly increased after stimulation by IFN-γ/CpG, IFN-γ/LPS, and Newcastle disease virus. The increase was particularly noticeable at 14–24 h. The levels of transcription factor II B (TFIIB), tested as a loading control, remained unchanged. Stimulation by CpG or LPS alone led to only a meager increase in

FIGURE 4. SUMO3 conjugation reduces genome-wide mobility of IRF8 and target-binding activity. (A) NIH3T3 cells were transduced with pMSCV vectors for GFP alone, GFP-SUMO3, wt GFP-IRF8, or GFP-IRF8-SUMO3, and localization of each construct was viewed by live cell imaging. (B) In FRAP analysis, a small region within the nucleus was bleached at indicated time, and the recovery of fluorescence (FL) signals was plotted against time (sec). The recovery curves represent the average of 15 separate cells. See reduced recovery of GFP-IRF8-SUMO1 in Supplemental Fig. 3. (C) CL2 cells were transduced with indicated pMSCV vectors and stimulated with IFN-γ or IFN-γ/CpG. Binding of these constructs to the indicated regions of the Il12b gene was tested by ChIP using anti-GFP Ab. Values represent the average of three determinations ± SD. The bottom diagram indicates the exon-intron organization of Il12b. The arrow marks the TSS.
SUMOylated proteins, whereas IFN-γ stimulation alone resulted in a more substantive increase, in line with the observations on ubiquitinated proteins (Supplemental Fig. 3A) (47). A similar, but less dramatic increase was detected for SUMO1-conjugated nuclear proteins after IFN-γ/TLR stimulation (Supplemental Fig. 3B). Together, a large number of nuclear proteins are conjugated to SUMO after RAW macrophage cell activation. The global shift in SUMOylated proteins observed in this study is reminiscent of stress-induced changes in SUMOylated proteins reported in other cell types, in which some proteins are newly SUMOylated, whereas others are deSUMOylated after stress (8–11).

**SENP1 is induced in macrophages after IFN-γ/CpG stimulation and deconjugates SUMO3 from IRF8**

In light of the results that IRF8 SUMOylation decreased after IFN-γ/CpG stimulation, despite the global increase in SUMOylated nuclear proteins (Fig. 1A), we considered the possibility that IRF8 is one of those proteins that are deSUMOylated upon macrophage activation, examples found in other types of stress (8–11). Before testing whether SUMO was enzymatically removed from IRF8, we asked whether expression of Senp genes that encode SUMO-deconjugating enzymes alters after macrophage activation (3, 4, 48). We tested all seven Senp transcripts expressed in RAW cells. As shown in Supplemental Fig. 3C, that among all seven Senp transcripts, Senp1 mRNA was most robustly induced in RAW cells after IFN-γ/CpG stimulation: other Senps (Senp2–Senp6) were also induced, although weakly, although Senp7 and Senp8 were not. Immunoblot results shown in Supplemental Fig. 3D indicated that SENP1 protein was not detectable before stimulation, but induced after stimulation. In BM-derived macrophages, Senp1 mRNA was also induced along with other Senps after stimulation (Supplemental Fig. 3E). Given the global increase in SUMOylated proteins after macrophage activation (Fig. 5A), we also tested expression of genes involved in SUMO conjugation. As shown in Supplemental Fig. 3E, IFN-γ/TLR stimulation strongly induced all four SUMO E3 ligases of the PIAS family, Pias1, Pias2, Pias3, and Pias4. Furthermore, ubiquitin-conjugating enzyme E21 and all three Sumo genes were also induced after IFN-γ/TLR stimulation.

**FIGURE 5.** Macrophage activation triggers a global change in SUMOylated nuclear proteins, and the SENP1 enhances IRF8 transactivation. (A) RAW cells were stimulated with IFN-γ overnight, followed by stimulation by Newcastle disease virus, LPS, or CpG for indicated time. A total of 30 μg nuclear extracts was resolved in 4–12% Nu PAGE and blotted with anti-SUMO2/3 Ab. 0, indicates no treatment. (B) 293T cells were cotransfected with 2 μg pcDNA vector for Flag-IRF8 and 2 μg HA-Senp1 and 500 μg lysates were immunoprecipitated with anti-Flag Ab and blotted with anti-HA Ab (upper panel) or Flag (middle panel). In the lower panel, WCE (20 μg) were blotted with Flag for Flag-IRF8 and 2 μg HA-Senp1 and 500 μg lysates were immunoprecipitated with anti-Flag Ab and blotted with anti-HA Ab (upper panel) or Flag (middle panel). In the lower panel, WCE (20 μg) were blotted with Flag for Flag-IRF8 and 2 μg HA-Senp1 and 500 μg lysates were immunoprecipitated with anti-Flag Ab and blotted with anti-HA Ab (upper panel) or Flag (middle panel). In the lower panel, WCE (20 μg) were blotted with Flag for Flag-IRF8 and 2 μg HA-Senp1 and 500 μg lysates were immunoprecipitated with anti-Flag Ab and blotted with anti-HA Ab (upper panel) or Flag (middle panel).

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Thus, IFN-γ/TLR stimulation enhanced expression of genes encoding factors that catalyze both SUMOylation and deSUMOylation.

SENP1 has been shown to interact with a substrate protein, as detected by communoprecipitation assays (49). We performed communoprecipitation experiments with cells expressing Flag-IRF8 and HA-SENP1. As shown in Fig. 5B, IRF8 and SENP1 were coprecipitated when expressed together, supporting the possibility that IRF8 may serve as a SENP1 substrate. We next tested whether SENP1 catalyzes deSUMOylation of IRF8. Flag-IRF8 was coexpressed with wt HA-SENP1 or C599S, a SENP1 mutant lacking the catalytic activity, along with V5-SUMO3, and communoprecipitation experiments were performed with anti-Flag Ab (37). As shown in Fig. 5C, in the absence of SENP1, Flag-IRF8 was readily conjugated to SUMO3. However, SUMO3 conjugation was virtually absent when SENP1 was coexpressed. Consistent with an enzymatic basis of this effect, the C599S mutant did not affect Flag-IRF8 SUMO3 conjugation (upper two panels). Immunoblot data in lower three panels showed that SENP1 and IRF8 were properly expressed and that SENP1, but not the C599S mutant, globally reduced SUMOylated proteins in the cells. Thus, SENP1 can remove SUMO3 from its substrate, IRF8. Reinforcing deSUMOylating activity of SENP1, polymeric SUMO chains linked to the IRF8-SUMO3 conjugate were also removed from IRF8 by SENP1, but not by the mutant (Supplemental Fig. 4A).

**Ectopic SENP1 expression enhances IRF8 transactivation**

To study whether SENP1 alters repressive activity of SUMOylated IRF8, we performed reporter assays with the Il12b and Ifnb promoters (Fig. 5D). The activity of both promoters was enhanced by transfection of IRF8, as expected. Cotransfection of wt SENP1 further enhanced activity of both reporters in a dose-dependent manner. However, the C599S mutant did not enhance activity of either reporter, indicating that SENP1 enhanced reporter activity by reducing SUMO conjugation. Supporting this idea, SENP1 had no effect on reporter activity by the K310R mutant (Supplemental Fig. 4E). To further investigate the effect of SENP1 expression on IRF8 function, SENP1 was stably introduced in IRF8−/− CL2 macrophages along with IRF8 by retroviral transduction, and expression of endogenous IRF8 target genes was tested. Results in Fig. 5E showed that IRF8 alone enhanced Il12b and Cc9 transcripts, and cotransduction of SENP1, but not C599S, further increased their expression. The effect of SENP1 coexpression was highly reproducible, and consistently seen with samples prepared separately. Cotransduction of IRF8 and SENP1, but not C599S in IRF8−/− DCs also led to increased expression of Il12b and Ifnb mRNAs (Fig. 5F). Thus, SENP1 increases IRF8’s ability to transactivate target genes, presumably by deSUMOylating IRF8.

**SENP1 knockdown weakens IRF8 transactivation**

To ascertain whether SENP1 promotes IRF8 transactivation, we knocked down Senp1 expression by shRNA and tested its effect on IRF8 target gene expression. Immunoblot analysis in Supplemental Fig. 4B showed that Senp1 protein levels were markedly reduced when Senp1 shRNA vector, but not scrambled shRNA (ctrl), was coexpressed in NIH3T3 cells. Senp and control shRNA were then stably expressed in CL2 macrophages along with IRF8, and tested for their effects on Il12b and Cc9 induction. Results in Fig. 5G showed that Senp1 shRNA reduced induction of both genes by ~30–45% relative to control shRNA or vector alone. Similarly, Senp1 shRNA reduced levels of Il12b and Ifnb induction approximately by half in DCs (Fig. 6B). Induction of additional IRF8 targets, Ciita, Cc9, and Irf7, was also reduced in the presence of Senp1 shRNA (Supplemental Fig. 4C, 4D). These results further support the idea that SENP1 relieves IRF8 of SUMO-mediated transcriptional repression.

**Discussion**

IRF8 plays a pivotal role in eliciting innate immune responses in activated macrophages and DCs (22, 29, 30, 32). We show that, in unstimulated RAW macrophages, IRF8 was SUMOylated, in a state unable to transactivate target genes. IRF8 became deSUMOylated after IFN-γ/TLR stimulation, coinciding with trans-
activation. The change in IRF8 SUMOylation took place in the wake of larger changes in SUMO-conjugated nuclear proteins, which was set off upon macrophage activation: RAW cell activation caused a global increase in SUMOylated proteins, accompanying induction of Sumo 1, 2, 3, the E2 enzyme ubiquitin-conjugating enzyme E2I, and all four Pias E3 ligases. Parallelizing with these changes, however, there was a dramatic induction of multiple Senp deSUMOylating enzymes. The simultaneous induction of enzymes that catalyze opposite reactions indicates that SUMO modifications occur in a bidirectional fashion during macrophage activation, where some proteins are newly SUMOylated, whereas others are deSUMOylated. Clearly, IRF8 belongs to the latter. Both events are likely to be relevant to the execution of innate immune responses. Somewhat analogous to our observations, Deyrieux and Wilson (34) reported that cellular SUMOylation dynamic plays a critical role in keratinocyte differentiation. During Ca2+-induced keratinocyte differentiation enzymes of SUMOylation pathways, E1, E2, and E3 as well as SENP1 are induced to globally alter SUMOylation of many proteins that is required for proper keratinocyte differentiation. In contrast, a variety of stress, including heat shock, oxidative stress, and genotoxic stress, prompts changes in SUMOylation of many proteins, which most likely contribute to cells’ stress responses (8, 10, 11, 50). Macrophage activation may share aspects of stress response, a view supported by the production of reactive oxygen species and NO in activation macrophages (23–25).

**SUMOylation of IRF8**

In RAW cells, a portion of IRF8 was conjugated to SUMO, mostly SUMO2/3 through the K310 residue. Coinmunoprecipitation analysis indicated that IRF8 can conjugate both monomeric and polymeric SUMO peptides, similar to other proteins (40, 51). SUMOylated IRF8 was associated with transcription repression, as judged by enhanced transactivation by the K310R mutant and diminished activation by the IRF8-SUMO3 conjugate, a conclusion consistent with the large body of evidence that SUMOylation impairs genome-wide mobility of IRF8, an in vivo parameter that signifies functional integrity of a transcription factor (38, 46). The reduced IRF8 mobility may be partly accounted for by the property of SUMO3 itself, because GFP-SUMO3 exhibited a very slow mobility as well. The slow mobility of SUMO3 noted in this work is in agreement with a report on SUMO1 (44). In ChIP analysis, binding of the IRF8-SUMO3 conjugate to the II12b target site was weaker than that of wt IRF8. The weakened target binding may be accounted for by the assembly of multiple repressive factors and poor genome-wide mobility (6, 7). In addition, epitope sequestration by assembled factors cannot be ruled out for reduced target binding.

**IRF8 SUMO deconjugation by SENP1**

SUMOylated IRF8 levels fell following macrophage activation, coinciding with marked induction of SENP1. Supporting the role for this enzyme in processing SUMOylated IRF8, SENP1 removed SUMO3 from IRF8 in a manner dependent on its catalytic activity. SENP1 could reduce monomeric and polymeric SUMO3 from IRF8. Moreover, SENP1 altered the functional property of IRF8, because SENP1 ectopic expression enhanced promoter activity and mRNA expression of IRF8 target genes. Supporting a direct link between deSUMOylation and IRF8 transactivation, SENP1 enhancement of IRF8 activity was also dependent on the intact catalytic activity. Moreover, the enhancing effect of SENP1 was reversed by shRNA-based Senp1 knockdown. These results led us to suggest a model in which SENP1, by virtue of its deSUMOylating activity, switches IRF8 from a repressor to an activator, a change critical for functional macrophage activation by IRF8 (Fig. 6C). At present, the mechanism by which SENP1 selectively deSUMOylates IRF8 among other proteins remains elusive. It is possible that a ready physical interaction between the two proteins as observed by our communoprecipitation assays may be a basis of substrate selectivity. Given that other SENPs besides SENP1 were also induced after IFN-γ/TLR stimulation, albeit less strongly, it is conceivable that they also take part in deSUMOylation of IRF8 or more likely other substrates. SENP members have different activities, ranging from SUMO maturation to chain editing to SUMO deconjugation (3, 4). SENPs are also differentially distributed in the nucleus. Some of these SENPs are shown to relieve SUMO-mediated transcriptional repression. For example, SENP1 deconjugates SUMO from HDAC1 and p300 to relieve repression (48, 49). Sir2uin 1 is constitutively SUMOylated, the state that confers full deacetylase activity. Sir2uin 1 is deSUMOylated by SENP1 after UV radiation or by oxidative stress, leading to inactivation of its histone deacetylase activity and increased apoptosis (49). SENPs are involved in other regulatory processes. Senp1−/− mice are embryonic lethal due to the dysregulation of HIF-1a stability, which is controlled by SUMOylation (48, 52). Moreover, Huang et al. (7, 53) recently demonstrated that SENP3 plays a role in regulating inflammatory gene expression in a model in which SUMOylated liver X receptor transrepresses NF-κB-activated genes by recruiting the NCOR complex. This transrepression was relieved by SENP3 recruited to the target genes by interacting with liver X receptor. Another recent study showed that SENP3 is also involved in p300 deSUMOylation and HIF-1α transactivation (54).

Our proposition that IRF8 deSUMOylation is causally linked to its functional activation in macrophages gains further credence in view of an emerging paradigm in which SENPs play an integral role in promoting maturation of hematopoietic cells. Tillmanns et al. (35) showed that the maintenance of myeloid progenitor cells depends on MafB repression through its SUMOylation. In that report, a MafB mutant lacking SUMOylation site prompts premature macrophage differentiation. Furthermore, GATA1, a transcription factor that regulates the globin gene switch, is deSUMOylated by SENP1 during RBC development (36). GATA1 deSUMOylation is shown to be a critical step for erythocyte formation, as evidenced by the absence of erythropoiesis in Senp1−/− fetal liver cells (36).

In summary, by its deSUMOylating activity, SENP1 switches repressive IRF8 to a potent transactivator, an event fundamental to the onset of innate immune responses in macrophages.

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**Disclosures**

The authors have no financial conflicts of interest.

**References**

Supplementary Data: Supplemental Figures and Legends

The SUMO deconjugating enzyme SENP1 switches IRF8 from a repressor to an activator during macrophage activation

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Figure S1

(A) **Nuclear localization of K310R.** NIH3T3 cells were transfected with Flag-wt IRF8 or Flag-K310R along with GFP-SUMO3, and nuclear localization of the two
proteins was examined by immunostaining using anti-Flag antibody (Ab).

(B) **Search for IRF8 SUMOylation sites.** 293T cells were transfected with indicated Flag-IRF8 mutants and V5-SUMO3, and immunoprecipitated with anti-Flag Ab and blotted with anti-V5 Ab as Figure 2A.

(C) **A summary of IRF8 SUMOylation site analysis.** Underlined K residues were mutated to R and tested for conjugation of SUMO1 or SUMO3. Only K310R failed to conjugate SUMO.

(D) **Stability of SUMO3 conjugates.** Flag-IRF8 fused to SUMO3 at the N- or C-terminus, or wt Flag-IRF8 was transfected into 293T cells, and extracts were immunoblotted with anti-IRF8 antibody 24 h later.

(E) The N-terminus SUMO3-IRF8 conjugate (SUMO3-IRF8) was constructed by inserting an EcoR1 IRF8 fragment and PCR amplified SUMO3 fragment into the HindIII and NotI site of 3X Flag CMV7.1 (Sigma). The C-terminus SUMO3 conjugate (IRF8-SUMO3) was constructed by ligation of the XbaI/ XhoI digested IRF8 fragment lacking the last stop codon to SUMO3 cDNA in pcDNA-Flag vector. 293T cells were transfected with Flag-IRF8, Flag-SUMO3-IRF8 or FlagIRF8-SUMO3 for 36 h and extracts were immunoblotted with anti-Flag antibody. A cleavage product, Flag-SUMO3 was generated from SUMO3-IRF8 but not IRF8-SUMO3 (bottom arrow). The bulk of SUMO3 was retained on IRF8-SUMO3.

(F) **The IRF8 K310R mutant enhances IRF8 transactivation.** IRF8-/- macrophages (CL2) were transduced with pMSCV vectors for wt IRF8 and K310R for 5 days and stimulated with (filled bars) or without (open bars) IFNγ and CpG. Levels of Pml and Ciita transcripts were measured by qRT-PCR. Values represent the average of three determinations +/- S.D. Statistical significance was tested by Student’s t-test, \( P \)-values \(^*<0.01, **<0.005.\)
(G) IRF8-/- macrophages (CL2) were transduced with the above vectors for wt IRF8 and IRF8-SUMO3 for 5 days and stimulated with (filled bars) or without (open bars) IFNγ and LPS, and Tnfa transcript levels were measured as above.

(H) RAW 264.7 macrophages were transduced with pMSCV vectors for wt IRF8 and K310R for 5 days and stimulated with (filled bars) or without (open bars) IFNγ and CpG. Levels of Il12p40 and Ifnb transcripts were measured by qRT-PCR. Values represent the average of three determinations +/- S.D. Statistical significance was tested by Student’s t-test, P-values **<0.005.
Figure S2

(A) Localization of indicated GFP proteins expressed in NIH3T3 cells was viewed on confocal microscope. Left and right panels depict images by lower or higher magnification. GFP signals were detected in living cells plated on chambered cover glass (NUNC) using a Zeiss 510 confocal laser-scanning microscope.

(B) FRAP analysis of SUMO1 conjugated IRF8. Photobleaching was performed under a 100X oil immersion lens with a 488 nm argon laser at a maximum laser power with a 285 milliseconds. Prior to the bleach pulse, 30 prebleach images were acquired. Recovery was monitored for 28 seconds (s) with 49 ms intervals. Prebleach and recovery images were collected at 0.5% laser power. A region marked by red circle in A was photobleached at indicated time (arrow) and fluorescence recovery was measured against time. The data represent the average of 15 separate analyses.
(A) **IFNγ treatment globally increases SUMO2/3 conjugated nuclear proteins in macrophages.** RAW cells stimulated with IFNγ overnight followed by NDV, LPS or CpG for indicated times (h) were lysed by a buffer containing 10 mM Tris HCl, pH 7.5, 0.05% Nonidet P-40, protease inhibitors at 4 °C. Nuclear pellets were lysed by high salt lysis buffer (50 mM Tris HCl, pH 7.5, 420 mM NaCl, 0.2 mM EDTA, protease inhibitors). Thirty μg of extracts were resolved in 4 to 12% Nu PAGE (Invitrogen) and blotted with anti-SUMO2/3 Ab (Santa Cruz).

(B) **IFNγ/TLR stimulation increases SUMO1 conjugated nuclear proteins in RAW macrophages.** Cytosolic and nuclear extracts from RAW cells stimulated by IFNγ and CpG or LPS were blotted with anti-SUMO1 Ab.

(C) **Induction of SUMO modification-related genes upon macrophage activation.** RAW cells were treated with or without IFNγ overnight followed by stimulation by CpG for 4 h. Levels of indicated Senp mRNAs were measured by qRT-PCR.

(D) Nuclear extracts from RAW cells stimulated with IFNγ, IFNγ/CpG, or NDV were immunoblotted with Ab for SENP1 (Abcam) or TFIIB (Sigma).

(E) BM derived macrophages were stimulated with IFNγ overnight followed by treatment with indicated TLR ligands for indicate time (h), and Senp, Pias and Sumo transcripts were detected by qRT-PCR. The values represent the average of three determinations +/- SD.
Figure S4

(A) SENP1 removes SUMO polymers from IRF8-SUMO3. 293T cells were transfected with indicated plasmids, and lysates were precipitated with anti-GFP Ab and blotted with anti-IRF8 Ab as in Figure 5C. WCE were blotted with indicated Abs.

(B) Senp1 shRNA reduces SENP1 protein expression. NIH3T3 cells were

Figure S4

(A) SENP1 removes SUMO polymers from IRF8-SUMO3. 293T cells were transfected with indicated plasmids, and lysates were precipitated with anti-GFP Ab and blotted with anti-IRF8 Ab as in Figure 5C. WCE were blotted with indicated Abs.

(B) Senp1 shRNA reduces SENP1 protein expression. NIH3T3 cells were
cotransfected with 2 μg of plasmid vector for control shRNA or Senp1 shRNA with 0.5 μg of pcDNA-Senp1 for 30 h and 20 μg of WCE were analyzed by immunoblotted with anti-SENP1 Ab (Abcam).

(C) **SENP1 shRNA reduces IRF8 transactivation.** CL2 macrophages were transduced with pMSCV vectors for IRF8 and pSuper Senp1 shRNA and control shRNA for 5 days and stimulated with IFNγ/CpG. Expression of Ciita and Irf7 transcripts was tested by qRT-PCR as above.

(D) **IRF8-/- DCs (10⁶ cells)** were transduced with indicated vectors for 5 days and stimulated with CpG for 4 h and expression of indicated transcripts was tested as above. The values represent the average of three determinations +/- S.D. Statistical significance was tested by Student’s t-test, P-values *<0.01, **<0.005.

(E) **RAW cells (1x10⁶)** were transfected with 600 ng of indicated luciferase reporters, 800 ng of pcDNA-IRF8, IRF8K310 or SENP1 vectors and 60 ng of pRL-TK vector for 30 h and stimulated with IFNγ overnight and CpG for 6 h. Reporter activity was normalized by Renilla luciferase activity. Values represent the average of three assays +/- S.D.