

Selective blockade of the inhibitory Fc γ receptor (Fc γ RIIB) in human dendritic cells and monocytes induces a type I interferon response program

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The ability of dendritic cells (DCs) to activate immunity is linked to their maturation status. In prior studies, we have shown that selective antibody-mediated blockade of inhibitory Fc γ RIIB receptor on human DCs in the presence of activating immunoglobulin (Ig) ligands leads to DC maturation and enhanced immunity to antibody-coated tumor cells. We show that Fc γ receptor (Fc γ R)-mediated activation of human monocytes and monocyte-derived DCs is associated with a distinct gene expression pattern, including several inflammation-associated chemokines, as well as type 1 interferon (IFN) response genes, including the activation of signal transducer and activator of transcription 1 (STAT1). Fc γ R-mediated STAT1 activation is rapid and requires activating Fc γ Rs. However, this IFN response is observed without a detectable increase in the expression of type I IFNs themselves or the need to add exogenous IFNs. Induction of IFN response genes plays an important role in Fc γ R-mediated effects on DCs, as suppression of STAT1 by RNA interference inhibited Fc γ R-mediated DC maturation. These data suggest that the balance of activating/inhibitory Fc γ Rs may regulate IFN signaling in myeloid cells. Manipulation of Fc γ R balance on DCs and monocytes may provide a novel approach to regulating IFN-mediated pathways in autoimmunity and human cancer.

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Abbreviations used: Fc γ R, Fc γ receptor; GEP, gene expression profile; IC, immune complex; IDC, immature DC; IFI, IFN- α inducible; IFNAR, IFN- α receptor; IRG, IFN response gene; STAT, signal transducer and activator of transcription.

The Fc γ R system comprises both activating and inhibitory receptors, and the balance of these two types of receptors determines the outcome of immune complex (IC)-mediated inflammation, immunity, and antibody-based immunotherapy (1). Altering this balance by using a selective blocking antibody against the human inhibitory Fc γ RIIB receptor in the presence of activating Ig ligands in human plasma leads to enhanced generation of antitumor T cell responses (2). Mice deficient in the inhibitory Fc γ R Fc γ RII also show enhanced T cell immunity to model antigens (3). However, the mechanisms by which activating Fc γ Rs mediate maturation of human DCs and enhance adaptive immunity remain to be clarified.

IFNs are pleiotropic cytokines with potent antiviral, antitumor, growth suppressive, and

immunomodulatory properties (4). The cellular effects of both type I (IFN- α and - β) and type II (IFN- γ) IFNs are mediated via activation of the STAT family of transcription factors and downstream activation of a distinct set of "IFN response genes" (IRGs) (5). IFNs play an important role in the regulation of both innate and adaptive immunity (6). For example, IFNs play a critical role in T cell-dependent antibody responses to antigens delivered with the classical complete Freund's adjuvant, DNA vaccines, and immunostimulatory DNA (7–9), and they promote the induction of cytotoxic T cells in vivo (10, 11). IFN-mediated signaling pathways also play an important role in immune surveillance and protection from tumors (12). Dysregulation of IFN signaling has been observed in patients with several autoimmune diseases (6, 13). Therefore, pathways that regulate

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IFN signaling in myeloid cells, particularly DCs, may have a major impact on immunity to tumors and pathogens, as well as autoimmunity. An important aspect of the biology of IFN signaling is that the level of constitutive signaling in the absence of pathogens determines the strength of IFN signaling in response to pathogens (14). Therefore, there is a need to identify the factors that regulate the level of this constitutive or basal IFN signaling.

We show that Fc γ R-mediated maturation of human DCs is associated with a distinct pattern of gene expression. This includes the expression of several inflammation-associated cytokines and chemokines, and the induction of several typical IRGs. These data suggest that the balance of activating/inhibitory Fc γ Rs can regulate the IFN response program in human DCs and monocytes.

RESULTS

A distinct gene expression profile (GEP) of DCs treated with anti-Fc γ RIIB antibody

We have previously shown that treatment of monocyte-derived immature DCs (IDCs) with an anti-Fc γ RIIB-blocking antibody in the presence of Ig ligands in normal human plasma leads to DC maturation and enhancement of anti-tumor T cell immunity (2). To further characterize Fc γ R-mediated enhancement of DC function, we analyzed the GEPs of pure populations of monocyte-derived DCs (MoDcs) from healthy donors ($n = 5$) using Affymetrix Human Genome U133 Plus 2.0 microarrays. IDCs cultured in 1% plasma were treated for 24 h with either anti-Fc γ RIIB or isotype control antibody. To test whether Fc γ R-mediated DC maturation was distinct from other maturation stimuli, we also compared DCs matured using the inflammatory cytokine cocktail (TNF- α , IL-1 β , IL-6, and PGE₂) that is commonly used in DC immunotherapy trials (15). To first validate the GEP data at the protein level, we compared the gene expression data for some of the genes associated with DC maturation (CD83 and CD80) with the detection of corresponding proteins by flow cytometry (Fig. 1 A). As expected, mRNA expression, as well as protein levels of CD80 and CD83, increased in DCs matured with the Fc γ RIIB blocking antibody and in cytokine-matured DCs compared with isotype-treated IDCs. Of the 24,296 expressed genes on the array, 1,801 were differentially regulated in DCs treated with anti-Fc γ RIIB antibody (RIIB DCs) versus those treated with isotype control antibody (Iso DCs). Most of these genes were also up-regulated in DCs matured with inflammatory cytokines (Cyt DCs), suggesting that these reflected changes in gene expression shared with other DC maturation stimuli. However, we also identified a distinct set of 95 genes that were differentially expressed only in RIIB DCs (Fig. 1 B and Table S1, available at <http://www.jem.org/cgi/content/full/jem.20062545/DC1>). Thus, the GEP of RIIB DCs includes genes shared with Cyt DCs, as well as a distinct subset of genes specifically overexpressed only in RIIB DCs.

Changes in type I IFN-induced genes after treatment with anti-Fc γ RIIB antibody

Interestingly, a majority of the genes expressed selectively in RIIB DCs were known to be induced by type I IFNs. To better characterize the IFN-responsive genes in DCs, we treated IDCs ($n = 3$) with 1,000 U/ml of IFN- α 2b and identified 167 genes that were up-regulated by greater than fivefold at 24 h in IFN-treated DCs compared with untreated DCs (Fig. 1 C and Table S2, available at <http://www.jem.org/cgi/content/full/jem.20062545/DC1>). Indeed, 54 of 95 genes in the RIIB DC-specific signature were IFN-inducible genes (Table S1). Up-regulation of IFN-induced genes by anti-Fc γ RIIB antibody was not unique to DCs, but also observed in monocytes treated with this antibody (Fig. 1 C). The IFN-induced genes in these cells included several well-known type I IFN response signature genes, such as IFN- α inducible (IFI) protein 27, myxovirus resistance 1 (Mx1), Mx2, 2'-5' oligoadenylate synthase, cig5, GIP3, IFI16, serpin E, IFI44, STAT1, and IRF-7 (Table S1). To validate the data for expression of IFN-induced genes in the microarray, the expression of two of these genes (IFI27 and Mx1) was also analyzed by real-time PCR, which confirmed the microarray findings (Fig. 1 D). Surprisingly, this increase in IRGs was not associated with an increase in the levels of any of the known type I IFNs themselves (IFN- α , - β , - ω , - τ , and including newer members of the family such as IL28A, IL-28B, and IL-29; Fig. 1 E) and only a modest increase in IFN- γ at the mRNA and protein level as analyzed in the supernatants using Luminex (Table I). It is notable that a low level signal for one of the type I IFN genes (IFN- α 17) was detectable by microarray in DCs treated with isotype control mAb, and consistent with this, low levels of IFN- α were also detectable in the supernatants by Luminex (Fig. 1 E and Table I). Importantly, however, treatment with anti-Fc γ RIIB mAb did not lead to a change in the expression of any of the type I IFNs when compared with isotype mAb-treated cells. Therefore, a distinct component of Fc γ R-mediated activation of DCs is the strong and rapid activation of IFN-induced genes without a concurrent increase in the expression of type I IFNs themselves.

Expression of inflammation-associated chemokines and cytokines in Fc γ R-matured DCs

In addition to the IFN-inducible genes, several other genes involved in inflammation were also differentially up-regulated in RIIB DCs relative to Iso DCs or Cyt DCs. This included several CC chemokines (CCL2/MCP-1, CCL19, CCL3, CCL5/Rantes, and CCL4/MIP1 β), cytokines (IL-1 α), FcR, and complement-related genes. To validate the microarray data at the protein level, the DC culture supernatants were also analyzed for the presence of cytokines and chemokines by Luminex analysis. When compared with Iso DCs, RIIB DCs secreted greater amounts of MIP1 α , IL1 α , IL1 β , IP10, IL-12p70, and CCL5/Rantes, which were also consistent with the microarray data (Table I and Fig. 2). Therefore, DC maturation by selective signaling via activating Fc γ Rs is characterized by a distinct pattern of inflammation-associated genes relative to DCs matured using inflammatory cytokines.

Table I. Expression of cytokines and chemokines secreted by DCs by Luminex analysis

	Iso DC	RIIB DC	Cyt DC	RIIB DC versus Iso DC (p-value)	RIIB DC versus Cyt DC (p-value)
IL-1 α	57.19 (29.4)	699.5 (184.4)	87.6 (80.8)	<0.001	<0.001
IL-1 β	11 (15.6)	668.9 (780)	NE	0.071	NE
IL-2	10.2 (4.1)	14.3 (4.7)	21.9 (14)	0.120	0.170
IL-3	102.2 (16.2)	176 (56.5)	137.6 (35.5)	0.023	0.147
IL-5	0 (0)	0.0	0 (0)	0.000	0.000
IL-6	229.6 (142.6)	7941.8 (4116.4)	NE	0.005	NE
IL-7	0.1 (0.2)	27 (3.8)	16.4 (12.7)	0.000	0.080
IL-8	628.1 (408.1)	10000 (0)	5727 (4992.1)	<0.001	0.069
IL-10	22.5 (2.8)	1200.9 (987.5)	108.6 (111.5)	0.027	0.035
IL-12p40	40.5 (28.4)	4713.3 (5019.4)	3191 (4483.3)	0.056	0.333
IL-12p70	15.9 (16.6)	148.2 (148.5)	29 (20.1)	0.064	0.082
IL-13	8.9 (3.4)	28.9 (21.1)	39.7 (47.2)	0.055	0.345
IL-15	3.8 (7.7)	0.9 (1.8)	0 (0)	0.242	0.178
IFN- γ	116.5 (8.8)	264.2 (61.1)	198.3 (81.6)	0.002	0.122
TNF- α	5.3 (6.3)	1059.8 (911.3)	NE	0.030	NE
Eotaxin	15.5 (19.2)	35 (11)	43.1 (14.5)	0.065	0.204
MCP1	1173.1 (1074)	2776.34 (2655.6)	1314.8 (1908.4)	0.153	0.203
Rantes	27.8 (16.5)	1531 (682)	264.8 (356.3)	0.002	0.008
MIP1a	710.6 (351.3)	9746.9 (388)	3067.8 (2733.7)	<0.001	0.001
IP10	395.3 (185.4)	8629.8 (2364.7)	637.9 (664.9)	<0.001	<0.001
IFN- α	10.2 (9.9)	10.3 (12.5)	28.9 (15.6)	0.493	0.056

IDCs ($n = 4$) were treated with anti-Fc γ RIIB antibody (RIIB DC), inflammatory cytokine cocktail (Cyt DC), or isotype control antibody (Iso DC). 24 h later, the culture supernatant was collected and examined for cytokines and chemokines by Luminex assay. NE, not evaluable, as these cytokines were added exogenously to Cyt DCs.

Mechanism of Fc γ R-mediated induction of type 1 IFN response

Because of the prominence of the IFN signature in RIIB DCs, we focused on further characterizing the mechanism for induction of these responses. Type 1 IFN-mediated activation of its receptor activates Jak protein tyrosine kinases, which in turn phosphorylate STAT proteins, including STAT1 (16). Therefore, we used the detection of phosphorylated STAT1 (P-STAT1) protein by flow cytometry as a marker for IFN-induced response at a single-cell level. Treatment of IDCs with the anti-Fc γ RIIB was associated with an increase in P-STAT1 detectable at 24 h after anti-Fc γ RIIB mAb treatment (Fig. 3 A). The increase in P-STAT1 observed in these experiments was also confirmed by Western blot analysis (Fig. 3 B). Anti-Fc γ RIIB-mediated up-regulation of P-STAT1 was not unique to DCs, but was also seen in monocytes (Fig. 3 A). In prior studies, we have shown that DC maturation induced by anti-Fc γ RIIB requires the presence of activating Ig ligands in normal human plasma (2). Consistent with this, anti-Fc γ RIIB enhanced P-STAT1 in DCs cultured in 1% plasma, but not those cultured in serum-free media (Fig. 3 C).

Next, we studied the effects of IFN- α blockade on anti-Fc γ RIIB-mediated STAT1 phosphorylation. As expected, the addition of IFN- α to IDCs led to an increase in P-STAT1, which was blocked by preincubation of DCs with IFN- α

and IFN- α receptor (IFNAR) blocking antibodies (Fig. 4 A). However, preincubation with the IFN- α and IFNAR blocking antibodies did not abrogate the up-regulation of P-STAT1 in anti-Fc γ RIIB-treated DCs (Fig. 4 B). It is notable, however, that although these mAbs inhibit the increase in P-STAT1 in response to exogenous IFN- α , they do not abolish the constitutive levels of P-STAT1 (Fig. 4 A and not depicted). Similarly, pretreatment of monocytes with anti-IFNAR mAb inhibited the increase in P-STAT1 in response to exogenous IFN- α , but not after treatment with anti-Fc γ RIIB (Fig. 4 C).

As there was a mild, but detectable, increase in IFN- γ in the DC supernatants, we also analyzed the effect of an IFN γ blocking mAb on anti-Fc γ RIIB-induced P-STAT1. The IFN γ blocking antibody was also unable to inhibit the up-regulation of P-STAT1 in anti-Fc γ RIIB-treated DCs (Fig. 4 D). Similar findings were observed in monocytes (unpublished data). Anti-Fc γ RIIB-mediated P-STAT1 up-regulation was rapid and detectable within an hour of mAb treatment in both monocytes and DCs (Fig. 4 E). Together, these data suggest that selective engagement of activating Fc γ Rs leads to rapid activation of IFN signaling without a concurrent increase in the expression of type I IFNs themselves.

Role of activating Fc γ Rs

Induction of DC maturation by anti-Fc γ RIIB antibody requires the presence of activating Ig ligands present in normal

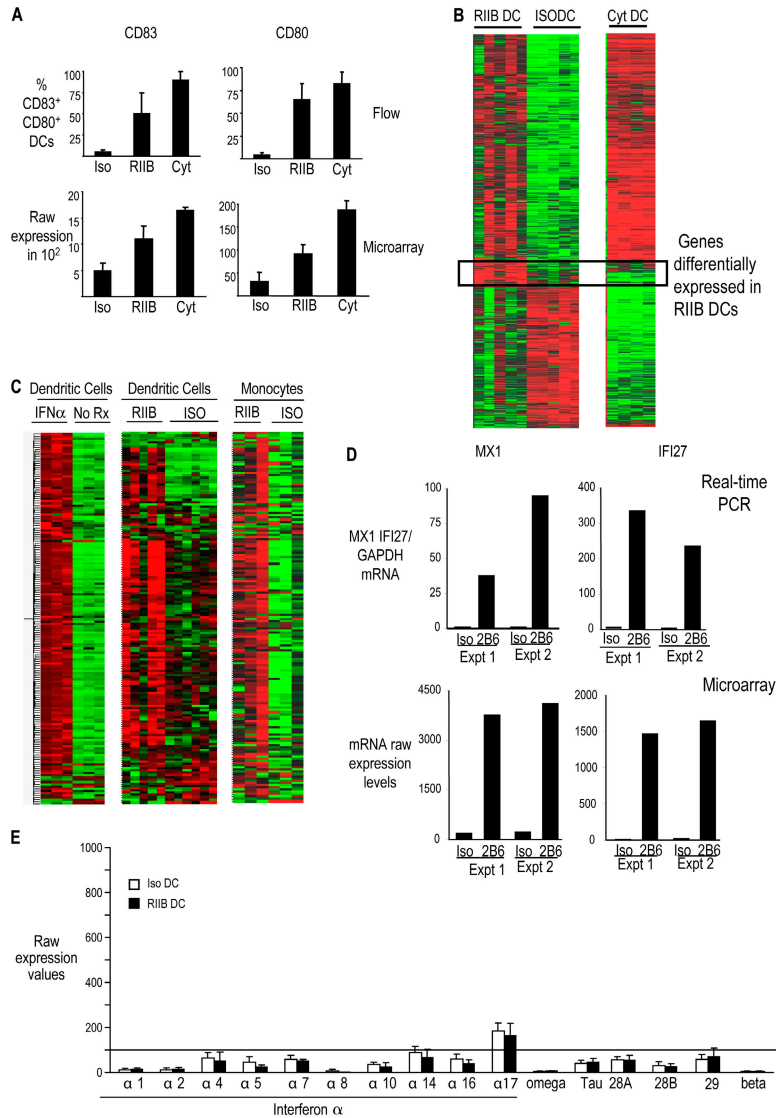


Figure 1. Changes in GEPs of human DCs treated for 24 h with isotype control antibody, anti-Fc γ RIIB antibody, or an inflammatory cytokine cocktail. (A) Validation of microarray. Immature monocyte-derived DCs from healthy donors ($n = 4$) were treated with 20 μ g/ml mouse IgG1 (Iso), 20 μ g/ml anti-Fc γ RIIB blocking antibody (RIIB), or inflammatory cytokine cocktail (IL-1 β , TNF- β , PGE2, and IL-6; Cyt). 24 h later, some DCs from each of the three conditions were analyzed for the expression of DC maturation markers (CD83 and CD80) by flow cytometry. RNA was extracted from the rest of the DCs and gene expression was examined using Human Genome U133 Plus 2 Affymetrix chips. These graphs show the changes in the expression of DC maturation markers (CD80 and CD83) at the level of mRNA (by microarray) and protein (by flow cytometry). (B) Heatmap of 1,801 genes differentially expressed between Iso DC and RIIB DC. Day 5 immature Mo-DCs were treated with either 20 μ g/ml mouse IgG1 (Iso DC), 20 μ g/ml anti-Fc γ RIIB antibody (RIIB DC), or inflammatory cytokine cocktail (IL-1 β , TNF- β , PGE2, and IL-6; Cyt DC). 24 h later, the DCs were harvested and RNA was extracted and analyzed using the Human Genome U133 Plus 2 Affymetrix gene array chips and the GeneSpring software. Genes that were marked as present and expressed above a raw level of 100 were included in the analysis. Iso DC, RIIB DC, and Cyt DC were analyzed to detect genes that were

differentially expressed between the three groups using a parametric test with variance assumed equal (ANOVA) with a p-value cut off of 0.05, followed by the Benjamin and Hochberg false discovery rate multiple correction. 4,759 genes were found to be differentially expressed between Iso DC, RIIB DC, and Cyt DC. Of these 4,759 genes, 1,801 genes were found to be differentially expressed in RIIB DCs compared with Iso DCs (twofold difference; $P < 0.05$). (left) The expression of these 1,801 genes in RIIB DC and Iso DC. (right) The expression of the same 1,801 genes in DCs treated with inflammatory cytokines (Cyt DC). The box shows a subset of 95 genes that are overexpressed only in RIIB DCs compared with both Iso DCs and Cyt DCs. Details of these genes are noted in Table S1. (C) Heat map of IRGs in monocytes and DCs treated with anti-Fc γ RIIB antibody or isotype control antibody. Day 5 monocyte derived IDCs ($n = 3$) were treated with IFN- α 2b (1,000 U/ml) or left untreated. 24 h later, RNA was extracted and analyzed using the Human Genome U133 Plus2 Affymetrix chips and the GeneSpring software (version 7.2). 167 genes were found to be up-regulated by more than fivefold in IFN-treated DCs compared with untreated DCs. (left) Expression of the 167 IFN- α -induced genes in DCs treated with IFN- α and untreated DCs (No Rx). Expression of the 167 IRGs was then compared in DCs ($n = 5$) and monocytes ($n = 3$) treated with anti-Fc γ RIIB mAb (RIIB) or isotype control antibody (Iso). (D) Expression of two

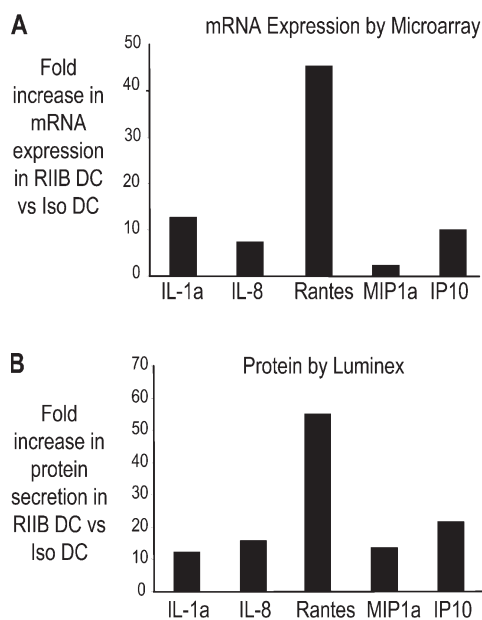


Figure 2. Up-regulation of chemokines/cytokines in Fc γ R-matured DCs. Validation of microarray data at the protein level. Monocyte-derived IDCs (four different donors) were treated with 20 μ g/ml anti-Fc γ RIIB antibody (RIIB DC) or 20 μ g/ml isotype control antibody (mouse IgG1; Iso DC). 24 h later, DCs were harvested and RNA was extracted and analyzed using the U133 Plus2 Affymetrix chips (as in Fig. 1). The culture supernatant was analyzed for the expression of cytokines/chemokines by Luminex analysis. The graphs show the fold increase in mRNA expression (A) or protein secretion (B) for the cytokines and chemokines in RIIB DC compared with Iso DC.

human plasma (2). The Mo-DCs used in these experiments were generated from CD14⁺ monocytes purified using immunomagnetic beads and cultured in 1% plasma. These DCs express higher levels of CD32, as well as CD16, compared with DCs generated from plastic adherent monocytes (Fig. 5 A and not depicted) (17). To evaluate the role of specific activating Fc γ Rs, the DCs were preincubated with antibodies against CD16 and CD32A before blockade of CD32B. Prior incubation with a cocktail of antibodies against activating Fc γ Rs, Fc γ RIIA (CD32A), and Fc γ RIIIA (CD16) inhibited anti-Fc γ RIIB-mediated up-regulation of DC maturation markers (CD80 and CD83; Fig. 5 B).

Preincubation of DCs with a cocktail of antibodies against both activating Fc γ Rs (CD32a and CD16) inhibited anti-Fc γ RIIB-mediated induction of P-STAT1 (Fig. 5 C). These antibodies also blocked anti-Fc γ RIIB-mediated up-regulation of STAT1 in monocytes (Fig. 5 D). This was also associated

IFN-induced genes (Mx1 and IFI27) in DCs treated with anti-Fc γ RIIB mAb or isotype control was analyzed by TaqMan, and expression data was compared with the data obtained by microarray analysis. RNA from two donors (used in the microarray analysis) was analyzed by TaqMan to verify the expression of two IFN-induced genes (Mx1 and IFI27). The figure shows the Taqman expression data compared with the expression data

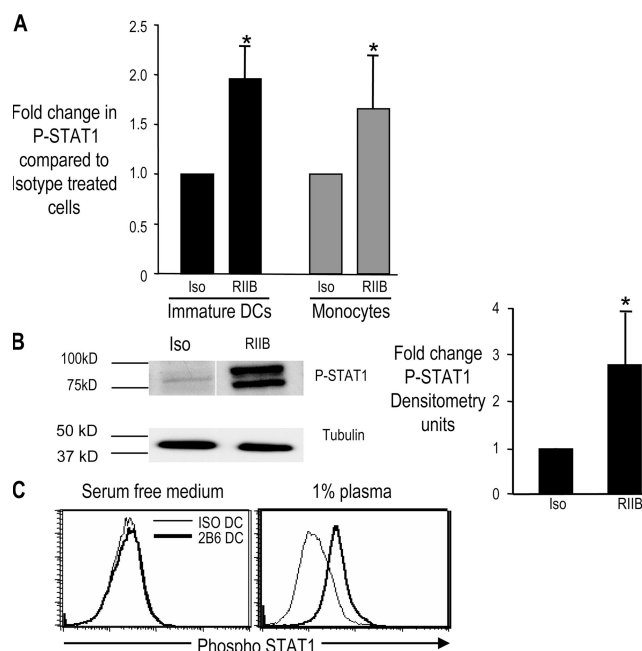


Figure 3. Mechanism of Fc γ R-mediated induction of IFN response. (A) Up-regulation of P-STAT1 in DCs or monocytes treated with anti-Fc γ RIIB mAb versus isotype control. Immature Mo-DCs ($n = 8$) or freshly isolated monocytes ($n = 7$) were treated with anti-Fc γ RIIB antibody (RIIB) or isotype control antibody (Iso). 24 h later, P-STAT1 expression was examined by flow cytometry. The histogram shows fold change in expression of P-STAT1 in anti-Fc γ RIIB antibody-treated versus isotype control antibody-treated cells. *, $P < 0.05$. (B) Western blot confirmation of up-regulation of phosphorylated STAT1 in DCs treated with anti-Fc γ RIIB antibody versus isotype control antibody. Monocyte-derived IDCs were treated with isotype control mouse IgG1 antibody (Iso) or anti-Fc γ RIIB antibody (RIIB). 24 h later, the DCs were harvested and the protein was analyzed on a 7.5% polyacrylamide gel to detect the presence of phosphorylated STAT1. (left) Representative of four separate experiments. (right) The summary of data for quantitative densitometry. Value for P-STAT1 was first normalized against data for α -tubulin in that sample, before comparison between RIIB DCs and Iso DCs. Data shown are the summary of four separate experiments. *, $P < 0.05$. (C) Expression of phosphorylated STAT1 (P-STAT1) in DCs treated with anti-Fc γ RIIB antibody or isotype control antibody in serum-free medium compared with 1% plasma. Immature monocyte-derived DCs were treated with anti-Fc γ RIIB antibody (RIIB DC) or isotype control antibody (Iso DC) in serum-free medium or in medium supplemented with 1% human plasma. 24 h later, the expression of P-STAT1 was examined by flow cytometry. Data are representative of three similar experiments. Data represent the mean \pm the SD.

with down-regulation of the IFN-induced genes on microarray analysis (unpublished data). To examine the relative contribution of specific activating Fc γ Rs in anti-Fc γ RIIB

obtained by microarray analysis. (E) Expression of type I IFNs by microarray. Gene expression data obtained from DCs treated with either isotype antibody or anti-Fc γ RIIB antibody, as in Fig. 1 B, was analyzed for the expression of type I IFN genes. The histogram shows the mean \pm the SD for the mRNA expression from four different donors. Table S1 is available at <http://www.jem.org/cgi/content/full/jem.20062545/DC1>.

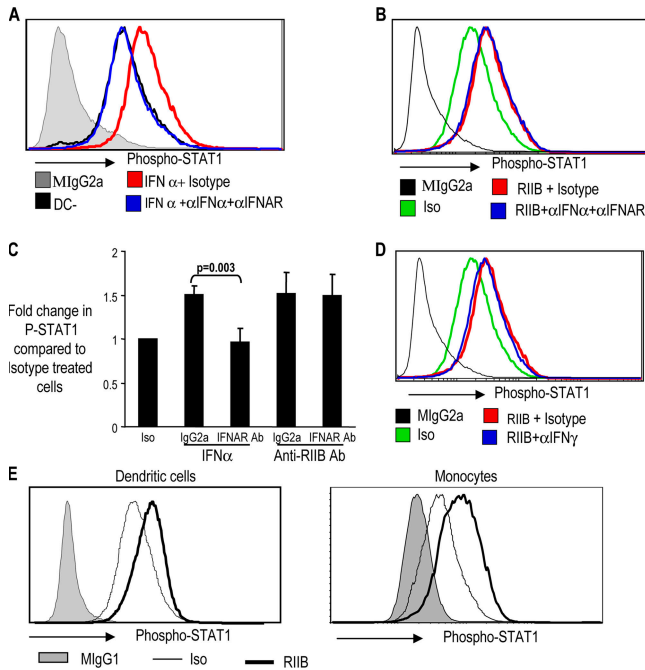


Figure 4. Induction of IFN response by anti-FcγRIIB antibody is not inhibited by blocking antibodies against IFN-α and IFN-γ. (A) Up-regulation of P-STAT1 by exogenous IFN-α is blocked by antibodies against IFN-α. Monocyte-derived IDCs were either left untreated (DC-) or treated with IFN-α2b (1,000 U/ml intron A). The DCs were either treated with a combination of blocking antibodies against IFN-α, as well as IFNAR (αIFNAR; both at 10 μg/ml) or with their isotype control antibodies (Isotype; mouse IgG1 and mouse IgG2a, respectively; both at 10 μg/ml) for 45 min before treatment with IFN-α. The DCs were analyzed for their expression of P-STAT1 by flow cytometry. Gray area of the histogram shows staining of the DCs with isotype control (Mouse IgG2a) for P-STAT1 antibody. The graph represents one of two similar experiments. (B) Effect of blocking antibodies against IFN-α on anti-FcγRIIB-mediated induction of P-STAT1. Monocyte-derived IDCs were treated with anti-FcγRIIB antibody (5 μg/ml RIIB) or mouse IgG1 isotype control antibody (Iso). DCs treated with anti-FcγRIIB antibody were either treated with blocking antibodies against IFN-α and IFNAR (10 μg/ml RIIB + αIFNα + αIFNAR) or isotype control antibody (RIIB + Isotype; 10 μg/ml mouse IgG2a and 10 μg/ml IgG1, respectively) for 45 min before the addition of anti-FcγRIIB antibody. 24 h later, flow cytometry was performed to examine the expression of P-STAT1. Some DCs were also stained with mouse IgG2a, which is isotype control for the P-STAT1 antibody. One of two similar experiments. (C) Effect of anti-IFNAR blocking antibody on anti-FcγRIIB-mediated induction of P-STAT1. Freshly isolated PBMCs were treated with IFN-α2b (1,000 U/ml intron A) or anti-FcγRIIB antibody and an isotype control antibody either in the presence of 20 μg/ml IFNAR antibody (IFNAR Ab) or isotype control antibody (mouse IgG2a). 1 h later, flow cytometry was performed to examine the expression of P-STAT1 on CD14+ monocytes. Data shown are the summary of three similar experiments. *, P < 0.05. Data represent the mean ± the SD. (D) IFN-γ blocking antibodies do not inhibit the up-regulation of P-STAT1 by anti-FcγRIIB antibody. IDCs were treated with either anti-FcγRIIB antibody (5 μg/ml RIIB) or isotype control antibody (Iso). DCs treated with anti-FcγRIIB antibody were either treated with blocking antibodies against IFN-γ and IFNAR (10 μg/ml RIIB + αIFNγ) or isotype control antibody (10 μg/ml RIIB + Isotype; mouse IgG1) for 45 min before the addition of anti-FcγRIIB antibody. 24 h later, flow cytometry was performed to examine the expression of

mAb-mediated STAT1 up-regulation, we also evaluated the effect of individual blockade of FcγRIIIA and FcγRIIA on up-regulation of P-STAT1. Blockade of FcγRIIIA on monocytes with clone 3G8 led to greater inhibition of P-STAT1 up-regulation, compared with FcγRIIA blocking antibody (clone IV.3), suggesting that FcγRIIIA may be the major activating FcγR contributing to STAT1 phosphorylation in monocytes (Fig. 5 E). Aggregation of FcγRs bearing ITAM motifs leads to the activation of Syk family of tyrosine kinases (18, 19). Tassiulas et al. recently demonstrated that Syk and associated adaptor proteins can enhance IFN-α signaling by direct phosphorylation of STAT1 (20). Consistent with this, pretreatment of DCs with a Syk inhibitor, piceatannol, inhibited the up-regulation of P-STAT1, as well as maturation induced by anti-FcγRIIB (Fig. 5 F). Collectively, these data suggest that the signals via activating FcγRs contribute to the induction of P-STAT1 under these experimental conditions.

Role of STAT1 in FcγR-mediated DC maturation

Prior studies have shown that type I IFNs can mediate phenotypic and functional activation of human DCs. We hypothesized that the activation of IFN pathway/STAT1 by anti-FcγRIIB mAb may be essential for FcγR-mediated DC maturation. To test this directly, we knocked down STAT1 protein in IDCs using RNAi. IDCs were electroporated with STAT1 siRNA or nontargeted control siRNA. The expression of STAT1 protein was markedly inhibited at 48 and 72 h in DCs treated with STAT1 siRNA, but not in nontargeted control siRNA (Fig. 6 A). At 72 h after treatment with siRNA, DCs were treated with anti-FcγRIIB or isotype control mAb. DCs treated with STAT1 siRNA showed decreased up-regulation of maturation markers (CD80 and CD83; Fig. 6, B and C). However, these DCs were capable of maturation in response to an inflammatory cytokines cocktail. Therefore, STAT1 plays an important role in FcγR-mediated induction of DC maturation.

DISCUSSION

ICs and antibodies play an important role in autoimmune and allergic inflammation and in protection from infectious pathogens (21). The immunologic effects of ICs depend on the balance between activating and inhibitory FcγRs, including in DCs (3, 21). These data demonstrate that manipulating this balance via antibody-mediated blockade of inhibitory FcγRIIB in the presence of activating ligands in the human

P-STAT1. Some DCs were also stained with mouse IgG2a, which is isotype control for the P-STAT1 antibody. The graph shows one of two similar experiments. (E) Anti-FcγRIIB induced up-regulation of P-STAT1 is rapid. Immature monocyte derived DCs (left) or monocytes (right) were treated with 10 μg/ml anti-FcγRIIB antibody (RIIB) or mouse IgG1 isotype control antibody (Iso). 1 h later flow cytometry was performed to detect the expression of P-STAT1. Gray histogram shows staining with mouse IgG2a isotype control antibody for P-STAT1. The graphs show one of five similar experiments.

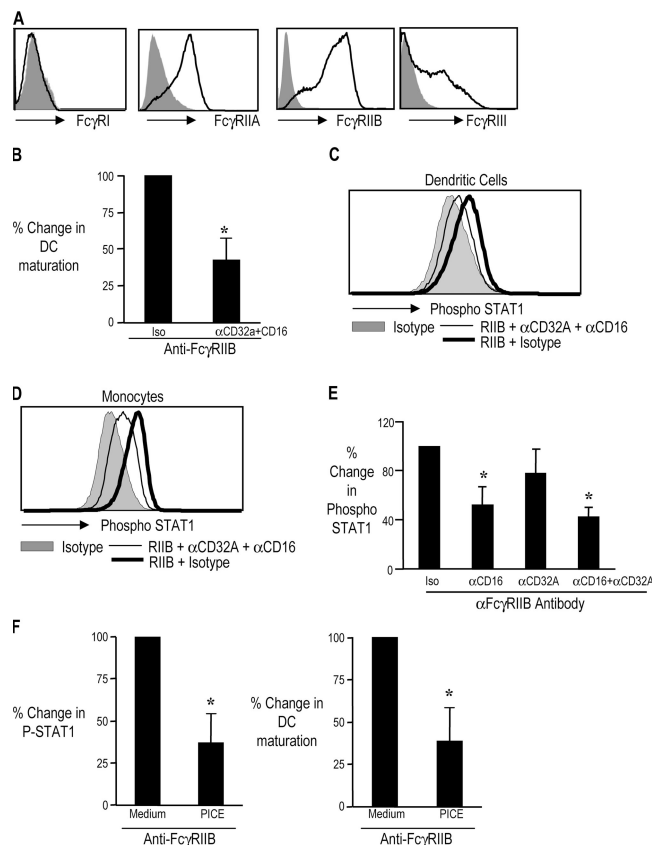


Figure 5. Role of activating Fc γ R in the FcR-mediated DC maturation and induction of P-STAT1. (A) Expression of Fc γ R1, RIIA, RIIB, and RIIIA on DCs generated from CD14⁺ monocytes by flow cytometry. Day 5 monocyte-derived IDCs were examined for the expression of Fc γ R1, Fc γ RIIA, Fc γ RIIB, and Fc γ RIII by flow cytometry. The area in gray represents staining with isotype control antibodies. Figure represents one of six similar experiments. (B) Down-regulation of anti-Fc γ RIIB antibody induced DC maturation by blocking antibodies against activating Fc γ R. IDCs ($n = 4$) were treated with isotype control antibody or anti-Fc γ RIIB antibody either with blocking antibodies against CD32A and CD16 antibody (α CD32A + α CD16; clone IV.3 and 3G8, both at 10 μ g/ml) or their isotype control antibodies (Isotype; mouse IgG2a and IgG1, respectively). 24 h later, the expression of CD80 and CD83 was monitored by flow cytometry, and the double-positive cells were used to assess DC maturation. Change in maturation between isotype-treated and anti-Fc γ RIIB-treated DCs was considered as 100%. The figure shows the percentage of decrease in maturation of DCs treated with blocking antibodies against the activating Fc γ R (CD32A and CD16). Data are a summary of four similar experiments. *, $P < 0.05$. (C and D) Down-regulation of anti-Fc γ RIIB antibody induced P-STAT1 by blocking antibodies against activating Fc γ R. IDCs (C) or PBMCs (D) were treated with mouse IgG1 isotype control antibody (Isotype) or anti-Fc γ RIIB antibody (RIIB). Some of the DCs and PBMCs treated with anti-Fc γ RIIB antibody were pretreated with blocking antibodies against CD32A and CD16 (α CD32A + α CD16; clone IV.3 and clone 3G8, respectively, both 10 μ g/ml) or isotype control antibodies for the CD32A and CD16 blocking antibodies (mouse IgG2a and mouse IgG1, respectively; 10 μ g/ml) for 45 min. Expression of P-STAT1 on CD11c⁺ DCs or CD14⁺ monocytes was examined by flow cytometry. Data are representative of four similar experiments for DCs and three for monocytes. (E) PBMCs were treated with MlgG1 isotype control antibody or anti-Fc γ RIIB antibody. Some of the anti-Fc γ RIIB-treated PBMCs were pretreated

serum has major effects on gene expression and activation status of human monocytes and DCs. Signaling via activating Fc γ R on DCs provides a distinct form of maturation stimulus that is characterized by the expression of P-STAT1 and several IRGs, as well as inflammation-associated chemokines/cytokines. Activation of P-STAT1 is critical to Fc γ R-mediated effects on DCs, because suppression of STAT1 inhibits Fc γ R-mediated DC maturation.

The finding that Fc γ R signaling can alter the expression of IRGs has implications for the regulation of IFN signaling in Fc γ R-expressing myeloid cells. Typically, induction of IFNs in response to viral pathogens is mediated by cytosolic receptors (e.g., RIG1 and MDA5) in infected cells, or endosomal receptors (e.g., Toll receptor family) in antigen-presenting cells (22). Downstream signaling in response to IFNs and activation of Jak-STAT pathway is also highly regulated (e.g., by the PIAS or SOCS family of proteins) (23). However, studies by Taniguchi et al. have shown that the strength of IFN response to pathogens depends on the level of constitutive signaling in the absence of pathogens (14). Our data suggest that the balance of activating/inhibitory Fc γ R signaling in DCs may have a direct impact on this basal level of IFN signaling in myeloid cells in the absence of pathogens. This balance may, in principle, be altered both in the context of specific Fc γ R polymorphisms, or the ability of ICs to bind activating versus inhibitory Fc γ R.

Aggregation of Fc γ R bearing ITAM motifs leads to the activation of Syk family of tyrosine kinases (18, 19). Tassiulas et al. recently showed that Syk and associated adaptor proteins can enhance IFN- α signaling by direct phosphorylation of STAT1 (20). Our data extend these observations and suggest a role for activating Fc γ R on human monocytes and

with blocking antibody to CD16 (3G8), CD32A (IV.3), or a combination of blocking antibodies against CD16 and CD32A (3G8 + IV.3) or their isotype control antibodies (mouse IgG1 and mouse IgG2a, respectively; Iso). 1 h later, the expression of P-STAT1 in CD14⁺ monocytes was analyzed by flow cytometry. The change in expression of P-STAT1 between MlgG1-treated cells and anti-Fc γ RIIB-treated cells was considered as 100%. The figure shows the percentage of decrease in this P-STAT1 phosphorylation by blocking antibodies against CD16 (α CD16) and CD32A (α CD32A), either alone or in combination (α CD16 + α CD32A). The histogram shows the summary of experiments on three separate donors (*, $P < 0.05$). (F) Inhibition of Syk tyrosine kinase abrogates anti-Fc γ RIIB induced up-regulation of P-STAT1. Immature Mo-DCs were treated with medium alone (medium) or Syk tyrosine kinase inhibitor piceatannol (PICE) at 5 μ M concentration. 30 min later, the DCs were treated with either anti-Fc γ RIIB antibody or isotype control mouse IgG1 antibody (MlgG1). 24 h later, the DCs were analyzed for their P-STAT1 expression and maturation (CD80 and CD83) by flow cytometry. Change in expression of P-STAT1, as well as maturation between MlgG1-treated cells and anti-Fc γ RIIB-treated cells, was considered as 100%. The histogram on the left shows the percentage of decrease in this P-STAT1 phosphorylation by piceatannol treatment. The histogram on the right shows the percentage of decrease in DC maturation by piceatannol treatment. The graphs are a summary of experiments on three separate donors (*, $P < 0.05$). Data represent the mean \pm the SD.

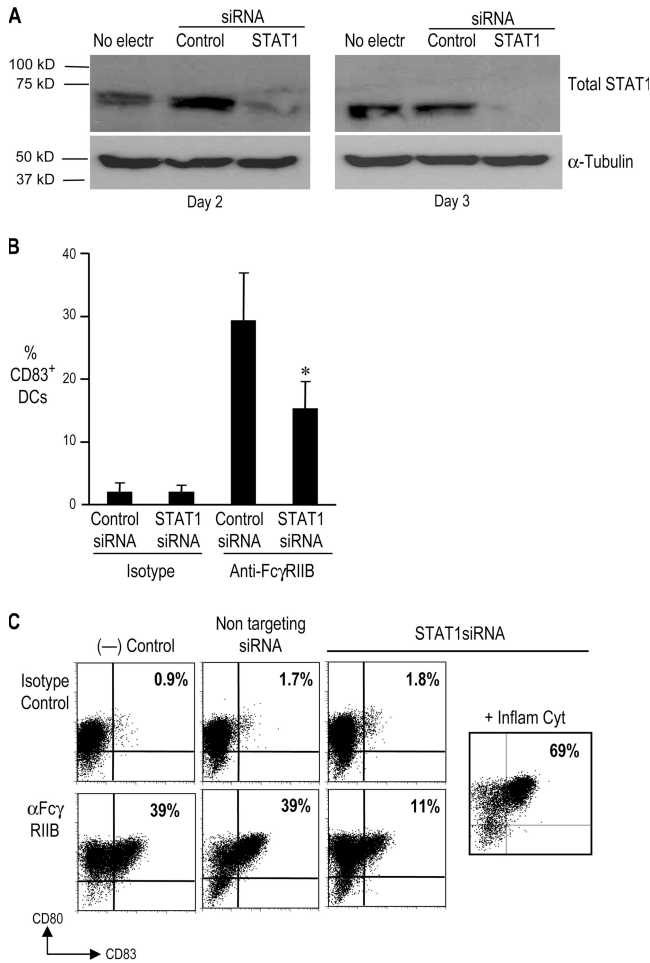


Figure 6. FcγR-mediated DC maturation can be inhibited by STAT1 knockdown. (A) Day 4 IDCs were electroporated with 10 μg of STAT1 siRNA (STAT1) or nontargeting siRNA (Control). Some DCs were cultured without electroporation (No electr). The DCs were harvested 48 and 72 h after electroporation, and a Western blot was performed to detect total STAT1 protein. Data are representative of two separate experiments. (B and C) DCs were harvested 72 h after electroporation with either STAT1 siRNA or nontargeting control siRNA. Some nonelectroporated DCs were also harvested. The DCs were treated with anti-FcγRIIB antibody or isotype control antibody. Some of the DCs electroporated with STAT1 siRNA were also treated with the inflammatory cytokine cocktail. 24 h later, DC maturation was determined by examining the expression of CD80, CD83, and CD11c by flow cytometry. (B) Summary of data in four independent experiments (mean ± the SD; *, P < 0.05). (C) Data from a representative experiment.

DCs in enhancing the expression of IRGs. Further studies are needed to evaluate this pathway.

Importantly, the FcγR-mediated induction of IRGs was not associated with a detectable increase in the expression of any of the type I IFNs, and it did not require the addition of exogenous IFNs to these cultures. Blocking mAbs against IFNAR and IFN-α, which block the increase in P-STAT in response to exogenous IFN-α, also fail to block this response. The induction of STAT1 phosphorylation by anti-FcγRIIB was rapid and detectable within 1 h of treatment with anti-

FcγRIIB blocking antibodies, which also argues against the IFN signature being caused by synthesis of new IFNs. It is also important to emphasize that the IFN signature is not observed in DCs treated specifically with inflammatory cytokines (Cyt DCs). Collectively, these data suggest a novel link between FcγR signaling and IFN response in human DCs and monocytes. However, it is important to note that low levels of type I IFNs (IFN-α17) were expressed in control DCs and detectable in supernatants (although unchanged in response to anti-FcγRIIB mAb). Furthermore, the IFNAR blocking antibodies were unable to block the constitutive level of P-STAT1. Therefore, the observed effects are consistent with the ability of FcγR-mediated signals to modify the constitutive IFN signaling in these cells (14).

The nature of specific activating FcγRs and other signals that are important for regulating IFN signaling in DCs requires further study and may be cell/tissue type specific. In our experiments, the signal was delivered via both FcγRIIIA and FcγRIIA. Recent studies have shown that FcγRIIIA is expressed on distinct subsets of human monocytes and DCs (24, 25), although their functional significance remains to be clarified. Polymorphisms in both FcγRIIA and FcγRIIIA may therefore have a significant impact on immune activation mediated by ICs and antibody-coated tumor cells (26, 27).

Selective blockade of inhibitory FcγR signaling on DCs also leads to the up-regulation of several other genes not in the type I IFN pathway, but implicated in autoimmune and allergic inflammation. These include several CC chemokines, such as CCL5, CCL4, and CCL3, which were documented at both mRNA and protein level. Expression of these chemokines may allow FcγR-matured DCs to attract several immune cells, including T cells, monocytes, DCs, and eosinophils, which play an important role in inflammation (28).

These data also have several clinical implications. Dysregulation of FcγR signaling may contribute to the altered IFN signaling observed in several autoimmune states (13, 27). Therefore, either restoring the FcγR balance or inhibiting STAT1 may help suppress immune activation in lupus. Indeed, recent studies have shown that only small changes in inhibitory FcγR can have major effects on immunopathology in lupus (29). Altered IFN signaling and STAT1/STAT3 ratio in myeloid DCs and other myeloid cells has also been implicated in the loss of protective antitumor immunity and immune surveillance in several models (12, 30, 31). Interestingly, recent studies have also implicated a more direct role for ICs in promoting tumor growth (32). Our observations may serve to link these findings together. Manipulating FcγR balance on tumor-infiltrating DCs (e.g., via anti-FcγRIIB antibodies) may be of value as a general approach to regulate endogenous antitumor immunity. Modulation of IFN signaling may also contribute to the efficient induction of T cell immunity observed by selectively targeting ICs or antibody-coated tumor cells to activating FcγRs on human and murine DCs (2, 3, 33, 34); therefore, manipulating the FcγR balance may also be targeted for improving the efficacy of antibody-based therapy (35) or DC-based immunotherapy of cancer (36).

MATERIALS AND METHODS

Generation of DCs. PBMCs were isolated from blood of healthy donors via density gradient centrifugation using Ficoll-Hypaque (GE Healthcare). CD14⁺ cells were obtained using CD14 microbeads and LS columns (Miltenyi Biotec) and used to generate DCs. DCs were generated by culturing CD14⁺ cells in RPMI-1640 medium with L-glutamine (Mediatech) supplemented with 1% single donor plasma, gentamicin (20 µg/ml; BioWittaker), and 0.01 M Hepes (Cambrex). For some experiments, DCs were cultured in AIM-V serum free medium (Invitrogen), 20 ng/ml GM-CSF (Immunex) and 12.5 ng/ml IL-4 (R&D Systems) were added to the culture on days 0, 2, and 4. On day 5 of culture, IDCs were either treated with 10–20 µg/ml anti-FcγRIIB antibody (clone 2B6; MacroGenics), 10–20 µg/ml mouse IgG1 isotype control (Sigma-Aldrich), or inflammatory cytokine cocktail (10 ng/ml IL-1β, 1,000 U/ml IL-6, 10 ng/ml TNF-α (all from R&D Systems), and prostaglandin E₂ (1 µg/ml, Sigma-Aldrich). As R/H polymorphisms of the activating receptor CD32A can potentially impact FcR-mediated DC maturation, they were monitored by flow cytometry using specific antibodies (17, 37). All donors in this study had HH/HR genotype (not depicted). Purity of the DC preparations was evaluated by flow cytometry and was >95%. The percentage of contaminating NK cells was <0.1%.

Microarray analysis. IDCs on day 5 of culture ($n = 5$) or CD14⁺ monocytes isolated from immunomagnetic bead selection ($n = 3$) were treated with 20 µg/ml anti-FcγRIIB mAb, isotype control (mouse IgG1 mAb), or inflammatory cytokine cocktail. After 24 h of culture, cells were pelleted in RLT buffer. RNA was extracted from the DCs using the RNeasy kit (QIAGEN) as per the manufacturer's protocol. Total RNA (1–2 µg) was labeled and hybridized using the Enzo T7 labeling kit (Life Sciences) on Affymetrix Human Genome U133 Plus 2.0 microarray (Affymetrix). Washing and scanning was done with Fluidics Station 400 and GeneChip Scanner 3000 (both from Affymetrix) following the manufacturer's protocol. Affymetrix GCOS 1.2 software was used to obtain the raw signal and present/absent gene data. Further analysis of the data was performed using GeneSpring software version 7.2 (Silicon Genetics).

Data analysis of microarray data. The methodology for analysis of microarray data was similar to that used by Napolitani et al. (38). The microarray data was normalized as follows. The signal values below 0.01 were set to 0.01. The percentile for all the measurements in each sample was calculated using the values for all genes not marked "absent." Each measurement was divided by the 50th percentile of all measurements in that sample. Each gene was divided by the median of its measurements in all samples. Genes that were marked as present and expressed a raw level of >100 were included in the analysis. 29,499 genes passed this filter. Gene expressions in DCs treated with inflammatory cytokines, FcγRIIB blocking antibody, and isotype control antibody were compared, and genes with statistically different expression between the groups based on the values of the replicates were calculated using a parametric test with variance assumed equal (ANOVA) using a p-value cut off of 0.05, followed by Benjamin and Hochberg false discovery rate multiple-test correction. A gene list of 4,759 significant genes was obtained and used to determine fold changes in expression between the three different conditions. Of these 4,759 genes, 1,801 genes were differentially regulated between FcγRIIB antibody-treated DCs by twofold or higher, compared with isotype control antibody-treated DCs. Microarray data are available in the National Center for Biotechnology Information Gene Expression Omnibus (<http://www.ncbi.nih.gov/geo/>) under accession no. GSE7509.

Real time PCR (TaqMan) analysis. RNA was isolated with RNeasy Mini kit (Qiagen) and RT-PCR was conducted with Assays-on-Demand primer-probe for IFI27 and Mx1 using the ABI PRISM 7700 sequence detection system (both Applied Biosystems). Expression of GAPDH was monitored as a housekeeping gene. Reactions were set up in triplicates using EZ PCR Core Reagents (Applied Biosystems) according to the manufacturer's instructions with 20 ng of total RNA. Relative expression of target genes was calculated using the comparative threshold cycle method.

Luminex assay. Cell supernatants from 24-h DC cultures were analyzed for 20 cytokines and chemokines using the Protein Multiplex Immunoassay kit (Biosource International) as per the manufacturer's protocol. In brief, Multiplex beads (Biosource) were vortexed and sonicated for 30 s, and 25 µl was added to each well and washed two times with wash buffer. The samples were diluted 1:2 with assay diluent and loaded onto a Multiscreen BV 96-well filter plate (Millipore) with 50 µl of incubation buffer already added to each well. Serial dilutions of cytokine standards were prepared in parallel and added to the plate. Samples were then incubated on a plate shaker at 600 rpm in the dark at room temperature for 2 h. The plate was applied to a Multi-screen Vacuum Manifold (Millipore) and washed twice with 200 µl of wash buffer. 100 µl of biotinylated anti-human Multi-Cytokine Reporter (Biosource International) was added to each well. The plate was incubated on a plate shaker at 600 rpm in the dark at room temperature for 1 h. The plate was applied to a Multiscreen Vacuum Manifold (Millipore) and washed twice with 200 µl of wash buffer. Streptavidin-phycoerythrin was diluted 1:10 in wash buffer, and 100 µl was added directly to each well. The plate was incubated on a plate shaker at 600 rpm in the dark at room temperature for 30 min. The plate was then applied to the vacuum manifold, washed twice, and each well was resuspended in 100 µl wash buffer and shaken for 1 min. The assay plate was then transferred to the Bio-Plex Luminex 100 XYP instrument (Millipore) for analysis. Cytokine concentrations were calculated using Bio-Plex Manager 3.0 software with a 5-parameter curve-fitting algorithm applied for standard curve calculations.

Detection of STAT1 phosphorylation by flow cytometry. The methodology for the detection of P-STAT1 by flow cytometry was adapted from that described by Lesinski et al. (39). Freshly isolated PBMCs, purified CD14⁺ monocytes, or monocyte-derived IDCs were treated with either mouse IgG1 isotype control antibody or the FcγRIIB blocking antibody. After a 1–24-h culture, the cells were labeled with the mouse anti-human phospho STAT1 antibody (clone pY701; BD Biosciences) as per the manufacturer's protocol. In some experiments, cells were treated with 1,000 IU/ml IFN-α 2b (Schering-Plough). In some experiments, DCs or monocytes were pretreated with 10 µg/ml anti-IFNAR mAb (Fitzgerald Industries), 10 µg/ml anti-IFN α antibody (clone MMHA-1; PBL Laboratories), 10 µg/ml anti-IFN-γ antibody (clone MMHG-1; PBL Laboratories), 10 µg/ml anti-CD16 (clone 3G8; BD Biosciences), and 10 µg/ml antiCD32A (clone IV.3) or isotype controls.

Blocking inhibitory FcγR on human Mo-DCs and evaluation of their maturation. In brief, IDCs were harvested on day 5 of culture and treated with either 10 µg/ml anti-human FcγRIIB blocking antibody (clone 2B6) or 10 µg/ml mouse IgG1 isotype control antibody (Sigma-Aldrich). After overnight culture, DCs were harvested to assess DC maturation. In some experiments, DCs were pretreated with 10 µg/ml CD16 blocking antibody (clone 3G8; BD Biosciences), and 10 µg/ml CD32A blocking antibody (clone IV.3) or isotype controls for 1 h before treatment with anti-human FcγRIIB blocking antibody. The following antibodies were used for evaluating the surface changes associated with DC maturation: CD11c-APC, CD80-PE, CD83-FITC, CD86-PE, and HLA-DR-FITC (all obtained from Becton Dickinson).

Inhibition of STAT1 synthesis by RNA interference. CD14⁺ monocytes were cultured in 1% plasma supplemented with IL4 and GMCSF on days 0 and 2, as described in Generation of DCs. On day 4, the IDCs were harvested, washed with Opti-MEM I medium without phenol red (Invitrogen) and resuspended in Opti-MEM I at a concentration of 2.5×10^7 cells/ml. 4×10^6 IDCs were electroporated with 10 µg of STAT1 (siRNA; siGENOME SMARTpool; Thermo Fisher Scientific) or nontargeting siRNA (siCONTROL Nontargeted siRNA; Thermo Fisher Scientific) in a 4-mm electroporation cuvette using the ECM830 Square Wave Electroporator (Harvard Apparatus). The pulse conditions were a square wave pulse of 500V and 0.5 ms. The electroporated DCs were immediately resuspended in complete medium (RPMI 1% plasma) supplemented with IL4 and GMCSF.

Inhibition of STAT1 protein was examined by Western blot analysis 2 and 3 d after electroporation.

Western blot analysis to detect total and phosphorylated STAT1.

Nonelectroporated DCs, as well as DCs electroporated with either nontargeting RNAi or STAT1 RNAi, were harvested, washed in PBS, and lysed with the radioimmunoprecipitation assay buffer containing 150 mM NaCl, 10 mM Tris, pH 7.2, 0.1% SDS, 1% Triton X-100, 1% deoxycholate, 5 mM EDTA, 100 mM sodium orthovanadate, and protease inhibitors (protease inhibitor cocktail tablet; Boehringer Mannheim). 25–50 μ g of protein was resolved on a 7.5% polyacrylamide gel and transferred to a nitrocellulose membrane. Blots were probed with total STAT1 antibody (rabbit anti-human STAT1, 1:1,000; Cell Signaling Technologies) overnight according to the manufacturers protocol. The membrane was treated with goat anti-rabbit horseradish peroxidase secondary antibody (1:5,000; Southern Biotechnology). Detection was done using the ECL plus Western blot detection reagents (GE Healthcare).

For the detection of phosphorylated STAT1 protein, IDCs were treated with mouse IgG1 isotype control antibody or anti-Fc γ R1IB antibody. 24 h later, the cells were harvested and lysed with the radioimmunoprecipitation assay buffer containing 150 mM NaCl, 10 mM Tris, pH 7.2, 0.1% SDS, 1% Triton X-100, 1% deoxycholate, 5 mM EDTA, protease inhibitors, 10 mM sodium orthovanadate, and 10 mM β glycerol phosphate. Protein was resolved on a 7.5% polyacrylamide gel and transferred to a nitrocellulose membrane. Blots were probed with mAb specific for phosphorylated STAT1 (rabbit anti-human phospho-STAT1 antibody; 1:1,000, Cell Signaling Technologies) overnight, according to the manufacturers protocol. The membrane was treated with goat anti-rabbit horseradish peroxidase secondary antibody (1:5,000; Southern Biotechnology). Detection was done using the ECL plus Western blot detection reagents (GE Healthcare), and analyzed by quantitative densitometry using ImageJ software (W.S. Rasband, National Institutes of Health, Bethesda, MD; <http://rsb.info.nih.gov/ij/>). Data for P-STAT1 were first normalized against values for α -tubulin in that sample, before comparison to other samples.

Detection of DC maturation after STAT1 knockdown with the STAT1 RNAi.

Nonelectroporated DCs or DCs electroporated with either STAT1 siRNA or nontargeting control siRNA were harvested 72 h after electroporation. DCs were treated with anti-Fc γ R antibody or mouse IgG1 isotype control antibody. 24 h later, flow cytometry was performed to examine the surface expression of CD83, CD80, and CD11c.

Online supplemental material. Table S1 shows a list of 95 genes that are differentially expressed in DCs treated with anti-Fc γ R1IB antibody. Table S2 shows a list of genes up-regulated by more than fivefold at 24 h in immature Mo-DCs cultured in the presence of IFN- α 2b (1,000 U/ml), compared with untreated DCs. The MIAME checklist for microarray analysis is presented as supplemental text.

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