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Article

A Cluster of Interferon-γ-Inducible p65 GTPases Plays a Critical Role in Host Defense against *Toxoplasma gondii*

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SUMMARY

Interferon-γ (IFN-γ) is essential for host defense against intracellular pathogens. Stimulation of innate immune cells by IFN-γ upregulates ~2,000 effector genes such as immunity-related GTPases including p65 guanylate-binding protein (Gbp) family genes. We show that a cluster of *Gbp* genes was required for host cellular immunity against the intracellular parasite *Toxoplasma gondii*. We generated mice deficient for all six *Gbp* genes located on chromosome 3 (Gbp^{chr3}) by targeted chromosome engineering. Mice lacking Gbp^{chr3} were highly susceptible to *T. gondii* infection, resulting in increased parasite burden in immune organs. Furthermore, Gbp^{chr3}-deleted macrophages were defective in IFN-γ-mediated suppression of *T. gondii* intracellular growth and recruitment of IFN-γ-inducible p47 GTPase Irgb6 to the parasitophorous vacuole. In addition, some members of Gbp^{chr3} restored the protective response against *T. gondii* in Gbp^{chr3}-deleted cells. Our results suggest that Gbp^{chr3} play a pivotal role in anti-*T. gondii* host defense by controlling IFN-γ-mediated Irgb6-dependent cellular innate immunity.

INTRODUCTION

Interferon-γ (IFN-γ) is an important T helper 1 (Th1) cell cytokine that strongly suppresses the growth and survival of intracellular pathogens (Boehm et al., 1997). Stimulation of innate immune cells such as macrophages and dendritic cells by IFN-γ results in robust gene expression of a number of effector molecules. These include immunity-related GTPases such as the Mx proteins, immunity-related p47 GTPases (Irgs), and p65 guanylate-binding proteins (Gbps) (Shenoy et al., 2007; Taylor et al., 2004). Mx proteins have been shown to participate in host defense against RNA viruses such as influenza and vesicular stomatitis virus (Sadler and Williams, 2008). Among the Irgs, mice deficient in Irgm1 (Lrg-47) are highly susceptible to *Listeria*, *Salmonella*, and mycobacteria (Deretic, 2006; MacMicking, 2004). Furthermore, Gbps have recently been shown to induce antibacterial responses involving phagocytic oxidases, autophagic effectors, and inflammasome (Kim et al., 2011; Shenoy et al., 2012). Thus, IFN-γ-inducible immunity-related GTPases play pivotal roles in antiviral and antibacterial immune systems.

Toxoplasma gondii is an obligatory intracellular protozoan parasite that infects virtually all warm-blooded vertebrates including human and mouse (Boothroyd, 2009; Israelski and Remington, 1993). Infection of immunocompromised individuals such as those suffering from AIDS or those being treated with chemotherapy often leads to fatal toxoplasmosis encephalitis (Montoya and Remington, 2008). Innate immune cells, which recognize microbial components mainly via Toll-like receptors (TLRs) and the chemokine receptor CCR5, are essential in controlling *T. gondii* infection via the production of proinflammatory cytokines such as interleukin-12 (IL-12) (Aliberti et al., 2003; Hunter and Remington, 1995; Yarovinsky and Sher, 2006). IL-12 potentiates polarization of naive T cells to Th1 cells, from which IFN-γ is produced in an antigen-dependent fashion (Trinchieri, 2003; Whitmarsh et al., 2011). IFN-γ-inducible GTPases are also important for the inhibition of *T. gondii* growth by IFN-γ. Mice lacking Irgm1, Irgd (Irg-47), Irgm3 (Igtp), or Irga6 (Iigp1) are susceptible to acute and chronic infection (Collazo et al., 2001; Howard et al., 2011; Taylor et al., 2000). IRGs are recruited to the parasitophorous vacuole (PV), a membrane formed during invasion that is maintained to surround the intracellular

replicating parasites. Accumulation of Irgs eventually leads to disruption of the integrity of the PV membranes (Howard et al., 2011; Ling et al., 2006; Taylor et al., 2007; Zhao et al., 2008).

Not only Irgs but also Gbps are known to accumulate around the PV shortly after *T. gondii* invasion (Degrandi et al., 2007). Moreover, because virulent strains of *T. gondii* inhibit the recruitment of Gbps around the PV (Degrandi et al., 2007; Virreira Winter et al., 2011), Gbps are considered anti-*T. gondii* defensive factors. Among Gbps, Gbp1 and Gbp2 are reported to modulate cellular proliferation (Gorbacheva et al., 2002; Guenzi et al., 2001). In addition, Gbp1 is involved in the regulation of matrix metalloproteinase 1 in cancer cell lines (Guenzi et al., 2003; Li et al., 2011). Although in vitro studies have been reported, the physiological protective role of Gbps against *T. gondii* remains uncertain. The mouse genome carries 13 *Gbp* genes (11 active members and 2 pseudogenes) that are organized in clusters and share a high degree of homology (Kresse et al., 2008). Six and seven family members are tandemly aligned on chromosomes 3 and 5, respectively (Kresse et al., 2008). Such a complex configuration has hampered in vivo investigation of the *Gbp* genes through genetic approaches.

To elucidate the in vivo functional contribution of the Gbps to host defense against *T. gondii*, we have generated mice lacking the entire cluster of *Gbp*s on chromosome 3 (Gbp^{chr3}) by Cre-loxP-based chromosome engineering. Gbp^{chr3}-deleted mice were highly susceptible to *T. gondii* infection with a considerably increased parasite burden in tissues. Furthermore, Gbp^{chr3}-deleted macrophages showed defective suppression of parasite growth in response to IFN-γ. Although parasite infection-induced production of oxidants and proinflammatory cytokines as well as autophagy-related 4b (Atg4b) recruitment to the parasites were normal, IFN-γ-induced disruption of the PV membrane and localization of Irgs such as Irgb6 and Irgb10 to the PV were compromised in Gbp^{chr3}-deleted cells. Moreover, endogenous Gbps colocalized and interacted with Irgb6. The reintroduction of *Gbp1*, *Gbp5*, or *Gbp7* into Gbp^{chr3}-deleted cells *partially* restored the IFN-γ-dependent anti-*T. gondii* response. Taken together, these results demonstrate that this cluster of Gbps has a defensive function against *T. gondii* by positively regulating IFN-γ-inducible Irgb6-dependent cellular innate immunity.

RESULTS

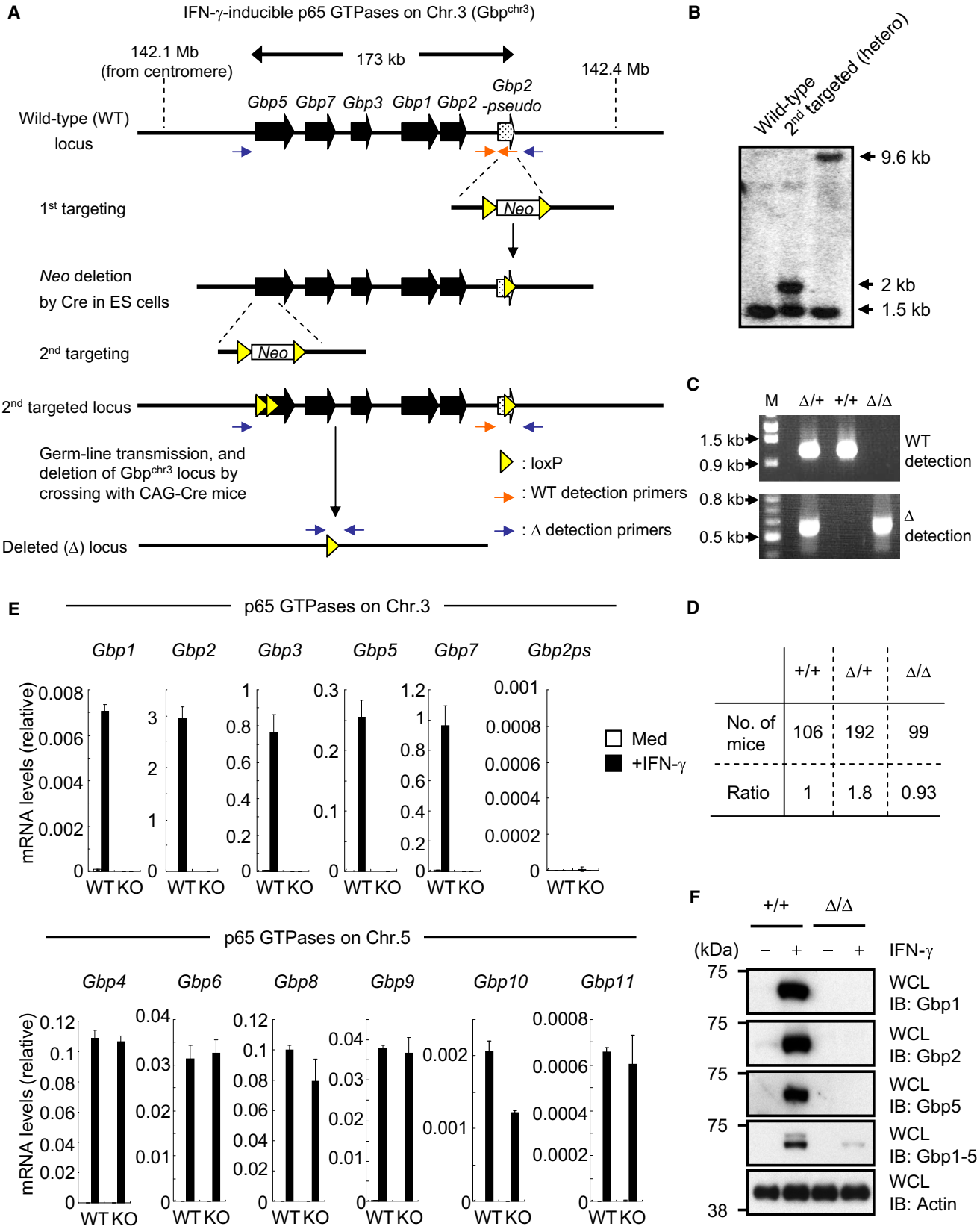
Generation of Mice Lacking the Entire *Gbp* Locus on Chromosome 3

To assess the anti-*T. gondii* immunity of Gbp^{chr3} in vivo, we generated embryonic stem (ES) cells possessing loxP sites at the most proximal and distal loci from the centromere in the gene cluster (*Gbp5* and *Gbp2ps*, respectively) by sequential conventional gene targeting methods (Figure 1A and Figure S1 available online). Deletion of the entire Gbp^{chr3} locus spanning 173 kb was achieved by crossing the F1 mice with CAG-Cre transgenic mice and was confirmed by Southern blotting and PCR (Figures 1B and 1C). Gbp^{chr3}-deleted mice were successfully obtained by intercrossing heterozygous mice, were born at the expected Mendelian ratio, and were healthy and normal in specific-pathogen-free conditions (Figure 1D). Under nonstimulated conditions, Gbp^{chr3}-deleted mice showed normal parameters of cellular immunity (Figure S1H). To test whether the expression of Gbp^{chr3} was correctly ablated in Gbp^{chr3}-deleted cells, we confirmed by quantitative RT-PCR that the mRNAs derived from Gbp^{chr3} (*Gbp1*, *Gbp2*, *Gbp3*, *Gbp5*, *Gbp7*, and *Gbp2ps*) were not induced in response to IFN-γ in Gbp^{chr3}-deleted cells (Figure 1E). On the other hand, mRNAs for the Gbps on chromosome 5 (*Gbp4*, *Gbp6*, *Gbp8*, *Gbp9*, *Gbp10*, and *Gbp11*) were normally induced. We further analyzed Gbp^{chr3} expression by protein immunoblotting and found that Gbp1, Gbp2, and Gbp5 proteins were not detected. A weak signal was observed when we performed blotting with Gbp1-5 monoclonal antibodies that are raised against amino acids 1–300 mapping at the N terminus of human GBP1 and that recognize a shared epitope among Gbp1, Gbp2, Gbp3, Gbp4, and Gbp5. Expression of the gene encoding Gbp4, which is not disrupted in Gbp^{chr3}-deleted cells, could account for this weak signal (Figure 1F).

Gbp^{chr3} Confer Resistance to *T. gondii* In Vivo

First, mice lacking *Gbp2ps* alone or deficient in both *Gbp2ps* and *Gbp5* were infected intraperitoneally with the avirulent *T. gondii* strain ME49 and shown to exhibit survival rates similar to those of wild-type mice (Figures S1C and S1F). In contrast, infection of Gbp^{chr3}-deleted mice revealed that these mutant mice were highly prone to die after *T. gondii* infection (Figure 2A). Because Gbp1 and Gbp5 were recently shown to be involved in host defense against *Listeria monocytogenes* (Kim et al., 2011; Shenoy et al., 2007), we challenged Gbp^{chr3}-deleted mice with *L. monocytogenes* by intraperitoneal injection. Wild-type and Gbp^{chr3}-deleted mice exhibited comparable bacterial burdens in tissues and survival rate (Figures S2A and S2B). To monitor the course of infection more accurately, we generated transgenic ME49 *T. gondii* expressing luciferase and monitored the kinetics by in vivo imaging in wild-type and Gbp^{chr3}-deleted mice infected with 10^2 parasites. Significant increases in parasite number were observed at days 7, 8, and 9 in Gbp^{chr3}-deleted mice compared with those in wild-type mice (Figures 2B and S2C). We next compared parasite burdens in organs from the infected mice. The parasite load showed an excellent correlation with luciferase counts in vitro (Figures S2D and S2E). We collected the spleens and mesenteric lymph nodes from wild-type or Gbp^{chr3}-deleted mice 9 days after a challenge with luciferase-expressing parasites and calculated the parasite numbers according to the luciferase signal. The parasite load in the tissues originating from Gbp^{chr3}-deleted mice was markedly elevated in comparison with that from wild-type mice (Figure 2C). Taken together, these findings demonstrate that Gbp^{chr3} protects against the spreading and proliferation of *T. gondii* in vivo.

Next we examined immune responses during parasite infection in wild-type and Gbp^{chr3}-deleted mice. IL-12 is important for the development of type I immunity, in which IFN-γ-producing $CD4^+$ and $CD8^+$ T cells play central roles for anti-*T. gondii* responses (Hunter et al., 1995). The concentrations of IL-12p40 and IFN-γ measured in sera were similarly increased in wild-type and Gbp^{chr3}-deleted mice infected with *T. gondii* (Figure 2D). Furthermore, cellularity in spleens and IFN-γ production from splenic $CD4^+$ and $CD8^+$ T cells in response to anti-CD3 was comparable in wild-type and Gbp^{chr3}-deleted mice (Figures 2E and S2F), suggesting that the high susceptibility to *T. gondii* in Gbp^{chr3}-deleted mice was not due to defects in production of IL-12 or IFN-γ or in T cell responses.


A
IFN-γ-inducible p65 GTPases on Chr.3 (Gbp^chr3)
142.1 Mb
(from centromere)
173 kb
142.4 Mb
Gbp5 Gbp7 Gbp3 Gbp1 Gbp2 Gbp2 -pseudo
Wild-type (WT) locus
1st targeting
Neo
Neo deletion by Cre in ES cells
2nd targeting
Neo
2nd targeted locus
Germ-line transmission, and deletion of Gbp^chr3 locus by crossing with CAG-Cre mice
Deleted (Δ) locus
: loxP
: WT detection primers
: Δ detection primers
B
Wild-type
2nd targeted (hetero)
9.6 kb
2 kb
1.5 kb
C
M Δ/+ +/+ Δ/Δ
1.5 kb
0.9 kb
WT detection
0.8 kb
0.5 kb
Δ detection
D
+/+ Δ/+ Δ/Δ
No. of mice 106 192 99
Ratio 1 1.8 0.93
E
p65 GTPases on Chr.3
Gbp1 Gbp2 Gbp3 Gbp5 Gbp7 Gbp2ps
mRNA levels (relative)
Med
+IFN-γ
WT KO
p65 GTPases on Chr.5
Gbp4 Gbp6 Gbp8 Gbp9 Gbp10 Gbp11
F
+/+ Δ/Δ
(kDa) – + – + IFN-γ
75
WCL IB: Gbp1
WCL IB: Gbp2
WCL IB: Gbp5
WCL IB: Gbp1-5
38
WCL IB: Actin

Gbp^{chr3} Are Essential for IFN-γ-Induced Reduction of *T. gondii* Infection in Macrophages

Macrophages play a vital role in IFN-γ-mediated cellular innate immunity against *T. gondii* (Suzuki et al., 1988). Peritoneal macrophages represent a major cell type targeted by the parasite at the early stage after intraperitoneal infection (Jensen et al., 2011). To analyze the impact of *Gbp*^{chr3} deficiency in this cell type, we infected wild-type or *Gbp*^{chr3}-deleted peritoneal macrophages with *T. gondii* expressing luciferase in the presence of IFN-γ and assessed the luciferase units at 1, 12, 24, 36, or 48 hr postinfection (Figure 3A). We observed higher luciferase emissions in *Gbp*^{chr3}-deleted cells than in wild-type cells at all time points tested except for 1 hr. Next, we analyzed the IFN-γ-dependent reduction of parasite burden by counting parasite numbers and the luciferase units in macrophages and mouse embryonic fibroblasts (MEFs) from wild-type or *Gbp*^{chr3}-deleted mice. The dose-dependent reduction of parasite numbers and luciferase signals in both cell types lacking *Gbp*^{chr3} was less pronounced than that in wild-type cells (Figures 3B and S3A). On the other hand, costimulation of tumor necrosis factor-α (TNF-α), which is known to strongly enhance antitoxoplasmal activity in macrophage in combination with IFN-γ (Sibley et al., 1991), abrogated the difference between wild-type and *Gbp*^{chr3}-deleted cells (Figure S3B). These data suggested a selective impairment in IFN-γ-mediated reduction of parasite burden.

IFN-γ leads to inhibition of *T. gondii* proliferation and promotes its clearance from macrophages (Ling et al., 2006). To measure clearance, we next used confocal microscopy to compare the degree of parasite infection and growth in macrophages isolated from wild-type or *Gbp*^{chr3}-deleted mice. At 5 hr postinfection, the percentage of cells infected with the parasites was comparable between wild-type and *Gbp*^{chr3}-deleted mice (Figure 3C). In contrast, the infection rate of IFN-γ-stimulated macrophages from *Gbp*^{chr3}-deleted mice was remarkably higher than that from wild-type cells after 20 hr (Figures 3C, 3D, and S3C). Next, we compared the rate of parasite replication in wild-type and *Gbp*^{chr3}-deleted macrophages in the presence of IFN-γ by counting the parasite numbers per PV. In *Gbp*^{chr3}-deleted cells, the number of parasites per PV was modestly increased compared with that in wild-type cells (Figures 3D and 3E), indicating that Gbp^{chr3} are not only required for clearance but also inhibit parasite replication in IFN-γ-stimulated macrophages.

Impact of Gbp^{chr3} Deficiency on IFN-γ-Mediated Anti-*T. gondii* Response

Among Gbp^{chr3}, Gbp1 and Gbp7 have been shown to participate in antibacterial host defense by inducing an oxidative response and recruiting autophagy effectors such as Atg4b to *L. monocytogenes* and *Mycobacterium bovis* (Kim et al., 2011). To test whether these mechanisms also apply to anti-*T. gondii* defense, we first measured IFN-γ-induced oxide ion (O^{2-}) production in wild-type or *Gbp*^{chr3}-deleted cells infected with *T. gondii* (Figure 4A). The O^{2-} production in *Gbp*^{chr3}-deleted macrophages was normally enhanced. Next, we examined Atg4b recruitment to the parasite by confocal microscopy. As observed for pathogenic bacteria, Atg4b colocalized with intracellular *T. gondii* in wild-type cells. The increased recruitment of Atg4b to parasites in *Gbp*^{chr3}-deleted macrophages was not altered during the course of infection (Figures 4B and S4A). Taken together, these results suggest that the anti-*T. gondii* action of Gbp^{chr3} operates independently of O^{2-} and Atg4b recruitment.

IFN-γ is also known to augment proinflammatory cytokine production in response to TLR ligands, and *T. gondii* possesses a profilin that can serve as a TLR11 ligand called profilin-like molecule (Plattner et al., 2008; Yarovinsky et al., 2005). We analyzed the production of proinflammatory cytokines in macrophages. Production of tumor necrosis factor α, IL-6, and IL-12p40 in IFN-γ-treated *Gbp*^{chr3}-deleted cells in response to *T. gondii* infection or lipopolysaccharide (LPS) was normal (Figure S4B), indicating that Gbp^{chr3} are dispensable for IFN-γ-mediated production of proinflammatory cytokines.

Because nitric oxide (NO) was previously reported to inhibit *T. gondii* replication in vitro (Adams et al., 1990), we assessed NO production in wild-type and *Gbp*^{chr3}-deleted cells. Nonstimulated, *T. gondii*-infected, or LPS-treated macrophages from *Gbp*^{chr3}-deleted mice produced similar concentrations of nitrite ion (NO^{2-}) compared with that produced by wild-type cells (Figure S4C). Furthermore, hepatic expression of iNOS mRNAs was unchanged between wild-type and *Gbp*^{chr3}-deleted mice in *T. gondii* infection (Figure S4D). In spite of normal NO production in *Gbp*^{chr3}-deleted macrophages (Figure S4C), the inhibition of the parasite replication was modestly impaired by the *Gbp*^{chr3} deficiency (Figure 3E). To examine whether NO is involved in inhibition of parasite replication in *Gbp*^{chr3}-deleted cells, we compared the replication in *Gbp*^{chr3}-deleted cells between those untreated and those treated with aminoguanidine (AG), an NO inhibitor (Figure 4C). Although the infection rate was not affected at all, AG-treated *Gbp*^{chr3}-deleted cells contained significantly larger numbers of the parasites per vacuole than control cells, indicating that NO indeed plays a major role in inhibition of *T. gondii* replication in *Gbp*^{chr3}-deleted macrophages in vitro. To further determine relative importance of Gbp^{chr3} in the IFN-γ-dependent anti-*T. gondii* response in vivo, we compared parasite burdens and the survival rate of wild-type and

Figure 1. Generation of *Gbp*^{chr3}-Deleted Mice

(A) The gene targeting strategy for *Gbp*^{chr3} locus by chromosome engineering.

(B) Southern blot analysis of offspring from the heterozygote intercrosses. Genomic DNA was extracted from mouse tails, digested with KpnI and PstI, electrophoresed, and hybridized with the radiolabelled probe indicated in Figure S1G. Southern blotting resulted in a 1.5 kb band for wild-type locus, a 2.0 kb band for deleted (Δ) locus, and 9.6 kb for second targeted locus.

(C) PCR detection of mice with homozygous deletion of *Gbp*^{chr3} locus. Primers used are denoted in (A).

(D) Numbers of offspring by intercross of heterozygous (Δ/+) mice.

(E) Quantitative PCR analysis of the expression of the indicated Gbp mRNA in wild-type (WT) or *Gbp*^{chr3}-deleted (KO) peritoneal macrophages unstimulated or stimulated with 100 ng/ml IFN-γ.

(F) Peritoneal macrophages treated with 100 ng/ml IFN-γ for 24 hr were lysed. The lysates were detected by protein immunoblot with the indicated Abs.

Data are representative of two (B, C, E) and three (F) independent experiments. See also Figure S1.

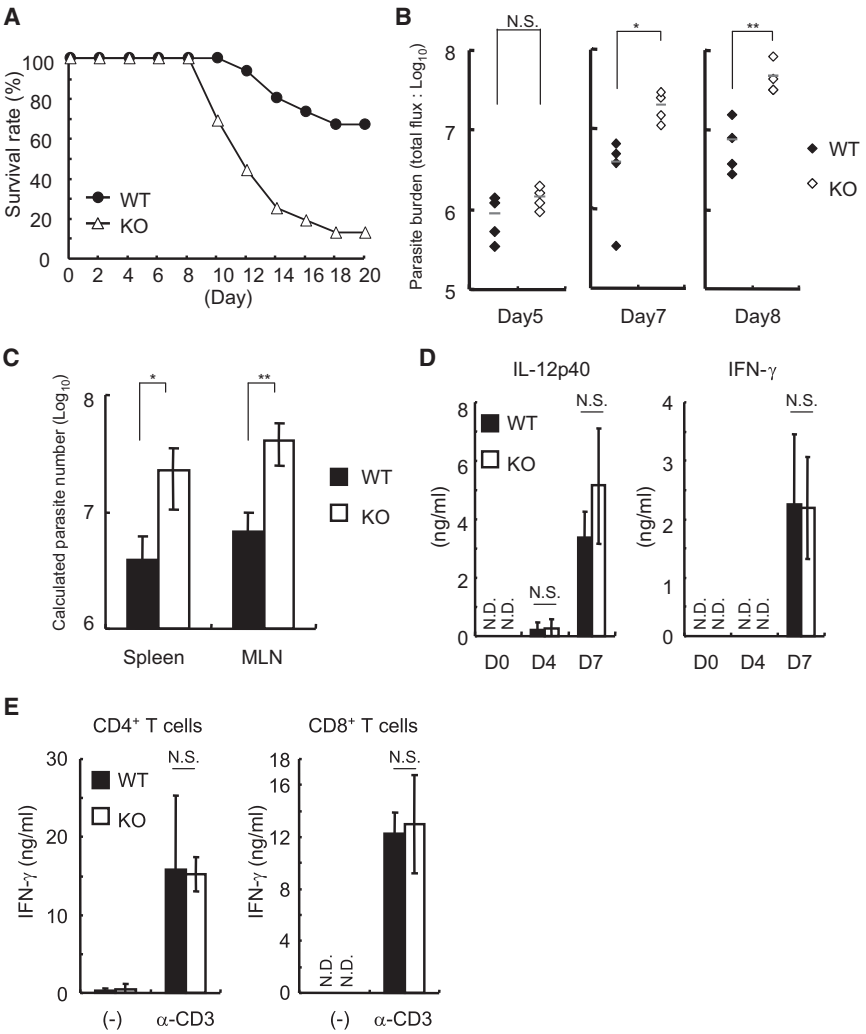


Figure 2. *Gbp*^{chr3}-Deleted Mice Are Highly Susceptible to *T. gondii*

(A) Wild-type (n = 15) or *Gbp*^{chr3}-deleted (KO) (n = 16) mice were infected with 1 × 10^2 *T. gondii*, and the survival rates were monitored for 20 days.

(B) Total photon emission analysis from wild-type or *Gbp*^{chr3}-deleted mice (n = 4 per group) infected with 1 × 10^2 ME49 *T. gondii* luciferase-expressing parasites at days 5, 7, or 8 after infection. The flux (photons/s/cm^2/sr) was determined as a measure of parasite burden. *p < 0.01, **p < 0.02.

(C) Quantification of parasites in indicated tissues from 4 mice per group at day 9 postinfection by the standard curve in Figure S2. Indicated values are means ± SD of quadruplicates. *p < 0.005, **p < 0.03.

(D) Sera were taken from indicated mice (n = 4 per group) at the indicated time postinfection of 1 × 10^2 *T. gondii*. Serum concentrations of the indicated cytokines were determined by ELISA.

(E) Splenic CD4$^+$ or CD8$^+$ T cells from indicated mice (n = 4 per group) at day 7 postinfection of 1 × 10^2 *T. gondii* were cultured in the presence of 5 μg/ml plate-bound anti-CD3 for 24 hr. Concentration of IFN-γ in the culture supernatants was measured by ELISA. Indicated values are means ± SD of triplicates.

N.D., not detected; N.S., not significant. Data are representative of three (C) and two (B, D, E) independent experiments. Data in (A) are pooled from three independent experiments with five or six mice per group. See also Figure S2.

Gbp^{chr3}-deleted mice treated with anti-IFN-γ (Figures 4D and 4E). Although the mortality was not increased by anti-IFN-γ treatment in *Gbp*^{chr3}-deleted mice (Figure 4D), *Gbp*^{chr3}-deleted mice treated with anti-IFN-γ displayed much higher parasite burdens than did control IgG-treated groups (Figure 4E). These results indicated that Gbp^{chr3} do not fully account for the IFN-γ-dependent anti-*T. gondii* immunity in vitro and in vivo.

Impaired Disruption of the PV Membrane in *Gbp*^{chr3}-Deleted Macrophages

Gbp^{chr3} deficiency resulted in compromised IFN-γ-induced clearance of *T. gondii* in macrophages (Figure 3). The killing by IFN-γ-activated macrophages is accompanied by blebbing of the PV membrane shortly after parasite entry (Ling et al., 2006; Zhao et al., 2008). Therefore, we examined by electron microcopy the morphology of the parasite PV membrane in wild-type and *Gbp*^{chr3}-deleted macrophages 6 hr postinfection. Blebbing and vesiculation were easily detectable in the vicinity of the parasite PV membrane in wild-type cells (Figure 5A). In sharp contrast, in *Gbp*^{chr3}-deleted cells the parasites were surrounded by a continuous intact PV membrane (Figure 5B), suggesting that Gbps affect IFN-γ-induced clearance of *T. gondii* shortly after invasion. Moreover, 24 hr postinfection, a markedly increased number of dead parasites with damaged parasite membranes or no PV membranes were observed in wild-type cells compared with those in *Gbp*^{chr3}-deleted cells (Figure 5C and data not shown).

Defective Irgb6 Recruitment to PVs in *Gbp*^{chr3}-Deleted Macrophages

A previous study demonstrated that macrophages lacking the autophagy protein Atg5 exhibit defective blebbing of the *T. gondii* PV (Zhao et al., 2008), which is reminiscent of the phenotype observed in *Gbp*^{chr3}-deleted cells (Figure 5B). Atg5 has been reported to be required for the recruitment of Irga6 to the PV (Zhao et al., 2008). Irgs including Irga6 (also known as IIGP1), Irgb6 (TGTP), and Irgb10 were shown to be phosphorylated and dampened by *T. gondii* ROP18, a parasite-secreted kinase that acts as a virulence effector molecule (Fentress et al., 2010; Steinfeldt et al., 2010). The CTG strain of *T. gondii* produces an extremely low *ROP18* mRNA expression (Boothroyd and Dubremetz, 2008; Saeij et al., 2006; Taylor et al., 2006); hence the CTG parasites as well as ME49 are permissive for Irg recruitment to the PV membrane (Saeij et al., 2006; Taylor et al., 2006). By using the CTG strain expressing luciferase, we found that the IFN-γ-dependent decrease in CTG parasites was impaired in *Gbp*^{chr3}-deleted macrophages (Figure 6A),

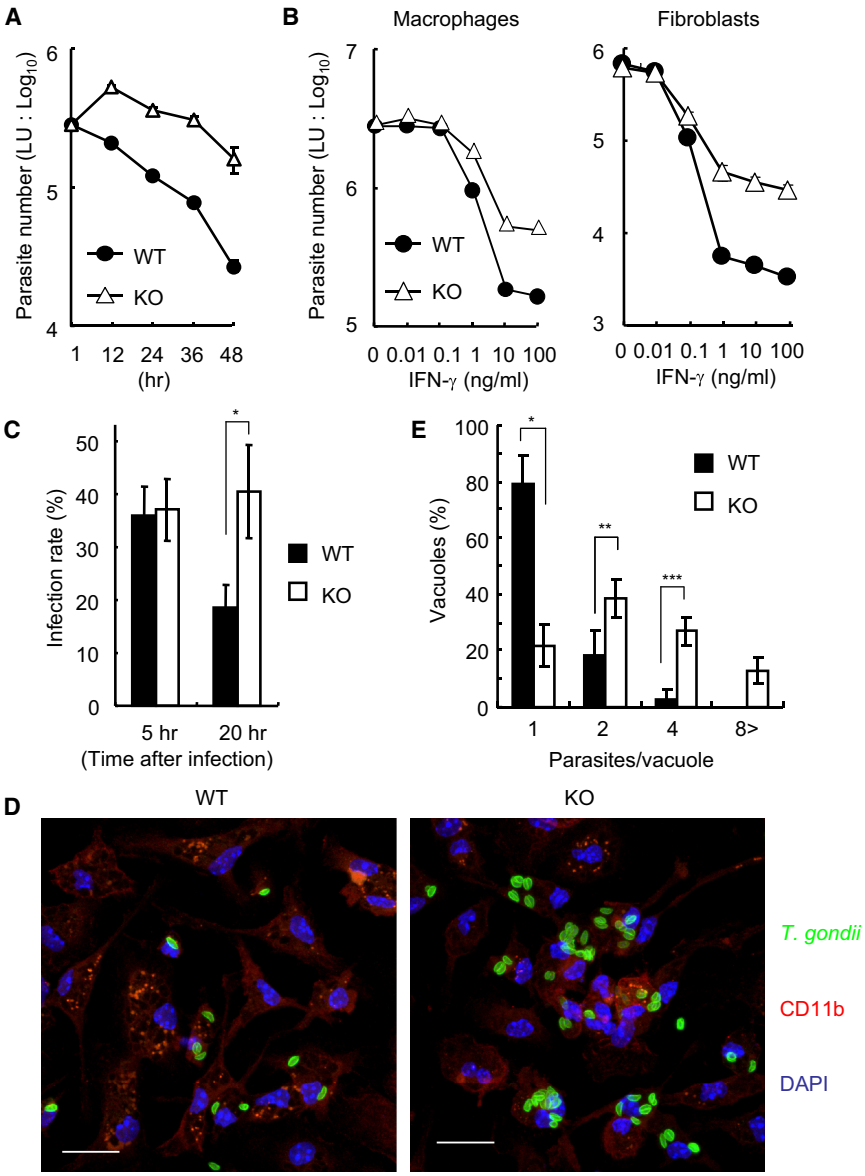


Figure 3. Impaired IFN-γ-Mediated Parasite Clearance in *Gbp*^{chr3}-Deleted Macrophages

(A) Wild-type (WT) and *Gbp*^{chr3}-deleted (KO) peritoneal macrophages were treated with 100 ng/ml IFN-γ for 24 hr. IFN-γ-treated cells were infected with ME49 *T. gondii* expressing lucifease (moi = 0.5) and harvested at the indicated point postinfection. The luciferase units (LU) were assayed with the lysates. Indicated values are means ± SD of triplicates.

(B) WT and KO peritoneal macrophages or MEFs were untreated or treated with the indicated concentrations of IFN-γ for 24 hr. Untreated or IFN-γ-treated cells were infected with ME49 *T. gondii* expressing luciferase (moi = 0.5) and harvested at 36 hr postinfection. The LU were assayed with the lysates. Indicated values are means ± SD of triplicates.

(C) The percentage of WT and KO macrophages containing at least one *T. gondii* parasite at the indicated points postinfection. Indicated values are means ± SD of triplicates. $^{**}p < 0.03$.

(D) WT and KO peritoneal macrophages were treated with 100 ng/ml IFN-γ for 24 hr. IFN-γ-treated cells were infected with ME49 *T. gondii* (moi = 0.5), fixed at 24 hr postinfection, and stained with rabbit anti-*T. gondii* (Alexa 488, green) or rat anti-Cd11b (Alexa 594, red). Scale bars represent 20 μm.

(E) The number of *T. gondii* parasites per vacuole in WT or KO macrophages at 24 hr postinfection. Indicated values are means ± SD of triplicates. $^{*}p < 0.02$, $^{**}p < 0.04$, $^{***}p < 0.002$.

Data are representative of five (A), three (B), and two (D) independent experiments. Data in (C) and (E) are pooled from three independent experiments in which almost 150 cells and 140 vacuoles were counted, respectively. See also Figure S3.

suggesting that Irg-mediated immunity could be affected by *Gbp*^{chr3} deficiency. To test this possibility, we first analyzed the amount of Irgb6 protein expression in IFN-γ-treated macrophages. Similar amounts of Irgb6 protein were induced in wild-type and *Gbp*^{chr3}-deleted cells upon stimulation by IFN-γ (Figure 6B). Next, we tested the recruitment of Irgb6 to the PV by confocal microscopy. At 6 hr postinfection, Irgb6 was detectable at the PV in wild-type cells. In contrast, the recruitment of Irgb6 was severely impaired in *Gbp*^{chr3}-deleted macrophages (Figure 6C). The extent of Irgb6 recruitment was analyzed at 2, 4, and 6 hr postinfection. Compared with that in wild-type cells, the percentage of Irgb6-positive parasites in *Gbp*^{chr3}-deleted cells was significantly lower at every time point analyzed (Figure 6D). In addition to Irgb6, we tested the recruitment of other Irgs such as Irgb10, Irga6, and Irgm3 (Figure S5). The recruitment of Irgb10 to the PV was also impaired in *Gbp*^{chr3}-deleted cells (Figure S5A), whereas that of Irga6 was comparable between wild-type and *Gbp*^{chr3}-deleted cells (Figures S5B and S5C). We failed to observe the accumulation of Irgm3 to the PVs even in IFN-γ-stimulated wild-type cells (data not shown). Taken together, these results suggest that Gbp^{chr3} are required for recruitment of Irgb6 and Irgb10 to the PV of *T. gondii* in IFN-γ-activated macrophages.

Gbps Interact with Irgb6

In a previous study, an overexpressed green fluorescent protein (GFP)-Gbp1 fusion protein was shown to colocalize with endogenous Irgb6 and *T. gondii* parasites (Virreira Winter et al., 2011). We confirmed these observations with endogenous proteins by costaining for anti-Gbp1-5 and anti-Irgb6 in wild-type macrophages infected with *T. gondii* (Figure 6E). Furthermore, intensity profile analysis revealed that the signals for both proteins were detected at almost the same sites (Figure 6F). Then, we assessed the interaction between Gbps and Irgb6 in endogenous settings. Immunoprecipitation with anti-Gbp1-5 coprecipitated Irgb6 in IFN-γ-stimulated macrophages, and the amount of Irgb6 coimmunoprecipitated was markedly increased upon *T. gondii* infection (Figure 6G). These results suggest that the

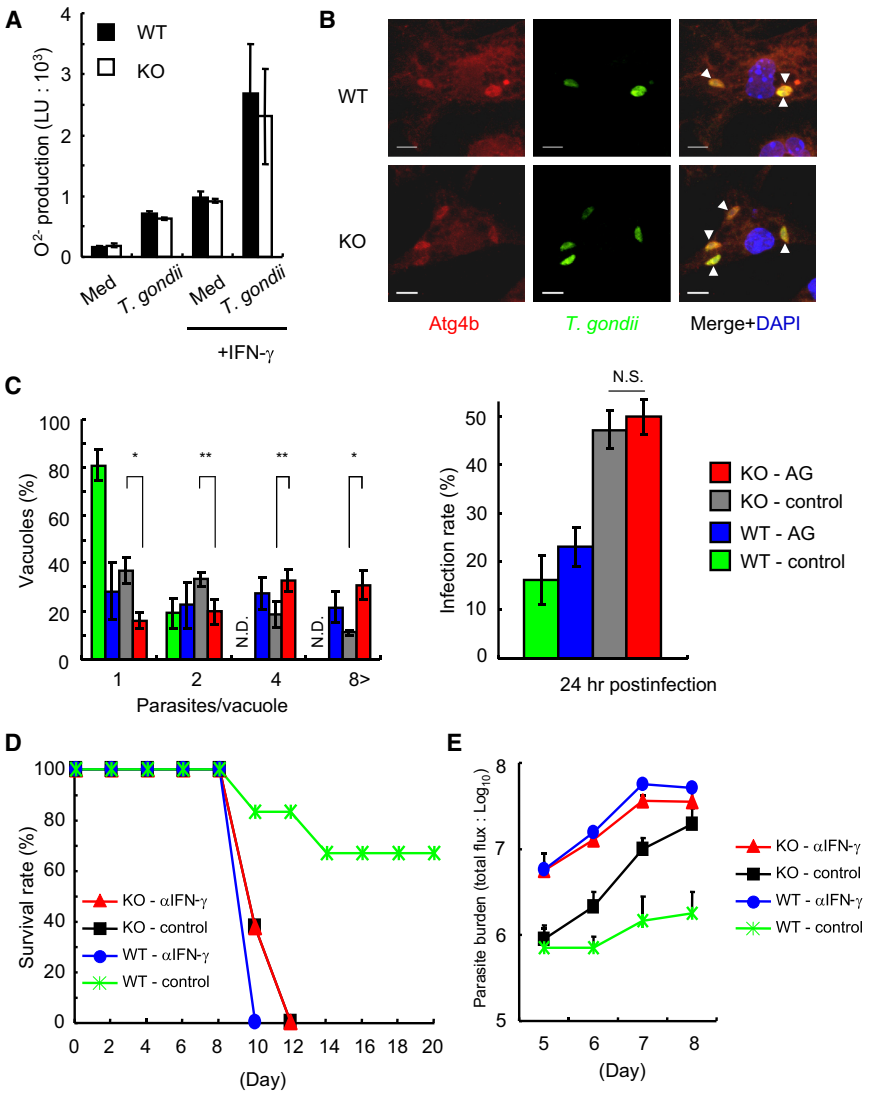


Figure 4. Assessment of IFN-γ-Dependent Responses in *Gbp*^{chr3}-Deleted Mice

(A) Wild-type (WT) and *Gbp*^{chr3}-deleted (KO) peritoneal macrophages were treated with 100 ng/ml IFN-γ for 24 hr. Untreated or IFN-γ-treated cells were uninfected or infected with ME49 *T. gondii* (moi = 0.5) for 24 hr. Concentration (light unit; LU) of O^{2-} in cells was measured with an O^{2-}-specific chemiluminescence reagent by luminometer. Indicated values are means ± SD of triplicates.

(B) WT and KO peritoneal macrophages were treated with 100 ng/ml IFN-γ for 24 hr. IFN-γ-treated cells were infected with ME49 *T. gondii* (moi = 0.5), fixed at 6 hr postinfection, and stained with mouse anti-*T. gondii* (Alexa 488, green) or rabbit anti-Atg4b (Alexa 594, red). Scale bars represent 5 μm.

(C) WT and KO peritoneal macrophages were untreated or treated with 100 ng/ml IFN-γ for 24 hr in the presence or absence of 100 μM AG. Cells were infected with ME49 *T. gondii* expressing luciferase (moi = 0.5). Left: The number of parasites per vacuole in WT or KO macrophages at 24 hr postinfection is shown. Indicated values are means ± SD of triplicates. *p < 0.01, **p < 0.03. Right: The percentage of WT and KO macrophages containing at least one parasite at 24 hr postinfection is shown. Indicated values are means ± SD of triplicates. N.S., not significant.

(D) Mice (WT intraperitoneally injected with 1 mg of anti-IFN-γ [n = 5] or with control IgG [n = 6] and KO with anti-IFN-γ [n = 8] or with control IgG [n = 8] before 1 day of the parasite challenge) were intraperitoneally infected with 1 × 10^2 *T. gondii*, and the survival rates were monitored for 20 days.

(E) Total photon emission analysis from indicated mice (n = 4 per group) infected with 1 × 10^2 ME49 *T. gondii* luciferase-expressing parasites at days 5, 6, 7, or 8 after infection. The flux (photons/s/cm^2/sr) was determined as a measure of parasite burden.

Data are representative of two (A, B, C, E) independent experiments. Data in (D) are pooled from two independent experiments. See also Figure S4.

association of Irgb6 with Gbps is fundamental to the defective Irgb6 recruitment to *T. gondii* in *Gbp*^{chr3}-deleted cells.

Gbp^{chr3} Participate Differentially in Anti-*T. gondii* Defense

Although the Gbps share a high degree of homology (Kresse et al., 2008), variation in the amino acid sequence among Gbp^{chr3} is greater than that among Gbp^{chr5}. It is uncertain whether the five active members of *Gbp*^{chr3} (*Gbp1*, *Gbp2*, *Gbp3*, *Gbp5*, and *Gbp7*) similarly or differentially contribute to the anti-*T. gondii* cellular immunity. To address this question, we cloned each Gbp into drug-responsive retroviral expression vectors and expressed them in a tightly doxycycline-dependent manner in *Gbp*^{chr3}-deleted primary MEFs (Figure 7A). Parasite clearance in the absence or presence of IFN-γ was then tested in these transduced cells. Reintroduction of *Gbp1*, *Gbp5*, or *Gbp7* into *Gbp*^{chr3}-deleted MEFs *partially* restored the IFN-γ-dependent clearance of *T. gondii* compared with wild-type cells (Figures 3B and 7B), indicating that these proteins possess common and compensatory functions in the anti-*T. gondii* immune response. Taken together, our results show that Gbp^{chr3} play a pivotal role in IFN-γ-mediated cellular innate immune response to *T. gondii*.

DISCUSSION

This study makes use of targeted chromosome engineering to provide genetic evidence for the critical role of Gbp^{chr3} in defense against *T. gondii* infection. Mice lacking *Gbp*^{chr3} were highly susceptible to *T. gondii* infection, and *Gbp*^{chr3}-deleted macrophages are defective in IFN-γ-mediated inhibition of intracellular parasite growth.

The Gbp-mediated cellular immune mechanism was previously reported to be pleiotropic and implicated IFN-γ-inducible phagocytic oxidative killing and the trafficking of antimicrobial peptides to autophagolysosomes by their interaction with NADPH oxidase subunits and autophagy-related molecules such as Atg4b, respectively (Kim et al., 2011). In this previous

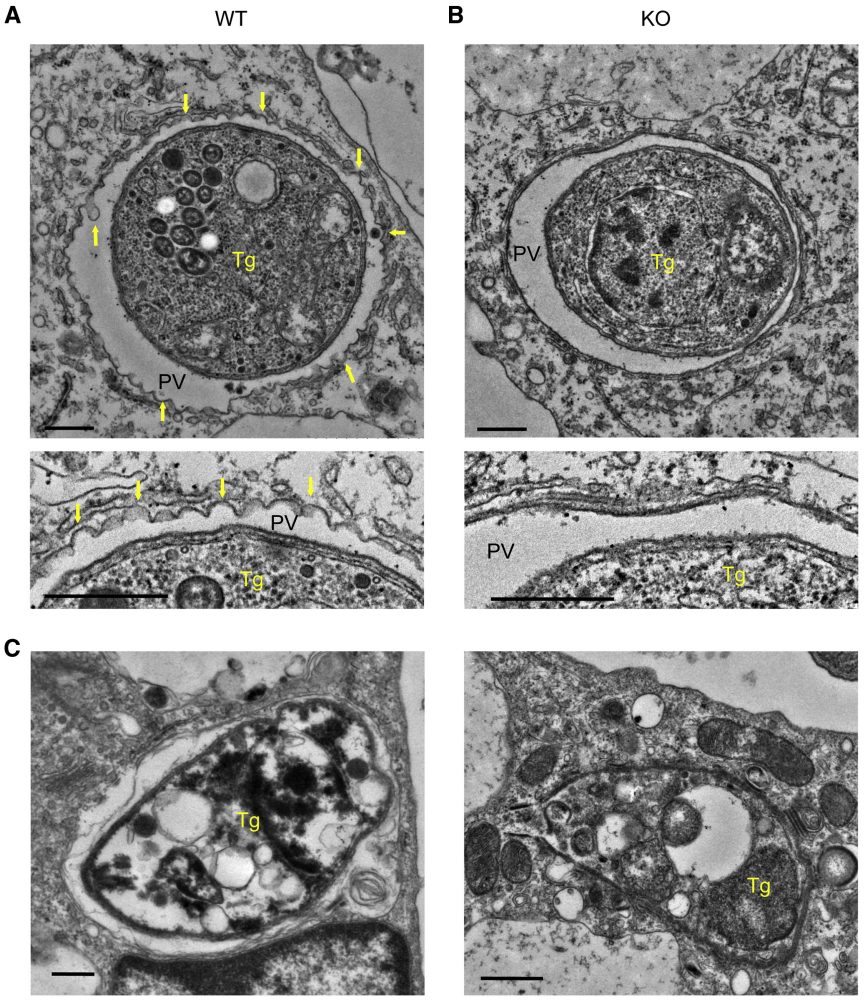


Figure 5. Electron Microscopic Analysis of IFN-γ-Activated Macrophages

(A and B) Wild-type (A) or *Gbp*^{chr3}-deleted (B) peritoneal macrophages were treated with 100 ng/ml IFN-γ for 24 hr. IFN-γ-treated cells were infected with ME49 *T. gondii* (moi = 0.5) for 6 hr and then analyzed by electron microscopy. The lower images are enlarged views from the upper images. Tg, *T. gondii*; PV, parasitophorous vacuole. Arrows indicate protrusive blebs inside or outside parasitophorous vacuoles (A).

(C) Images of dead parasites at 24 hr postinfection in IFN-γ-treated wild-type cells. Tg, *T. gondii*.

Scale bars represent 0.2 μm. Data are representative of two independent experiments.

study, Gbp function was mainly tested with dominant-negative forms of Gbps or small interfering RNA-mediated knockdown (Kim et al., 2011). Here, we observed normal O^{2-} production in response to IFN-γ stimulation and Atg4b recruitment to PVs in *Gbp*^{chr3}-deleted macrophages. Instead, we found that *Gbp*^{chr3} deficiency affected recruitment of some Irgs to *T. gondii*-infected macrophages. The discrepancy between the two studies might be explained by the different pathogens, or alternatively, it is possible that the other cluster of *Gbp*s on chromosome 5 (*Gbp*^{chr5}) may play a compensatory role in the oxidative and autophagy-related responses.

Blebbing of the PV membrane was not induced in IFN-γ-activated macrophages lacking *Gbp*^{chr3}, which is akin to that observed in Atg5-deficient cells (Zhao et al., 2008). The similar phenotypes of both mutant mice prompted us to examine the localization of Irgs during *T. gondii* infection. *Gbp*^{chr3} deficiency impaired accumulation of Irgb6 and Irgb10 but not of Irga6 around the parasite, suggesting that the Gbps play a major role in some Irg recruitment. This model contrasts with previous studies suggesting that Irgs control localization of Gbps based on the fact that IFN-γ prestimulation is required for the localization of overexpressed Gbps to *T. gondii* (Degrandi et al., 2007), and that Gbp2 localization is altered in Irgm-deficient cells (Traver et al., 2011). Considering that Gbps and Irgs participate in the anti-*T. gondii* response in close proximity, it is possible that the two families of IFN-γ-inducible GTPases mutually control their localization, at least in part, through physical association. Given that the number of IRGs and GBPs varies among species (2 IRGs and 7 GBPs in human versus 23 Irgs and 13 Gbps in mouse) (Kresse et al., 2008; Shenoy et al., 2007), evolutionary pressure to increase or decrease the number and variety of IRGs could influence the repertoire of GBPs. Furthermore, virulent *T. gondii* strains such as RH are capable of evading recognition by Irgs via the action of the virulence factor ROP18 that phosphorylates the Irgs (Degrandi et al., 2007; Fentress et al., 2010; Steinfeldt et al., 2010; Virreira Winter et al., 2011). In addition, virulent *T. gondii* also prevents the accumulation of Gbps (Virreira Winter et al., 2011). Taken together, ROP18 phosphorylation of Irgs could affect the interaction with Gbps, promoting their dissociation from the parasites. In terms of the kinetics of the recruitment of Irgb6, given that *Gbp*^{chr3} deficiency affected the later phase rather than the early stage, Gbp^{chr3} might play a role in persistence of Irgb6 on PVs in the later stage. Alternatively, other factors including Gbp^{chr5} might also participate in the Irgb6 recruitment in the early phase of infection. Whether deficiency of either *Gbp*^{chr3} or *Gbp*^{chr5} (or both) also affects the localization of other Irgs such as Irgm and Irgd deserves further investigation. Despite the fact that Irgb6 and Irgb10 are shown to be phosphorylated by ROP18 in vitro (strongly indicating the defensive roles of these Irgs) (Fentress et al., 2010), their in vivo functions in immunity to *T. gondii* should still be assessed and confirmed under physiological conditions via mice lacking these Irgs.

The impaired inhibition of *T. gondii* replication in *Gbp*^{chr3}-deleted cells prompted us to investigate the effect of NO on the *Gbp*^{chr3}-dependent resistance to *T. gondii* in vitro. We found that the presence or absence of NO did not affect parasite clearance; the rate of macrophages infected with *T. gondii* was not altered by addition of the NO inhibitor. Given the collaboration

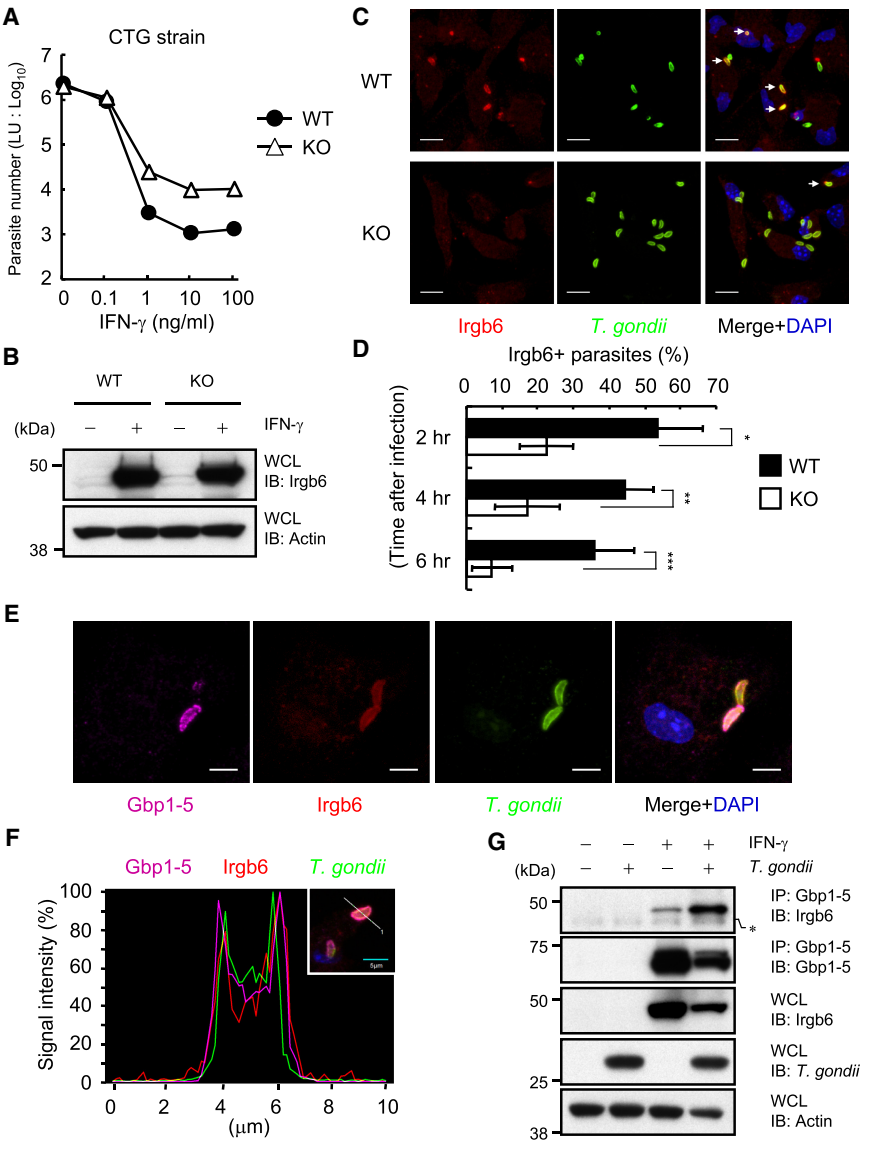


Figure 6. Defective Irgb6 Recruitment in *Gbp*^{chr3}-Deleted Macrophages

(A) Wild-type (WT) and *Gbp*^{chr3}-deleted (KO) peritoneal macrophages were untreated or treated with the indicated concentrations of IFN-γ for 24 hr. Untreated or IFN-γ-treated cells were infected with CTG *T. gondii* expressing luciferase (moi = 0.5) and harvested at 36 hr postinfection. The luciferase units (LU) were assayed with the lysates. Indicated values are means ± SD of triplicates.

(B) Peritoneal macrophages treated with 100 ng/ml IFN-γ for 24 hr were lysed. The lysates were detected by protein immunoblot with the indicated Abs.

(C) WT and KO peritoneal macrophages were treated with 100 ng/ml IFN-γ for 24 hr. IFN-γ-treated cells were infected with ME49 *T. gondii* (moi = 0.5), fixed at 6 hr postinfection, and incubated with rabbit anti-*T. gondii* (Alexa 488, green) or goat anti-Irgb6 (Alexa 594, red). Scale bars represent 10 μm. Arrows indicate colocalization of Irgb6 with *T. gondii*.

(D) The percentage of WT and KO macrophages positive for Irgb6 staining at the indicated points postinfection. Indicated values are means ± SD of triplicates. *p < 0.03, **p < 0.005, ***p < 0.004.

(E and F) Peritoneal macrophages were treated with 100 ng/ml IFN-γ for 24 hr. IFN-γ-treated cells were infected with ME49 *T. gondii* (moi = 0.5), fixed at 6 hr postinfection, and incubated with rabbit anti-*T. gondii* (Alexa 488, green), goat anti-Irgb6 (Alexa 594, red), or mouse anti-Gbp1-5 (Alexa 647, magenta). The samples were analyzed by confocal microscopy and subsequent intensity profile (F). Scale bars represent 5 μm. The signal intensities of each color on the 10 μm white line on the parasites are shown.

(G) Peritoneal macrophages treated with 100 ng/ml IFN-γ for 24 hr were lysed. The lysates were immunoprecipitated with anti-Gbp1-5 and detected by protein immunoblot with the indicated antibodies. Asterisk indicates nonspecific.

Data are representative of two (A, B) or three (C, D, E, F, G) independent experiments. Data in (D) are pooled from three independent experiments, in which almost 150 cells at each time point were counted. See also Figure S5.

of Irg and Gbp, the NO-independent role of Gbp^{chr3} may be parallel to the NO-independent effect of Irg on *T. gondii* clearance (Collazo et al., 2002). In contrast, the presence of Gbp^{chr3} had a modest effect on the inhibition of *T. gondii* replication, which critically involves NO, because *Gbp*^{chr3}-deleted macrophages contained larger numbers of the parasites per PV. Because there is no direct evidence indicating a role of Irg in the suppression of *T. gondii* replication to date, these data may be indicative of an unknown Irg-independent mechanism(s) of Gbp^{chr3} to promote the NO-mediated inhibition of the parasite replication in vitro. Nevertheless, considering that iNOS deficiency has a minor effect on early resistance in vivo (Scharton-Kersten et al., 1997), the high susceptibility in acute *T. gondii* infection in *Gbp*^{chr3}-deleted mice might be due to the defective parasite clearance dependent on some Irgs. Given the relative significance of Gbp^{chr3} in IFN-γ-dependent anti-*T. gondii* responses, the fact that anti-IFN-γ treatment in *Gbp*^{chr3}-deleted mice resulted in enhanced parasite burdens in vivo suggests that additional IFN-γ-inducible effector(s) controls early resistance to the parasite. On the other hand, *Gbp*^{chr3}-deleted mice treated with anti-IFN-γ displayed similar survival rate of mice with control IgG. The apparent disparity might be because anti-IFN-γ blocked IFN-γ-dependent anti-*T. gondii* response, leading not only to high parasite burdens but also to immune pathology mediated by massive and dysregulated doses of IFN-γ at the terminal phase of *T. gondii* infection (Nguyen et al., 2003). In addition, costimulation of TNF-α with IFN-γ resulted in incomparable inhibition of the parasite growth and abolished the Gbp^{chr3}-mediated effect. Although NO concentration in TNF-α and IFN-γ-treated macrophages was markedly higher than that in cells stimulated with IFN-γ alone, further investigation is required to determine whether only the difference of NO

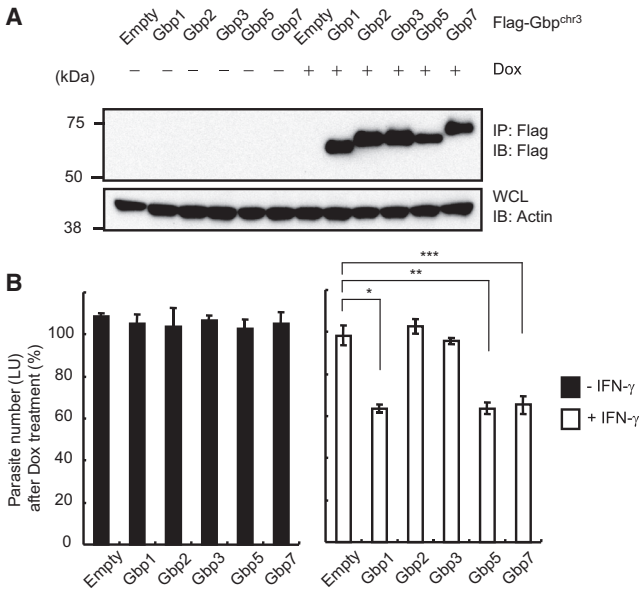


Figure 7. Individual Participation of Gbp^{chr3} in IFN-γ-Dependent Inhibition of *T. gondii* Growth

(A) Retroviral vectors encoding the indicated Flag-tagged Gbps were stably introduced in *Gbp*^{chr3}-deleted (KO) primary MEFs. The transfected MEFs were treated with or without 1 μg/ml doxycycline (Dox) for 24 hr and lysed. The lysates were immunoprecipitated with anti-Flag and detected by protein immunoblot with the indicated antibodies.

(B) The KO MEFs transfected with indicated Gbps were unstimulated or stimulated with 100 ng/ml of IFN-γ for 24 hr with or without 1 μg/ml Dox. Cells were infected with ME49 *T. gondii* parasites expressing luciferase (moi = 0.5) for 36 hr, and the luciferase units (LU) were analyzed. The percentages of the activities after each Gbp induction with Dox over those without Dox (no Gbp induction) in nonstimulated (left) or stimulated (right) cells were shown. Indicated values are means ± SD of triplicates.

Data are representative of two independent experiments. *, **, ***$p < 0.001$.

concentration or other NO-independent effects such as autophagy and augmentation of phagocytic activity accounts for the enhanced protective effect (Keller et al., 2011; Langermans et al., 1992; Leenen et al., 1994).

Gbp1, Gbp5, and Gbp7, but not Gbp2 or Gbp3, were able to restore, albeit not fully, the killing activity by IFN-γ in *Gbp*^{chr3}-deleted MEFs, indicating that Gbp^{chr3} contribute differentially to the anti-*T. gondii* host defense mechanism. This is consistent with a previous finding that knockdown of some Gbps abrogates IFN-γ-dependent suppression of bacterial growth (Kim et al., 2011). In the context of the functional redundancy within Gbp^{chr3}, we observed that mice lacking *Gbp5*, or *Gbp5* and *Gbp2ps*, were resistant to *T. gondii* infection, suggesting that *Gbp5* deficiency may be compensated for by other *Gbp*^{chr3}-encoded proteins such as Gbp1 and Gbp7. On the other hand, mice lacking Gbp1 alone are susceptible to *L. monocytogenes* and *M. bovis*. Interestingly, Gbp5 has recently been shown to play a key role in the host defense in *L. monocytogenes* infection, indicating the nonredundant function of Gbp1 and Gbp5 in the antibacterial response (Kim et al., 2011; Shenoy et al., 2012). In contrast, our *Gbp*^{chr3}-deleted mice displayed resistance to *L. monocytogenes*. This discrepancy may reflect different modes of infection (e.g., oral versus intraperitoneal infections in the previous and current studies, respectively) or utilization of different mice strains (*Gbp1* or *Gbp5* singly deficient mice in previous studies and *Gbp*^{chr3}-deleted mice in this study) (Kim et al., 2011; Shenoy et al., 2012). Moreover, we could not exclude the possibility that Gbp^{chr3} other than *Gbp1* and *Gbp5* play a negative role in Gbp1- and Gbp5-mediated anti-*Listeria* response. It remains to be seen whether deficiency of Gbp1 alone would be sufficient to disrupt the anti-*T. gondii* defense system, but it is nevertheless plausible that each member of Gbp^{chr3} may play a differential role to combat distinct types of pathogens. Given that Gbp2 associates with Gbp1 (Virreira Winter et al., 2011), Gbp2 (and Gbp3) might play a role in host defense against other pathogens or have an additional anti-*T. gondii* effect in the presence of other Gbp^{chr3}. Considering that *Gbp*^{chr3}-deleted cells lack a large genetic region, there is an inherant risk that the phenotype observed in *Gbp*^{chr3}-deleted mice could be unrelated to Irg. Although we adopted in vitro retroviral transfection for the restoration of each Gbp in this study, none of Gbp^{chr3} failed to fully restore the effect of IFN-γ. It remains unclear whether the failure is due to limit of the transfection method or due to nonredundancy among Gbp^{chr3} except for Gbp5. In vivo transgenic reconstitution of the region by an artificial chromosome such as BAC and YAC should be utilized to reveal a role of each Gbp specifically participating in antiparasite in vivo responses in the future (Copeland et al., 2001).

In conclusion, our genetic study with *Gbp*^{chr3}-deleted mice has established that Gbp^{chr3} are required for IFN-γ-mediated host defense against *T. gondii* by regulating Irgb6 recruitment to the parasite. To understand the functions of this family of IFN-γ-inducible p65 GTPases fully, future studies will have to include the second cluster, Gbp^{chr5}, and will need to investigate the effect of these proteins in defense against a broader range of parasites, as well as against other pathogens including viruses and bacteria.

EXPERIMENTAL PROCEDURES

Cells, Mice, and Parasites

C57BL/6 mice were obtained from SLC. ME49 and CTG derivatives of *T. gondii* were maintained in Vero cells by biweekly passage in RPMI1640 (Nacalai Tesque) supplemented with 2% heat-inactivated fetal calf serum (JRH Bioscience), 100 U/ml penicillin, and 0.1 mg/ml streptomycin (GIBCO). MyD88-deficient mice were kindly provided by S. Akira. Animal experiments were conducted with the approval of the Animal Research Committee of the Graduate School of Medicine and of the Research Institute for Microbial Diseases, Osaka University.

Reagents

Antibodies against *T. gondii* (sc-73210), TGTP (Irgb6; sc-11079), Atg4b (sc-130968), Gbp1 (sc-10586), Gbp2 (sc-10588), Gbp5 (sc-160356), Gbp1-5 (sc-166960), and actin (sc-8432) were purchased from Santa Cruz. Anti-CD11b (M1/70) was obtained from Becton Dickinson. Aminoguanidine hydrochloride, LPS (a TLR4 ligand) from *Salmonella minnesota* Re 595, and anti-Flag were purchased from Sigma. Anti-GAP45 rabbit and anti-MIC2 mouse antibodies were as described previously (Frénal et al., 2010). Anti-Irga6 (165/3) and Irgb10 (940/6) rabbit antibodies and anti-Irgm3 (BD Transduction Laboratories) mouse antibodies were kindly provided by J. Howard. Anti-IFN-γ (BE0055) and control Rat IgG1 (BE0088) were obtained from BioXcell. Recombinant IFN-γ and TNF-α were obtained from Peprotech.

In Vivo Measurement of Parasites by Imaging

Mice were intraperitoneally infected with 1 × 10^2 freshly egressed ME49 tachyzoites expressing luciferase resuspended in 100 μl PBS, and bioluminescence was assessed on the indicated days after infection. Treatment of anti-IFN-γ and control IgG was performed by the intraperitoneal injection 1 day before *T. gondii* infection. For the detection of bioluminescence emission, mice were intraperitoneally injected with 3 mg of D-luciferin in 200 μl PBS (Promega), maintained for 5 min to allow adequate dissemination of the luciferin, then anesthetized with isoflurane (Dainippon Sumitomo Pharma). At 10 min postinjection of D-luciferin, abdominal photon emission was assessed during a 60 s exposure by an in vivo imaging system (IVIS 100; Xenogen) and analyzed as described previously (Yamamoto et al., 2011).

Immunofluorescence

Peritoneal macrophages (1 × 10^6) infected with *T. gondii* (moi = 0.5) were fixed for 10 min in PBS containing 3.7% formaldehyde. Cells were permeabilized with PBS containing 0.1% Triton X-100 and then blocked with 8% fetal calf serum in PBS. Subsequently, cells were incubated with anti-CD11b rat antibody (1:200) and anti-GAP45 rabbit antibody (1:1,000) in Figure 3D; anti-Atg4b rabbit antibody (1:200) and anti-*T. gondii* mouse antibody (1:50) in Figure 4B; anti-GAP45 rabbit antibody (1:1,000), anti-Irgb6 goat antibody (1:50), and anti-Gbp1-5 mouse antibody (1:200) in Figure 6; and anti-Irgb10 or anti-Irga6 rabbit antibody (1:1,000) and anti-MIC2 mouse antibody (1:1,000), anti-Irgm3 mouse antibody (1:250), and anti-GAP45 rabbit antibody (1:1,000) in Figure S5, for 1 hr at 37°C, followed by incubation with donkey IgG antibodies (1:10,000): Alexa Fluor 488-conjugated anti-rabbit IgG, Alexa Fluor 594-conjugated anti-goat, or Alexa Fluor 647 or Alexa Fluor 594-conjugated anti-mouse (Molecular Probes) for 1 hr at room temperature in the dark. Finally, the immunostained cells were mounted with PermaFluor (Thermo Scientific) on glass slides and analyzed by confocal laser microscopy (FV1000-D IX-81; Olympus); the images were analyzed with Fluoview (Olympus).

Transmission Electron Microscopy

Peritoneal macrophages (1 × 10^6) untreated or treated with 100 ng/ml IFN-γ for 24 hr were infected with *T. gondii* (moi = 0.5) for 6 or 24 hr. After washing with PBS, the cells were fixed with 2.5% glutaraldehyde in 0.1 M phosphate buffer at 4°C overnight. The cells were postfixed with 1% OsO_4 in the same buffer at 4°C for 1 hr, dehydrated in a graded series of ethanol, and embedded in Quetol 812 (Nissin EM). Silver sections were cut with an ultramicrotome, stained with lead citrate and uranyl acetate, and observed with an H-7650 electron microscope (Hitachi).

Statistical Analysis

The unpaired Student's t test was used to determine the statistical significance of the experimental data.

SUPPLEMENTAL INFORMATION

Supplemental Information includes Supplemental Experimental Procedures, five figures, and one table and can be found with this article online at http://dx.doi.org/10.1016/j.immuni.2012.06.009.

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