Amplified Host Defense by Toll-Like Receptor-Mediated Downregulation of the Glucocorticoid-Induced Leucine Zipper (GILZ) in Macrophages

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Activation of toll-like receptors (TLRs) plays a pivotal role in the host defense against bacteria and results in the activation of NF-κB-mediated transcription of proinflammatory mediators. Glucocorticoid-induced leucine zipper (GILZ) is an anti-inflammatory mediator, which inhibits NF-κB activity in macrophages. Thus, we aimed to investigate the regulation and role of GILZ expression in primary human and murine macrophages upon TLR activation. Treatment with TLR agonists, e.g., Pam3CSK4 (TLR1/2) or LPS (TLR4) rapidly decreased GILZ mRNA and protein levels. In consequence, GILZ downregulation led to enhanced induction of pro-inflammatory mediators, increased phagocytic activity, and a higher capacity to kill intracellular bacteria (Salmonella enterica serovar typhimurium), as shown in GILZ knockout macrophages. Treatment with the TLR3 ligand polyinosinic: polycytidylic acid [Poly(I:C)] did not affect GILZ mRNA levels, although GILZ protein expression was decreased. This effect was paralleled by sensitization toward TLR1/2- and TLR4-agonists. A bioinformatics approach implicated more than 250 miRNAs as potential GILZ regulators. Microarray analysis revealed that the expression of several potentially GILZ-targeting miRNAs was increased after Poly(I:C) treatment in primary human macrophages. We tested the ability of 11 of these miRNAs to target GILZ by luciferase reporter gene assays. Within this small set, four miRNAs (hsa-miR-34b*,−222,−320d,−484) were confirmed as GILZ regulators, suggesting that GILZ downregulation upon TLR3 activation is a consequence of the synergistic actions of multiple miRNAs. In summary, our data show that GILZ downregulation promotes macrophage activation. GILZ downregulation occurs both via MyD88-dependent and -independent mechanisms and can involve decreased mRNA or protein stability and an attenuated translation.

Keywords: inflammation, MyD88, TRIF, NF-κB, cytokine, nitric oxide, phagocytosis, microRNA
INTRODUCTION

Macrophages are strategically distributed all over the body and represent the first line of defense to invading pathogens. In response to bacterial or viral infections, these phagocytic cells engulf and destroy the infectious agent. Macrophages recognize a variety of microbial products, such as bacterial cell wall components and nucleic acids, via pathogen recognition receptors (PRRs). These receptors include Toll-like receptors (TLRs), as well as cytosolic NOD-like receptors, RIG-I-like receptors, and DNA sensors (1, 2).

Once a pathogen is recognized and phagocytosed, macrophages generate a wide range of biologically active molecules, e.g., pro-inflammatory cytokines, chemokines, and growth factors. Cytokines released by macrophages, such as interferons (IFNs), tumor necrosis factor (TNF)-α, and interleukin (IL)-10/β, stimulate the activity of both the innate and adaptive immune response. Pathogen uptake usually also results in antimicrobial responses such as reactive oxygen species (ROS) and nitric oxide (NO) production, thus enabling a rapid response to infection (3, 4).

Activated macrophages can be deactivated by various mediators, including IL-10, prostaglandins, G-protein coupled receptor ligands, or glucocorticoids (GCs) (3, 4). Anti-inflammatory factors such as IL-10 are also released by macrophages themselves, and failure to produce these mediators can lead to non-resolving inflammation (3, 4).

The Glucocorticoid-induced leucine zipper (GILZ) is an endogenous inhibitor of immune responses. GILZ exerts its anti-inflammatory activity mainly by inhibition of the pro-inflammatory transcription factors nuclear factor (NF)-κB and activator protein (AP)-1 (5–8). In addition, GILZ has been implicated in the negative regulation of mitogen-activated protein kinase (MAPK) signaling (9–11).

Early studies on GILZ focused on its effects on thymocytes and T lymphocytes because of its strong induction by GCs in the thymus (12). Several reports indicated that the pro-apoptotic effects of GILZ overexpression in T lymphocytes mimic GC treatment (5, 8, 12–14). Moreover, GILZ depletion attenuates GC-induced apoptosis in B lymphocytes (15, 16).

In monocytes and macrophages, GILZ can be upregulated by various anti-inflammatory factors, such as GCs, IL-4 or IL-10, and the natural compound curcumin (6, 17–20). In accordance with the GC-like actions of GILZ in lymphocytes, GILZ overexpression in macrophage-like THP-1 cells results in decreased expression of chemokines and macrophage activation markers, as well as NF-κB activity upon treatment with the TLR4 ligand lipopolysaccharide (LPS) (6).

We previously reported that primary human in vitro differentiated and ex vivo pulmonary macrophages express high constitutive levels of GILZ (11, 21). Both siRNA-mediated GILZ knockdown in human macrophages and GILZ knockout in murine bone marrow-derived macrophages (BMMs) increased the responsiveness toward LPS, suggesting that repression of endogenous GILZ expression represents a positive feedback loop in macrophage activation.

Little is known about the role and regulation of GILZ after stimulation with other TLR ligands, e.g., activators of TLR1/2 or TLR3. The extracellular TLR1/2 heterodimer recognizes bacterial triacylated lipopeptides and their mimic, the synthetic compound Pam3CSK4. In contrast, intracellular TLR3 detects double-stranded RNA, i.e., an intermediate in viral replication, as well as its synthetic analog polynosinic:polycytidylic acid [Poly(I:C)]. TLRs differentially activate transcription factors due to the varying involvement of the adapter molecules MyD88 (myeloid differentiation primary response gene 88) and TRIF (TIR domain-containing adapter inducing IFN-β). All TLRs except TLR3 can initiate MyD88-dependent signaling, and MyD88-independent signaling via TLR3 or TLR4 utilizes TRIF for signal transduction (1, 2).

In the present study, we provide evidence for dual regulation of GILZ upon MyD88- and TRIF-dependent TLR activation and link GILZ expression levels with pivotal macrophage defense mechanisms, such as phagocytosis and bac terial activity.

MATERIALS AND METHODS

Materials

Cell media (RPMI1640, #R0883; DMEM, #D6546), fetal calf serum (FCS, #F7524), penicillin/streptomycin (#P433), and glutamine (#G7513) were from Sigma-Aldrich. Anti-GILZ antibodies were obtained from either Santa Cruz Biotechnology (polyclonal goat anti-GILZ Ab, #sc-26518) or eBioscience (CFLKG15, #14-4033-82). The anti-tubulin antibody (#T9026) was obtained from Sigma-Aldrich. Anti-rabbit IRDye 680- and anti-mouse IRDye 800-conjugated secondary antibodies were from LI-COR Biosciences (#926-68071, #926-32210). The anti-rabbit antibody (#926-132-120). Anti-p44/42 (ERK1/2) mouse antibody (L34F12, #4696S) and anti-phospho-p44/42 MAPK (Thr202/Tyr204) rabbit mAbs (20G11, #4376S) were obtained from Cell Signaling Technology. TLR ligands, i.e., ultrapure LPS from Escherichia coli K12 (#tlrl-pelk), Pam3CSK4 (#tlrl-pms), lipoteichoic acid (LTA, #tlrl-plea), and Poly(I:C) (#tlrl-plt) were purchased from InvivoGen. Phorbol 12-myristate 13-acetate (PMA, #524400) and BAY 11-7082 (Cay10010266-10) were from Cayman Chemical. BAY 11-7085 (#196872) was
obtained from Calbiochem. Human M-CSF (#M6518), MTT (#M5655), actinomycin D (#A9415), and aminurinicularboxylic acid (ATA, #A1895) were obtained from Sigma-Aldrich. Murine GM-CSF (#130-095-735), M-CSF (#130-101-704), IFN-γ (#130-105-782), IL-4 (#130-097-761), and TNF-α (#130–101–689) were obtained from Miltenyi Biotec. Primers and dual-labeled probes were from Eurofins MWG Operon. Taq polymerase (5 U/µL, #E00007), Taq buffer (#B0005) and the dNTP mix (#D0056) were from Genscript. Restriction enzymes (BamH1, #R3136S; EcoR1, #R0101S; Sac1, #R0156S; Spe1, #R0133L) were purchased from New England Biolabs. Other chemicals were obtained from either Sigma-Aldrich or Carl Roth unless stated otherwise.

**Mice**

Mice were housed in a 12:12 h light-dark cycle with food and water ad libitum. Mice expressing Cre recombinase under the control of endogenous Lys2 (lysozyme 2) promoter/enhancer elements (LysMcre mice, The Jackson Laboratory, #B6.129P2-Lys2tm1(cre)Ifo1) were crossed with C57BL/6J mice bearing LoxP sites upstream and downstream of Gilt exon 6 (11, 22) to obtain myeloid-specific GILZ KO mice. Genotyping was performed according to protocols provided by The Jackson Laboratory and as described by Bruscoli et al. (22).

**Cell Culture**

**Cell Lines**

THP-1 (#TIB202), U937 (#CRL-1593.2), and L929 cells (#CRL-6364) were obtained from ATCC and grown in standard medium (RPMI 1640, 10% FCS, 100 U/mL penicillin G, 100 µg/mL streptomycin, 2 mM glutamine). THP-1 and U937 were differentiated into macrophage-like cells by treatment with PMA (100 nM) for 48 h. HEK293T cells (ATCC, #CRL-3216) were cultured in high glucose DMEM medium with supplements (10% FCS, 100 U/mL penicillin G, 100 µg/mL streptomycin, 2 mM glutamine). HEK-BlueTM Null1 cells (Invivogen, #hkb-null1) were grown in high glucose DMEM medium supplemented with 10% FCS, 2 mM glutamine, 50 U/mL penicillin G, 50 µg/mL streptomycin, 100 µg/mL Normocin (Invivogen, #ant-nr-1), and 100 µg/mL Zeocin (Invivogen, #ant-zn-1).

**Human Alveolar Macrophages (AMs)**

AMs were isolated from human non-tumor lung tissue obtained from patients undergoing lung resection. The use of human material for isolation of primary cells was reviewed and approved by the local Ethics Committees (permission no. 213/06; State Medical Board of Registration, Saarland, Germany). The informed consent of all participating subjects was obtained. Isolation was performed according to a previously described method (23) with minor modifications. After visible bronchi were removed, the lung tissue was cut into pieces (~1 cm³) and washed at least three times with RS (balanced salt solution; 137 mM NaCl, 5 mM KCl, 0.7 mM Na₂HPO₄, 10 mM HEPES, 5.5 mM glucose, pH 7.4). The washing buffer was collected and centrifuged (15 min, 350 x g). Erythrocytes were lysed by incubation with hypotonic buffer (155 mM NH₄Cl, 10 mM KHCO₃, 1 mM Na₂EDTA) for 2–5 min at 37°C, and the cell suspension was washed with PBS (137 mM NaCl, 2.7 mM KCl, 10.1 mM Na₂HPO₄, 1.8 mM KH₂PO₄, pH 7.4). Cells were resuspended in AM medium (RPMI 1640, 5% FCS, 100 U/mL penicillin G, 100 µg/mL streptomycin, 2 mM glutamine), seeded at a density of 0.5–1 x 10⁶ cells/well into a 12- or 6-well plate and incubated at 37°C for 2 h. Adherent cells were washed at least 5 times with PBS, and cells were cultured in AM medium overnight before further use. AM preparations were 95% pure as assessed by flow-cytometric analysis of intracellular CD68 expression (23, 24).

**Human Monocyte-Derived Macrophages**

Monocytes were isolated from healthy adult blood donors (Blood Donation Center, Saarbrücken, Germany) and differentiated with M-CSF as described in Dembek et al. (25). The use of the human material was reviewed and approved by the local Ethics Committees (permission no. 130/08; State Medical Board of Registration, Saarland, Germany). Peripheral blood mononuclear cells (PBMCs) were isolated by density gradient centrifugation using Lymphocyte Separation Medium 1077 (Promocell, #C-44010) and Leucosep tubes (Greiner Bio-One, #227290) as recommended by the suppliers. After washing with PBS, monocytes were purified from PBMCs by magnetic cell sorting using anti-CD14 microbeads (Miltenyi, #130-050-201) according to the manufacturer’s instructions, except that 10% of the recomended bead amount was used (25). Monocyte purity was > 95% as indicated by CD14 expression (data not shown). Monocytes were seeded into 12 well plates (5 x 10⁶ cells/well) and differentiated into macrophages in RPMI 1640 supplemented with 10% FCS, 100 U/mL penicillin G, 100 µg/mL streptomycin, 2 mM glutamine, and 20 mg/mL M-CSF at 37°C and 5% CO₂ for 7 d. The medium was changed every other day.

**Murine Bone Marrow-Derived Macrophages (BMMs)**

BMMs were obtained from 8 to 12-week-old male wild-type (WT, LysMCre+/floxed GILZ−/−) or GILZ knockout (KO, LysMCre+/floxed GILZ+/+) mice as described previously with minor modifications (11). Femurs and tibias were flushed with standard medium (RPMI 1640, 10% FCS, 100 U/mL penicillin G, 100 µg/mL streptomycin, 2 mM glutamine). After centrifugation (10 min, 250 x g), erythrocytes were lysed by incubation in hypotonic buffer (155 mM NH₄Cl, 10 mM KHCO₃, 1 mM Na₂EDTA) for 2 min at 37°C. Cells were washed with PBS, resuspended in standard medium containing M-CSF (50 ng/mL, 30 mL per preparation), transferred into a 75 cm² culture flask and allowed to adhere overnight. Non-adherent cells were collected and cultured in a 150 cm² culture flask for another 5 d in M-CSF-containing medium. Differentiated cells were detached with accutase (Sigma-Aldrich, #A6964), suspended in standard medium supplemented with 10 ng/mL M-CSF, and seeded into 96-well plates (7.5 x 10⁴ cells/well) for TNF measurements or into 12-well plates (5 x 10⁵ cells per well) for all other applications unless indicated otherwise. BMMs were > 95% pure as indicated by flow cytometric analysis using an antibody against F4/80 (data not shown).
**MPI Cells**

MPI cells, i.e., non-transformed self-renewing primary murine macrophages, were obtained from WT or GILZ KO mice based on a previously described method (26). MPI cells were prepared from fetal livers of 15-d-old mouse embryos by pushing the tissue through 70 µm cell strainers while washing with PBS. Cells were washed once with PBS and then resuspended in standard medium (RPMI 1640, 10% FCS, 100 U/mL penicillin G, 100 µg/mL streptomycin, 2 mM glutamine) supplemented with 30 ng/mL GM-CSF. Proliferating cells were subcultured by splitting them 1:5 after 6–8 d. Cell preparations were > 95% positive for macrophage markers F4/80 and CD68, as indicated by flow cytometric analysis (data not shown). Cells were seeded into 96-well plates (4 × 10⁴ cells/well) for determination of NO release or into 12 well plates (2 × 10⁵ cells/well) for phagocytosis assays.

**Determination of Cell Viability**

The MTT colorimetric assay was performed to ensure that non-toxic concentrations of U0126, BAY-11-7082, BAY-11-7085, and ATA were used. Cells were incubated for 24 h with the test compounds diluted in a cell-type specific medium. After that, cells were incubated with MTT solution (5 mg/mL in medium) for 2 h. The supernatant was discarded, and cells were lysed in 100 µL DMSO. Absorbance measurements were carried out at 550 nm with 630 nm as the reference wavelength using a microplate reader (Tecan Sunrise). The cell viability obtained from at least two independent experiments was calculated relative to untreated and solvent controls (data not shown).

**Cytokine Quantification**

Cytokines were quantified in AM supernatants using a Milliplex MAP Human Cytokine Kit (Millipore, #HCYTOMAG-60K) as detailed in Hoppstädter et al. (23). AMs were kept at a density of 1 × 10⁵ cells per well in 96 well plates in a total volume of 100 µL medium in the presence or absence of Pam3CSK4 (100 ng/mL) or Poly(I:C) (10 µg/mL) for 6 h. The supernatants were collected and stored at −80°C until they were used in the multiplex cytokine assay. The immunoassay procedure was performed using a serial dilution of the 10,000 pg/mL human cytokine standard according to the manufacturer’s instructions, and the plate was read at the Luminex 200 System (Millipore).

Murine TNF-α was quantified by bioassay as previously described (1). L929 cells were seeded at a density of 3 × 10⁴ cells per well into a 96-well plate. After 24 h, the medium was replaced by 100 µL of actinomycin D solution (1 µg/mL in standard medium) and cells were incubated for 1 h at 37°C. Subsequently, BMM supernatants (100 µL per well) were added. Dilution series of recombinant murine TNF-α (100–2,500 pg/mL) were run alongside the samples to generate a standard curve. The plates were incubated for an additional 24 h at 37°C. The MTT assay (see Determination of Cell Viability) was used to assess cell viability.

**RNA Isolation, Reverse Transcription, and Quantitative RT-PCR**

Total RNA was isolated using the RNeasy Plus Mini Kit (Qiagen, #74134), and 200 ng RNA were reverse transcribed using the High-Capacity cDNA Reverse Transcription Kit (Applied Biosystems, #4368813) according to the manufacturer’s recommendations. The cDNA was diluted with 80 µL TE buffer (Applichem, #A0368) before use. The CFX36 Touch™ Real-Time PCR Detection System (Bio-Rad) was used for real-time RT-PCR. For TNF, IL6, CXCL8, GILZ, and ACTB, one 25 µL reaction mix contained 2.5 U Taq polymerase, 500 nM sense and antisense primers, 60–100 nM probe, 200 µM dNTPs, 3–4 mM MgCl2, 2.5 µL Taq buffer, 3 µL Template, and molecular biology grade water (Applichem, #A7398). The reaction conditions were 95°C for 5 min followed by 40 cycles of 15 s at 95°C, 15 s at a reaction dependent temperature varying from 57 to 60°C, and 15 s at 72°C. For ZFP36 detection, the 5x HOT FIREPol® EvaGreen® qPCR Mix Plus (Solis Biodyne, #08-25) was used according to the manufacturer’s recommendations. Primer and probe sequences as well as specific reaction conditions are given in Supplementary Table 1.

**Western Blotting**

Cells were lysed in lysis buffer [50 mM Tris-HCl, 1% (m/v) SDS, 10% (v/v) glycerol, 5% (v/v) 2-mercaptoethanol, 0.004% (m/v) bromphenol blue] supplemented with a protease inhibitor mix (cOmplete; Roche Diagnostics, #04693124001) and stored at −80°C until further use. After sonication, lysates were boiled for 5 min at 95°C. Proteins were separated by SDS-PAGE (12–15% gels using the Mini-Protean Tetra Cell system (BioRad) and transferred onto Immobilon FL-PVDF membranes (Millipore, #IPFL00010) using the Tetra Blotting Module (BioRad). The membranes were blocked in blocking buffer for near-infrared Western Blotting (Rockland, MB070) for 1 h, incubated with primary antibody dilutions (1:500–1:2,000 in Rockland blocking buffer) for 3 h at room temperature or at 4°C overnight and with IRDye 680 or IRDye 800 conjugated secondary antibodies (1:5,000–1:10,000) for 1.5 h at room temperature. Blots were scanned with an Odyssey Infrared Imaging System (LI-COR Bioscience), and relative protein amounts were determined using either Odyssey or ImageJ software.

**Griess Assay**

MPI cells were cultured in 96-well plates (4 × 10⁴ cells per well) and treated as indicated. After 20 h, the concentration of nitrite was measured in the supernatants by Griess assay as previously described (20). In brief, 90 µL 1% sulfanilamide in 5% H₃PO₄ and 90 µL 0.1% N-(1-naphthyl)ethylenediamine dihydrochloride in H₂O were added to 100 µL of the cell culture supernatant, followed by absorbance measurement at 550 nm and the reference wavelength 630 nm using a Tecan Sunrise microplate reader. A standard curve of sodium nitrite dissolved in the medium was run along with the samples. Total cellular protein concentrations used for data normalization.
were determined by Pierce BCA protein assay (ThermoFisher Scientific, #23225) according to the manufacturer's instructions.

**Phagocytosis Assay**

Particle uptake was quantified after incubation of macrophages with 1.75 μm latex beads (Fluoresbrite Carboxylated YG microspheres; Polysciences, #17687) at a 50:1 bead/cell ratio in full medium. Subsequently, cells were washed 5 times with ice-cold PBS to remove remaining fluorophores and detached from plates using ice-cold TEN buffer (40 mM Tris, 1 mM EDTA, 150 mM NaCl). Cells were resuspended in ice-cold PBS (137 mM NaCl, 2.7 mM KCl, 10.1 mM Na₂HPO₄, 1.8 mM KH₂PO₄, pH 7.4) and particle-associated fluorescence was determined using an LSR Fortessa flow cytometer (BD Biosciences). Particle uptake was verified by quenching the fluorescence of particles that were merely attached with trypan blue solution (1 mg/mL in PBS, data not shown).

**Salmonella Infection and Determination of Colony Forming Units (CFU)**

The *Salmonella* enterica serovar typhimurium (*S. typhimurium*) wild-type strain 12023 (resp. 14028s) was a gift from D. Monack (Stanford University). *S. typhimurium* was grown to stationary phase overnight in LB medium (Carl Roth, #X968.1) at 37 °C and 20 rpm. After several washing steps, the microarrays were inserted into the respective cloning site of pMIR-REPORT (Ambion, #AM5795). Plasmid DNA was purified from overnight cultures with the Genopure Plasmid Midi Kit (Roche, #3143414001). Reporter gene assays were performed by transfecting HEK293T cells with the pMIR-REPORT reporter construct, a pre-miRNA vector, and phRG-TK Renilla (Promega, #E6291). HEK 293T cells were seeded at a density of 2 × 10⁴ cells per well into a 96-well-plate, grown for 24 h and transfected using PolyFect (Qiagen, #301105) according to the manufacturer's recommendations. Twenty-four hours after transfection, cells were lysed with 5x Passive Lysis Buffer (Promega, #E1941). Luciferase activity was determined after the addition of firefly luciferase substrate (470 μM D-luciferin, 530 μM ATP, 270 μM coenzyme A, 33 mM DTT, 20 mM Tricine, 2.67 mM MgSO₄, and 0.1 mM EDTA, pH 7.8) or renilla substrate (0.1 M NaCl, 25 mM Tris-HCl, pH 7.5, 1 mM CaCl₂, 1 μM coelenterazine) by luminescence measurement using a Glomax Discover multiplate reader (Promega). Firefly luciferase activity was normalized to renilla luciferase activity to correct for variations in transfection efficiency, and relative luminescence was calculated relative to P5G5 control transfected cells. Data were also normalized to the impact of each miRNA on the luciferase mRNA without GILZ 3′UTR to adjust for GILZ 3′UTR-specific effects.

**miRNA Detection**

Total RNA for miRNA detection was isolated from AMs using the miRNEasy Kit (Qiagen, #217004). The RNA quality was assessed using the Agilent Bioanalyzer 2100 and RNA 6000 Pico Chips (Agilent Technologies, #5067-1513) as recommended by the supplier. For miRNA expression analysis, the Agilent microarray platform and Human miRNA 8x60K microarrays (Release 16.0, Agilent Technologies, #G4870A) were used according to manufacturer's instructions. In brief, 100 ng total RNA and the miRNA Complete Labeling and Hyb Kit (Agilent Technologies, #5190-0456) were used to generate fluorescently labeled miRNA. The microarrays were loaded and incubated for 20h at 55°C and 20 rpm. After several washing steps, the microarrays were scanned with the Agilent Microarray Scanner at 3 microns in double path mode. The Total Gene Signal provided by the Agilent Feature Extraction software was used for quantitative data analysis. Normalization between arrays was carried out using quantile normalization. The data have been deposited in NCBI’s Gene Expression Omnibus (27) and are accessible through GEO Series accession number GSE123756.

**Luciferase Reporter Gene Assay**

miRNA overexpression vectors were generated by cloning the miRNA precursor sequences into the BamH1/EcoR1 sites of pSG5 (Agilent Technologies, #216201) using the primers given in Supplementary Table 2 or as previously described (28–30). GILZ 3′UTR-luciferase reporter constructs were generated by coupling the GILZ 3′UTR sequence to a firefly luciferase reporter gene (21). Human GILZ 3′UTR cDNA was amplified using the Expand High fidelity PCR System (Sigma-Aldrich, #11732641001) and the following primers: 5′-GCC TAC TAG TGC AGA GCC ACT AAA CTT G-3′ and 5′-AAT AGA GCT CAC TCT CAC AAA ACC CGC TAC-3′. The SacI/SpeI digested PCR product was inserted into the respective cloning site of pMIR-REPORT (Ambion, #AM5795). Plasmid DNA was purified from overnight cultures with the Genopure Plasmid Midi Kit (Roche, #3143414001). Reporter gene assays were performed by transfecting HEK293T cells with the pMIR-REPORT reporter construct, a pre-miRNA vector, and phRG-TK Renilla (Promega, #E6291). HEK 293T cells were seeded at a density of 2 × 10⁴ cells per well into a 96-well-plate, grown for 24 h and transfected using PolyFect (Qiagen, #301105) according to the manufacturer's recommendations. Twenty-four hours after transfection, cells were lysed with 5x Passive Lysis Buffer (Promega, #E1941). Luciferase activity was determined after the addition of firefly luciferase substrate (470 μM D-luciferin, 530 μM ATP, 270 μM coenzyme A, 33 mM DTT, 20 mM Tricine, 2.67 mM MgSO₄, and 0.1 mM EDTA, pH 7.8) or renilla substrate (0.1 M NaCl, 25 mM Tris-HCl, pH 7.5, 1 mM CaCl₂, 1 μM coelenterazine) by luminescence measurement using a Glomax Discover multiplate reader (Promega). Firefly luciferase activity was normalized to renilla luciferase activity to correct for variations in transfection efficiency, and relative luminescence was calculated relative to P5G5 control transfected cells. Data were also normalized to the impact of each miRNA on the luciferase mRNA without GILZ 3′UTR to adjust for GILZ 3′UTR-specific effects.

**miRNA Mimic Transfection**

miRNA mimics (hsa-miR-34b, −222, −320d, −484, and scrambled control; MISSION miRNA, Sigma-Aldrich, #HMI0510, #HMI0400, #HMI0475, #HMI0593, #HMC0002) were reverse transfected into HEK293T cells (2.5 × 10⁵ cells in a 6-well plate) using the Lipofectamine RNAiMAX transfection reagent (ThermoFisher Scientific, #13778075) as recommended by the supplier. Mimics were used at a final concentration of 50 nM. To examine potential synergistic effects, a combination of all four miRNA mimics (12.5 nM each) was compared with scrambled controls (50 nM) or miR-34b mimic (12.5 nM) only. The latter was co-transfected with scrambled control mimics (37.5 nM) to ensure comparable transfection conditions.

**NF-κB/AP-1 Reporter Gene Assay**

HEK-Blue cells (HEK-Blue Blue™ Null, Invivogen, #hkb-null1) expressing secreted embryonic alkaline phosphatase (SEAP) under the control of the IFN-β minimal promoter fused to five NF-κB and AP-1 binding sites were used to determine NF-κB/AP-1 activity. Cells were seeded into 96-well plates (5 × 10⁵ cells/well) and treated immediately as indicated. After 24 h,
supernatants were collected, and SEAP activity was determined using the QuantiBlue reagent (Invivogen, #REP-QB2) according to the supplier’s instructions. For miRNA mimic experiments, HEK-Blue cells (2 x 10^5 cells/well in 96-well plates) were reverse transfected with single miRNA mimics (50 nM), scrambled control (50 nM), or a mix of miR-34b*,−222,−320d, and−484 mimics (12.5 nM, each) using the Lipofectamine RNAiMAX transfection reagent as recommended by the supplier (see miRNA Mimic Transfection for details). Twenty-four hours later, cells were treated as indicated and incubated for another 24 h. Subsequently, SEAP activity was determined as described above.

**TAT-Fusion Proteins**

The cell-permeable trans-activator of transcription peptide (TAT)-GILZ fusion protein and the respective control protein (TAT-Co) were generated and purified as previously (13, 31). In brief, TAT and TAT-GILZ sequences were inserted into the pGEX-4T2 plasmid (GE Healthcare, #28-9545-50) to produce an in-frame fusion protein. The GST fusion protein expression was induced in *E. coli* BL21 (GE Healthcare, #27-1542-01) with 0.1 mM isopropyl β-d-thiogalactopyranoside (Sigma-Aldrich, #I5502). After lysis by sonication, proteins were purified with glutathione-sepharose 4B beads (GE Healthcare, #17-0756-01) according to the manufacturer’s instructions. Protein purity was evaluated by SDS-PAGE and Coomassie blue staining.

**Statistics**

All experiments were performed at least three times, and at least two biological replicates were analyzed for all in vitro experiments unless stated otherwise. Data distribution was determined by the Shapiro-Wilk test. For normally distributed data, means of two groups were compared with non-paired *t*-test. For data that were not normally distributed, means of two groups were compared using the Mann-Whitney test. Means of more than two groups were compared by one-way ANOVA with Bonferroni’s post hoc test (normal distribution) or Kruskal-Wallis ANOVA followed by Mann-Whitney test (no normal distribution). Statistical significance was set at *p* < 0.05, *p* < 0.01, or *p* < 0.001. Data analysis was performed using Origin software (OriginPro 8.6G; OriginLab).

**RESULTS**

**Downregulation of GILZ by TLR Agonists**

Because most studies on GILZ focused on its pharmacological induction, there are only limited data on the regulation of endogenous GILZ during the immune response. To investigate macrophage responses toward TLR1/2 and TLR3 ligands, we treated human AMs, human monocyte-derived macrophages, as well as non-differentiated and PMA-differentiated U937 and THP-1 cells with Pam3CSK4 and Poly(I:C). Out of these human macrophages and macrophage-like cells, AMs were the only cell type that showed a robust activation by Poly(I:C) regarding cytokine production (Figure 1A, Supplementary Figure 1). Thus, we used these cells to further examine GILZ expression. As reported previously (21), GILZ mRNA levels were highly reduced upon TLR1/2 and TLR4, but not TLR3 stimulation (Figure 1B). In contrast, GILZ protein levels decreased both after MyD88-dependent stimulation by Pam3CSK4 and TRIF-dependent activation by Poly(I:C) (Figures 1C–E), implicating a dual regulation of GILZ.

During the late stages of sepsis, macrophages adjust to the prolonged stimulation by LPS or other TLR ligands and enter a hyporesponsive state termed endotoxin or LPS tolerance. LPS-tolerant macrophages are characterized by reduced secretion of pro-inflammatory cytokines and upregulation of anti-inflammatory genes. We previously showed that GILZ represents a key mediator of LPS tolerance (11). In contrast, pretreatment of AMs with low-dose Poly(I:C) for 24 h resulted in reduced GILZ levels (Figures 2A,B) and an enhanced LPS- or Pam3CSK4-induced pro-inflammatory response, as shown by qPCR analysis of TNF, IL6, and CXCL8 expression (Figures 2C,D). This observation is in line with a recent report showing that macrophages generate a type of innate immune memory by activating the TLR3 pathway, thereby enhancing the immune response to a subsequent PAMP exposure (32). Our data suggest that this might, at least in part, be due to GILZ downregulation.

**Functional Implications of Gils Downregulation**

We then sought to characterize the functional implications of GILZ downregulation. To examine whether lack of GILZ modulates macrophage activation after Pam3CSK4 and Poly(I:C), we treated murine WT and GILZ KO macrophages with increasing dosages of both stimuli, and TNF-α secretion was quantified. Compared with their WT counterparts, GILZ KO cells released higher levels of TNF-α after stimulation with either TLR ligand (Figures 3A,B). Loss of GILZ has been shown to enhance LPS-induced ERK-signaling in macrophages and thereby promote inflammatory actions, including TNF-α production (11). Therefore, we investigated Pam3CSK4- and Poly(I:C)-mediated ERK phosphorylation. We detected ERK activation after treatment with both TLR ligands, and GILZ KO BMMs showed higher levels of pERK after 15 min for Pam3CSK4 and after 30 min for Poly(I:C) when compared with equally treated WT cells (Figures 3C–F).

Since ERK activity might play a role in increased cytokine expression concomitant with GILZ KO, we blocked ERK activation with the inhibitor U0126 and quantified TNF-α after Pam3CSK4 or Poly(I:C) treatment. Inhibition of ERK signaling abolished the increased responsiveness of GILZ KO cells toward both ligands, indicating that loss of GILZ enhances the inflammatory response by modulating ERK activity (Figures 3G,H).

Apart from the production of inflammatory cytokines, macrophage-dependent host defense involves the release of nitric oxide (NO) and phagocytosis of the invading pathogen. Thus, we examined the influence of GILZ expression on these parameters. NO production was elevated in GILZ KO cells when compared with equally treated WT cells upon TLR4 activation by LPS alone or in combination with IFN-γ (Figure 4A). TLR2 ligands, i.e., Pam3CSK4 or lipoteichoic acid (LTA), induced NO only when...
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FIGURE 1 | GILZ is downregulated in AMs upon MyD88-dependent and MyD88-independent TLR activation. (A) Responsiveness of AMs toward the TLR1/2 ligand Pam3CSK4 (Pam, 100 ng/mL, 6 h) and the TLR3 ligand Poly(I:C) (PIC, 10 µg/mL, 6 h). Cytokine production was determined by Luminex bead assay (n = 4, triplicates; #p < 0.05, ##p < 0.01, ###p < 0.001 compared with Co; *p < 0.05, **p < 0.01, ***p < 0.001 compared with Pam-treated AMs). (B) AMs were left untreated or treated with LPS (1 µg/mL), Pam (1 µg/mL) or PIC (10 µg/mL) for 2 h and GILZ mRNA was quantified by qRT-PCR using ACTB as a housekeeping gene (n = 4, duplicates; ***p < 0.001 compared with Co). (C–E) AMs were treated with Pam (C,E; 1 µg/mL) or PIC (D,E; 10 µg/mL) for the indicated time points and GILZ levels were determined by Western blot. Tubulin served as a loading control. (C,D) Representative blots. (E) GILZ signal intensities were quantified and normalized to tubulin values [n = 3; *, #p < 0.05 compared with 0 h for Pam (*) and PIC (#)-treated cells].

combined with IFN-γ. As seen for LPS, the response was higher in GILZ KO macrophages (Figure 4B).

To assess their phagocytic capacity, WT and GILZ KO macrophages were incubated with fluorescent latex particles. The total percentage of cells with particle-associated fluorescence and the percentage of cells that engulfed 0, 1, 2, or more than 2 particles was determined by flow cytometry after 15 and 30 min (Figures 4C–E). GILZ KO macrophages phagocyted more particles than WT cells, which was especially evident in the fraction that took up more than two particles.

To determine the influence of GILZ expression on the phagocytic activity of differentially polarized macrophages, WT, and GILZ KO macrophages were either left untreated or pretreated with LPS/IFN-γ, IL-4, or dexamethasone for 20 h. Subsequently, latex beads were added, and uptake was quantified by flow cytometry. Phagocytosis was enhanced by LPS/IFN-γ and impaired by dexamethasone treatment. Regardless of the treatment scheme, GILZ KO macrophages had a higher uptake efficiency than equally treated WT cells (Figures 4F,G and Supplementary Figure 2).

Next, we investigated whether the loss of GILZ modulated the bactericidal capacity. WT and GILZ KO BMMs were exposed to Salmonella typhimurium, and extracellular bacteria were removed after 30 min. After 8 h the number of viable bacteria within the cells was determined. GILZ KO macrophages were more efficient regarding the killing of intracellular bacteria, as indicated by lower CFU counts when compared with WT macrophages (Figure 4H).

Mechanisms of GILZ Downregulation

We then aimed to address the mechanisms underlying GILZ downregulation in activated macrophages. Both the MyD88- and TRIF-dependent pathways can lead to the activation of the pro-inflammatory transcription factor NF-κB. Thus, we blocked NF-κB signaling in AMs by pretreatment with the inhibitors BAY-11-7082 or BAY-11-7085, followed by treatment with the TLR ligands LPS, Pam3CSK4, and Poly(I:C), and found that GILZ downregulation was abrogated in inhibitor-treated cells (Figures 5A,B). The efficiency of the inhibitor treatment was verified by the impairment of LPS-induced TNF production (Supplementary Figure 3). As seen previously, LPS treatment also resulted in downregulation of GILZ on the mRNA level. The effect was reversible by inhibition of NF-κB (Figure 5C), suggesting an involvement of an NF-κB-inducible factor in GILZ mRNA regulation. We previously reported that downregulation of GILZ upon MyD88-dependent TLR activation required the presence of the RNA-binding protein tristetraprolin (TTP, encoded by ZFP36) (11, 21). Therefore, we speculated that TTP
FIGURE 2 | Pretreatment with low-dose Poly(I:C) reduces GILZ expression paralleled by sensitization toward LPS and Pam3CSK4. (A,B) AMs were left untreated (Co) or treated with LPS (100 ng/mL) or Poly(I:C) (PIC, 1 µg/mL) for 24 h. GILZ levels were determined by Western blot using tubulin as a loading control. (A) Representative blot. (B) GILZ signal intensities were quantified and normalized to tubulin values (n = 3, duplicates; **p < 0.01 compared with Co).

(C,D) AMs were either left untreated (NT) or pretreated with PIC (1 µg/mL), followed by stimulation with LPS (1 µg/mL, C) or Pam3CSK4 (Pam, 1 µg/mL, D) for another 2 h. TNF, IL6, and CXCL8 mRNA expression was determined by qRT-PCR using ACTB as a housekeeping gene (n = 4, duplicates; **p < 0.01, ***p < 0.001).

might be, at least in part, regulated via NF-κB. We found indeed that ZFP36 mRNA was induced upon LPS treatment in an NF-κB-dependent fashion (Figure 5D).

In line with the finding that Poly(I:C) was unable to reduce GILZ mRNA levels, we observed no induction of TTP after Poly(I:C) treatment (Figure 5E), suggesting an entirely different mode of action.

miRNAs are endogenous small noncoding RNAs that can facilitate translational repression (33). A bioinformatics approach implicated more than 250 miRNAs as potential GILZ regulators. Interestingly, there were more predictions for GILZ than for well-characterized miRNA targets, such as CXCL8, IL6, and TNF (34) (Figure 5F). To evaluate whether miRNAs are involved in Poly(I:C)-induced GILZ repression, AMs were either left untreated or treated with Poly(I:C), aurantricarboxylic acid (ATA), an inhibitor of ribonuclease activities which also affects miRNA processing (35), or a combination of both for 8 h, followed by quantification of GILZ protein levels. We observed that ATA treatment was able to prevent Poly(I:C)-mediated GILZ downregulation, suggesting that miRNAs indeed affect GILZ expression (Figure 5G).

Microarray analysis revealed that the expression of several potentially GILZ-targeting miRNAs was increased in Poly(I:C)-treated AMs, although these effects were in some cases rather donor-dependent (Figure 6A). We tested the ability of 11 of these miRNAs predicted to target GILZ and expressed by Poly(I:C)-treated AMs to regulate GILZ by luciferase reporter gene assays. Within this small set, four miRNAs decreased the activity of the luciferase reporter encoded by a gene fused to the GILZ 3′UTR (Figure 6B). To determine the influence of these miRNAs on endogenous GILZ expression, we transfected HEK293T cells with mimics of the four candidate miRNAs, i.e., miR-34b*, −222, −320d, and −484, and observed that all mimics were able to reduce GILZ expression, although to a different extent (Figures 6C,D). To examine whether these miRNAs exert synergistic effects, HEK293T cells were transfected with the most potent mimic, i.e., miR-34b*, or a mix of all four miRNA mimics. As shown in Figures 6E,F, repression of GILZ expression was more efficient when using the miRNA mimic mix, suggesting that GILZ downregulation upon TLR3 activation is a consequence of the synergistic actions of multiple miRNAs.

We then wondered whether reduced GILZ expression upon miRNA mimic treatment might affect downstream effectors. To this end, HEK-Blue cells expressing a reporter gene under the control of the transcription factors NF-κB and AP-1 were used. Since HEK-Blue cells express endogenous TLR3, reporter gene activity was inducible by Poly(I:C). This effect was profoundly impaired by co-treatment with a cell-permeable GILZ peptide, indicating that GILZ can inhibit NF-κB/AP-1 activation in this system (Figure 6G). In line with these findings, Poly(I:C)-induced NF-κB/AP-1 activity was enhanced in HEK Blue cells transfected with the mir-34b* mimic or a mix of all four mimics, i.e., under conditions that resulted in the highest degree of GILZ repression (Figure 6H).

DISCUSSION

To date, only a few studies have addressed the regulation of endogenous GILZ during the immune response in the
absence of glucocorticoids. In both epithelial (36) and endothelial cells (37), inflammatory cytokines have been demonstrated to attenuate GILZ levels. GILZ expression has also been shown to be decreased in activated macrophages from patients with Crohn’s disease or tuberculosis (6), and nasal explants from patients suffering from chronic rhinosinusitis lack GILZ (38).
Anti-bacterial host defense is enhanced in GILZ knockout macrophages. (A,B) Wildtype- (WT) and GILZ Knockout- (KO) MPI macrophages were treated with LPS (10 or 100 ng/mL) Pam3CSK4 (Pam, 1 µg/mL), or lipoteichoic acid (LTA, 5 µg/mL) with or without co-stimulation by IFN-γ (20 ng/mL) for 20 h. NO-production was measured by Griess assay. Data were normalized to total protein concentration and expressed as x-fold of untreated cells (n = 4, triplicates).

(C–E) Phagocytic activity of WT and GILZ KO MPI cells. MPI cells were incubated with fluorescent latex particles (diameter 1.75 µm, 50 particles per cell) for 15 or 30 min and particle-associated fluorescence was quantified by flow cytometry (15 min: n = 2, 30 min: n = 6, triplicates). (C) Representative histogram for 15 min. 0 P , cells without particles; 1 P , 1 particle; 2 P , 2 particles; > 2 P , more than 2 particles. (G) Phagocytic activity as a total percentage of cells with particles. (E) The (Continued)
**FIGURE 4** | Percentage of cells that engulfed 0, 1, 2 or more than 2 particles was quantified as shown in (C), (F,G) WT and GILZ KO BMMs were left untreated (Co) or treated with LPS (1 µg/mL) and INF-γ (IFN, 20 ng/mL), IL-4 (20 ng/mL) or dexamethasone (Dex, 1 µM) for 20 h, followed by incubation with fluorescent latex particles (diameter 1.75 µm, 50 particles per cell) for 1 h. Particle uptake was quantified by flow cytometry. (F) Total percentage of cells with particle-associated fluorescence. (G) Percentage of cells with 0, 1, 2 or more than 2 particles. (H) Relative number of viable bacteria in *Salmonella typhimurium*-infected WT and GILZ KO BMMs. Extracellular bacteria were removed after 30 min, and the remaining bacteria were killed using gentamicin. Eight hours following infection, monolayers were lysed, and the number of intracellular bacteria was determined as Colony Forming Units (CFU) (*n = 8, triplicates*). *p < 0.05, **p < 0.01, ***p < 0.001 as indicated or compared with equally treated WT cells (E,G).

**FIGURE 5** | Mechanisms of GILZ downregulation upon TLR activation. (A,B) AMs pretreated with BAY-11-7082 (BAY82, 5 µM), BAY-11-7085 (BAY85, 5 µM), or the solvent control DMSO (0.1%) for 1 h, followed by treatment with LPS (100 ng/mL), Pam3CSK4 (Pam, 1 µg/mL), Poly(I:C) (PIC, 10 µg/mL), or medium (Co) for 4 h. GILZ expression was determined by Western blot. Tubulin served as a loading control. (A) Representative blot. (B) GILZ signal intensities were quantified and normalized to tubulin values (*n = 7*). Values for unstimulated DMSO controls were set as 100%. (C,D) After preincubation with BAY-11-7082 and BAY-11-7085 (5 µM, 1 h), solvent (0.1% DMSO) or medium only (Co), AMs were treated with LPS (100 ng/mL) for 2 h. GILZ and ZFP36 mRNA expression was determined by qRT-PCR using ACTB as a housekeeping gene (*n = 3, duplicates*). (E,F) AMs were either left untreated (Co) or treated with LPS (100 ng/mL), Pam3CSK4 (Pam, 1 µg/mL), or Poly(I:C) (PIC, 10 µg/mL) for 4 h. TTP levels were determined by Western blot using tubulin as a loading control, GILZ signal intensities were normalized to tubulin and are shown as a percentage of untreated cells (*n = 2, triplicates*). (F) The number of miRNAs predicted to target GILZ, CXCL8, IL6, and TNF was assessed via the microRNA Data Integration Portal (mirDIP, accession date 02/02/2018). (G) AMs were either left untreated (Co) or treated with Poly(I:C) (PIC, 10 µg/mL), aminocarboxylic acid (ATA, 25 µM), or a combination of both for 8 h. GILZ expression was quantified by Western blot. Signal intensities were normalized to tubulin and expressed as a percentage of untreated cells (*n = 3,* p < 0.05, **p < 0.01, ***p < 0.001.
FIGURE 6 | miRNA-mediated GILZ downregulation. (A) miRNA induction in Poly(I:C)-treated AMs (10 µg/mL for 2 h or 1 µg/mL for 24 h) was assessed by microarray analysis. Data are presented as fold change compared with untreated cells from the same donor. Individual changes per donor (n = 3) are shown. (B) The interaction (Continued)
Furthermore, reduced GILZ mRNA levels were observed in liver tissue from patients with alcoholic hepatitis (17).

Besides the high abundance of GILZ in endothelial cells (21, 37), we previously demonstrated high GILZ levels in primary human monocyte-derived and pulmonary macrophages (21). GILZ expression was diminished in human alveolar macrophages as well as in vivo in mouse lungs upon TLR4 activation.

Further investigations indicated that GILZ mRNA was destabilized upon MyD88-mediated TLR activation, since LPS and Pam3CSK4, but not Poly(I:C), the activator of MyD88-independent TLR3, reduced GILZ mRNA expression. Downregulation of GILZ on the mRNA level required the presence of both the RNA-binding protein TTP and the GILZ 3′-untranslated region (21).

Our results presented within this study suggest that NF-κB activation is also required, which might at least in part be due to NF-κB-mediated TTP induction. Indeed, NF-κB has been previously shown to regulate TTP on the transcriptional level in LPS-stimulated RAW264.7 macrophages (39). However, the composition of the entire RNA-protein complex rather than the binding of TTP alone might dictate the fate of GILZ mRNA. The regulatory networks driving mRNA decay are considered to consist of multiple RNA-binding proteins, whose binding activity may be further modulated by miRNAs (40, 41). Thus, we cannot rule out the possibility that TTP-mediated GILZ mRNA destabilization requires accessory factors that may also be regulated by NF-κB.

Although Poly(I:C) did not decrease GILZ mRNA levels in macrophages, GILZ protein expression was reduced. Furthermore, Poly(I:C)-induced GILZ downregulation was dependent on both NF-κB activation and the presence of miRNAs. Our investigations revealed that multiple miRNAs act as GILZ regulators, suggesting that GILZ downregulation upon TLR3 activation is instead a consequence of synergistic miRNA actions. Promoter analysis using the miRGen v3.0 database (42) revealed that only mir-222 might be a direct target of NF-κB, whereas the promoters of mir-34b, mir-320d-1, and mir-484 lack NF-κB binding sites. NF-κB activation might influence miRNA expression indirectly, e.g., by modulation of miRNA processing via induction of the RNase Dicer (43). Finally, it is also possible that additional miRNAs are induced that were not identified by our approach.

Investigations on the functional significance of GILZ downregulation revealed an enhanced sensitivity toward LPS in human macrophages after siRNA-mediated GILZ knockdown, as indicated by increased cytokine expression and NF-κB activity (21). Within a follow-up study (11) we observed an increased response toward LPS in GILZ KO BMMs, such as enhanced NF-κB and AP-1 activity, suggesting repression of GILZ expression as a regulatory mechanism that amplifies macrophage activation. These findings are in line with our observation that treatment with a cell-permeable GILZ fusion protein can prevent PIC-induced NF-κB/AP-1-dependent activation in HEK-Blue cells.

In addition to its direct interaction with pro-inflammatory transcription factors, GILZ has also been reported to attenuate MAPK signaling in general and ERK signaling in particular (9–11). In the present study, we showed that the absence of GILZ increased both TLR1/2 and TLR3 activation, as indicated by augmented ERK activity, and, as a consequence, increased production of TNF-α. These findings suggest a role for GILZ repression in innate immune memory/trained immunity. Both terms describe an adaption of innate immune cell functions due to previous pathogen exposure, resulting in cross-protection between infections with different pathogens (32, 44).

The increased production of cytokines, such as TNF-α, promotes an acute inflammatory response that helps to clear invading organisms but also contributes to tissue damage. Another factor that plays a similarly ambivalent role in response to infections is nitric oxide (NO), a molecule with antimicrobial and proinflammatory functions. NO is generated from arginine in a reaction catalyzed by the enzyme nitric oxide synthase (NOS). Due to its inducible expression the principal isoform expressed in macrophages, i.e., NOS2, is also known as iNOS. iNOS expression can be elevated by activating cytokines, such as IFN-γ, and repressed by anti-inflammatory mediators, e.g., IL-4 or IL-10. Various PAMPs of bacterial, viral, or fungal origin have also been reported to enhance iNOS expression and function, with LPS as the most prominent example. LPS usually synergizes with IFN-γ to activate both NF-κB and STAT1, thereby inducing high levels of iNOS (45). Our data show that LPS- as well as LPS/IFN-γ-induced NO production can be increased by GILZ depletion. This finding is in line with the observation that LPS-induced NF-κB activity is increased in GILZ KO macrophages (11). In addition, we showed that treatment with TLR2 ligands, i.e., Pam3CSK4 and LTA, results in similar effects when co-administered with IFN-γ. TLR2 ligands did not, however, induce NO on their own, which might be attributed to the lack of host defense.
of TRIF-dependent IFN-β production and subsequent STAT1 activation upon TLR2 activation (46).

The ability to engulf particles, including pathogens and apoptotic cells, is a significant characteristic of macrophages. The phagocytic capacity is not necessarily associated with a specific macrophage phenotype, but rather constitutes a general attribute of these cells (24, 47). In this study, we show that lack of GILZ promotes phagocytosis in untreated as well as differentially polarized macrophages, including GC-treated cells. GCs have only recently been shown to impair phagocytosis by inhibiting the expression of genes that are required for phagosome formation (48). Whether or not these genes, such as CD14, CD48, and MARCKS are regulated in a GILZ-dependent manner presently remains elusive.

Although GILZ knockout macrophages showed a higher phagocytic activity, the number of viable bacteria in S. typhimurium-infected cells was reduced. This finding suggests that GILZ depletion enhances the bactericidal activity of macrophages, which is also supported by the observation that NO production is increased in GILZ knockout macrophages.

In conclusion, our data show that GILZ expression can be downregulated on different levels depending on the nature of the stimulus. In return, GILZ downregulation can promote essential macrophage functions, such as cytokine production, phagocytosis, and bactericidal activity. Thus, GILZ repression can help to clear pathogens, but might also be detrimental to the host by promoting excessive inflammatory disorders. Therefore, GILZ might represent a potential target for therapeutic interventions, both in order to restrict or to stimulate innate immune responses depending on the nature of the disease.

AUTHOR CONTRIBUTIONS

JH, BD, RL, NH, SF, MM, PL, and CB designed, performed, and analyzed the experiments. JH wrote the paper. All authors contributed to drafting the manuscript. SB, CR, FG, EM, and HH provided materials and discussed the data. AK initiated the study and participated in data interpretation and manuscript preparation. All authors read and approved the final manuscript.

FUNDDING

This work was funded, in part, by the Deutsche Forschungsgemeinschaft, Bonn, Germany (KI702).

ACKNOWLEDGMENTS

The authors thank Theresa Stadter, Katharina Sauer, Anke Gückeisen, and Sarah Bender for support in generating murine macrophage data, and Eva Dilly for support in animal maintenance.

SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fimmu.2018.03111/full#supplementary-material

REFERENCES

1. Leifer CA, Medvedev AE. Molecular mechanisms of regulation of Toll-like receptor signaling. J Leukocyte Biol. (2016) 100:927–41. doi: 10.1189/jlb.2MR0316-117RR
2. Brubaker SW, Bonham KS, Zanoni I, Kagan JC. Innate immune pattern recognition: a cell biological perspective. Ann Rev Immunol. (2015) 33:257–90. doi: 10.1146/annurev-immunol-032414-112240
3. Zhou D, Huang C, Lin Z, Zhan S, Kong L, Fang C, et al. Macrophage polarization and function with emphasis on the evolving roles of coordinated regulation of cellular signaling pathways. Cell Signal. (2014) 26:192–7. doi: 10.1016/j.cellsig.2013.11.004
4. Mosser DM, Edwards JP. Exploring the full spectrum of macrophage activation. Nat Rev Immunol. (2008) 8:958–69. doi: 10.1038/nri2448
5. Ayrolde E, Migliorati G, Bruscoli S, Marchetti C, Zollo O, Cannarile L, et al. Modulation of T-cell activation by the glucocorticoid-induced leucine zipper factor via inhibition of nuclear factor kappaB. J Biol Chem. (2007) 35:517–28. doi: 10.1093/nar/gkl1080
6. Berrebi D, Bruscoli S, Cohen N, Fousat A, Migliorati G, Bouchet-Delbos L, et al. Synthesis of glucocorticoid-induced leucine zipper (GILZ) by macrophages: an anti-inflammatory and immunosuppressive mechanism shared by glucocorticoids and IL-10. Blood (2003) 101:729–38. doi: 10.1182/blood-2002-02-0538
7. Di Marco B, Massetti M, Bruscoli S, Macchiariulo A, Di Virgilio R, Velardì E, et al. Glucocorticoid-induced leucine zipper (GILZ)/NF-kappaB interaction: role of GILZ homo-dimerization and C-terminal domain. Nucleic Acids Res. (2007) 35:517–28. doi: 10.1093/nar/gkl1080
8. Mittelstadt PR, Ashwell JD. Inhibition of AP-1 by the glucocorticoid-inducible protein GILZ. J Biol Chem. (2001) 276:29603–10. doi: 10.1074/jbc.M101522200
9. Ayrolde E, Zollo O, Macchiariulo A, Di Marco B, Marchetti C, Riccardi C. Glucocorticoid-induced leucine zipper inhibits the Raf-extracellular signal-regulated kinase pathway by binding to Raf-1. Mol Cell Biol. (2002) 22:7929–41. doi: 10.1128/MCB.22.22.7929-7941.2002
10. Ayrolde E, Zollo O, Bastianelli A, Marchetti C, Agostini M, Di Virgilio R, et al. GILZ mediates the antiproliferative activity of glucocorticoids by negative regulation of Ras signaling. J Clin Investig. (2007) 117:1605–15. doi: 10.1172/JCI30724
11. Hoppstädter J, Kessler SM, Bruscoli S, Huer H, Riccardi C, Kiemer AK. Glucocorticoid-induced leucine zipper: a critical factor in macrophage endotoxin tolerance. J Immunol. (2015) 194:6057–67. doi: 10.4049/jimmunol.1403207
12. D’Adamo F, Zollo O, Moraca R, Riccardi C, Kieler AK. Glucocorticoid-induced leucine zipper: a critical factor in macrophage endotoxin tolerance. J Immunol. (2015) 194:6057–67. doi: 10.4049/jimmunol.1403207
13. Delfino DV, Agostini M, Spinicelli S, Vacca C, Riccardi C. Inhibited cell immune responses depending on the nature of the disease. J Immunol. (2015) 98:4134–1. doi: 10.1074/jbc.M114.584481
14. Delfino DV, Agostini M, Spinicelli S, Vacca C, Riccardi C. Inhibited cell immune responses depending on the nature of the disease. J Immunol. (2015) 98:4134–1. doi: 10.1074/jbc.M114.584481
15. Bruscoli S, Biagioli M, Sorcini D, Frammartino T, Cimino M, Sportoletti P, et al. Lack of Glucocorticoid-induced leucine zipper family protect T lymphocytes from TCR/CD3-activated cell death. Immunity (1997) 7:803–12.
16. D’Adamo F, Zollo O, Moraca R, Riccardi C, Kieler AK. Glucocorticoid-induced leucine zipper: a critical factor in macrophage endotoxin tolerance. J Immunol. (2015) 194:6057–67. doi: 10.4049/jimmunol.1403207
17. D’Adamo F, Zollo O, Moraca R, Riccardi C, Kieler AK. Glucocorticoid-induced leucine zipper: a critical factor in macrophage endotoxin tolerance. J Immunol. (2015) 194:6057–67. doi: 10.4049/jimmunol.1403207
18. Bruscoli S, Biagioli M, Sorcini D, Frammartino T, Cimino M, Sportoletti P, et al. Lack of Glucocorticoid-induced leucine zipper (GILZ) deregulates B cell survival and results in B cell lymphocytosis in mice. Blood (2015). 126:1790–801. doi: 10.1182/blood-2015-03-631580
19. Bruscoli S, Sorcini D, Flamini S, Gagliardi A, Adamo F, Ronchetti S, et al. Glucocorticoid-induced leucine zipper inhibits interferon-gamma production in B cells and suppresses colitis in mice. Front Immunol. (2018) 9:1720. doi: 10.3389/fimmu.2018.01720.
17. Hamdi H, Bigorgne A, Naveau S, Balian A, Bouchet-Delbos L, Cassard-Doulcier AM, et al. Glucocorticoid-induced leucine zipper: a key protein in the sensitization of monocytes to lipopolysaccharide in alcoholic hepatitis. Hepatology (2007) 46:1986–92. doi: 10.1002/hep.21880
18. Vago JP, Tavares LP, Garcia CC, Lima KM, Perucci LO, Vieira EL, et al. The role and effects of glucocorticoid-induced leucine zipper in the context of inflammation resolution. J Immunol. (2015) 194:4940–50. doi: 10.4049/jimmunol.1401722
19. Hoppstädter J, Kiemer AK. Glucocorticoid-induced leucine zipper (GILZ) in immunosuppression: master regulator or bystander? Oncotarget (2015) 6:38446–57. doi: 10.18632/oncotarget.6197
20. Hoppstädter J, Hachenthal N, Valbuena-Perez JV, Lampes S, Astanina K, Kunze MM, et al. Induction of glucocorticoid-induced leucine zipper (GILZ) contributes to anti-inflammatory effects of the natural product curcumin in macrophages. J Biol Chem. (2016) 291:22949–60. doi: 10.1074/jbc.M116.733253
21. Hoppstädter J, Diesel B, Eisler UK, Schmid T, Brüne B, Kiemer AK. Glucocorticoid-induced leucine zipper is downregulated in human alveolar macrophages upon Toll-like receptor activation. Eur J Immunol. (2012) 42:1282–93. doi: 10.1002/eji.201142081
22. Bruscoli S, Velardi E, Di Sante M, Bereshchenko O, Venanza A, Coppedo M, et al. Long glucocorticoid-induced leucine zipper (L-GILZ) protein interacts with ras protein pathway and contributes to spermagenesis control. J Biol Chem. (2012) 287:1242–51. doi: 10.1074/jbc.M111.316372
23. Hoppstädter J, Diesel B, Zarbock R, Breinig T, Monz D, Koch M, et al. Differential cell reaction upon Toll-like receptor 4 and 9 activation in human alveolar and lung interstitial macrophages. Respir Res. (2010) 11:124. doi: 10.1186/1465-9921-12-11
24. Hoppstädter J, Seif M, Dembek A, Cavelius C, Huwer H, Kraegeloh A, et al. Hoppstädter et al. Lack of GILZ Amplifies Host Defense expression and hybridization array data repository. Nucleic Acids Res. (2017) 45:6353–63. doi: 10.1093/nar/gkw1193
25. Dembek A, Laggai S, Kessler SM, Czepukojc B, Simon Y, Kiemer AK, et al. Downregulation of the glucocorticoid-induced leucine zipper (GILZ) promotes vascular inflammation. Atherosclerosis (2014) 234:391–400. doi: 10.1016/j.atherosclerosis.2014.03.028
26. Eddleston J, Herschbach J, Bagel-Steven AL, Christiansen SC, Zuraw BL. The anti-inflammatory effect of glucocorticoids is mediated by glucocorticoid-induced leucine zipper in epithelial cells. J Allergy Clin Immunol. (2007) 119:115–22. doi: 10.1016/j.jaci.2006.08.027
27. Hahn RT, Hoppstädter J, Hirschfelder K, Hachenthal N, Diesel B, Kessler SM, et al. Induction of glucocorticoid-induced leucine zipper (GILZ) contributes to anti-inflammatory effects of the natural product curcumin in macrophages. J Biol Chem. (2016) 291:22949–60. doi: 10.1074/jbc.M116.733253
30. Liu B, Liu Q, Yang L, Palaniappan SK, Bahar I, Thiagarajan PS, et al. The role and effects of glucocorticoid-induced leucine zipper in the context of inflammation resolution. J Immunol. (2015) 194:4940–50. doi: 10.4049/jimmunol.1401722
31. Cannarile L, Cuzzocrea S, Santucci L, Agostini M, Mazzon E, Esposito E, et al. Glucocorticoid-induced leucine zipper is protective in Th1 mediated models of colitis. Gastroenterology (2010) 138:530–41. doi: 10.1053/j.gastro.2008.09.024
32. Liu B, Liu Q, Yang L, Palaniappan SK, Bahar I, Thigaranjan PS, et al. Innate immune memory and homeostasis may be conferred through crosstalk between the TLR3 and TLR7 pathways. Sci Signal. (2016) 9:a97a0. doi: 10.1126/scisignal.aac9340
33. Iwakawa HO, Tomari Y. The functions of MicroRNAs: mRNA decay and translational repression. Trends Cell Biol. (2015) 25:651–65. doi: 10.1016/j.tcb.2015.07.011
34. Rebane A, Akdis CA. MicroRNAs: Essential players in the regulation of inflammation. J Allergy Clin Immunol. (2013) 132:15–26. doi: 10.1016/j.jaci.2013.04.011
35. Tan GS, Chiu CH, Garchow BG, Metzler D, Diamond SL, Kirikidou M. Small molecule inhibition of RISC loading. ACS Chem Biol. (2012) 7:403–10. doi: 10.1021/cb200253b
36. Eddleston J, Herschbach J, Bagel-Steven AL, Christiansen SC, Zuraw BL. The anti-inflammatory effect of glucocorticoids is mediated by glucocorticoid-induced leucine zipper in epithelial cells. J Allergy Clin Immunol. (2007) 119:115–22. doi: 10.1016/j.jaci.2006.08.027
37. Hahn RT, Hoppstädter J, Hirschfelder K, Hachenthal N, Diesel B, Kessler SM, et al. Induction of glucocorticoid-induced leucine zipper (GILZ) promotes vascular inflammation. Atherosclerosis (2014) 234:391–400. doi: 10.1016/j.atherosclerosis.2014.03.028
38. Zhang XH, Lu X, Long XB, You Y, Gao QX, Cui YH, et al. Chronic rhinosinusitis with and without nasal polyps is associated with decreased expression of glucocorticoid-induced leucine zipper. Clin Exp Allergy (2009) 39:647–54. doi: 10.1111/j.1365-2228.2008.03198.x
39. Chen YL, Jiang YW, Su YL, Lee SC, Chang MS, Chang CJ. Transcriptional regulation of tristetraprolin by NF-kappaB signaling in LPS-stimulated macrophages. Molecular biology reports (2013) 40:2867–77. doi: 10.1007/s11033-012-2302-8
40. Doyle P, Tenenbaum SA. Trans-regulation of RNA-binding protein motifs by microRNA. Front Genet. (2014) 5:79. doi: 10.3389/fgen.2014.00079
41. Wu X, Brewer G. The regulation of mRNA stability in mammalian cells: 2.0. Gene (2012) 500:10–21. doi: 10.1016/j.gene.2012.03.021
42. Georgakilas G, Vlachos IS, Zagganas K, Vergoulis T, Paraskevopoulos MD, Kanellos I, et al. DIANA-miRGen v3.0: accurate characterization of microRNA promoters and their regulators. Nucleic Acids Res. (2016) 44:D190–5. doi: 10.1093/nar/gkv1254
43. Guan Y, Yao H, Wang J, Sun K, Cao L, Wang Y. NF-kappaB-DICER-miRs axis regulates TNF-alpha expression in responses to endothox stress. Int J Biol Sci. (2015) 11:1257–68. doi: 10.7150/ijbs.12611
44. Netea MG, Joosten LA, Latz E, Mills KH, Natoli G, Stunnenberg HG, et al. Trained immunity: a program of innate immune memory in health and disease. Science (2016) 352:aaf1098. doi: 10.1126/science.aaf1098
45. Bogdan C. Nitric oxide synthase in innate and adaptive immunity: an update. Trends Immunol. (2015) 36:161–78. doi: 10.1016/j.it.2015.01.003
46. Toschakov V, Jones BW, Perera PY, Thomas K, Cody MJ, Zhang S, et al. TLR4, but not TLR2, mediates IFN-beta-induced STAT1alpha/beta-dependent gene expression in macrophages. Nat Immunol. (2002) 3:392–8. doi: 10.1038/nij774
47. Mills CD, Ley K. M1 and M2 macrophages: the chicken and the egg of immunity. J Innate Immun. (2014) 6:716–26. doi: 10.1159/000364945
48. Olivares-Morales MJ, De La Fuente MK, Dubois-Camacho K, Parada D, Diaz-Jimenez D, Torres-Riquelme A, et al. Glucocorticoids impair phagocytosis and inflammatory response against crohn’s disease-associated adherent-invasive Escherichia coli. Front Immunol. (2018) 9:1026. doi: 10.3389/fimmu.2018.01026

Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.