Novel Majeed Syndrome–Causing LPIN2 Mutations Link Bone Inflammation to Inflammatory M2 Macrophages and Accelerated Osteoclastogenesis

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Objective. To identify novel heterozygous LPIN2 mutations in a patient with Majeed syndrome and characterize the pathomechanisms that lead to the development of sterile osteomyelitis.

Methods. Targeted genetic analysis and functional studies assessing monocyte responses, macrophage differentiation, and osteoclastogenesis were conducted to compare the pathogenesis of Majeed syndrome to interleukin-1 (IL-1)–mediated diseases including neonatal-onset multisystem inflammatory disease (NOMID) and deficiency of the IL-1 receptor antagonist (DIRA).

Results. A 4-year-old girl of mixed ethnic background presented with sterile osteomyelitis and elevated acute-phase reactants. She had a 17.8-kb deletion on the maternal LPIN2 allele and a splice site mutation, p.R517H, that variably spliced out exons 10 and 11 on the paternal LPIN2 allele. The patient achieved long-lasting remission receiving IL-1 blockade with canakinumab. Compared to controls, monocytes and monocyte-derived M1-like macrophages from the patient with Majeed syndrome and those with NOMID or DIRA had elevated caspase 1 activity and IL-1β secretion. In contrast, lipopolysaccharide-stimulated, monocyte-derived, M2-like macrophages from the patient with Majeed syndrome released higher levels of osteoclastogenic mediators (IL-8, IL-6, tumor necrosis factor, CCL2, macrophage inflammatory protein 1α/β, CXCL8, and CXCL1) compared to NOMID patients and healthy controls. Accelerated osteoclastogenesis in the patient with Majeed syndrome was associated with higher NFATc1 levels, enhanced JNK/MAPK, and reduced Src kinase activation, and partially responded to JNK inhibition and IL-1 (but not IL-6) blockade.

Conclusion. We report 2 novel compound heterozygous disease-causing mutations in LPIN2 in an American patient with Majeed syndrome. LPIN2 deficiency drives differentiation of proinflammatory M2-like macrophages and enhances intrinsic osteoclastogenesis. This provides a model for the pathogenesis of sterile osteomyelitis which differentiates Majeed syndrome from other IL-1–mediated autoinflammatory diseases.

INTRODUCTION

Genetically defined autoinflammatory bone diseases are caused by dysregulation of innate immune responses resulting in systemic inflammation (1) and osteomyelitis, which includes Majeed syndrome, caused by recessive loss-of-function mutations in LPIN2 (2), and deficiency of interleukin-1 (IL-1) receptor antagonist (DIRA), caused by loss-of-function mutations in IL1RN.
encoding the IL-1 receptor (IL-1R) antagonist (3,4). Majeed syndrome presents in early childhood with osteitis and osteomyelitis, dyserythrocritinaemia, and elevation of acute-phase reactants. Since its initial description in 1989 (5), 23 patients from 10 families/kindreds of Jordanian, Indian, Arabic, Turkish, and Chinese ethnic backgrounds have been described as having 7 autosomal-recessive, disease-causing loss-of-function mutations (summarized in Supplementary Table 1, available on the Arthritis & Rheumatology website at http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract). Clinical similarities to DIRA and a rapid resolution of the inflammatory disease manifestations upon treatment with short- and long-acting IL-1 inhibitors suggest a prominent role of IL-1 in disease pathogenesis (6,7).

**LPIN2 encodes lipin 2, a magnesium-dependent phosphatidate phosphatase enzyme that associates with membrane lipids and localizes to organelles. Lipin 2 catalyzes the conversion of phosphatidic acid to diacylglycerol (DAG) and acts as a branch point for the synthesis of triacylglycerol and phospholipids (8–12). Recently, a link between lipin 2 deficiency and inflammation was described (13). Lipin 2 negatively regulates MAPK phosphorylation and contributes to the activation of the NLRP3 inflammasome. Bone marrow–derived macrophages (BMMs) from Lpin2−/− mice or small interfering RNA–induced silencing of Lpin2 in murine and human macrophages showed increased IL-1 production in response to stimulation with lipopolysaccharide (LPS) and ATP (13). Reduced lipin 2 levels, as seen in patients with Majeed syndrome with disease-causing loss-of-function mutations, result in a decrease in cellular cholesterol levels, which increases ion currents through the ATP-activated P2X7 receptor and subsequent NLRP3 inflammasome activation with mature IL-1β release (13).

Despite insights that link LPIN2 mutations to increased pro–IL-1β transcription and to NLRP3 inflammasome activation, pathomechanisms that provide insight into the differential development of osteomyelitis in Majeed syndrome and DIRA (but not in the NLRP3 inflammasomopathy, neonatal-onset multisystem inflammatory disease [NOMID]) remain unresolved (Supplementary Figure 1, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract) and suggest NLRP3 inflammasome–independent effects of LPIN2 in regulating bone homeostasis at the growth plate of long bones. In the present study we demonstrated that LPIN2 mutations differentially affect monocytes, monocyte-derived M1-like macrophages (M1-MDMs) and M2-MDMs, and osteoclastogenesis in the various IL-1-mediated diseases. In contrast to the NLRP3 inflammasomopathy, NOMID, LPS-stimulated M2-MDMs from patients with Majeed syndrome and DIRA are inflammatory, and the mutations promote accelerated osteoclastogenesis. These effects were mediated through changes in phosphorylation patterns in macrophages in the patient with Majeed syndrome and could largely be reversed through IL-1 inhibition. Our data suggest a novel role of lipin 2 in macrophage polarization and osteoclastogenesis, and we propose a disease model for sterile osteomyelitis.

**PATIENTS AND METHODS**

**Patients.** The parents of the patient with Majeed syndrome and the adult healthy control donors provided written informed consent. Patients were enrolled in an institutional review board–approved National Institutes of Health (NIH) natural history protocol (ClinicalTrials.gov identifier: NCT02874595). Blood samples were collected. Further details about all methods described in this section are presented in Supplementary Methods (http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract).

**Genetic analyses.** Targeted sequencing. Clinical genetic testing of ELANE, LPIN2, MEFV, MVK, NLRP3, PSTPIP1, and TNFRSF1A (GeneDx) was inconclusive.

Array-based comparative genomic hybridization (aCGH). Genomic DNA from the mother of the patient with Majeed syndrome was examined by aCGH using the current version of ExonArrayDx (GeneDx).

Deletion breakpoint delineation by polymerase chain reaction (PCR) and Sanger sequencing. Serial forward and reverse primers close to the 5’ and 3’ ends of the deletion, respectively, were designed to obtain new amplicons. PCR products were sequenced in both directions using an ABI 3100 Genetic Analyzer (Applied Biosystems).

Detection of LPIN2 splicing abnormalities by complementary DNA (cDNA) PCR and sequencing. Complementary DNA primers were designed using Primer3Plus to amplify the region between exon 8 and exon 12 of LPIN2 NM_014646.2.

**LPIN2 transcript quantification in RNA-Seq data using StringTie.** Raw reads were mapped to the reference human genome (hg19) by TopHat (version 2.0.8; https://ccb.jhu.edu/software/tophat/index.shtml), and we assembled transcripts and determined normalized expression values for each spliced event using StringTie (version 1.3.3).

**Cell culture experiments.** Differentiation of human monocytes and monocyte-derived macrophages. Human peripheral blood–derived monocytes and macrophages were prepared from heparinized venous blood as previously described (14). Peripheral blood mononuclear cells (PBMCs) were suspended in RPMI 1640 medium containing a low concentration (1%) of fetal bovine serum (FBS). Adherent monocytes were seeded in multiwell plates or dishes; they were differentiated into M1-MDMs and M2-MDMs by culturing with RPMI 1640 supplemented with 10% FBS containing either 10 ng/ml recombinant human granulocyte–macrophage colony-stimulating factor (GM-CSF) (for M1-MDMs) or 100 ng/ml recombinant human M-CSF (for M2-MDMs) (both from PeproTech). After 3 days, media were replaced with either
GM-CSF or M-CSF, and fresh complete media and cells were incubated for another 2 or 3 days.

Osteoclast differentiation and pit formation assay. Osteoclasts were prepared from peripheral blood–derived monocytes (15). Briefly, PBMCs from the patient with Majeed syndrome and the controls were suspended in α-minimum essential medium (α-MEM) containing 1% FBS. Monocytes were enriched by allowing attachment to plates and were cultured in α-MEM containing 10% FBS with M-CSF (100 ng/ml) for 6 days; floating cells were removed, and the attached MDMs were used as osteoclast precursors. To generate osteoclasts, M-CSF macrophages were cultured with combinations of M-CSF (100 ng/ml) and RANKL (30 ng/ml). After an additional 3–6 days of culture, cells were fixed and stained for tartrate-resistant alkaline phosphatase (TRAP) (Cosmo Bio). TRAP-positive multinucleated cells containing >3 nuclei were considered TRAP-positive multinuclear osteoclasts.

For the pit assay, harvested cells were inoculated on fluorescein-labeled CaP-coated 24-well plates at a density of 1 × 10⁴ cells/well (Cosmo Bio), and were cultured for an additional 3–6 days with M-CSF (100 ng/ml) and RANKL (30 ng/ml) according to the manufacturer’s recommendation. After 6 days, plates were washed with phosphate buffered saline and treated with 5% sodium hypochlorite for 5 minutes. After washing the plates, 5 different regions in each well were photographed by microscopy, and the pit areas were measured with ImageJ software.

Stimulation/inhibition assays. Adherent monocytes, M1-MDMs, or M2-MDMs were stimulated in the presence of LPS (1 μg/ml) for 2 hours and 30 minutes followed by 30-minute stimulation with ATP (1 mmole) (Sigma-Aldrich). After 24 hours, culture supernatants were collected. For blocking assays, the following were used: IL-1 inhibitor anakinra (Sobi), potent JNK inhibitor SP600125 or phosphorylation of JNK (InvivoGen), macrophage migration inhibitory factor (MIF) inhibitor ISO-1 (R&D Systems), potent tumor necrosis factor (TNF) inhibitor SPD304 (Sigma-Aldrich), and IL-6 inhibitor tocilizumab (Genentech).

Cytokine/chemokine production and enzyme-linked immunosorbent assay (ELISA). Relative levels of multiple cytokines and chemokines in the supernatants of macrophages were analyzed using a proteome profiler human cytokine array according to the instructions of the manufacturer (R&D Systems). Briefly, culture supernatants (200 μl) of macrophages, which were collected after centrifugation, were added to dot-bLOTS onto which the captured antibodies had been spotted in duplicate. After incubation with the secondary antibody mixture, the resultant signals were detected using a Bio-Rad image analyzer. A total of 36 cytokines, chemokines, and acute-phase proteins can be screened (16,17). The intensity of the spots was quantified using ImageJ software. In order to assess concentrations of IL-1α, IL-1β, and IL-10, the supernatants of macrophages stimulated in the presence or absence of LPS and ATP were collected and analyzed by ELISA (R&D Systems).

Phosphokinase array. A human Phospho-Kinase Array Kit (no. ARY003B; R&D Systems) was used to simultaneously detect the relative site-specific phosphorylation of 43 kinases. Briefly, cells were plated and cultured for M2-MDM differentiation with M-CSF (100 ng/ml), and 0.5 × 10⁶ cells were lysed and assayed according to the manufacturer’s instructions. The array results were quantified using ImageJ software.

Immunofluorescence staining. Stimulated and unstimulated macrophages were fixed in 4% paraformaldehyde, permeabilized with 0.1% Triton X-100, and stained with a single antibody or combinations of anti-gasdermin D antibody (no. LS-B4537; LifeSpan Biosciences), anti–caspase 1 antibody (no. ab-1872; Abcam), anti–lipin 2 antibody (no. HPA017857; Sigma-Aldrich), anti–RANK antibody (no. ab222215; Abcam), and anti–IL-1R1 antibody (no. ab106278; Abcam) for 12 hours, followed by Alexa Fluor 568–conjugated anti-mouse IgG antibody (Invitrogen) or Alexa Fluor 488–conjugated anti-rabbit IgG antibody (Abcam). DAPI (Molecular Probes) was used to visualize nuclei. Signals were visualized with a confocal laser scanning microscope (Leica SP8). Image processing was performed with Imaris 9.2.1 software.

Statistical analysis. Statistical analyses and graphing were performed using GraphPad Prism 6. Differences in unpaired data with parametric or nonparametric distributions were analyzed by Student’s t-test and Mann-Whitney U test, respectively, assuming the same variance for each group. P values were not corrected for multiple comparisons and must be viewed as exploratory.

RESULTS

Majeed syndrome in 4-year-old patient caused by 2 novel compound heterozygous mutations in LPIN2.

A 4-year-old female patient of Puerto Rican and African American descent was evaluated at the NIH for suspected Majeed syndrome (Supplementary Results, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract). A radiograph of the right knee obtained at 26 months to evaluate recurrent right knee pain revealed horizontal lucencies at the distal femur and proximal tibia. Laboratory examinations showed systemic inflammation with elevated erythrocyte sedimentation rate (ESR) (55 mm/hour) and C-reactive protein (CRP) level (61 mg/liter), and a knee magnetic resonance imaging demonstrated an abnormal bone marrow signal in the bilateral distal femoral metaphyses (Figure 1B).

The patient had anemia, and a bone marrow biopsy showed 70–90% marrow cellularity with trilineage hematoipoiesis and dyserythropoietic, binuclear erythrocyte precursors (Supplementary Figure 2, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract). At the first NIH visit at the age of 4 years, the laboratory examination showed mild anemia with the following test results: hemoglobin 10.1 gm/dl (normal 10.2–12.7), mean corpuscular
volume 65.2 fl (normal 72.3–85), thrombocytosis with platelets 556 K/μl (normal 189–394), and systemic inflammation (ESR 77 mm/hour [normal <15] and CRP 104 mg/liter [normal <5]). After intermittent short courses of nonsteroidal antinflammatory drugs, the patient was started on treatment with the monoclonal anti–IL-1β antibody, canakinumab, at 2 mg/kg every 8 weeks. She has continued to receive treatment for the past 6.5 years, with significant clinical and laboratory improvement (Supplementary Table 2, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract).

The clinical features and treatment response suggested an autoinflammatory bone disease, and genetic testing identified a previously unreported mutation in LPIN2, c.1550G>A, p.R517H, a variant that was initially thought to be homozygous. The presence of the missense variant in the patient’s father but not her mother prompted a search for a large deletion that revealed a multiexon deletion spanning exons 7–18, which was confirmed by aCGH of the maternal DNA (Figure 1C). PCR and Sanger sequencing of the approximate breakpoint regions that were identified by aCGH demonstrated a 17,770-bp deletion (Chr18: 2,921,149–2,938,919) (Supplementary Figures 3A–C, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract).

To investigate aberrant splicing by the paternally inherited mutation in exon 10, we sequenced LPIN2 cDNA between exons 8 and 12 and detected 2 alternatively spliced transcripts with deleted exon 10 alone (17.2% of the transcript) or exons 10 and 11 (26.7% of the transcript). LPIN2 mRNA expression in human cell subsets and tissues was quantified by real-time quantitative polymerase chain reaction. Fold changes in LPIN2 expression (relative to 18S) are shown; bars show the mean ± SD of 3 technical replicates. Levels of LPIN1, LPIN2, and LPIN3 mRNA were quantified by RNA-Seq analysis in human monocytes from healthy controls; bars show the mean ± SD of 3 samples. PBMC = peripheral blood mononuclear cell; NK = natural killer; HUVEC = human umbilical vein endothelial cell.
of transcripts lacked exons 10 and 11 (Figure 1D). LPIN2 was widely expressed, including in myeloid cells (Figure 1E), and of the 3 lipins, monocytes almost exclusively expressed LPIN2 (Figure 1F). Using commercial antibodies that are directed to the C-terminal domain (which was deleted in the mutation from the patient’s mother), we detected ~50% of normal lipin 2 protein expression in patient cells, which was likely from the paternal splice variants (data not shown).

**Lipin 2 mutations cause constitutive inflammasome activation and high IL-1β production in patient monocytes and M1-MDMs.** Previously, serum IL-1β levels were reported to be elevated in patients with Majeed syndrome who had lipin 2 mutations (6) and in macrophages from *Lpin2−/−* mice (13). We assessed IL-1β production in LPS- and ATP-stimulated monocytes and in M1-MDMs and M2-MDMs, comparing the patient with Majeed syndrome to disease controls and healthy controls. Increased inflammasome activation and IL-1β secretion were observed on 2 separate visits in the patient with Majeed syndrome, despite receiving canakinumab. IL-1β release in monocytes was elevated to a level similar to that observed in patients with the IL-1–mediated autoinflammatory osteomyelitis syndrome (DIRA) but lower than that observed in patients with the IL-1–mediated neutrophilic osteomyelitis (NOMID), which does not present with osteomyelitis (Figure 2A). IL-1β release was blocked by the recombinant IL-1 receptor antagonist, anakinra (shown in monocytes) (Figure 2B). In contrast to monocytes, unstimulated M1-MDMs and M2-MDMs from the patient with Majeed syndrome and patients with DIRA exhibited elevated IL-1β levels, and LPS- and ATP-stimulated IL-1β secretion was also increased in M1-MDMs, but not in M2-MDMs, in the patient with Majeed syndrome (Figure 2C).

Consistent with these observations, caspase 1 and gasdermin D expression were up-regulated in M1-MDMs in the patient with Majeed syndrome, but gasdermin D expression in M2-MDMs was comparable to that in healthy controls (Supplementary Figure 4 and Supplemental Movie 1, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract). In contrast, LPS- and ATP-stimulated M1-MDMs and M2-MDMs from NOMID patients secreted significantly more IL-1β than healthy controls (Figures 2C and D). While LPS- and ATP-stimulated IL-1α secretion was variable in monocytes and M1-MDMs, IL-1α tended to be higher in stimulated M2-MDMs from the patient with Majeed syndrome and those with DIRA, but not in NOMID M2-MDMs (Supplementary Figure 5, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract).

**Proinflammatory M2-MDMs from the patient with Majeed syndrome compared to patients with NOMID and healthy controls.** Monocytes from the patient with Majeed syndrome differentiated normally into M1-MDMs and M2-MDMs. While M2-MDMs expressed low levels of CD80 (M1-MDM marker), similar to controls, they had lower expression of

**Figure 2.** Effect of disease-causing *Lpin2* mutations on interleukin-1β (IL-1β) production in monocytes and macrophages. IL-1β concentration in supernatant was analyzed by enzyme-linked immunosorbent assay. A, Adherent monocytes from peripheral blood mononuclear cells were stimulated with lipopolysaccharide (LPS; 1 μg/ml) for 2.5 hours and ATP (1 mmole) for 30 minutes. Each symbol represents an individual subject: healthy controls (n = 3), patient with Majeed syndrome (n = 1), patients with neonatal-onset multisystem inflammatory disease (NOMID) (n = 3), and patients with deficiency of the IL-1 receptor antagonist (DIRA) (n = 3). Bars show the mean ± SD. Values from the patient with Majeed syndrome represent the mean of 3 technical replicates. B, LPS- and ATP-stimulated monocytes were cultured in the presence or absence of anakinra (10 mg/ml). Each symbol represents an individual subject: healthy controls (n = 6), patient with Majeed syndrome (n = 1), and patients with NOMID (n = 3), and patients with DIRA (n = 4). Bars show the mean ± SD. Values from the patient with Majeed syndrome were obtained from 3 independent experiments, using samples from separate visits, with the mean of 2 technical replicates shown. C and D, M1-like macrophages (C) and M2-like macrophages (D) were stimulated with LPS (1 μg/ml) and ATP (1 mmole) for 24 hours. Each symbol represents an individual subject: healthy controls (n = 6), patient with Majeed syndrome (n = 1), patients with NOMID (n = 3), and patients with DIRA (n = 4). Bars show the mean ± SD. Values for the patient with Majeed syndrome represent the mean of the technical replicates. * = P < 0.05; ** = P < 0.001, by 2-sample *t*-test assuming equal variances to compare disease groups, or by Student’s unpaired *t*-test to compare stimulation with anakinra to no stimulation with anakinra.

**CD163** (M2-MDM marker) compared to M2-MDMs from healthy controls (Supplementary Figure 6, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract). To assess responses of M1-MDMs and M2-MDMs to inflammatory stimuli, cytokine and chemokine profiles were compared among LPS- and ATP-stimulated M1-MDMs and M2-MDMs from the patient with Majeed syndrome and those with NOMID or DIRA (Figures 3A and B). IL-8 was constitutively expressed in the M2-MDMs but not the M1-MDMs from the patient with Majeed syndrome and from 1 of 4 DIRA patients.
Compared to healthy controls and NOMID patients, the M2-MDMs from the patient with Majeed syndrome secreted more IL-6 and IL-8 in response to LPS stimulation, which was most similar to the DIRA patients. TNF elevation was higher in M1-MDMs than in M2-MDMs from the patient with Majeed syndrome compared to healthy controls; however, levels in M2-MDMs from the patient with Majeed syndrome and patients with NOMID or DIRA were similar. Higher secretion of chemokines, including CXCL1/growth-related oncogene α (GROα), macrophage inflammatory protein 1α/β (MIP-1α/β), CCL5/RANTES, and CXCL10, was observed in LPS- and ATP-stimulated M2-MDMs from the patient with Majeed syndrome compared to unstimulated macrophages as well as stimulated disease control macrophages (Figures 3A and B). IL-10 production by M2-MDMs from the patient with Majeed syndrome and 3 of 4 DIRA patients was lower compared to healthy controls and NOMID patients (Figure 3C).

Enhanced osteoclastogenesis in the patient with Majeed syndrome compared to healthy controls and patients with NOMID. Although many of the proinflammatory cytokine/chemokines that are up-regulated in Majeed syndrome promote osteoclastogenesis (18–21), we hypothesized that the LPIN2 mutations may promote osteoclastogenesis independently of external cytokine supplementation. M2-MDMs are osteoclast precursors (22,23) that differentiate into osteoclasts upon RANKL stimulation. In fact, higher numbers of monocyte-derived TRAP-positive osteoclasts were derived from M2-MDMs from the patient with Majeed syndrome compared to cells from healthy controls and NOMID patients; more heterogeneity in osteoclast formation was observed in the DIRA patients (Figure 4A). Osteoclast function, assayed by bone resorption using a pit assay, showed higher bone resorption in the patient with Majeed syndrome and DIRA patients compared to the healthy controls and NOMID patients (Figure 4B). RANK expression that accelerates osteogenesis (24) was increased in M2-MDMs from the patient with Majeed syndrome compared to healthy controls (Figure 4C), while overall IL-1R1 expression was comparable to that in healthy controls, with higher cell surface expression of IL-1R1 in the M2-MDMs from the patient with Majeed syndrome (Figure 4D). RANKL stimulation in healthy controls increased IL-1R1 surface expression (Supplementary Figure 7, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract), suggesting a role of RANKL in regulating surface expression of IL-1R1. Induction of IL-1R1 expression by c-Fos enabled IL-1–induced osteoclastogenesis in murine BMMs.
REGULATION OF INFLAMMATORY PROPERTIES OF M2- MDMs BY A PHOSPHOLIPASE ENCODED BY LPIN2

in a RANKL/RANK-independent manner (24,25). In human cells, osteoclast differentiation was not induced by recombinant IL-1β, independent of RANKL, nor did IL-1β enhance RANKL-induced osteoclastogenesis (Supplementary Figure 8, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract). Binding of RANKL to its receptor (RANK) activates and/or induces expression of key transcription factors such as NF-κB, c-Fos, melanocyte-inducing transcription factor, PU.1, and NFATc1, which is considered a master regulator of osteoclast differentiation in vitro and in vivo (26). NFATc1 orchestrates a signaling cascade that stimulates activator protein 1 and costimulatory signal–mediated intracellular Ca2+ oscillation (24,26,27). Consistent with these observations, NFATc1 was significantly up-regulated in the supernatant of LPS- and ATP-stimulated M2-MDMs and in osteoclasts from the patient with Majeed syndrome compared to healthy controls (Supplementary Figure 9, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract).

IL-1 inhibition effectively blocks proinflammatory cytokine production in M2-MDMs and accelerated osteoclastogenesis in the patient with Majeed syndrome. The clinical benefit of IL-1 blocking treatments in Majeed syndrome prompted investigation into the effect of IL-1 inhibition on stimulated cytokine and chemokine production in M2-MDMs and on osteoclastogenesis. IL-1 inhibition with anakinra significantly reduced the production of inflammatory cytokines and osteoclastogenic chemokines in LPS- and ATP-stimulated M2-MDMs from the patient with Majeed syndrome and healthy controls, including the neutrophil-recruiting chemokines CXCL1/GROα and IL-8, as well as IL-6, interferon-γ–inducible protein 10, TNF, and CCL5/RANTES (Figure 5A). In contrast, IL-10 levels remained low (data not shown). IL-1 inhibition with anakinra effectively reduced the number of TRAP-positive osteoclasts in a dose-dependent manner (Figure 5B), and the inhibitory effect was more pronounced in the patient with Majeed syndrome compared to controls. Although IL-1β production increased during osteoclastogenesis, levels did not significantly differ between healthy controls and the patient with Majeed syndrome (Supplementary Figure 10A, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract). In contrast, the osteoclast differentiation factor MIF and serpin/plasminogen activator inhibitor (PAI) were induced in Majeed syndrome osteoclasts compared to healthy control osteoclasts (Supplementary Figures 10B and C). Similarly, osteoclastogenic factors CXCL1, CCL2/monocyte chemotactic protein 1 (MCP-1), and CXCL8, which may function as autocrine factors in amplifying osteoclastogenesis, increased during osteoclast differentiation in the patient and controls (with higher levels in the patient with Majeed syndrome on day 8) (Supplementary Figures 10D–F). Osteoclastogenesis was inhibited by ISO-1 and anakinra, but it was less inhibited by tocilizumab, SPD304, and SP600125.
Anakinra reduced MIF, serpin/PAI, and intercellular adhesion molecule 1 production during osteoclastogenesis more effectively than IL-6 inhibition (Supplementary Figure 11, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract), and may block osteoclastogenesis by reducing autocrine production of the osteoclastogenic factors MIF and serpin/PAI.

**LPIN2 mutations affect M2-MDM activation and osteoclast differentiation by modification of phosphokinase activity of JNK/MAPKs and Src kinases.** Lipin 2 depletion leads to accumulation of phosphatidic acid, which activates the JNK 1/c-Jun pathway to induce transcription of proinflammatory cytokine genes including IL-6, CCL-2/MCP-1, and TNF in macrophage cell lines (28). Activation of several JNK/MAPKs is known to promote osteoclast differentiation (28–34). We hypothesized that the chemokine dysregulation and/or osteoclast differentiation may be mediated by up-regulation of JNK/MAPK pathways and screened for differences in site-specific phosphorylation in 43 kinases in M2-MDMs. Of those screened, 9 were differentially phosphorylated (Figure 6B). Site-specific phosphorylation was increased in JNK1/2/3, MSK1/2, FAK, PRAS40, mechanistic target of rapamycin (mTOR), and STAT5b in the M2-MDMs from the patient with Majeed syndrome compared to healthy controls. Phosphorylation was unchanged in the Src family kinase member Hck and the multifunctional Ser/Thr protein kinase glycogen synthase kinase 3α/β. In contrast, c-Src kinase phosphorylation at position p.Y419, which is essential for full activation (35), was markedly reduced in the patient with Majeed syndrome compared to healthy controls. Phosphorylation was also lower in Pyk-2 and WNK1 in the patient compared to healthy controls (Figure 6B).

Overall, our data suggest that disease-causing LPIN2 mutations alter bone homeostasis by modulating phosphokinase activity in M2-MDMs, thus accelerating osteoclastogenesis. The proinflammatory effect of the LPIN2 mutation on M2-MDMs and osteoclasts was inhibited by IL-1 blockade with anakinra.

**DISCUSSION**

Majeed syndrome is caused by loss-of-function mutations in LPIN2, and disease-causing mutations are recessive and include missense mutations at positions p.S734L or p.R736H or truncating mutations at positions p.C181*, p.S439Wfs*, p.R564Kfs*3, p.Y747*, and p.R776Sfs*66 (Supplementary Table 2, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract). This is the first report of an American patient with Majeed syndrome who is compound heterozygous, exhibiting a previously undescribed 17.8-kb deletion of exons...
7–18 and a novel missense mutation at position 517 that affects a splice site that alternatively splices out exon 10, or exons 10 and 11. The 17.8-kb deletion may be missed with targeted and whole-exome sequencing techniques. Treatment with IL-1 blocking agents reverts the inflammatory bone disease and normalizes the acute-phase reactants, emphasizing the need for a rapid diagnosis.

While IL-1–mediated autoinflammatory diseases including NOMID and DIRA share significant systemic inflammation with granulocytosis and elevated acute-phase reactant levels, only patients with DIRA or Majeed syndrome (but not cryopyrinopathies such as NOMID) develop osteomyelitis (Supplementary Figure 1, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract). The inflammatory bone lesions in patients with Majeed syndrome primarily localize to the growth plates of long bones, which prompted the evaluation of the role of disease-causing LPIN2 mutations in molecular mechanisms that regulate bone homeostasis, specifically the polarization of regulatory M2-DMCs. Consistent with other IL-1–mediated autoinflammatory diseases, Majeed syndrome M1-DMCs secreted more IL-1β compared to M1-DMCs from healthy controls but less than NOMID M1-DMCs.

In contrast, antiinflammatory or regulatory M2-DMCs from the patient with Majeed syndrome secreted high levels of proinflammatory cytokine IL-6 and chemokines IL-8, CCL2/MCP-1, CXCL1/GROα, CCL5/RANTES, and CXCL10. Many were also increased in DIRA patients, but not in NOMID and healthy control M2-DMCs, suggesting that the inflammatory M2-DMC phenotype cannot be attributed to inflammasome activation. In fact, IL-8 (36), CCL2/MCP-1 (37,38), MIP-1α/β (39), and CCL5/RANTES (21), which were elevated in LPS- and ATP-stimulated Majeed syndrome M2-DMCs, accelerate osteoclast differentiation and have been associated with the progression of metastatic bone lesions. These osteoclastogenic factors may act in a paracrine manner, promoting osteoclastogenesis at the growth plate. CCL2/MCP-1 recruits monocytes that differentiate into tissue macrophages, and IL-8 and CXCL1, both prominent neutrophil-attracting chemokines, may orchestrate the recruitment of neutrophils into the growth plate, which is the hallmark of sterile osteomyelitis and is absent in conditions of exclusive osteoclast activation (such as osteoporosis).

Osteoclastogenesis was enhanced in the patient with Majeed syndrome, independently of the aforementioned cytokines and chemokines. The enhanced osteoclastogenesis in this patient and those with DIRA was significantly reduced by IL-1 blockade with anakinra. Mechanisms by which IL-1 blockade controls bone inflammation remain incompletely understood. However, the
increase in RANKL and in the induction of NFATc1, a master regulator of osteoclastogenesis, as well as the increased surface distribution of IL-1R1, suggest a role of intracellular signaling pathways that accelerate osteoclastogenesis independently of autocrine production of osteoclastic factors. These factors also illustrate a complex role of lipin 2 deficiency in regulating bone inflammation that includes the induction of RANK in M2-MDMs. This was consistent with the relatively small effect of culture supernatant from the patient with Majeed syndrome on enhancing osteoclastogenesis in vitro (data not shown).

Lipin 2 has emerged as a metabolic checkpoint that links fatty acid metabolism to proinflammatory signaling in macrophages (28), by controlling the cellular concentrations of DAG and phosphatic acid in saturated fatty acid–overloaded macrophages (40–42). In models of Lipin2 deficiency, the reduced phosphatic acid phosphatase activity results in decreased DAG production and phosphatic acid accumulation, which can activate ERK/ JNK signaling pathways (13). Consistent with these findings, JNK and MSK1/2 phosphorylation, which promote proinflammatory chemokine/cytokine production in adaptive and innate immune cells (43,44), was increased in the M2-MDMs from the patient with Majeed syndrome. In contrast, the profound decrease in Src kinase phosphorylation at p.Y419 in the patient’s M2-MDMs had not previously been implicated with lipin 2 deficiency. Src activation is essential for regulating the antinflammatory property of M2-MDMs and for promoting M2-MDM polarization by increasing IL-10 production (45) and by promoting the production of IL-4–induced STAT6 and Jak1-dependent arginase 1 (46). The low IL-10 secretion and CD163 expression in Majeed syndrome M2-MDMs, which were not corrected by IL-1 inhibition (data not shown), are likely caused by the profound suppression of Src phosphorylation (47) and are consistent with low levels of Pyk-2 phosphorylation (48,49) also observed in the patient with Majeed syndrome.

Src is also a central regulator of osteoclast resorption (50–53), but we did not assess Src phosphorylation in mature osteoclasts. In contrast, FAK, PRAS40, mTOR, and STAT5b kinases (which play a role in cell spreading, migration, and survival in tumor cells [47,54–59], in regulating mTORC1-dependent polarization of macrophages [60,61], and in osteoclastogenesis [62]) were increased in the M2-MDMs from the patient with Majeed syndrome. Although we did not assess the effect of IL-1 blockade on modulating kinase phosphorylation, the significant suppression of proinflammatory cytokine production and bone differentiation factors MIF and serpin/PAI with IL-1 blockade is likely regulated through modulation of phosphokinases (63). Our findings add to recently published data on a novel genetic cause of multifocal osteomyelitis that illustrate the relevance of another Src kinase, Fgr, in promoting osteoclastogenesis (64).

Bone-resident tissue macrophages recently referred to as “osteomacs” (65) have been proposed to have a pivotal role in regulating bone homeostasis. Osteomacs form canopy-like structures at bone remodeling sites, are CD68-positive, and TRAP-negative (66,67). Their distinctive stellate and spindle-shaped morphology makes it intriguing to speculate that Lpin2 mutations influencing the polarization of the spindle-like M2-MDMs in our in vitro culture system may regulate the inflammatory polarization of osteomacs in vivo, thus regulating bone inflammation in patients with Majeed syndrome. Since bone biopsies were not available and Lpin2−/− murine models do not develop osteomyelitis, this hypothesis has not been further investigated.

Limitations of our study include the evaluation of only 1 patient with Majeed syndrome; however, the inclusion of mono- genic controls, particularly NOMID and DIRA patients, and normal controls demonstrates disease-specific clustering. The absence of osteomyelitis in Lpin2−/− mice further limits the ability to evaluate the bone phenotype.

Taken together, our data suggest that lipin 2 modulates bone homeostasis independently of inflammasome activation by driving osteoclastogenesis and altering phosphorylation of JNK and Src kinases that render regulatory M2-MDMs inflammatory and accelerate osteoclastogenesis. We propose a model for sterile osteomyelitis that illustrates the important role of phosphokinases in modulating the inflammatory properties of M2-MDMs and osteoclastogenesis (Supplementary Figure 12, http://onlinelibrary.wiley.com/doi/10.1002/art.41624/abstract). Our data may assist in characterizing pathways and identifying genetic causes for clinically or genetically uncharacterized conditions presenting with chronic recurrent multifocal osteomyelitis and suggest additional targets for treatment.

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AUTHOR CONTRIBUTIONS

All authors were involved in drafting the article or revising it critically for important intellectual content, and all authors approved the final version to be published. Dr. Goldbach-Mansky had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

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Acquisition of data. Bhuyan, de Jesus, Mitchell, Leikina, VanTries, Herzog, Onel, Oler, Montalegre Sanchez, Johnson, Bichell, Marrero, Fernandez de Castro, Huang, Ganesan.

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