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Macrophage-derived IL-1 β enhances monosodium urate crystal-triggered NET formation

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Abstract

Objective and design—Arthritic gout is caused by joint inflammation triggered by the damaging effects of monosodium uric acid (MSU) crystal accumulation in the synovial space. Neutrophils play a major role in mediating joint inflammation in gout. Along with neutrophils, other immune cells such as macrophages are present in inflamed joints and contribute to gout pathogenesis. Neutrophils form neutrophil extracellular traps (NETs) in response to MSU crystals. In presence of MSU crystals, macrophages release IL-1 β , a cytokine crucial to initiate gout pathogenesis and neutrophil recruitment. Our research investigated interactions between human macrophages and neutrophils in an *in vitro* model system and asked how macrophages affect NET formation stimulated by MSU crystals.

Materials or subjects—Human neutrophils and PBMCs were isolated from peripheral blood of healthy volunteers. PBMCs were differentiated into macrophages *in vitro* using human M-CSF.

Treatment—Human neutrophils were pretreated with macrophage-conditioned media, neutrophil-conditioned media or recombinant human IL-1 β prior to stimulation by MSU crystals.

Method—Interaction of neutrophils with MSU crystals was evaluated by live imaging using confocal microscopy. The presence of myeloperoxidase (MPO) and neutrophil elastase (NE) was measured by ELISA. NET formation was quantitated by Sytox Orange-based extracellular DNA release assay and NE-DNA ELISA. AggNET formation was assessed by macroscopic evaluation.

Results—We found that crystal- and cell-free supernatants of macrophages stimulated with MSU crystals promote MSU crystal-stimulated NET formation in human neutrophils. This observation was confirmed by additional assays measuring the release of MPO, NE and the enzymatic activity of NE. MSU crystal-induced NET formation remained unchanged when neutrophil supernatants were tested. IL-1 β is a crucial cytokine orchestrating the onset of inflammation in gout and is known to be released in large amounts from macrophages following MSU crystal stimulation. We found that recombinant IL-1 β strongly promoted MSU crystal-induced NET formation in human neutrophils. Interestingly, IL-1 β alone did not induce any NET release. We also found that clinical grade anakinra, an IL-1 receptor blocker, strongly reduced the NETosis-enhancing effect of macrophage supernatants indicating that IL-1 β is mainly responsible for this effect.

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Conclusions—Macrophage-derived IL-1 β enhances MSU crystal-induced NET release in neutrophils. We identified a new mechanism by which macrophages and IL-1 β affect neutrophil functions and could contribute to the inflammatory conditions present in gout. Our results also revealed a new anti-inflammatory mechanism of anakinra.

1. Introduction

Gout is a sterile inflammation of the joints mediated mainly by infiltrated macrophages and polymorphonuclear neutrophilic granulocytes (PMN) driven by monosodium urate (MSU) crystal deposition [1]. These innate immune cells mistakenly recognize MSU crystals as dangers and undergo unnecessary activation resulting in painful, tissue-damaging inflammation [2]. While macrophages are the initiators of the inflammatory cascade, PMN infiltration and activation play a major role in bringing inflammation to its full expansion in gout [3–6]. IL-1 β is a crucial and potent proinflammatory cytokine which also orchestrates gouty inflammation [7–9]. IL-1 β is abundant in the synovial fluid of gout patients [3]. Its central role in gout inflammation is shown by successful IL-1 β -inhibitory therapies [9–11]. Anakinra, a recombinant IL-1Ra protein, rilonacept, an IL-1 receptor fusion protein and canakinumab, an anti-IL-1 β antibody have all been shown to successfully treat acute gout attacks [9]. The primary role of IL-1 β in mediating the inflammatory cascade in gout is thought to be its potent effect to recruit PMNs to the synovial space [9, 12]. MSU crystals activate the NLRP3 inflammasome and induce release of IL-1 β in macrophages [7]. We and others have shown that MSU crystal-induced NLRP3 inflammasome activation in macrophages requires calcium and NADPH oxidase-independent production of reactive oxygen species (ROS) [13–15]. Caspase-1 activated by the inflammasome cleaves pro-IL-18 and pro-IL-1 β forming bioactive cytokines [16, 17].

PMNs are the most abundant leukocytes during gout inflammation reaching very high concentrations in the synovial fluid [5, 18]. PMNs are known to engulf MSU crystals triggering ROS production, cytokine release and neutrophil extracellular trap (NET) formation [5, 6, 18–20]. In a previous study, we described robust NET release of human PMNs in response to calcium pyrophosphate dihydrate (CPPD) microcrystals, the causative agents of pseudogout, a condition similar to gout [21]. NETs are composed of extracellular DNA network associated with histones and PMN granule proteins including myeloperoxidase (MPO) and neutrophil elastase (NE) [22]. NET formation in gout is mistakenly initiated and leads to the release of dangerous PMN cargo into the synovium. In gout, the primary antimicrobial function of NETs (trapping extracellular microbes) manifests in MSU crystal immobilization [5, 6]. MSU crystal-induced NET release requires autophagy, NADPH oxidase-dependent ROS production and RIP3K-MLKL signaling pathway [5, 18, 23]. A recent study shed new light on a potential anti-inflammatory role of NETs in gout [5]. At high PMN densities, NETosed PMNs and MSU crystals form ‘aggregated NETs’ (aggNETs), the proposed basis of gouty tophi, a characteristic white material that appears at the beginning of the resolution phase of acute gout flares [5, 24]. Although our view on the role of NETs in gout pathogenesis is improving, less is known how the two most important cell types initiating and mediating inflammation in gout, macrophages and PMNs, interact.

Our study addressed the question how macrophages influence NET formation stimulated by gout-causing MSU crystals. Our data show that proinflammatory macrophages stimulated with MSU crystals release mediators that cause neutrophils to have over-exaggerated release of NETs induced by the crystals. We also show that IL-1 β is a potent enhancer of NET formation and it is mainly responsible for the NETosis-promoting effect of macrophages. We propose that macrophage- and IL-1 β -enhanced NET formation can be a novel contributor to gout inflammation.

Materials and Methods

2.1. Human subjects

De-identified healthy human volunteers were recruited at the University of Georgia to donate blood. The studies were performed according to the guidelines of the World Medical Association's Declaration of Helsinki. Enrolled blood donors signed consent forms as described previously [25, 26]. The human blood protocol (UGA# 2012-10769) and the associated consent form were reviewed and approved by the Institutional Review Board (IRB) of the University of Georgia.

2.2. Human neutrophils and autologous serum

Human neutrophils were isolated as described previously [21, 26]. Briefly, coagulation was prevented with heparin and red blood cells were removed by Dextran sedimentation (GE Healthcare). Neutrophils were isolated using 5-step Percoll gradient centrifugation. Cell viability was >98% (Trypan Blue) and neutrophil purity was > 95% (cytospin and flow cytometry). Autologous serum was prepared by centrifugation and sterile filtration. Calcium- and magnesium-containing HBSS (Mediatech, Manassas, VA, USA) supplemented with 1% autologous serum, 5 mM glucose and 10 mM HEPES was used as assay buffer.

2.3. PBMC isolation and in vitro macrophage differentiation

PBMCs were isolated from 65% Percoll gradient interphase, washed twice with PBS and seeded at 2–3 million cells/well concentration in RPMI medium supplemented with Glutamax (Gibco Thermo Fischer, Waltham, MA). The next day the media was supplemented with 50 ng/ml M-CSF and 1% autologous serum or FBS and macrophages were differentiated for 5–7 days into M1 phenotype as described previously [13].

2.4. Macrophage supernatant collection (Macrophage-conditioned medium)

Differentiated macrophage cultures were primed with 10 μ g/ml LPS for 30 mins to mimic proinflammatory conditions found in the gout synovium. Macrophages were then washed 4X HBSS to remove LPS and were subsequently incubated with or without 250 μ g/ml MSU crystals for 5 hours. Supernatants were collected and then spun at 10,000g for 5 mins. The pellet was discarded, cell-, crystal- and debris-free supernatants were frozen at –80 °C and called the 'macrophage-conditioned medium'.

2.5. Extracellular DNA release assay

NETs were quantified essentially as described previously [21, 27]. Briefly, PMNs were allowed to adhere to poly D-lysine coated 96-well black transparent bottom plates (Thermo Scientific, Rochester, NY) in assay buffer with 0.2% Sytox Orange (Life Technologies, Grand Island, NY, USA). Fluorescence (excitation: 530 nm, emission: 590 nm) was measured in a fluorescence microplate reader (Varioskan Ascent, Thermo Scientific) for 6 hours at 37 °C. Increase in fluorescence normalized on maximal value (saponin-treated PMNs) was referred to as “extracellular DNA release” and expressed as “% of max”. The obtained relative fluorescence units (RFU) were sometimes also normalized against PMA.

2.6. Confocal microscopy of NETs

As previously detailed [21, 26], 2×10^6 human neutrophils in assay medium were incubated for 4 hrs at 37 °C with or without 100 nM PMA and 100 µg/ml MSU crystals in 35 mm glass bottom dishes (MatTek, Ashland, MA) pre-coated for 1 hr with 1% human serum albumin (Sigma Aldrich, St Louis, MO, USA). 2.5 µM Sytox Orange (Invitrogen, Grand Island, NY, USA) was added prior to fluorescence imaging (exc/em: 547/570 nm). Images were collected using a Nikon A1R confocal microscope system equipped with a Nikon Eclipse Ti-E inverted microscope, built in Perfect Focus, high-speed motorization and NIS Elements software (Nikon Instruments, Melville NY). Live cell imaging was carried out using a Tokai Hit INY-G2A-TIZ incubator (Tokai Hit CO, Ltd, Shizuoka-ken with Japan) for temperature, humidity and CO₂ control. The Coherent Sapphire 561nm 20mW laser was used to excite Sytox Orange using a CFI Plan APO VC 60X Oil NA 1.4 WD 0.13 mm objective. Optical sections of neutrophils were analyzed, Z-stacks spanning 1 µm were acquired and 3-D image reconstructions were processed using the Nikon NIS Element Version 4 software. Final image preparation was carried out using Adobe Photoshop.

2.7. High-throughput live imaging of NET formation

A microplate-based NET imaging assay was established using a motorized stage support of a confocal microscope. PMNs were seeded at a concentration of 250,000 cells/well in a 96 well plate Optical Btm Pit Polymer Base black plate (Thermoscientific, Rochester, NY). PMNs were preincubated with 10 ng/ml IL-1β prior to imaging. PMNs were then stimulated with 200 µg/ml MSU crystals for 16 hours. Nikon A1R confocal microscope system with a 10X lens was used to capture transmitted light and fluorescence images. A field of interest was chosen to perform imaging and their position was defined using the NIS elements software. The chosen field was a true representation of the entire well. Images were taken every 15 min for 12 hours using automated capture component of the NIS elements software. Mean Sytox Orange intensities of the fluorescent images were quantified using the measure region of interest (ROI) feature of the Nikon A1 Elements software. The entire image which represented the field was selected as the ROI. These measurements were used to calculate changes in SYTOX orange intensities over time (Mean SYTOX orange intensity). Samples were always run as three biological replicates.

2.8. Measurement of AggNET formation

Visualization of aggregated NET formation was modified after adaptation [5]. 1,000,000 PMNs in assay medium were added in a 48-well plate with or without 1mg/ml MSU crystals at 37 °C with or without 10 ng/ml IL-1 β . Images of aggNETs were taken after 5 hr incubation using an inverted light microscope [5].

2.9. ELISA

Myeloperoxidase (MPO), IL-1 β or IL-8 levels were quantitated using commercial ELISA kits from R&D systems (Minneapolis, MN), Millipore and BioLegends, respectively. Manufacturer's instructions were followed for sample dilutions and processing. 250,000 PMNs/well were seeded in 24-well plates and stimulated with 250 μ g/ml MSU crystals in HBSS for up to 4 hours at 37 C. Cell supernatants collected were either immediately processed or stored (–80°C) for later analysis. Undiluted supernatants were used to measure IL-8 and IL-1 β , whereas supernatants were diluted 1:100 to determine MPO and HNE concentrations. HNE release was assessed as described [25, 26] by ELISA using anti-HNE rabbit polyclonal antibody (Merck Millipore, Billerica, MA). Supernatant samples diluted with coating buffer (25 mM carbonate, 25 mM bicarbonate, pH 9.6) were incubated overnight at 4°C in 96-well high-binding microtiter ELISA plates (Greiner bio-one, Frickenhausen, Germany). After blocking with 1% BSA for 1 hour, anti-rabbit mouse polyclonal Ab (1:500 in PBS [Calbiochem], 481001 [Merck Millipore]) was added for 2 hours at RT. After repeated washes, samples were incubated with horseradish peroxidase–linked donkey anti-rabbit Ab (1:2000 in PBS, NA934V; GE Healthcare) for 1 h. Yellow coloration of the TMB substrate (Thermo Scientific, Rockford, IL) after the addition of 1N hydrochloric acid (Sigma) was read at 450-nm wavelength with Eon microplate photometer (BioTek, Winooski, VT). Commercially available HNE (stock: 1 mg/ml; Cell Sciences, Canton, MA) was used as standard.

2.10. HNE-DNA ELISA

NETs were quantitated as described previously [25, 27]. Briefly, PMN supernatants were exposed to DNase I digestion for 15 min that was stopped by adding 2.5 mM EGTA. Diluted supernatants and “NET standard samples” were added to 96-well high binding microtiter ELISA plates (Greiner bio-one, Frickenhausen, Germany) coated anti-HNE rabbit antibody. Horse radish peroxidase-labelled anti-DNA secondary antibody (1:500 in PBS) was added for 2 hours followed by repeated washes. Absorbance of the TMB substrate was read after addition of 1N hydrochloric acid (Sigma) at 450 nm with Eon microplate photometer (BioTek, Winooski, VT). NET's were quantitated as percentage of the NET-standard in the undiluted supernatants.

2.11. Quantification of enzymatic activity

To measure HNE activity, the Neutrophil Elastase Activity Assay Kit (Cayman Chemical Ann Arbor, MI, USA) was used following manufacturer's protocol. Briefly, supernatants (50 μ l) were placed into 96-well black plates. Substrate (Z-Ala-Ala-Ala-Ala 2Rh110, cat no. 600613) solution was added to assess elastase activity by measuring production of the highly fluorescent product (compound R110) using 485 nm excitation and 525 nm emission

wavelengths in microplate fluorimeter (Varioskan Flash, Thermo Scientific). Data are expressed either as kinetics (RFU) or end-point values normalized on maximal (PMA-stimulated) data.

2.12. Statistical analysis

Data are represented as mean $\pm$ S.E.M. P value was calculated with ANOVA and Dunnett's or Tukey's post-hoc tests, and was marked as * when $p < 0.05$, ** when $p < 0.01$ and *** when $p < 0.001$

Results

3.1. Secreted neutrophil products do not affect MSU crystal-induced NET release

PMNs release NETs in response to gout-causing MSU crystals [5, 6, 18]. First, we exposed human PMNs to different doses of MSU crystals and measured NET formation using the extracellular DNA-binding dye, Sytox Orange. MSU crystals induced NET release in a dose-dependent manner (Fig. 1A). An MSU crystal concentration of 100 $\mu\text{g/ml}$ was required to trigger a minimal NET release that is higher than the signal without crystals (Fig. 1A). Throughout the manuscript 100 or 250 $\mu\text{g/ml}$ MSU crystal doses were used that are significant and detectable but still small enough to leave room to record any NETosis-enhancing effects.

To assess whether PMNs themselves promote NET formation, we first stimulated human PMNs with MSU crystals and collected supernatants after 5 hours. Cells, debris and crystals were removed by centrifugation resulting in a clean supernatant referred to as "PMN-conditioned medium" that was kept frozen. This conditioned medium was then added to unchallenged PMNs isolated from another donor in presence or absence of MSU crystals and DNA release was followed. As shown in Figure 1B, the PMN-conditioned medium did not affect NET formation indicating that PMNs do not intrinsically accelerate MSU crystal-induced NETs in a positive feedback fashion.

3.2. Macrophages release NETosis enhancing mediators upon MSU crystal stimulation

Since macrophages are present in the gout synovium in large numbers and are responsible for initiating inflammation [2], we repeated the same experiment as before using macrophages, not PMNs, to collect the conditioned medium. When this "macrophage-conditioned medium" was added to human PMNs, MSU crystal-induced DNA release was significantly enhanced (Fig. 1C). Spontaneous NET formation remained unaffected (Fig. 1C). These data were also confirmed by confocal microscopy showing the larger amount of three-dimensional extracellular DNA release in the presence of macrophage-conditioned medium than in its absence (Fig. 1D). These results indicate that MSU crystal-exposed macrophages release secretory products that strongly accelerate NET formation of PMNs stimulated by MSU crystals.

3.3. Macrophage-conditioned medium increases the release of primary granule markers and NETs from MSU crystal-stimulated PMNs

NETs are composed of an extracellular DNA meshwork associated with histones and PMN granule components, such as myeloperoxidase (MPO) and neutrophil elastase (NE) [22]. The definition of NETs requires the detection of DNA-protein complexes characteristic for NETs. First, we confirmed that we can detect MPO ($p=0.0469$) and HNE ($p=0.0694$) by ELISA in the supernatants of human PMNs after MSU crystal stimulation (Fig. 2A–B). Second, we observed that addition of macrophage-conditioned medium significantly enhanced MSU crystal-induced MPO ($p<0.0001$) and HNE ($p=0.0003$) release (Fig. 2A–B). To prove that released MPO and DNA are present in a complex defining NETs, we utilized an ELISA developed in our laboratory capable of quantitating NET-specific MPO-DNA complexes [25, 27]. MSU crystals initiated NET release that was further enhanced by mediators secreted by macrophages (Fig. 2C). Overall, our results presented so far describe a novel mechanism by which NET formation is enhanced in PMNs by macrophages.

3.4. IL-1 β potentiates MSU crystal-induced DNA release from PMNs

As the next step we aimed at identifying which macrophage-secreted component is responsible for the NETosis-enhancing effect. Macrophages have been described to release several cytokines and other inflammatory mediators upon MSU crystal exposure [2]. IL-1 β was our primary suspect potentially responsible for the NET-promoting effect of macrophages because IL-1 β is a central cytokine orchestrating gout inflammation and it is released in copious amounts from MSU crystal-stimulated macrophages [2, 7, 9]. We confirmed that differentiated macrophages produced IL-1 β upon MSU crystal activation in our hands, as well (Fig. 3A). Human PMNs failed to release any detectable IL-1 β (Fig. 3A). We stimulated human PMNs with MSU crystals in presence of recombinant human IL-1 β and measured extracellular DNA release. While IL-1 β had only a minimal NETosis-enhancing effect when 100 $\mu\text{g/ml}$ MSU crystals were used, it strongly amplified NET formation in a dose-dependent manner when 250 $\mu\text{g/ml}$ MSU crystal concentration was applied (Fig. 3B–C). The highest dose of IL-1 β enhanced NET formation to the extent observed with the positive control PMA (Fig. 3B–C). Interestingly, IL-1 β alone (without MSU crystals) did not induce NET release (Fig. 3B–C). Lower doses of MSU crystals were chosen to be used in these experiments that allow detection of any potential, NETosis-enhancing effect of IL-1 β . These results indicate a novel effect of IL-1 β , a central cytokine in gout inflammation, on PMN activation by MSU crystals.

3.5. IL-1 β enhances MPO, HNE and NET release following MSU crystal stimulation

To confirm that the DNA release-promoting effect of IL-1 β is also accompanied by enhanced levels of released granule markers and NETs, we quantitated MPO, HNE, HNE-DNA and HNE enzymatic activity levels in supernatants of MSU crystal-induced PMN in the absence or presence of IL-1 β . We found that IL-1 β significantly promoted the release of MPO and HNE from human PMNs after MSU crystal stimulation (Fig. 4A–B, $p<0.05$ in each combination between untreated and MSU crystal-treated samples). In addition, HNE remained enzymatically active and bound to DNA in PMN supernatants after MSU crystal activation (Fig. 4C–D). These data are in accordance with those presented in Figure 3

indicating that IL-1 β alone does not induce NET formation but it significantly promotes NET release triggered by MSU microcrystals.

3.6. Imaging the NET-promoting effect of IL-1 β by high-throughput confocal live microscopy

Next, we performed high throughput imaging in order to visualize NET formation real-time using confocal microscopy. This method developed in our laboratory enables simultaneous imaging of NET-forming live PMNs on a 96-well-based format in presence of Sytox Orange. Recorded videos allow spatiotemporal resolution of the NET-forming process while kinetic data enable concomitant quantitative comparisons. The recorded videos confirmed previous data. IL-1 β alone did not induce DNA release while MSU crystals triggered NET formation (Fig. 5A–B). Addition of IL-1 β to MSU crystals resulted in more robust NET release than MSU crystals alone (Fig. 5A–B). In addition, video recordings show that PMNs start to aggregate soon after MSU crystal exposure (Fig. 5A). This microaggregate formation is unique to MSU crystals among NET-inducing stimuli and forms the microscopic basis of later, macroscopic aggNET formation.

3.7. IL-1 β enhances aggregated NET formation

MSU crystals induce the formation of aggregated NETs (aggNET) at higher PMN densities that provide the structural basis of gout tophi [5]. Since aggNET formation depends on MSU crystal-induced NET release that is enhanced by IL-1 β , we wanted to study how IL-1 β affects aggNET formation. Human PMNs at higher cell density were exposed to enhanced MSU crystal dose (1mg/ml) modeling peak phases of gout inflammation [5]. As shown in figure 5C, MSU crystals initiated aggNET formation that was further enhanced by IL-1 β , as assessed by light microscopy. Thus, acceleration of MSU crystal-triggered NET release by IL-1 β also leads to faster and more robust aggregated NET formation *in vitro*.

3.8. Anakinra abolishes the NET-promoting effect of macrophages

To prove that IL-1 β is the main component found in macrophage supernatants promoting MSU crystal-induced NET release, we pretreated PMNs with clinical grade anakinra, an IL-1R antagonist, prior to addition of MSU crystals and macrophages supernatants. Supernatants were collected from macrophages obtained from nine independent volunteers (donors #4–12) and were tested in groups of three on PMNs isolated from three, additional blood donors (donors #1–3) (Fig. 6A). Although we experienced donor-to-donor variations (Fig. 6A), anakinra generally efficiently inhibited the NET-enhancing effect MSU crystal-stimulated macrophage supernatants (Fig. 6B). Anakinra treatment resulted in a 76.5+/-6.3% (mean+/-S.E.M., n=12) inhibition of the signal (Fig. 6B). Thus, we described a novel effect of anakinra that could be a significant component of its *in vivo* mechanism of action.

Discussion

In line with previous reports of other groups, we confirmed that gout-causing MSU crystals elicit robust NET formation in human PMNs *in vitro* [5, 18, 23, 28]. Concentrations of MSU crystals higher than 100 μ g/ml induce significant extracellular DNA release in PMNs. The 100–1000 μ g/ml MSU crystal dose range used here represents pathophysiological

concentrations found in the synovial fluid of gout patients [5]. Therefore, the *in vitro* findings described here likely have human *in vivo* clinical relevance.

PMN-conditioned media did not alter MSU crystal-induced NET release indicating that PMNs themselves do not secrete any inflammatory mediators that would promote NET formation in a positive feedback fashion. Macrophages, however, were found to significantly enhance NET release following MSU crystal exposure in PMNs. The interaction between PMNs and macrophages is an essential component of gout inflammation in joints [16, 29]. It is known that resident macrophages at the synovium represent the first cell type to come in contact with MSU microcrystals and release several cytokines that attract additional macrophages and PMNs to the site of inflammation [16, 29]. Previous reports described how macrophages react to already formed NETs [30–32] but their NETosis-enhancing effect described here is a new finding that adds to the complex interplay between macrophages and PMNs.

Results shown here suggest that IL-1 β is the major macrophage-derived molecule promoting NET formation. These results complement previously published data showing that gout synovial fluid supernatants promote spontaneous NET release in human PMNs that is partially inhibited by anakinra [18]. MSU crystal stimulation of macrophages leads to NLRP3 inflammasome activation and production of the pro-inflammatory cytokine IL-1 β [1, 7]. IL-1 β initiates PMN chemotaxis that is thought to be the main mediator of PMN influx to the synovium in gout [7]. Accumulating data suggest more complex mechanisms by which macrophages interact with PMNs in gout indicating that macrophage-derived IL-1 β not only attracts PMNs to the site of inflammation but it also accelerates the release of PMN-derived inflammatory mediators including NETs. Not only caspase-1 but neutrophil elastase can also cleave pro-IL-1 β producing bioactive IL-1 β [3, 33]. This mechanism would accelerate inflammation since IL-1 β would enhance its own activation by activating PMNs to release NET-related proteases. Data shown here also propose that targeting IL-1 β in gout *in vivo* fulfills its beneficiary effects not only through diminished PMN recruitment but also via attenuated PMN activation by MSU crystals.

Interestingly, IL-1 β alone does not induce NETs in our hands but it enhances NET release induced by MSU crystals. Whether this observation is unique for MSU crystals or it is also true for other NET-triggering stimuli such as bacteria, fungi or other microcrystals, remains to be investigated. Several potential mechanisms could underlie the NETosis-enhancing effect of IL-1 β . IL-1 β could increase NET formation by its known beneficiary effect on the activation of the NADPH oxidase [34, 35]. The NADPH oxidase is required for MSU crystal-induced NET formation [5]. IL-1 β could enhance the phosphorylation of the oxidase subunits or their expression. This mechanism is likely because human PMNs of IRAK4- or NEMO-deficient patients with impaired TLR/IL-1R signaling have reduced NADPH oxidase priming and activation [36]. IL-1 β is a known chemoattractant for PMNs [37]. Although it remains to be studied, a migratory component of MSU crystal-induced NET/aggNET formation is likely that could be promoted by IL-1 β .

At the same time, IL-1 β also promotes aggNET formation, a novel mechanism proposed to shut down acute inflammation in gout [5]. The highly concentrated protease activity of

aggNETs degrades several proinflammatory cytokines including IL-1 β putting a break on the inflammatory process [5]. Therefore our results implicate that promoting the aggNET-promoting mechanism of IL-1 β offers a novel anti-inflammatory strategy and could have therapeutic benefits.

Although IL-1 β is the major proinflammatory molecule released by macrophages promoting gout inflammation, other secretory products could also promote NET formation. IL-8, C5a, G-CSF, IFN- γ , and TNF α have all been reported to promote NET formation [5, 21, 38–40]. G-CSF is indispensable for neutrophil functions [41] and assists in the NETosis process [16, 28, 42]. TNF- α is released during the acute phase of gout and can trigger a NET response [16, 21, 28, 39, 42]. The potential contributions of these inflammatory mediators to the NETosis-enhancing effect of macrophages remain to be studied in the future.

In summary, our data presented here provide a new link between macrophages and PMNs, two important innate immune cell types mediating the general and pathological inflammatory pathways. IL-1 β is a major player in gout and NET formation promoted by IL-1 β could contribute to gouty inflammation.

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Abbreviations

NETs	Neutrophil Extracellular Traps
AggNET	aggregated Neutrophil Extracellular Traps
PMN	polymorphonuclear neutrophils
IL-1β	Interleukin-1 beta
PBMC	Peripheral Blood Mononuclear cells
NLRP3	NACHT, LRR and PYD domains-containing protein 3
ROS	reactive oxygen species
MSU	monosodium urate
MPO	myeloperoxidase
NE	neutrophil elastase
HNE	human neutrophil elastase

M-CSF macrophage colony stimulating factor

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Figure 1. Supernatants of macrophages, but not neutrophils, enhance extracellular DNA release from MSU crystal-stimulated human neutrophils

A) Dose-dependence of MSU crystal-stimulated NET release in human PMNs measured in the presence of the fluorescent, extracellular DNA-binding dye, Sytox Orange. Mean $\pm$ /–S.E.M., $n=3$. **B–D)** Human macrophages differentiated with M-CSF for 5–7 days or neutrophils were exposed to 250 μ g/ml MSU crystals for 5 hours. Collected supernatants were centrifuged to remove cells, crystals and debris and referred to as “PMN-conditioned medium” or “macrophage-conditioned medium”, respectively. The effects of conditioned media on NET formation were subsequently tested on human neutrophils stimulated in the absence (UT, untreated) or presence (MSU) of 250 μ g/ml MSU crystals by Sytox Orange-based fluorescence. Fluorescence data were normalized on the 100 nM PMA-triggered signal as the positive control for NET formation. **B)** “PMN-conditioned media” of neither MSU crystal-stimulated (MSU PMN) nor untreated neutrophils (UT PMN) had any significant effect on NET release (No Sup). Mean $\pm$ /–S.E.M., $n=4$, Dunnett’s test. **C)** “Macrophage-conditioned media” of MSU crystal-stimulated macrophages (MSU MAC) significantly increased NET formation in comparison to that seen in absence of conditioned media (No Sup). Conditioned media of macrophages without crystal stimulation (UT MAC, untreated) had no significant effect. Mean $\pm$ /–S.E.M., $n=11$, Dunnett’s test. **D)** Representative confocal microscopy images of MSU crystal-stimulated human neutrophils show enhanced DNA release (Sytox Orange) by the MSU crystal-stimulated macrophage conditioned media compared (lower panels) to that without addition of conditioned media (upper panels) ($n=3$). 1) Merged images of transmitted light and Sytox Orange fluorescence and 2) three-dimensional Z-stack fluorescent images are shown for both experimental conditions. Black bars indicate 10 μ m distance. RFU, relative fluorescence unit; PMA, phorbol myristate acetate; MAC, macrophage; PMN, neutrophil; MSU, monosodium urate; n.s., not significant; ****, $p<0.0001$.

Figure 2. Macrophage supernatants increase MPO, HNE and NET release from human neutrophils following MSU crystal stimulation

Differentiated human macrophages were exposed to 250 µg/ml MSU crystals for 5 hours to collect and centrifuge their supernatants. This “macrophage-conditioned medium” (MSU MAC) was added to MSU crystal-stimulated human neutrophils and releases of **A**) MPO (n=9), **B**) HNE (n=9) and **C**) NETs (MPO-DNA, n=6) were quantitated by ELISA assays (mean+/-S.E.M., Dunnett’s test). As comparisons, PMNs were also treated with mock only (No MAC) or supernatants of macrophages without crystal stimulation (UT MAC, untreated). MPO, myeloperoxidase; HNE, human neutrophil elastase; MAC, macrophage. *, p<0.05; **, p<0.01; ***, p<0.001.

Figure 3. IL-1 β promotes DNA release from human neutrophils stimulated with MSU crystals
A) MSU crystal-stimulation resulted in IL-1 β release in human macrophages (MF) but not in human PMNs as measured by ELISA. Mean $\pm$ S.E.M., n=4. Student's t-test. **B)** Following treatment with increasing doses of recombinant human IL-1 β , human neutrophils were left unstimulated or were stimulated with 100 or 250 μ g/ml MSU crystals and DNA release was measured in presence of the membrane-impermeable DNA-binding dye, Sytox Orange. DNA release over 4 hours was presented as endpoint values (mean $\pm$ S.E.M., n=3). **C)** Kinetics of the previous experiment (panel B) (one representative, n=4). Fluorescence data were normalized on the maximal signal obtained by stimulation with 100 nM PMA. RFU, relative fluorescence unit; MSU, monosodium urate; UT, untreated; PMA, phorbol myristate acetate; n.s., not significant; *, p<0.05; ***, p<0.001; ****, p<0.0001.

Figure 4. IL-1 β enhances MSU crystal-stimulated releases of MPO, HNE and NETs from human neutrophils

Human neutrophils were treated with recombinant human IL-1 β (0–5 ng/ml) prior to addition of 250 μ g/ml MSU crystals. Neutrophils were also left unstimulated (UT, no crystals). Five hours later the following parameters were quantitated in the supernatants: **A**) MPO by commercial ELISA (n=5), **B**) HNE by ELISA (n=9), **C**) NETs (HNE-DNA ELISA, n=7) and **D**) HNE activity by commercial kit (n=7). NET release values were normalized on PMA (100%). Data are shown as mean $\pm$ S.E.M. Significance was calculated using Dunnett's test. *, p<0.05; **, p<0.01. MPO, myeloperoxidase; HNE, human neutrophil elastase; PMA, phorbol myristate acetate.

Figure 5. IL-1 β increases MSU crystal-induced aggregated NET formation

250,000 human PMNs/well were incubated in wells of a 96-well microplate with IL-1 β (10 ng/ml) and stimulated with 100 μ g/ml MSU crystals or 100nM PMA for 15 hours in presence of the extracellular DNA-binding dye, Sytox Orange. Interactions of PMNs and crystals were followed using confocal microscopy every 15 minutes. **A)** Representative images taken at t=0, 2, 4, 6. hrs are shown (n=3). Black bars indicate 100 μ m distances. Insets indicated by white frames are enlarged on the right side. **B)** Representative kinetics of Sytox Orange fluorescence curves followed for 8 hours (n=3, mean $\pm$ S.D. at each time point). **C)** AggNET formation was evaluated by light microscopy after a 5-hour incubation of human PMNs with 1 mg/ml MSU crystal in presence or absence of 10 ng/ml human IL-1 β (One representative image, n=3). MSU, monosodium urate; aggNET, aggregated NETs; RFU, relative fluorescence unit.

Figure 6. Anakinra inhibits the NET-enhancing effect of macrophage supernatants

A) Macrophages were isolated from nine independent donors (donors #4–12) and macrophage-conditioned media were collected when cells were left untreated (UT, no crystals) or stimulated with 250 mg/ml MSU crystals for 5 hours (MSU). Supernatants were cleaned, frozen and used in groups of three on PMNs isolated from three independent blood donors as indicated. PMNs were pretreated with 30 mg/ml anakinra for 1 hr before addition of 100 mg/ml MSU crystals and/or macrophage supernatants. EcDNA release was measured in PMNs for 6 hours following stimulation by the crystals using Sytox Orange dye. Results are expressed as percentage of maximal DNA release and shown as mean $\pm$ S.D. of triplicates. B) Data shown in panel A) were further analyzed by calculating the MSU crystal-induced DNA release signal and its averages. Data are shown as mean $\pm$ S.E.M. (n=12) and were analyzed by one-way ANOVA and Tukey's post-hoc test. *, p<0.05; ***, p<0.001. MAC, macrophage; PMN, neutrophil; UT, untreated.