Neutrophil Inflammatory Response Is Downregulated by Uptake of Superparamagnetic Iron Oxide Nanoparticle Therapeutics

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Superparamagnetic iron oxide nanoparticles (SPION) are employed as diagnostics and therapeutics following intravenous delivery for the treatment of iron deficiency anemia (IDA) in adult patients with chronic kidney failure. Neutrophils are the first defense against blood borne foreign insult and recruit to vascular sites of inflammation via a sequential process that is characterized by adhesive capture, rolling, and shear resistant arrest. A primary chemotactic agonist presented on the glycocalyx of inflamed endothelium is IL-8, which upon binding to its cognate membrane receptor (CXCR1/2) activates a suite of responses in neutrophils. An early response is degranulation with accompanying upregulation of β2-integrin (CD11/CD18) and shedding of L-selectin (CD62L) receptors, which exert differential effects on the efficiency of endothelial recruitment. Feraheme is an FDA approved SPION treatment for IDA, but its effect on the innate immune response of neutrophils during inflammation has not been reported. Here, we studied the immunomodulatory effects of Feraheme on neutrophils freshly isolated from healthy human subjects and stimulated in suspension or on inflammatory mimetic substrates with IL-8. Cells treated with Feraheme exhibited reduced sensitivity to stimulation with IL-8, indicated by reduced upregulation of membrane CD11b/CD18 receptors, high affinity (HA) CD18, and shedding of CD62L. Feraheme also inhibited N-formyl-Met-Leu-Phe (fMLP) induced reactive oxygen species production. Neutrophil rolling, arrest, and migration was assessed in vascular mimetic microfluidic channels coated with E-selectin and ICAM-1 to simulate inflamed endothelium. Neutrophils exposed to Feraheme rolled faster on E-selectin and arrested less frequently on ICAM-1, in a manner dependent upon SPION concentration. Subsequent neutrophil shape change, and migration were also significantly inhibited in the presence of Feraheme. Lastly, Feraheme accelerated clearance of cytosolic calcium flux following IL-8 stimulation. We conclude that uptake of Feraheme by neutrophils inhibits chemotactic activation and downregulates normal rolling to arrest under shear flow. The mechanism involves increased calcium clearance following chemotactic activation, which may diminish the efficiency of recruitment from the circulation at vascular sites of inflammation.

Keywords: neutrophil recruitment, nanoparticle, iron oxide, inflammation, neutrophil degranulation, mechanosignaling, immunosuppression, innate immunity
INTRODUCTION

Superparamagnetic iron oxide nanoparticles (SPION) are used as contrast agents for magnetic resonance imaging (MRI) in the liver, central nervous system, gastrointestinal system, and in macrophages (1). SPION have also found clinical use as an FDA approved treatment of cancer (i.e., NanoTherm) and iron deficiency anemia (IDA) in patients with chronic kidney disease (CKD) (i.e., Feraheme) (1). While SPION have been used for various clinical applications, a primary consideration for any biomaterial based therapeutic is the host immune response elicited by the interaction of the biomaterial with blood cells. The increased use of SPION has raised concerns over their safety, with several formulations showing cytotoxic and immunotoxic effects (2). This motivated the current study to examine the systemic effects of Feraheme on neutrophils in human blood.

The innate immune system is by design able to recognize and eliminate foreign pathogens and particulate matter through pattern recognition receptors and opsonin receptors that sound the alarm upon detection of danger signals (3). Opsonization involves the binding of molecules such as antibodies or proteins of the complement system that enhances recognition and removal via endocytosis and phagocytosis of foreign matter by innate immune cells (4). As nanoparticles enter the blood stream, complement and other proteins immediately adsorb to the particle surface forming a corona (5). Physiochemical properties of nanoparticles have a direct influence on the composition and formation of the protein corona in such a manner to impact particle-cell interactions (6). Induction of immune hypersensitivity reaction in patients prompted FDA to cancel clinical use of the SPION Feridex and GastroMARK (7, 8). Further, studies have shown that SPION can induce oxidative stress and cell damage in vitro and in animal models (9, 10). Upon binding to the plasma membrane and endocytosis, nanoparticles can induce a variety of functional responses in neutrophils such as reactive oxygen species (ROS) and chemokine production that vary with different SPION formulations. While some SPION can induce these immune responses, others exert an inhibitory effect, thereby highlighting composition as a key factor when considering nanoparticle use as a therapeutic (2, 11). Conventional wisdom is that nanoparticle clearance is mediated by macrophages; however, recent studies indicate that neutrophils also play a key role in particle clearance and this process has been exploited to deliver therapeutic nanoparticles to tumors for cancer therapy (12, 13). Current literature on the immune effects of SPION have largely focused on monocytes and macrophages. This has resulted in an incomplete understanding of the full scope of SPION immune effects on neutrophil function (14).

Neutrophils are the most common leukocyte in blood circulation and are essential first responders during inflammation. Thus, intravenous administration of nanoparticles is likely to result in frequent neutrophil-particle interactions. Capture of SPION onto the plasma membrane and subsequent endocytosis can perturb the inflammatory response and recruitment of neutrophils to sites of inflammation. Dysregulation of neutrophil activation and recruitment is implicated in various autoimmune and inflammatory diseases (15). Failure of neutrophils to normally adhere to the endothelium and become activated can cause severe impairment of host defense against pathogens, while inappropriate levels of cell recruitment and activation leads to chronic inflammation and tissue damage (16, 17). There are reports of the capacity for coated (Polycrylic acid) and non-coated iron oxide nanoparticles and metal oxide nanoparticles TiO2, CeO2, and ZnO, to induce ROS production and activate neutrophil degranulation (18, 19). Excessive release of neutrophil granules, which contain chemokine and integrin receptors, elastase, collagenase, and myeloperoxidase, along with overproduction of ROS can lead to inflammatory tissue injury (20, 21). Uptake of gold nanoparticles has been linked to endoplasmic reticulum stress and cleavage of cytoskeletal proteins in human neutrophils leading to apoptosis (22). Further, nano and micro particles have been reported to inhibit the recruitment of neutrophils from the circulation to sites of inflammation in mice and to cause a reduction in neutrophil attachment to endothelial monolayers in vascular mimetic flow channels (23, 24). The fine balance between the protective functions of neutrophils that maintain immune competence versus exuberant response that can result in tissue and organ damage has prompted the current studies on neutrophil activation and inflammatory recruitment in the presence of Feraheme. Intravenous infusion with Feraheme for delivery of complexed iron is FDA approved at a higher single dose than other products on the market (25). A typical treatment regime consists of two 510-mg doses separated by 5–8 days as compared to other products that require between 5 and 10 individual doses, thereby limiting the need for repeated infusion which typically lowers patient compliance. Feraheme also is approved for infusion at higher rates than other iron products with comparable safety (25, 26). The increased application of Feraheme in anemia patients who can receive frequent intravenous injections as a course of therapy highlight a need to understand the effects of SPION on neutrophils and their capacity to maintain normal immunosurveillance.

In the current study, we examined the effects of Feraheme on neutrophil degranulation and alterations in receptor expression and conformation on the plasma membrane. We hypothesized that neutrophil uptake of Feraheme in blood alters the normal process of adhesion receptor activation, shear stress resistant neutrophil rolling to arrest and subsequent shape change that precedes cell migration. Vascular mimetic flow channels were employed to assess the kinematics of neutrophil interactions on a substrate of recombinant E-selectin and ICAM-1 in an established model of endothelial inflammation. Feraheme in suspension inhibited neutrophil activation and degranulation induced by IL-8 stimulation, resulting in alteration in the expression of adhesion molecules necessary for the efficient recruitment on inflamed endothelium.

MATERIALS AND METHODS

Small Molecules, Antibodies, and Other Reagents

Monoclonal antibodies for flow cytometric detection of high affinity β2-integrin (mAb24), L-Selectin (Dreg-55, Dreg-56), CD11b (M170), CD18 (1B4), CD66b (G10F5) CD11a (HI111), PSGL-1 (PL-2, KPL-1), CXCR1(8F1/CXCR1), and CXCR2 (5E8/CXCR2)
along with antibodies that block CD11b function (Mac-1 blocking, M1/70), fixation buffer, and IL-8 were purchased from Biolegend (San Diego, CA). Recombinant human ICAM-1-IgG and E-Selectin-IgG produced as Fc chimeric constructs were purchased from R&D Systems (Minneapolis, MN). Adenosine A₂A receptor agonist CGS-21680, N-formyl-Met-Leu-Phe (fMLP), and ROS indicator Dihydrorhodamine 123 were purchased from Millipore Sigma (Burlington, MA). Adenosine A₂A receptor antagonist ZM #235586-9. Neutrophils were isolated from whole blood as per manufacturers instruction (Fresenius Kabi). Briefly, by gradient based isolation, whole blood is layered on an equal volume of Polymorphoprep® as per manufacturers instruction (Fresenius Kabi). Briefly, by gradient based isolation, whole blood is layered on an equal volume of Polymorphoprep solution and centrifuged at 760g for 30 min at 25°C. The neutrophil layer is collected and washed in Phosphate-buffered saline (PBS) before resuspension in HBSS buffer containing 0.1% human serum albumin (HSA) without Ca²⁺ and Mg²⁺ and kept at 1 × 10⁷ cells/ml on ice after isolation and prior to experimentation. For EasySep neutrophil isolation, whole blood was diluted 1:1 with PBS then 100 µl of isolation cocktail and magnetic RapidSpheres each were added and incubated for 5 min. Cells were then placed in the EasySep magnet for 5 min. Cells were poured into a new tube and treated for an additional 5 min with 100 µl of magnetic RapidSpheres and placed on the EasySep magnet for 5 min twice more. Cells were then spun down and resuspended in HBSS containing 0.1% HSA without Ca²⁺ and Mg²⁺ and kept at 1 × 10⁶ cells/ml on ice after isolation and prior to experimentation. Cell purity was >90% as determined by Beckman cell coulter counter. Cell viability was ~99% as determined by flow cytometry detection of zombie violet fluorescent dye staining (Biolegend).

### Human Neutrophil Isolation

Whole blood was obtained from healthy donors consented through a University of California, Davis institutional review board protocol #235586-9. Neutrophils were isolated from whole blood via negative enrichment using EasySep™ direct human neutrophil isolation kit purchased from StemCell Technologies as per manufacturers instruction (Cambridge, MA). Some studies employed a percoll gradient separation using Polymorphoprep® as per manufacturers instruction (Fresenius Kabi). Briefly, for gradient based isolation, whole blood is layered on an equal volume of Polymorphoprep solution and centrifuged at 760g for 30 min at 25°C. The neutrophil layer is collected and washed in Phosphate-buffered saline (PBS) before resuspension in HBSS buffer containing 0.1% human serum albumin (HSA) without Ca²⁺ and Mg²⁺ and kept at 1 × 10⁷ cells/ml on ice after isolation and prior to experimentation. For EasySep neutrophil isolation, whole blood was diluted 1:1 with PBS then 100 µl of isolation cocktail and magnetic RapidSpheres each were added and incubated for 5 min. Cells were then placed in the EasySep magnet for 5 min. Cells were poured into a new tube and treated for an additional 5 min with 100 µl of magnetic RapidSpheres and placed on the EasySep magnet for 5 min twice more. Cells were then spun down and resuspended in HBSS containing 0.1% HSA without Ca²⁺ and Mg²⁺ and kept at 1 × 10⁶ cells/ml on ice after isolation and prior to experimentation. Cell purity was >90% as determined by Beckman cell coulter counter. Cell viability was ~99% as determined by flow cytometry detection of zombie violet fluorescent dye staining (Biolegend).

### Flow Cytometry Detection of Cell Surface Marker Expression

Neutrophils (1 × 10⁶ cells/ml) in HBSS buffer containing Ca²⁺ and Mg²⁺ at 1 mM were treated with IL-8 at a dose range of concentrations (0.01–100 nM) and Feraheme (1–6 mg/ml) for 5 min at 37°C before addition of antibody (1.5–5 µg/ml). Cells were stained with antibodies for high affinity β2-integrin (mAb24), L-Selectin (Dreg-56), CD11b (M1/70), CD18 (1B4), CD11a (H1111), PSGL-1(KPL-1), CXCR1/8F1(CXCR1), and CXCR2 (5E8/CXCR2) for 20 min at 37°C before fixation with 4% paraformaldehyde at room temperature. After two washes with PBS cells were analyzed using the Attune NxT flow cytometer (ThermoFisher). Neutrophils were gated by their characteristic forward scatter vs side scatter profile. Receptor expression in terms of sites/cell was determined by comparing the MFI of bound antibody to Quantum Simply Cellular beads (Bangs Laboratories, Inc., Fishers, IN) which contain five bead sets with increasing numbers of antibody binding sites on their surface. From this analysis, a linear relation between MFI and receptor expression was determined for each directly conjugated antibody bound to cells and the calibration bead set.

### Microfluidic Shear Flow Assay

Vascular mimetic flow chambers were utilized to record neutrophil rolling and arrest behavior on substrates of endothelial ligands under physiological shear stresses. Devices were prepared as described previously (27). Briefly, polydimethylsiloxane (PDMS) microfluidic flow chambers with a minimum feature size of 5 µm were produced by curing Sylgard 184 prepolymer (Dow Corning, Midland, MI, USA) over a patterned silicon wafer. Holes were punched into the PDMS for flow channel and vacuum port access and the device was reversibly vacuum sealed on a glass coverslip coated with E-selectin (1 µg/ml) alone or with ICAM-1 (1 µg/ml) in PBS for 60 min at room temperature. To limit non-specific adhesion of neutrophils, coverslips were treated with 1% casein for 10 min before washing with PBS and assembly of the microfluidic device.

Isolated neutrophils were treated with Ca²⁺ indicators 1 Fluor-4 AM or Rhod-2 AM (1 µg/ml) for 20 min at room temperature in the dark. Cells were spun down and resuspended at 1 × 10⁶ cells/ml and were treated with Mac-1 blocking antibody M1/70 and incubated with or without MNP in HBSS buffer containing Ca²⁺ and Mg²⁺ for 20 min at 37°C. Neutrophils were then treated with 0.5 nM IL-8 or vehicle control and immediately loaded into an open 100-µl reservoir and were drawn through the channel by negative pressure produced by a syringe pump at a shear of 2 dynes/cm².

Utilizing real time fluorescence microscopy, images of rolling neutrophils were acquired at 60 frames per minute for 4 min per field of view on an inverted microscope (Nikon) using a phase contrast 20× objective and recorded with 16-bit digital complementary metal oxide semiconductor (CMOS) camera (Andor ZYLA) connected to a PC (Dell) with NIS Elements imaging software (Nikon Instruments Inc.). Arrested neutrophils were defined as having a velocity less than 0.1 µm/s. Migrating neutrophils were identified by exhibition of a polarized shape defined as exceeding a length/width aspect ratio greater than 1.4 and phase dark contrast indicative of being outside the focal plane of rolling cells.

### L-Selectin and PSGL-1 Clustering

Total internal reflection fluorescence (TIRF) microscopy and quantitative dynamic foot printing (qDFP) were employed to record fluorescently tagged antibodies targeting L-selectin (non-blocking clone: DREG55) and PSGL-1 (non-blocking clone: PL2). Cluster area and frequency during rolling of isolated neutrophils over E-selectin in the absence of Feraheme was observed. A cluster was defined as an area of uniform fluorescence intensity two standard deviations above the mean fluorescence intensity of the cell. Additionally, a cluster was defined as having a surface area of 0.4 µm² or greater.

### Measurement of Reactive Oxygen Species

Neutrophils (1 × 10⁶ cells/ml) in HBSS buffer containing Ca²⁺ and Mg²⁺ at 1 mM were incubated with the ROS indicator Dihydrorhodamine 123 (2 µM) and treated with or without IL-8.
Feraheme (4 mg/ml) for 10 min at 37°C before addition of 1 μM fMLP. Cells were incubated with fMLP at 37°C for 5 min and then placed on ice to end reactions and ROS was then quantified through flow cytometry.

Measurement of Calcium Flux

Isolated neutrophils were treated with the Ca2+ indicator Fluo-4 AM (1 μg/ml) for 20 min at room temperature in the dark. Cells were spun down and resuspended at 1 × 10⁶ cells/mL in HBSS buffer containing Ca2+ and Mg2+ at 1 mM and treated with or without the adenosine A2A receptor antagonist ZM 241385 (2.5 μM) for 5 min at 37°C. Feraheme (4 mg/ml) and or adenosine A2a receptor agonist CGS 21680 (1 μM) were then added to the cell solutions for 10 min at 37C. After incubation, IL-8 (1 nM) was added to cell solutions and calcium fluorescence time course was immediately read on the Facsscan flow cytometer (BDbiosciences).

RESULTS

Feraheme Inhibits Neutrophil Activation and Degranulation Stimulated by IL-8

Stimulation with chemotactic factors activate within seconds neutrophil degranulation, which results from the fusion of granule membranes to the plasma membrane resulting in upregulation of additional CD11b/CD18 and CD66b receptors. Activation also results in a shift in the conformation of CD18 from a constitutive low affinity conformation at rest to high affinity upon CXCR ligation, as well as the shedding of L-selectin receptors through the action of the metalloprotease ADAM17 (28, 29). We first assessed the capacity of different concentrations of Feraheme in suspension to alter neutrophil responses to chemotactic activation via CXCR1/2 in cells stimulated at the Kᵣ of IL-8 stimulation (~1 nM). Neutrophils were incubated at 37°C in the presence and absence of Feraheme and IL-8 for 5 min before the addition of fluorescent antibodies for 20 min followed by cell fixation. The presence of Feraheme led to significant alterations in expression of CD62L, CD11b, and high affinity (HA) CD18 at each dose applied (~6 mg/ml) (Figure 1). A dose dependent increase in the inhibition of degranulation was observed, as well as diminished activation of integrin and proteolytic cleavage of L-selectin. Since the effect of Feraheme was observed to plateau at 4 mg/ml, subsequent studies were performed using this concentration.

We next evaluated the effect of Feraheme on adhesion receptor expression over a dose range of chemotactic stimulation with IL-8. A typical sigmoidal dose dependent decrease in L-selectin expression followed addition of IL-8 between 0.1 and 100 nM, which reached a maximum of ~90% loss at 5 nM IL-8 from baseline expression on quiescent neutrophils (Figure 2A). Upregulation in surface expression of CD11b and CD18 provided a sensitive measure of activation, increasing by 2-fold and 2.3-fold from baseline up to maximum expression at 10 nM IL-8 (Figures 2B, C). Another sensitive measure of neutrophil activation was provided by the increased binding of mAb24 that reports on the allosteric conversion of CD18 receptors from low to high affinity, which yielded a 3-fold increase in expression at maximum IL-8 stimulation (Figures 2B, D). It is noteworthy that the maximum extent of L-selectin shedding was lowered by Feraheme at 4 mg/ml compared with 0 mg/ml, as indicated by the elevated plateau at a dose of IL-8 >5 nM from 10% up to 25% total receptor expression (Figure 2A). Moreover, the extent of upregulation of CD11b/CD18 and conversion of CD18 to the HA conformation was significantly inhibited, as indicated by a ~2-fold increase in the EC50 of IL-8 stimulation. This resulted in a reduced sensitivity to activation that was significant at IL-8 doses of 0.5 and 1 nM for CD11b, total CD18, and HA CD18 (Figures 2B–D). A marker of the release of secondary granules is the increase in membrane expression of CD66b, which is transported to the plasma membrane along with CD11b/CD18 (20). The presence of Feraheme elicited a significant inhibition in upregulation of CD66b, increasing the EC50 of IL-8 stimulation by ~30% (Supplemental Figure 1). In contrast, CD11a and PSGL-1 adhesion receptors that are constitutively expressed on circulating neutrophils and typically do not register a change in receptor number in response to chemotactic stimulation, remained constant over the dose range of IL-8 stimulation in the presence of Feraheme (Supplemental Figure 1).

Neutrophils are reported to endocytose CXCR1/2 following ligation and signaling by IL-8 (30). This motivated experiments to determine if Feraheme altered the expression of CXCR1/2, consequently accounting for the diminished capacity to induce changes in adhesion molecule expression following stimulation. Unexpectedly, incubating unstimulated neutrophils in suspension with Feraheme at 4 mg/ml elicited a ~26% increase in expression of CXCR1 and a ~15% increase in CXCR2 (Figure 3). A significant amount of CXCR1 endocytosis following IL-8 simulation was not observed in the presence or absence of Feraheme (Figure 3A). In contrast, the small amount of CXCR2 endocytosis in response to IL-8 stimulation was not altered in the presence of Feraheme (Figure 3B). These results indicate that CXCR1/2 expression either remains constant or increases in the presence of Feraheme and IL-8 stimulation as compared to controls. Thus, it is unlikely that the mechanism left-shifting the IL-8 dose response for activation is diminished expression of chemotactic receptors. Feraheme is also not altering the endocytosis of CXCR2 receptors in cells treated with 1 nM IL-8 indicating it is not interfering with IL-8 ligation of CXCR2. Thus, it is probable that Feraheme is leading to dramatically reduced responses to IL-8 stimulation by affecting intracellular signaling downstream of receptor ligation.

Feraheme Inhibits the Activation of ROS Production in Neutrophils

To further explore the influence of Feraheme on neutrophil function, ROS production was quantified using flow cytometry to detect intracellular Dihydorhadamine 123 fluorescence. Production of ROS is a key antimicrobial mechanism of neutrophils and has also been implicated in recruitment to sites of inflammation by increasing vascular permeability (31). The bacterial tripeptide fMLP is a potent chemotactic factor that binds to G-protein coupled Formyl peptide receptors expressed on neutrophils and activates the respiratory burst and production of ROS. Feraheme led to a significant 30% reduction of ROS
production in neutrophils stimulated with 1 µM fMLP (Figure 4). Pre-treatment with IL-8 has previously been reported to prime neutrophils for enhanced fMLP induced ROS production through assembly of the NADPH oxidase components into lipid rafts on the plasma membrane (32). Feraheme significantly reduced ROS production by 48% in cells primed with low dose IL-8 and stimulated with fMLP. While uptake of SPION other than Feraheme have been reported to induce ROS production, the presence of Feraheme did not lead to detectable ROS production in unstimulated neutrophils. This data indicates that Feraheme inhibits formyl peptide receptor signaling of ROS production, including increased ROS induced by IL-8 priming.

**Neutrophil Recruitment on Endothelial Adhesion Molecules Is Altered in the Presence of Feraheme**

It has been reported that exposure to nanoparticles and microparticles reduced neutrophil recruitment to inflamed mesentery microvasculature in mice (23). Moreover, polystyrene nanoparticles have been shown to reduce adhesion of human neutrophils to inflamed human umbilical vein endothelial cells in parallel plate flow chamber studies (24). This motivated determination of the effect of Feraheme on the multistep process of recruitment in vitro using microfluidic flow channels (23, 24, 33). Fluorescence microscopy was employed to image the dynamics of...

**FIGURE 1** | Effect of Feraheme concentration on integrin and selectin expression following chemotactic stimulation of neutrophils. Isolated human neutrophils were incubated with 1 nM IL-8 and Feraheme magnetic nanoparticles (MNP) for 25 min and cell surface expression of (A) CD62L, (B) CD11b, and (C) HA CD18 was assessed by flow cytometry. Representative histograms depict fluorescent antibody detection for each adhesion receptor at baseline receptor expression and following IL-8 stimulation in presence and absence of MNP. Bivariate data are presented as the percent shedding from unstimulated baseline mean ± SEM for CD62L and percent of unstimulated baseline expression mean ± SEM for CD11b and HA CD18 (n ≥ 4 donors) with experimental replicates averaged for each donor. Paired T-test was performed comparing the average value at each concentration to the 0 mg/ml Feraheme condition of the same donor * and ** denote p value ≤ .05 and ≤ 0.01, respectively.
neutrophil rolling and arrest within vascular mimetic microfluidic channels containing substrates coated with E-selectin alone or in conjunction with ICAM-1. Neutrophil suspensions were incubated for 20 min with Feraheme at 1 mg/ml or 4 mg/ml and compared with vehicle control in response to stimulation with 0.5 nM IL-8 prior to perfusion through the flow channel. This dose of IL-8 corresponded with the lowest concentration of stimulus in which a significant inhibition of CD18 activation and L-selectin shedding was elicited by Feraheme. Capture and rolling on E-selectin is primarily mediated by recognition of sLex presenting ligands on L-selectin and PSGL-1 expressed on neutrophils. The velocity of neutrophil rolling on E-selectin at a venular shear stress of 2 dyne/cm² increased ~37% in the presence of Feraheme at 4 mg/ml compared to vehicle alone, whereas no alteration in rolling velocity was detected at 1 mg/ml (Figures 5A, B). This increased rolling velocity was accompanied by a 40% increase in the variance of mean velocity, which is consistent with interruption of adhesive bond formation resulting in an unsteady trajectory of neutrophil rolling on the substrate under shear flow. Neutrophils require ICAM-1 on the substrate to provide an anchor for activated integrins to bind in order to achieve shear resistant cell arrest (28). Perfusion through flow channels derivatized with E-selectin and ICAM-1 were used to study Feraheme’s effect on the kinematics of neutrophil arrest and transition to a migratory phenotype. In the absence of IL-8 chemotactic stimulation, Feraheme lead to a dose dependent increase in rolling velocity on E-selectin coated substrates, which increased by ~30% and ~60% when neutrophils were exposed to 1 and 4 mg/ml Feraheme, respectively (Figures 5C, D). Moreover, an ~80% increase in variance of mean velocity corresponded to treatment at 4 mg/ml Feraheme, while 1 mg/ml increased variance by 17%. Stimulation with IL-8 leads to the activation of HA CD18 that binds ICAM-1 and mediates deceleration to arrest (28). Consequently, stimulation with IL-8 on E-selectin/ICAM-1 coated substrates resulted in a 50% decrease in rolling velocity and within ~40 s most neutrophils exhibited a rapid transition to arrest (Figures 5C, F). Neutrophils treated with Feraheme at 4 mg/ml exhibited ~40% higher rolling velocity compared to the 1 mg/ml or vehicle control conditions (Figure 5D). In the absence of IL-8 stimulation, outside-in signaling via E-selectin binding and clustering of L-selectin is sufficient to activate ~40% of cells to convert from rolling to arrest that is supported by activation of HA CD18 to bind ICAM-1 (34). Feraheme at 1 and 4 mg/ml concentrations inhibited this L-selectin mediated signaling of arrest, as well as decreased transition to a migratory state from 12% of cells to <1% of cells assuming a polarized shape (Figure 5E). In contrast, in the presence of IL-8 stimulation, we did not detect significant inhibition in the frequency of neutrophils rolling to arrest in presence of Feraheme (Figure 5E). This is likely due to the fact that coupled with shear force mediated L-selectin signaling, even very low concentrations of IL-8 can lead to activation of HA CD18 receptors sufficient to mediate conversion from rolling to arrest (35). However, in the presence of Feraheme, a significant...
increase was detected in the duration of rolling before the onset of cell arrest for both IL-8 stimulated and unstimulated neutrophil suspensions. In the absence of IL-8, rolling time to arrest increased from 42.1 seconds in the 0 mg/ml condition to 55.1 and 91.9 seconds in the 1 mg/ml and 4 mg/ml conditions, respectively (Figure 5F). In IL-8 treated cells only the 4 mg/ml condition resulted in a slight reduction in rolling velocity and decreased outside-in signaling (via E-selectin in the presence of Feraheme and indicate this effect may be a function of diminished L-selectin bond formation despite the observed higher frequency and area of L-selectin within sites of adhesive contact.

**Feraheme Accelerates Clearance of Cytosolic Calcium After Flux**

Calcium serves as a secondary messenger downstream of GPCR signaling that mediate neutrophil inflammatory responses including degranulation, integrin activation and adhesion, shape change, and ROS production (36). IL-8 ligation of CXCR leads to a release of endoplasmic reticulum (ER) stored calcium into the cytosolic space which is subsequently sequestered by the ER calciosomes to replenish stores. High concentrations of extracellular adenosine which binds to the adenosine A2A receptor is known to dampen inflammatory responses (37). The mechanism, as depicted in Figure 7A, is initiated by ligand binding to the adenosine A2A receptor which induces dissociation and activation of the Gα subunit of the A2A receptor linked heterotrimeric G protein which then activates adenylyl cyclase. Adenylyl cyclase activation leads to production of cyclic AMP (cAMP), which in turn activates cAMP-dependent protein kinase (PKA) and leads to accelerated sequestration of calcium through PKA-activated endo-membrane Ca2+-ATPases. To determine if Feraheme’s inhibitory effects on intracellular signaling are mediated through accelerated clearance of IL-8 induced calcium flux, the time course of cytosolic calcium stimulated by IL-8 in Feraheme treated neutrophils was compared in the presence and absence of adenosine A2A receptor agonist CGS 21680 (CGS). Following IL-8 induced release of ER Ca2+, Feraheme elicited an accelerated decrease in cytosolic calcium as compared to untreated cells. The level of accelerated Ca2+ clearance was equivalent and not additive with CGS (Figure 7B). The duration required for a decrease to 20% of the maximum Ca2+ flux elicited by IL-8 was 60 seconds in untreated cells compared to 36, 22, and 22 seconds for Feraheme, CGS, and Feraheme + CGS, respectively (Figure 7C). Neither Feraheme nor CGS alone reduced the maximal Ca2+ flux elicited by IL-8, while added together they resulted in a slight reduction (Figure 7D).

To determine if Feraheme mediated inhibition acts through release and binding of extracellular adenosine, the clearance of

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**FIGURE 3** | Feraheme alters CXCR expression in neutrophils. Isolated human neutrophils were incubated with IL-8 or vehicle and Feraheme for 25 min and cell surface expression of (A) CXCR1 and (B) CXCR2 was assessed by flow cytometry. The data are presented as mean ± SEM (n ≥4 donors) with experimental duplicates averaged for each donor. Paired T-tests were performed comparing the average value of the 4 mg/ml to the 0 mg/ml Feraheme conditions of the same donor * and ** denote p value ≤ 0.05 and ≤ 0.001, respectively. Paired T-tests were performed comparing the average value for a condition to the 0 nM IL-8 condition of the same donor $ denotes p value ≤ 0.05.

**FIGURE 4** | Feraheme alters CXCR expression in neutrophils. Isolated human neutrophils were incubated with IL-8 or vehicle and Feraheme for 25 min and cell surface expression of (A) CXCR1 and (B) CXCR2 was assessed by flow cytometry. The data are presented as mean ± SEM (n ≥4 donors) with experimental duplicates averaged for each donor. Paired T-tests were performed comparing the average value of the 4 mg/ml to the 0 mg/ml Feraheme conditions of the same donor * and ** denote p value ≤ 0.05 and ≤ 0.001, respectively. Paired T-tests were performed comparing the average value for a condition to the 0 nM IL-8 condition of the same donor $ denotes p value ≤ 0.05.
calcium after IL-8 stimulated Ca$^{2+}$ flux was compared to that activated by the adenosine A$_{2A}$ receptor agonist CGS 21680 (CGS) while blocking signaling via the A$_{2A}$ receptor with the adenosine A$_{2A}$ receptor antagonist ZM 241385 (ZM) as depicted in Figure 8A. As expected, in neutrophils preincubated with ZM the sequestration effect of CGS was completely abrogated. In contrast, Feraheme treatment remained effective in accelerating the decrease in cytosolic Ca$^{2+}$ following IL-8 stimulation (Figure 8B). In untreated cells the duration to 20% of maximum Ca$^{2+}$ flux was 65 s compared to 46, 63, and 42 s for Feraheme, CGS, and Feraheme+ CGS, respectively (Figure 8C). All conditions registered the same maximal Ca$^{2+}$ flux (Figure 8D).

**DISCUSSION**

Neutrophil recruitment and attachment to inflamed endothelium is mediated by a multistep process involving ligation and mechanosignaling from the outside-in through selectins and integrins following bond formation (38). CXCR ligation of IL-8 by neutrophils interacting with inflamed endothelium under shear flow induces degranulation and the activation of CD18 to a high affinity state, as well as the rapid shedding of L-selectin (20, 29). Here, we demonstrated that, treating neutrophils in suspension over a dose range with Feraheme elicited reduced responsiveness to stimulation with 1 nM IL-8 at each dose tested (1–6 mg/ml). This was manifest as increased inhibition of CD11b upregulation, HA CD18 activation, and L-selectin shedding with increasing particle concentration of Feraheme. Treating with a Feraheme dose of 4 mg/ml resulted in a consistent ~2-fold right shift in the EC50 of stimulation with IL-8 corresponding to diminished upregulation of CD11b/CD18, activation of HA CD18, and L-selectin shedding. Chemokine receptor expression in unstimulated cells was significantly elevated in the presence of Feraheme, but these changes in expression did not explain the inhibitory effects observed. Feraheme also did not alter endocytosis of CXCR2 in response to IL-8 signaling, suggesting that its mechanism of action is not through down regulation or interference with CXCR ligation. Feraheme also significantly reduced fMLP induced ROS production in the presence or absence of IL-8 priming. Taken together, we conclude that the inhibitory effects of Feraheme are not exclusive to CXCR1/2 signaling by IL-8, but include signaling through other GPCR namely, formyl peptide receptors.

Uptake of nanoscale poly(styrene) and liposomal particles by neutrophils was previously shown not to enhance apoptosis, activation, or cell death. Ingested nanoparticles were reported to reside in intracellular compartments that are retained during degranulation (39). Uncoated SPION are reported to aggregate and induce the formation of neutrophil extracellular traps (NETs) that are extracellular fibers consisting of expelled DNA. This pro-inflammatory host response is mitigated by coating SPION with layers of dextran or HSA (40). In our experiments, Feraheme did not reduce cell viability, which exceeded 99%. Although we did not specifically test for netosis in our studies of neutrophil arrest and migration, no visible changes in cell morphology that is characteristic of netosis was observed. In-vivo studies have demonstrated that uptake of nanoparticles can diminish the efficiency of neutrophil recruitment to inflamed endothelium within the lungs of mice. However, the mechanism that may involve cell-particle uptake and competitive binding to adhesion receptors, which could interfere with chemotactic or adhesive ligand binding during recruitment remained elusive (23, 24). Here, we report that following 20 min of exposure to Feraheme in suspension, we observed diminished efficiency of neutrophil arrest and migration as measured in an established model of endothelial inflammation in vascular mimetic shear flow channels. We have recently reported that deceleration of neutrophils within the vascular mimetic substrate in microfluidic flow channels is mediated by as few as 200 HA CD18 binding ICAM-1 in order to transition from rolling to arrest (34). In the absence of IL-8 stimulation and in the presence of Feraheme, a ~50% reduction was measured in the frequency of neutrophils rolling to arrest on E-selectin/ICAM-1 at both 1 and 4 mg/ml. Moreover, neutrophils rolled over twice the distance before activation and binding of HA CD18 integrin bonds formed stable arrest. Inhibition of neutrophil CD11b/CD18 mediated migration after arrest was also observed at 1 and 4 mg/ml Feraheme. Thus, Feraheme uptake in a concentration dependent manner alters both selectin and GPCR mediated signaling of HA CD18 necessary to transition from rolling on E-selectin to arrest on ICAM-1. It has been previously shown that L-selectin catch-bonds, characterized by a prolonged bond lifetime as tensile force is increased, is necessary and sufficient to signal CD18 transition to a HA state (34). We show that Feraheme reduced both L-selectin shedding in response to IL-8 and L-selectin clustering during rolling on E-selectin, but it had no effect on PSGL-1 expression. Noteworthy was the finding that cluster area of L-selectin bonds during rolling on E-selectin in the presence of
shear stress and IL-8 stimulation was increased, while the density of bond formation within these clusters decreased. This indicates that Feraheme may alter L-selectin signaling, but not recognition by E-selectin on the surface of rolling neutrophils. This is consistent with the significant rise in the mean rolling velocity on E-selectin that was observed to increase with Feraheme concentration. This provides one potential explanation as to how Feraheme reduces the efficiency of the transition to cell arrest, in that it may alter the capacity of E-selectin/L-selectin to form catch-bonds. Increased availability of L-selectin to form bond clusters in the presence of Feraheme coincided with reduced density of bonds on E-selectin and also with a reduced frequency of cell arrest suggesting that the normal process of neutrophil deceleration at sites of inflammation may be perturbed. Reduced levels of L-selectin shedding may potentially contribute to the inhibition of arrest and polarization observed in Feraheme treated cells. Consistent with the latter observation are reports that L-selectin shedding

![FIGURE 5](https://example.com/figure5.png)
has been reported to amplify integrin-mediated outside-in signaling-dependent processes, including neutrophil migration, production of ROS, and phagocytosis (41). L-selectin shedding induces phosphorylation of phospholipase C (PLC)γ2, Akt, and Syk, critical regulators of intracellular calcium release that are important for E-selectin mediated slow rolling and CD18 integrin activation in neutrophils (41, 42). Thus, inhibition of L-selectin shedding may also interfere with normal activation of HA CD18 during rolling on E-selectin, despite the elevated expression of L-selectin on the surface. While it is likely that Feraheme is endocytosed by neutrophils in suspension, we cannot rule out that a fraction remain bound to the plasma membrane and sterically interfere with the recognition and mechanics of selectin bond formation. Due to optical limitations in real-time imaging of SPION bound to the plasma membrane at the nanoscale, we could not directly image whether Feraheme specifically bound to L-selectin receptors and sterically interfered with catch-bond formation.
associated with mechanotransduction of CD18 activation. What is clear is that Feraheme altered the efficiency of L-selectin’s capacity to mechanosignal activation of HA CD18, but not necessarily its ability to bind E-selectin and form focal clusters.

To elucidate a possible mechanism for Feraheme inhibition of GPCR signaling the effects of Feraheme on calcium flux after IL-8 stimulation was recorded. Feraheme significantly accelerated the clearance of calcium after flux and this response was comparable to that of the adenosine A2A receptor agonist CGS 21680 known to inhibit degranulation and ROS production. The effect of Feraheme was not dependent on adenosine as treatment with the adenosine A2A receptor antagonist ZM 241385 did not perturb the acceleration of calcium clearance in response to Feraheme. Calcium signaling plays a key role in GPCR mediated degranulation, L-selectin shedding, integrin activation, and ROS production which are all inhibited by Feraheme. Calcium is also necessary for selectin mechanosignaling induced activation of HA LFA-1 and for integrin signaling of shape change and polarization after arrest (38). Accelerated clearance of calcium could potentially result in the decreased arrest and polarization observed in Feraheme treated cells by inhibiting selectin mechanosignaling. These experiments highlight a consistent effect of Feraheme uptake on neutrophil intracellular signaling, but the mechanism which may involve Feraheme steric influence.
on normal cell surface receptor affinity or avidity in binding ligand needs to be confirmed in future studies. The non-additive inhibition in the presence of Feraheme and CGS could indicate a shared downstream mechanism for the accelerated clearance of Ca^{2+}. We propose that Feraheme may induce sequestration of calcium through activation of endo-membrane Ca^{2+}-ATPases rather than by efflux of calcium through activation of plasma membrane Ca^{2+}-pump, the proposed mechanism of action for CGS (Figure 9).

In summary, Feraheme exerted an immunosuppressive effect on neutrophil activation in response to IL-8 and fMLP, as well as outside-in signaling via E-selectin during neutrophil recruitment in shear flow. A potential mechanism may involve the observed Feraheme induced acceleration of calcium clearance resulting in a reduced capacity for cell signaling of immune responses. Further studies are warranted to elucidate the mechanism by which Feraheme induces accelerated calcium clearance, potentially though elevation of cAMP leading to downstream activation of endo-membrane Ca^{2+}-ATPases, and whether this is the primary mechanism for the inhibition of neutrophil inflammatory responses observed herein. Whether these effects are seen with other SPION formulations is also a relevant question to pursue.
These immunosuppressive effects could lead to impaired immune responses to infection or sterile tissue insult in patients treated with Feraheme. The current studies also point to the expanded use of SPION to downregulate the innate immune response as a therapy for ameliorating chronic inflammation and/or autoimmune diseases.

**DATA AVAILABILITY STATEMENT**

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

**ETHICS STATEMENT**

The studies involving human participants were reviewed and approved by UC Davis Institutional Review Board. The patients/participants provided their written informed consent to participate in this study.

**AUTHOR CONTRIBUTIONS**

GG designed the study and performed flow cytometry experiments. VM performed microfluidic experiments. GG and VM performed data analysis. GG wrote initial draft with aid of VM and SS. All authors contributed to the article and approved the submitted version.

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**SUPPLEMENTS MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fimmu.2020.571489/full#supplementary-material
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**Conflict of Interest:** 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.

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