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Effects of PVA coated nanoparticles on human immune cells

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Abstract: Nanotechnology provides new opportunities in human medicine, mainly for diagnostic and therapeutic purposes. The autoimmune disease rheumatoid arthritis (RA) is often diagnosed after irreversible joint structural damage has occurred. There is an urgent need for a very early diagnosis of RA, which can be achieved by more sensitive imaging methods. Superparamagnetic iron oxide nanoparticles (SPION) are already used in medicine and therefore represent a promising tool for early diagnosis of RA. The focus of our work was to investigate any potentially negative effects resulting from the interactions of newly developed amino-functionalized amino-polyvinyl alcohol coated (a-PVA) SPION (a-PVA-SPION), that are used for imaging, with human immune cells. We analyzed the influence of a-PVA-SPION with regard to cell survival and cell activation in human whole blood in general, and in human monocytes and macrophages representative of professional phagocytes, using flow cytometry, multiplex suspension array, and transmission electron microscopy. We found no effect of a-PVA-SPION on the viability of human immune cells, but cytokine secretion was affected. We further demonstrated that the percentage of viable macrophages increased on exposure to a-PVA-SPION. This effect was even stronger when a-PVA-SPION were added very early in the differentiation process. Additionally, transmission electron microscopy analysis revealed that both monocytes and macrophages are able to endocytose a-PVA-SPION. Our findings demonstrate an interaction between human immune cells and a-PVA-SPION which needs to be taken into account when considering the use of a-PVA-SPION in human medicine.

Keywords: nanoparticle, cell viability, cytokine, monocyte, macrophage

Introduction

Rheumatoid arthritis (RA) is a chronic, systemic, inflammatory autoimmune disease of connective tissues, which mostly affects synovial joints. The inflammatory process leads to a local release of pro-inflammatory cytokines, including TNF α and IL-1, that activate enzymes involved in the degradation of articular cartilage and subchondral bone. Other tissues and organs may also be harmed as a result of the disease.¹⁻⁹

Between 0.5% and 1% of the human population within industrialized countries suffer from RA (three times as many women as men), with the average age of disease onset between 40 and 50 years.¹⁰ The causes of RA are not completely clarified, but the development of the disorder seems to result from a combination of genetic factors and environmental triggers.¹¹⁻¹⁴ The typical clinical signs of RA (symmetric joint swellings, morning stiffness, pain, and asthenia) are important for diagnosing this disease. Additionally, imaging techniques like X-ray, ultrasound and magnetic resonance imaging (MRI) as well as laboratory tests (such as inflammatory markers, rheumatoid factors, and anti-citrullinated protein antibodies) complete the diagnosis and permit surveillance of disease activity, progress and therapy success.¹⁵⁻¹⁸ There is no cure for RA, but the treatment can improve symptoms and quality of life, and can

reduce or even stop the progress of the disease. Typically, disease-modifying anti-rheumatic drugs, biological agents, glucocorticoids and non-steroidal anti-inflammatory drugs are used for treatment.^{5,19–21} Nevertheless, some patients do not respond adequately to the prescribed drug, requiring 3 months of treatment before a lack of response becomes apparent. Only after this period of time has elapsed can treatment strategy be modified, with further progress of the disease potentially occurring due to the loss of this treatment time. Additionally, all treatment strategies involve some side effects.^{8,9,22–24}

It has been shown that the earlier patients with RA are treated the more effective is the treatment.^{25,26} Thus there is an urgent need for a very early diagnosis of RA in order to enable earlier treatment and to reduce or even to stop the progress of the disease. Furthermore, an early diagnosis can prevent joint damage, thereby lowering the additional therapeutic and disability costs which arise from reduced work capacity of the patients: nearly 30% of patients give up work within 1 year as a result of their condition and almost 60% within 6 years of diagnosis.²⁷ Thus, there is a considerable need for highly sensitive and specific diagnostic tools to detect early disease. Such tools could also facilitate further drug development and enable health professionals to distinguish between responders and non-responders before and after onset of therapy.

One promising approach to generate such a tool is the use of nanotechnology employing superparamagnetic iron oxide nanoparticles (SPION). These have been intensively investigated over the last 20 years for various *in vitro* and *in vivo* biomedical applications.^{28,29} Promising studies have been performed with SPION as contrast enhancers for MRI.^{30–35} Currently, SPION are used as contrast agents for liver function. For example, the product Feridex[®] was approved by the US Food and Drug Administration (FDA) in 1996.^{36,37} MRI has been used for many years in cross-sectional and observational studies in RA and in controlled clinical trials to identify early cartilage loss in osteoarthritis and RA.^{38,39} This method is also useful to assess and monitor disease activity in order to detect and follow pre-erosive features of RA (such as synovitis, bone marrow edema or osteitis, and tendon and ligament abnormalities) that other methods like radiography cannot detect.^{17,40,41} Detection of early cartilage and bone erosion is essential so that therapy can be initiated as early as possible following reliable diagnosis.⁴²

The contrast agents that are normally used in MRI are based on either particles with paramagnetic (like gadolinium) or superparamagnetic behavior (like iron oxide

nanoparticles).⁴³ As a gold standard for synovial imaging, gadolinium-enhanced MRI is used for the detection of synovial hypervascularity of arthritic joints and early cartilage degeneration without structural change. Gadolinium-enhanced MRI has limitations, as the synovial contrast enhancement is nonspecific and the enhancement patterns of normal and inflamed joints often overlap. The need for selective tissue-, cell-, and receptor-specific contrast agents is obvious since they should i) provide much improved detection of earlier degenerative events and responses to therapy, ii) better define and characterize the inflammatory and degradative processes, and iii) be more efficient ie, at lower doses. Targeted SPION offer a significant opportunity for improved diagnostic MRI imaging.⁴⁴

Similar to gadolinium, which is highly toxic as an element so needs to be given as a complex, iron oxide nanoparticles also need to be covered by a coating to guarantee safe application. The size of the particles and their coating characteristics determine pharmacokinetics, biodistribution and specificity, and possibly toxicity. SPION-based contrast agents have been developed as liver specific contrast agents because of their specificity for reticuloendothelial cells. Depending on their size, SPION of a larger hydrodynamic size are preferentially taken up by liver Kupffer cells. In contrast, smaller sizes of SPION are mostly taken up by macrophages of the lymph nodes, or are directed to liver, lung, or spleen, or may cross the blood–brain barrier.⁴⁵ SPION are mostly taken up intracellularly by endocytosis.^{46–48} Various studies show that dextran coated SPION uptake is driven by scavenger receptors, which are then metabolized in the lysosomes into ferritin or hemoglobin and as a result of this become part of the normal blood pool.^{43,47,49}

SPION coated with vinyl-alcohol/vinyl-amine copolymer (amino-polyvinyl alcohol [a-PVA]), as used in our study, are taken up *in vitro* by synovial cells, chondrocytes, and osteoblasts after a 2 hour incubation.⁵⁰ Furthermore, a very recent study revealed that amino-polyvinyl alcohol coated superparamagnetic iron oxide nanoparticles (a-PVA-SPION) were also taken up by human mesenchymal stromal cells (MSCs), which are of considerable interest in cell-based therapies. The study also demonstrated that these high contrast particles are non-toxic and suitable for MSC visualization in MRI, *in vitro* as well as *in vivo*, in rats.⁵¹

The colloidal stability and, therefore, cell uptake and cytotoxicity are strongly influenced by the composition of the SPION polymer coat (eg, polyvinyl alcohol, vinyl alcohol/vinyl amine copolymer, or polyethylenimine). Although these polymer-coated SPION are stable in water

and phosphate buffered saline (PBS), the choice of medium largely influences the cellular uptake of these particles.^{52,53}

Although various industrially produced SPION are being investigated in clinical trials and seem to produce very low side effects, the potential for cytotoxicity of these particles is still an important issue of debate. Investigation of the immune responses of dendritic cells to PVA coated SPION show a certain degree of immunomodulation.⁵⁴ Such properties – but also the impact of SPION on cell proliferation and cell functionality – depend upon the physicochemical properties of the particle surfaces and the cells used for such tests.^{55,56}

Therefore, we have investigated the interactions of amino functionalized SPION, developed for early diagnosis in arthritis, with different human immune cells obtained from patients suffering from RA and from healthy donors (HD). We investigated potential effects on specific immune cell viability as well as cytokine secretion as an indicator of immune cell activation.

Materials and methods

Materials

In this study we used a-PVA-SPION which were synthesized and characterized following a protocol previously described.⁵¹ The mean diameter of the magnetic cores (d_{core} in nm) of the a-PVA-SPION was determined by counting 400 diameters observed in transmission electron microscopy (TEM; CM12 from Philips Electron Optics, Zürich, Switzerland). Hydrodynamic size (d_{H} in nm) and Zeta potential (Z potential in mV) were measured at around pH 7 by dynamic laser scattering using a Malvern NanoZS (Nanosizer 7.2 software) using the theoretical refractive index of 2.42 of magnetite.⁵⁷ The saturation magnetization (Sat mag in emu/g) was obtained from freeze dried powder of SPION before PVA coatings using a superconducting quantum interference device.⁵⁸

Antibodies and reagents

Lipopolysaccharide (LPS), lectin from *Phaseolus vulgaris* (PHA), Brefeldin A, Saponin, and carboxyfluorescein diacetate N-succinimidyl ester (CFSE) were from Sigma-Aldrich Co., St Louis, MO, USA. Paraformaldehyde was from Carl Roth GmbH + Co. KG (Karlsruhe, Germany). Recombinant human macrophage colony stimulating factor (rhM-CSF) was from ImmunoTools GmbH (Friesoythe, Germany).

For flow cytometry, Priviligen® human immunoglobulin from CSL Bhering, (King of Prussia, PA, USA) was used. Anti-human-IL1 β -PE, anti-human-CD3-Pacific-blue, anti-human-CD14-APC-Cy7, anti-human-CD19-PE-Cy7, and

Annexin V/7AAD apoptosis kit were from BD Biosciences (San Jose, CA, USA) and anti-human-CD15-APC was from Miltenyi Biotech GmbH (Bergisch Gladbach, Germany).

Whole blood survival analysis, sample preparation, and stimulation

Venous blood (obtained from RA patients or HD) was collected in heparinized tubes. All patients met the American Rheumatism Association criteria (1987) for RA.⁵⁹ The characteristics of RA patients are summarized in Table S1. The study protocol was approved by the responsible local administrative body and ethics committee. RA patients as well as HD provided written informed consent before enrollment.

Immediately after retrieval of the blood samples, 100 μ L of whole blood was diluted with 100 μ L Roswell Park Memorial Institute (RPMI) 1640 culture medium (Thermo Fisher Scientific, Waltham, MA, USA) supplemented with 100 U/mL penicillinG, 100 μ g/mL streptomycin (both from PAA Laboratories) and 50 μ M β -mercaptoethanol (Sigma-Aldrich Co.) in deep-well-plates (Sarstedt AG & Co., Nuembrecht, Germany). Cells were stimulated with LPS (1 μ g/mL), PHA (5 μ g/mL), a-PVA-SPION (1 μ g/mL, 10 μ g/mL, 100 μ g/mL, 1,000 μ g/mL) or left untreated and incubated for 20 hours in a humidified incubator at 37°C (18% O₂/5% CO₂). Afterwards, supernatants were collected, immediately frozen, and stored at –80°C for cytokine secretion analysis and cells were prepared for flow cytometry (see section *Flow cytometric analysis*). For intracellular IL1 β analysis, secretion was blocked by adding 10 μ g/mL Brefeldin A followed by an additional incubation for 3 hours at 37°C.

Quantification of secreted cytokines

Secreted cytokines were quantified by Bio-Plex® Pro Cytokine 27-Plex Panel Human Group I on a Bio-Plex® 200 system with high-throughput fluidics (all Bio-Rad Laboratories Inc., Hercules, CA, USA). Intracellular IL1 β concentration was determined via flow cytometry as described below.

Isolation and stimulation of human CD14⁺ monocytes

Peripheral blood mononuclear cells were isolated from leukocyte apheresis filters of healthy blood donors by density gradient centrifugation using the Ficoll-Paque™ Plus technique (Amersham Biosciences Europe GmbH, Freiburg, Germany). CD14 positive monocytes were enriched up to 99% purity and >95% viability (data not shown) by MACS® Technology (Miltenyi Biotec GmbH) using anti-human CD14 conjugated magnetic beads as described by the manufacturer.

Fresh isolated CD14 positive monocytes were labeled with 0.3125 mM CFSE for 3.5 minutes at room temperature (RT) and cultured at 2×10^6 cells/mL in RPMI 1640 culture medium supplemented with 10% (v/v) heat-inactivated fetal calf serum (FCS) (Sigma-Aldrich Co.), 100 units/mL penicillinG, 100 $\mu\text{g/mL}$ streptomycin, and 50 μM β -mercaptoethanol for 20 hours in 24-well cell culture plates. Cells were stimulated with LPS (1 $\mu\text{g/mL}$), a-PVA-SPION (1 $\mu\text{g/mL}$, 10 $\mu\text{g/mL}$, 100 $\mu\text{g/mL}$, and 1,000 $\mu\text{g/mL}$) or left untreated for 20 hours in a humidified incubator at 37°C (18% O_2 /5% CO_2). Afterwards, cells were detached with Accutase (PAA Laboratories GmbH, Cölbe, Germany) and further prepared for flow cytometry (see section *Flow cytometric analysis*).

Generation of monocyte-derived macrophages (MDMs)

Freshly isolated CD14 positive monocytes (see section *Isolation and stimulation of human CD14⁺ monocytes*) as 6 mL of 2×10^6 cells/mL PBS were added to a covered 10 cm diameter Petri dish. Cells were incubated for 2 hours at 37°C (18% O_2 /5% CO_2). Afterwards, supernatants were carefully removed and 6 mL fresh RPMI 1640 medium containing 50 ng rhM-CSF was added to the adherent cells. After 24 and 72 hours at 37°C (18% O_2 /5% CO_2) medium was refreshed by adding 6 mL of RPMI 1640 containing 50 ng rhM-CSF. After a total of 6 days incubation, the differentiated MDMs were detached with Accutase and the remaining adherent cells were removed with a cell scraper.

Stimulation of monocytes/MDMs with a-PVA-SPION

Two different approaches have been followed: I) MDMs were labeled with 0.3125 mM CFSE for 3.5 minutes at RT and 5×10^4 cells/200 μL RPMI 1640 medium/well were incubated for 20 hours in 48-well cell culture plates. MDMs were stimulated with LPS (1 $\mu\text{g/mL}$), a-PVA-SPION (1 $\mu\text{g/mL}$, 10 $\mu\text{g/mL}$, 100 $\mu\text{g/mL}$, and 1,000 $\mu\text{g/mL}$) or left untreated in a humidified incubator at 37°C (18% O_2 /5% CO_2) and afterwards prepared for flow cytometry (see section *Flow cytometric analysis*). II) Monocytes were labeled with 0.3125 mM CFSE for 3.5 minutes at RT and 2×10^5 cells/100 μL PBS/well were incubated for 1 hour in 48-well cell culture plates for attachment. Afterwards, the PBS-buffer was removed and 200 μL RPMI 1640 medium containing 50 ng rhM-CSF was added. The cells were stimulated with 1 $\mu\text{g/mL}$ LPS or left untreated for 6 days. Stimulation with a-PVA-SPION during the differentiation process to

MDMs was performed as follows: a-PVA-SPION (1 $\mu\text{g/mL}$, 10 $\mu\text{g/mL}$, 100 $\mu\text{g/mL}$, and 1,000 $\mu\text{g/mL}$) were added at day zero within the first addition of RPMI 1640 medium containing 50 ng rhM-CSF or 24 hours later, when refreshing the medium or 72 hours later, when refreshing the medium again. Addition of fresh medium was performed as described in the section *Generation of monocyte-derived macrophages (MDMs)*. Cells were incubated in a humidified incubator at 37°C (18% O_2 /5% CO_2) until MDMs could be harvested after 6 days and prepared for flow cytometry.

Flow cytometric analysis

Whole blood extracellular staining

After lysis of erythrocytes, with a solution of potassium bicarbonate (KHCO_3) 0.01 M pH 7.5, ammonium chloride (NH_4Cl) 0.155 M, ethylenediaminetetraacetic acid (EDTA) 0.1 mM, cells were washed with PBS and transferred to 96-well V-bottom assay plates. Non-specific binding was blocked by pre-incubation with human immunoglobulin (5 mg/mL) for 10 minutes at 4°C . After washing the cells with PBS, an antibody-mix (anti-CD3, anti-CD14, anti-CD19, and anti-CD15) was added (all antibodies diluted 1/50 in PBS or 1/10 when additionally stained intracellularly) and incubation was performed for 15 minutes at 4°C in the dark. Cells were washed twice and cells were analyzed within 1 hour by flow cytometry using the MACSQuant analyzer equipped with plate MACSQuant MiniSampler. The acquired data were analyzed using FlowJo 7.6.1 software. For the investigation of survival of positive cells for the surface markers (CD3, CD14, CD19 or CD15), Annexin V-PE/7-AAD staining was performed according to the manufacturer's instructions after cell washing (to determine apoptosis). After data acquisition by flow cytometry, these populations were analyzed for the percentage of Annexin V (apoptotic) or 7-AAD (dead) positive cells.

Whole blood intracellular staining

After extracellular antibody staining, cells were fixed with 2% paraformaldehyde and subsequently cell membranes were permeabilized by washing twice with 0.5% Saponin-PBA (PBS/BSA/Acid). Non-specific binding was blocked with human immunoglobulin (5 mg/mL) for 10 minutes at RT. The washed (Saponin-PBA) cell pellets were resuspended in 10 μL of undiluted anti-human-IL1 β -PE antibody and incubated for 15 minutes at RT in the dark. After two final washing steps cells were analyzed by flow cytometry using the MACSQuant analyzer equipped with plate MACSQuant MiniSampler. The acquired data were analyzed using FlowJo

7.6.1 software, where positive cells for the according surface marker (CD3, CD14, CD19 or CD15) were selected and these populations were analyzed for the percentage and number of IL1 β positive cells.

Monocyte/MDM survival analysis

After detachment of the cells with Accutase and cell scraper, Annexin V-PE/7-AAD staining was performed according to the manufacturer's instructions and cells were analyzed within 1 hour by flow cytometry using the MACSQuant analyzer equipped with plate MACSQuant MiniSampler.

TEM analysis

Human monocytes (2×10^6 cells/mL) and MDMs (1×10^6 cells/mL) were incubated for 24 hours with PVA-SPION (1,000 $\mu\text{g/mL}$) or left untreated. Cells were prepared for TEM analysis as previously described.⁶⁰ Ultrathin sections were analyzed using routine TEM (EM 906; Carl Zeiss AG Oberkochen, Germany) techniques; micrographs were prepared and further processed using Adobe Photoshop CS software.

Statistical analysis

Statistical tests for the flow cytometry were performed using Graph Pad Prism Software. Data are presented as box and whiskers, whiskers from minimum to maximum or mean with standard deviation of at least four independent experiments. Multiple comparisons were analyzed by one-way or two-way analysis of variance (ANOVA) as indicated with Bonferroni's multiple comparison post hoc tests for normally distributed data. Probability values of $P < 0.05$ were considered to be statistically significant ($***P < 0.001$; $**P < 0.01$; $*P < 0.05$).

Results

Physicochemical characterizations of a-PVA-SPION

The d_{core} of the a-PVA-SPION (Table 1) is approximately 7 nm which is small enough to have superparamagnetic properties confirmed by Sat mag of 54 emu/g and the absence of hysteresis.^{58,61} The hydrodynamic size of the a-PVA-SPION is approximately 30 nm at pH 7 and the Zeta potential is approximately 22 mV indicating the positive initial charges of these nanoparticles at physiological pH.

Table 1 Physicochemical characterizations of a-PVA-SPION

Nanoparticles	d_{core} (nm)	d_{H} (nm)	Z potential (mV)	Sat mag (emu/g)
a-PVA-SPION	7.2 ± 2.5	31 ± 10	22 ± 6	54

Abbreviations: d_{core} , mean diameter of the magnetic cores; d_{H} , hydrodynamic diameter; Z potential, Zeta potential; Sat mag, the saturation magnetization; a-PVA-SPION, amino-polyvinyl alcohol coated superparamagnetic iron oxide nanoparticles.

a-PVA-SPION do not affect the survival of human immune cells within whole blood

We obtained venous blood from 19 patients with RA and 18 HD. To analyze the influence of a-PVA-SPION on human immune cells, we quantified the number of Annexin V (apoptotic) or 7AAD (dead) positive cells in combination with the cell specific extracellular markers (CD-antigens). For both RA (Figure 1A) and HD (Figure 1B), and for all cell types analyzed, we did not find any significant influence of a-PVA-SPION on apoptosis and viability. Furthermore, we found that CD3 positive T cells show the lowest apoptosis and necrosis rates as demonstrated by the low percentages of Annexin V and 7AAD positive cells (median values between 20% and 42%) whereas CD19 positive B cells showed the highest percentages of apoptosis and necrosis within their cell population (median values of Annexin V and 7AAD positive cells between 52% and 78%).

For RA patients (Figure 1A) we observed a non-significant a-PVA-SPION concentration-dependent increase in Annexin V positive CD14 monocytes (median value for a-PVA-SPION at 1 $\mu\text{g/mL}$ =45.9% and at 1,000 $\mu\text{g/mL}$ =68.2%). This was not observed in HD (Figure 1B), where we found a minimal non-significant a-PVA-SPION concentration-dependent decrease of Annexin V positive CD15 granulocytes (median value for PVA-SPION at 1 $\mu\text{g/mL}$ =60.5% and at 1,000 $\mu\text{g/mL}$ =45.5%).

a-PVA-SPION trigger cytokine secretion by dose-dependent cell activation in whole blood

From whole blood viability experiments, supernatants were collected prior to sample preparation for flow cytometry and secreted cytokines were analyzed using the Bio-Plex[®] Suspension array system. We examined supernatants from nine RA patients and four HD and found significantly increased secretion of the following cytokines in the presence of a-PVA-SPION (at 100 $\mu\text{g/mL}$) when compared to the untreated control (Figure 2): RA patients – IL1 β , IL4, IL6, IL8, IL9, IFN- γ , MCP-1, MIP1a, MIP1b, and PDGF; HD – IL1 β , IL4, IL6, IL8, MIP1b, and RANTES. One example is given in Figure 2A:

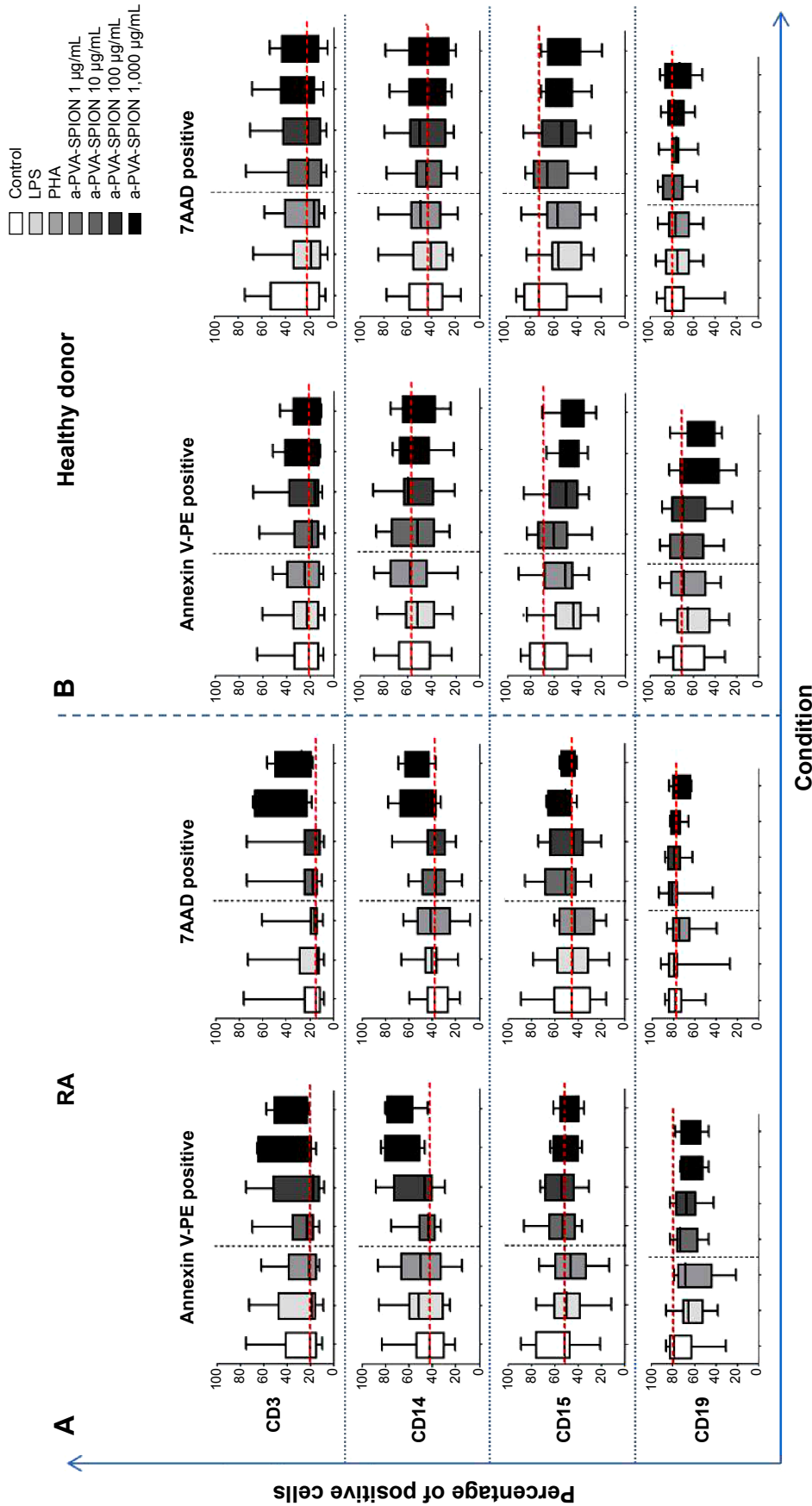


Figure 1 Influence of a-PVA-SPION on the viability of human immune cells.

Notes: Whole blood survival analysis was performed with blood samples obtained from patients with rheumatoid arthritis (**A**) n=19; or healthy donors (**B**) n=18. Cells were analyzed for specific surface markers, Annexin V (apoptosis marker) and 7AAD (necrosis marker). Data are given as box and whiskers; whiskers represent minimum to maximum; one-way ANOVA.

Abbreviations: RA, rheumatoid arthritis; ANOVA, analysis of variance; LPS, lipopolysaccharide; PHA, Phaseolus vulgaris; a-PVA-SPION, amino-polyvinyl alcohol coated superparamagnetic iron oxide nanoparticles; 7AAD, 7-amino-actinomycin D; PE, phycoerythrin.

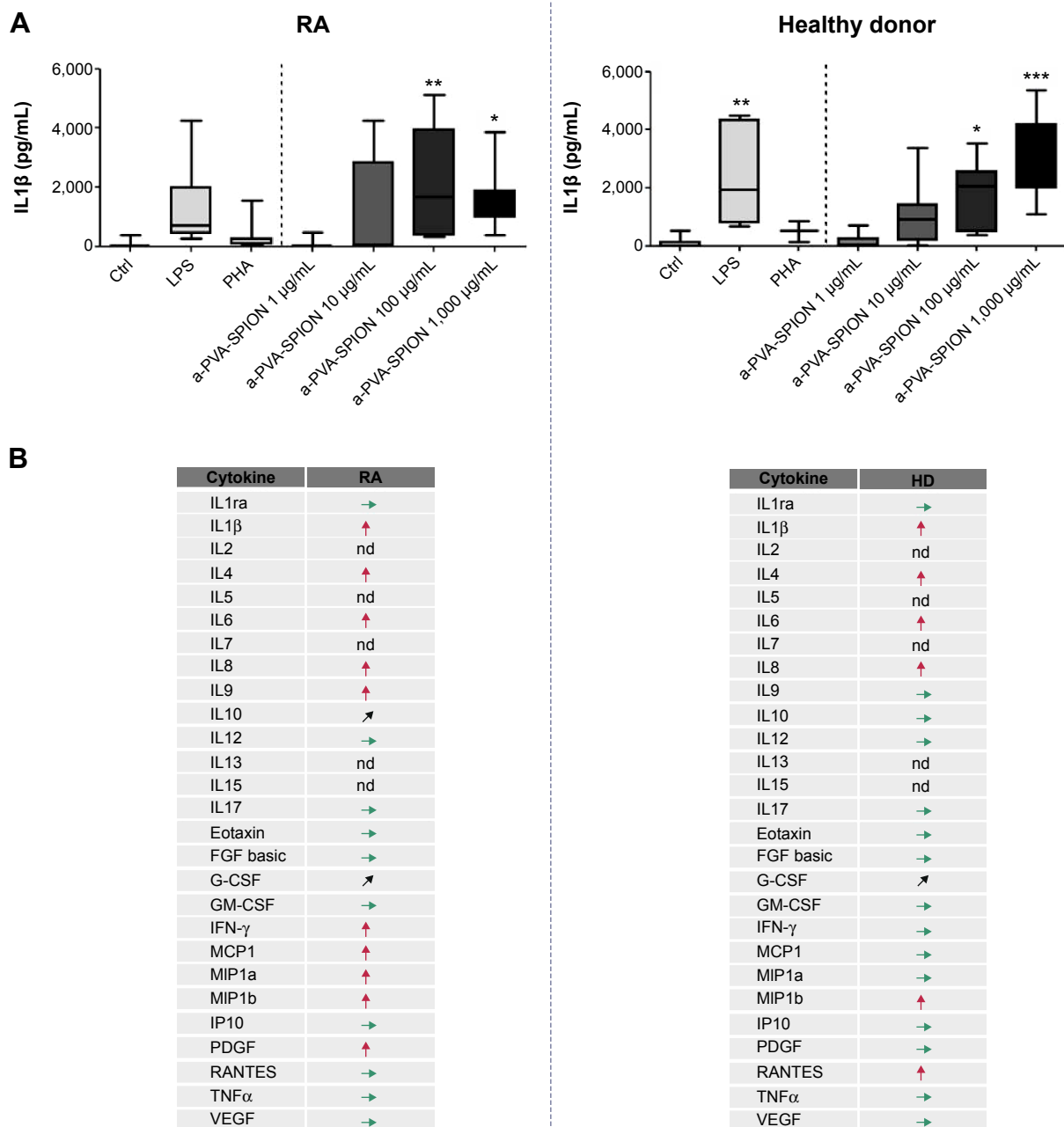


Figure 2 Comparison of the influence of a-PVA-SPIION on whole blood cytokine secretion by immune cells of RA patients and healthy donors.

Notes: Supernatants were collected from whole blood survival assays (blood samples obtained from patients with RA [n=9, left] or healthy donors [n=4, right]). One example for IL1β secretion is given in detail in (A), and a tabular overview for all measured cytokines is presented in (B). Red arrow: cytokines that were significantly increased at SPIION concentration 100 μg/mL compared to untreated control; black arrow: cytokines significantly increased at any other SPIION concentration than 100 μg/mL compared to untreated control; green arrow: cytokines that remained unchanged; nd: not detectable; data are given as box and whiskers; whiskers represent minimum to maximum; one-way ANOVA: ***P<0.001; **P<0.01; *P<0.05.

Abbreviations: RA, rheumatoid arthritis; ANOVA, analysis of variance; Ctrl, control; LPS, lipopolysaccharide; PHA, *Phaseolus vulgaris*; HD, healthy donors; a-PVA-SPIION, amino-polyvinyl alcohol coated superparamagnetic iron oxide nanoparticles.

unstimulated cells secrete IL1β at a median concentration of 55 pg/mL whereas 20 hours incubation with a-PVA-SPIION led to median values of 67.2 pg/mL (at 1 μg/mL), 1,075 pg/mL (at 10 μg/mL), 2,076 pg/mL (at 100 μg/mL), and 1,573 pg/mL (at 1,000 μg/mL) in blood samples obtained from RA patients. For HD we found similar results: control

cells secrete IL1β at median concentration of 96.3 pg/mL whereas a-PVA-SPIION incubation resulted in IL1β median values of 155.4 pg/mL (at 1 μg/mL), 1,105 pg/mL (at 10 μg/mL), 1,860 pg/mL (at 100 μg/mL), and 3,025 pg/mL (at 1,000 μg/mL) (Figure 2A; Tables S2 and S3). The secretion of G-CSF was found to be significantly increased at

1,000 $\mu\text{g/mL}$ of a-PVA-SPION for both, RA patients and healthy subjects, whereas IL10 was found to be significantly increased at 1,000 $\mu\text{g/mL}$ of a-PVA-SPION in RA patients. All other cytokines were unaffected by a-PVA-SPION treatment (see Tables and Figure 2B; all median concentrations are listed in Tables S2 and S3).

IL1 β secretion is increased due to a-PVA-SPION treatment, with human monocytes and granulocytes being the primary sources of induced IL1 β

To identify the cell population(s) responsible for the increased IL1 β secretion, we separated CD4 positive T cells, CD14 positive monocytes, and CD15 positive granulocytes from whole blood obtained from HD and incubated them for 20 hours at varying a-PVA-SPION concentrations followed by subsequent IL1 β measurement. Surprisingly, there was no influence of a-PVA-SPION on IL1 β secretion in each of the single cell populations (data not shown), ie, IL1 β was not detectable (except within the corresponding controls). This was in contrast to findings with whole blood experiments. So we repeated the whole blood assay and stained for both specific extracellular cell identity markers and for intracellular IL1 β that accumulated in the presence of Brefeldin A. We found a significant or numerical a-PVA-SPION-induced increase in the percentage of IL1 β positive cells for all cell types (Figure 3). Granulocytes represented the most commonly stained cell type followed, in descending order, by T cells, monocytes, and B cells, the latter representing the smallest cell population in the blood. The results were normalized to the respective cell counts. Thus both CD15 positive granulocytes and CD14 positive monocytes were identified as the major populations of IL1 β producing cells (Figure 3). A low a-PVA-SPION concentration (1 $\mu\text{g/mL}$) did not influence IL1 β secretion, whereas medium a-PVA-SPION concentrations led to a strong numerical (10 $\mu\text{g/mL}$) or significant (100 $\mu\text{g/mL}$, one-way ANOVA, $P < 0.05$) increase in the number of IL1 β positive cells compared to untreated control. A decline in the number of IL1 β positive cells was found for the highest a-PVA-SPION concentration (1,000 $\mu\text{g/mL}$) compared with concentrations of 100 $\mu\text{g/mL}$ or 10 $\mu\text{g/mL}$ (Figure 3).

a-PVA-SPION do not affect the viability of human monocytes but induce increased survival of MDMs in a dose dependent manner

The main task of monocytes and macrophages is the clearance of pathogens and toxins from the human body. Thus,

we analyzed the effects of a-PVA-SPION on the survival of i) isolated monocytes and ii) MDMs (Figure 4). In general, monocytes as well as MDMs exhibit a decrease in numbers over 20 hours. Thus, we have normalized the results to the unstimulated control (considered to be 100%). When adding a-PVA-SPION to human CD14 positive monocytes at varying concentrations for 20 hours ($n=8$), the viability was not affected by nanoparticles compared to the unstimulated control. However, we found a significant decrease in the percentage of viable human monocytes when stimulated with a positive control, namely LPS (38.9% Annexin V/7AAD negative cells) (Figure 4A). Interestingly, MDMs ($n=10$) showed a significant increase in the percentage of viable cells at medium and high a-PVA-SPION concentrations compared to the unstimulated control (311.5% Annexin V/7AAD negative cells for a-PVA-SPION at 100 $\mu\text{g/mL}$ and 450.7% Annexin V/7AAD negative cells for a-PVA-SPION at 1,000 $\mu\text{g/mL}$) (Figure 4B).

a-PVA-SPION affect the survival of MDMs primarily when added very early during the differentiation process

Next we analyzed the influence of nanoparticles on the naturally occurring differentiation process by adding

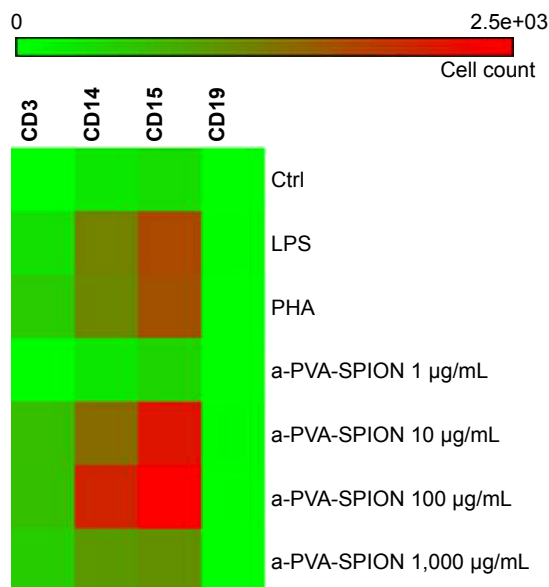


Figure 3 Impact of a-PVA-SPION on the number and distribution of IL1 β positive cells.

Notes: Whole blood samples obtained from healthy donors ($n=6$) were incubated for 20 hours at varying a-PVA-SPION concentrations. Cellular transport was blocked by adding Brefeldin A and cells were stained for extracellular markers as well as for intracellular IL1 β . Normalization according to cell count was performed and data were visualized with the help of a heat map. Data are given as number of IL1 β positive cells.

Abbreviations: Ctrl, control; LPS, lipopolysaccharide; PHA, *Phaseolus vulgaris*; a-PVA-SPION, amino-polyvinyl alcohol coated superparamagnetic iron oxide nanoparticles.

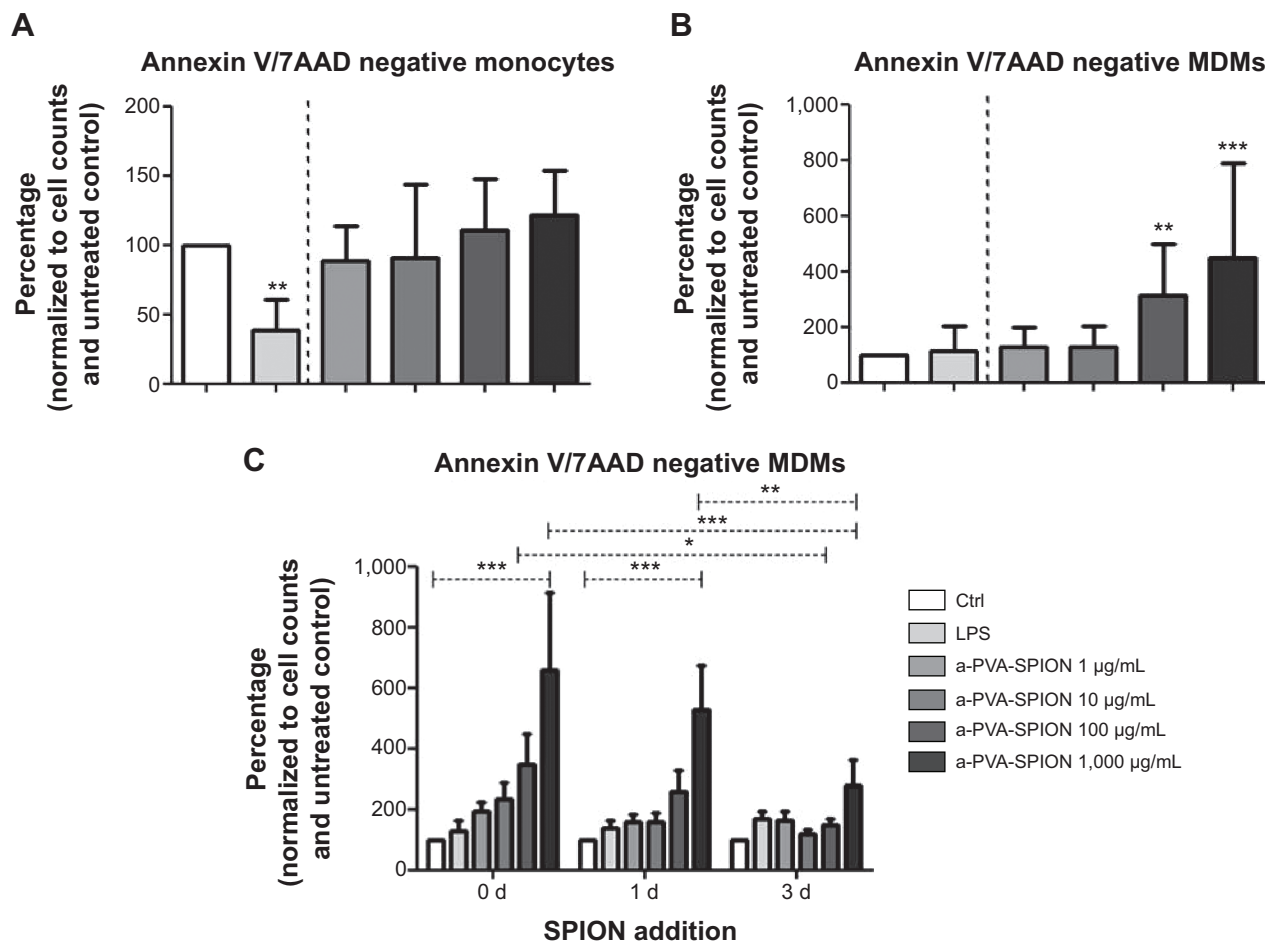


Figure 4 Influence of a-PVA-SPION on the viability of isolated monocytes and monocyte-derived macrophages (MDMs).

Notes: Human monocytes (**A**) $n=8$ or MDMs (**B**) $n=10$ were incubated for 20 hours with or without SPION at varying concentrations and the percentage of Annexin V/7AAD negative cells was determined. In another experiment the SPION were added during the differentiation process from monocytes to MDMs at the beginning (0 d), after 24 hours (1 d) or after 72 hours (3 d) and the percentage of Annexin V/7AAD negative cells was determined after a total of 6 days (**C**) $n=7$. Results are normalized to cell counts and untreated control. Data are given as mean \pm standard deviation; two-way ANOVA: *** $P<0.001$; ** $P<0.01$; * $P<0.05$.

Abbreviations: d, day(s); ANOVA, analysis of variance; Ctrl, control; LPS, lipopolysaccharide; 7AAD, 7-amino-actinomycin D; a-PVA-SPION, amino-polyvinyl alcohol coated superparamagnetic iron oxide nanoparticles.

a-PVA-SPION at different time points during this process. In summary, we found that the survival of MDMs ($n=7$) was significantly increased in a dose dependent manner, especially when a-PVA-SPION were added very early in the differentiation process (Figure 4C [0 days]). The later the nanoparticles were added, the weaker the observed increase in cell survival (Figure 4C).

a-PVA-SPION at 1,000 µg/mL are internalized by human monocytes and MDMs

Monocytes and macrophages are professional phagocytes. Thus, we expected that the a-PVA-SPION are internalized within these cells. However, we aimed to investigate the exact intracellular location of the SPION within the cells, and the extent of their accumulation. To this end, we performed TEM analyses of both untreated (Figure 5A, C) and

a-PVA-SPION (1,000 µg/mL) treated monocytes and MDMs (Figure 5B, D). We found a-PVA-SPION to be present in phagosomes within both monocytes (Figure 5B) and MDMs (Figure 5D). Interestingly, we observed phagosomes that were only partly filled, whereas others were tightly filled with a-PVA-SPION.

Discussion

There is considerable interest in the use of nanoparticulate materials in the fields of pharmaceuticals and health care industries, as well as in clinical medicine, either for soft materials like liposomes (mostly used for drug delivery) or inorganic hard materials (developed mainly for in vitro or in vivo diagnostics) like SPION. RA is a chronic, systemic, inflammatory autoimmune disease that in large part is diagnosed only after irreversible joint damage is observed using existing imaging modalities. One promising opportunity to

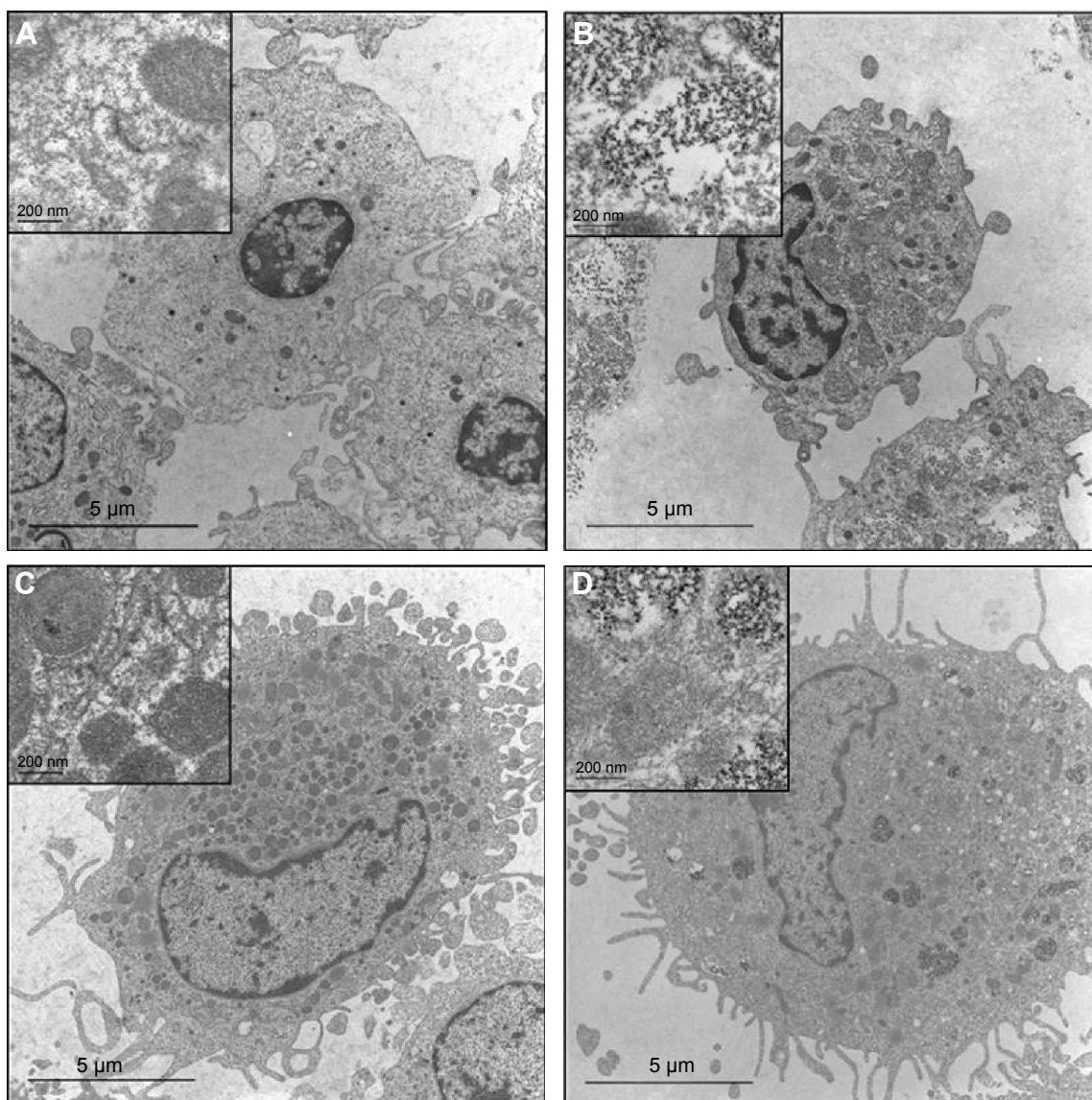


Figure 5 Transmission electron microscopy (TEM) of human monocytes and monocyte-derived macrophages (MDMs) incubated with a-PVA-SPIION.

Notes: Human monocytes were incubated for 24 hours without (A) or with (B) a-PVA-SPIION (1,000 μg/mL). (C, D) Human MDMs were incubated for 24 hours without (C) or with (D) a-PVA-SPIION (1,000 μg/mL). Both cell types showed uptake of SPIION in phagosomes; some are scattered but others were tightly filled with nanoparticles (scale indicated on the figure).

Abbreviation: a-PVA-SPIION, amino-polyvinyl alcohol coated superparamagnetic iron oxide nanoparticles.

achieve an early diagnosis of RA and other inflammatory and degenerative joint diseases, before significant joint destruction occurs, is represented by the use of SPIION as MRI contrast agents. SPIION are internalized in cells of the mononuclear phagocyte system, in tumor cells as well as in MSCs, which makes these particles of interest for use in MRI.^{51,62,63} The main advantage is that SPIION are taken up by immune cells like monocytes, which differentiate to macrophages while migrating into the inflamed tissue. Thus, tissue inflammation can be visualized by MRI in very early disease which enables early therapeutic intervention prior to onset of significant joint damage. This approach

is very much in agreement with current treat-to-target guidelines that recommend early and rigorous treatment to achieve low disease activity or remission targets as rapidly as possible.

However, prior to routine usage of novel engineered SPIION formulations in medical applications, it is essential to determine any potential toxic effects of these particles.^{64,65} Therefore, in this study we investigated *in vitro* the interaction between a-PVA-SPIION and human immune cells. We aimed to clarify possible toxic effects of a-PVA-SPIION on human immune cells using a whole blood system to retain the interaction between the different blood cells and/or soluble

factors, thereby mimicking as far as possible the *in vivo* situation. Moreover, the influence of a-PVA-SPION was further analyzed in greater detail using isolated pure cell populations of monocytes or MDMs.

The following major findings emerged from this study: i) a-PVA-SPION had no effect on the survival of human immune cells in the whole blood analysis but ii) the cytokine secretion was dose-dependently increased in this short term study. iii) Using the example of IL1 β , we were able to show that monocytes and granulocytes were the primary source for this cytokine. Furthermore, when analyzing monocytes and MDMs as pure cell populations in more detail, we found that iv) a-PVA-SPION improved the survival of MDMs in a dose dependent manner and that v) an early addition of a-PVA-SPION during the differentiation process led to the strongest impact on macrophage survival. Additionally, we were able to show that vi) a-PVA-SPION were internalized in human monocytes and MDMs. The main conclusion of these findings is that there is indeed a potentially clinically relevant interaction of a-PVA-SPION with human immune cells.

Immune cells represent the primary defense system against pathogens. Cells bearing phagocytic functions (especially professional phagocytes such as monocytes, macrophages, and polymorphonuclear granulocytes or neutrophils) are equipped with receptors that recognize foreign invaders.⁶⁶ One reaction after the detection of external substances within the human body is the secretion of cytokines and chemokines to alert and attract further immune cells that migrate into the affected area. In this study we demonstrated that the secretion of different cytokines is increased due to a-PVA-SPION treatment in a whole blood system. This is in contrast to the results on isolated human MDMs, which found no difference in the production of several cytokines (IL12, IL6, TNF- α , and IL1 β) in MDMs that were stimulated with dextran coated ultrasmall superparamagnetic iron oxide particle (Ferumoxtran-10) compared to untreated controls.⁶⁷ Whether or not the effect of the SPION in our study is persistent or short-lived is not known. Further studies are required to examine this.

Another interesting study revealed that LPS pre-treated monocyte-derived dendritic cells co-cultured with CD4 T cells show a SPION dependent decrease in IL1 β secretion.⁵⁴ When analyzing the IL1 β secretion in more detail in our study, we first incubated isolated T cells, monocytes and granulocytes with a-PVA-SPION in order to quantify the supernatants with regard to this cytokine. Interestingly, with the exception of the respective control, there was no measurable amount of IL1 β in the supernatants after a-PVA-SPION

stimulation (data not shown). But for both the whole blood cytokine secretion assay and the whole blood intracellular IL1 β staining the results on cytokine production were precise and reproducible. Thus, we assume that analysis of isolated single cell populations is not representative of the complex human *in vivo* situation where interactions between immune cells and other cells and soluble factors exist. Nevertheless, the question arises whether this effect on cytokine secretion (especially IL1 β) is of importance for the assessment of SPION toxicology. In freshly drawn blood, IL1 β serum levels of 5.5 pg/mL for RA patients and 2.6 pg/mL for HD have been reported.⁶⁸ After incubation, we found median values of 96.3 pg/mL and 54.9 pg/mL IL1 β for RA patients and HD in the untreated control, respectively. The difference with regard to freshly drawn blood may result from the incubation time of 20 hours, when apoptosis and consequently cell lysis may take place (see results of the whole blood survival assay), which mediate the release of soluble factors. These serum values were increased up to 2,000 pg/mL in our positive control, which is represented by LPS at high dosages. Similar values could be observed for a-PVA-SPION used at concentrations >10 μ g/mL. These results show that the batch of a-PVA-SPION used does indeed trigger a dose-dependent effect that is comparable to an inflammatory process. Thus, for some *in vitro* studies these SPION are usable but they are not feasible for *in vivo* imaging techniques that exceed local a-PVA-SPION concentrations of 10 μ g/mL. Further modulation of the formulation may possibly reduce this cytokine secretion thereby overcoming this problem, since we were able to show that there is no toxic effect with regard to survival and since we do not expect our immune cells to be totally inert when in contact with a-PVA-SPION. Furthermore, when thinking of the *in vivo* situation, one has to keep in mind that the properties of the peripheral blood and the tissues, especially in the case of inflammation, vary strongly. One example is oxygen availability, which is sufficient in the blood stream but deficient in inflamed tissues. Thus, immune cells have to adapt to these changes when infiltrating the inflamed area.⁶⁹ These environmental changes could result in a completely different behavior of immune cells, compared to our findings for peripheral blood cells. Another important consideration is the concentration of SPION that is attained *in vivo* when used as a contrast agent in clinical studies. Both questions remain to be analyzed in future studies.

A key immune cell, whose principal task is the recognition, uptake and lysis of foreign invaders, is the professional phagocyte.⁶⁶ Such cells do not distinguish between microbes,

foreign cell debris, apoptotic cells or nanoparticles. Our results indicate that a-PVA-SPION are endocytosed by monocytes and MDMs, which is in accordance with other studies.^{67,70} When nanoparticles enter the cell, they can sometimes directly interact with the cytosol while during phagocytosis these particles are retained within vesicles.^{71–73} Hence, the intracellular location of nanoparticles depends on the uptake mechanism, which in turn depends on the type and the charge of the nanoparticle. For example Schweiger et al reported that negatively charged SPION were found firstly in endosomes and later in lysosomes, whereas positively charged particles were found exclusively in lysosomes.⁷¹ Since we used a-PVA-SPION in RPMI media without FCS, the Zeta potential of the particles is slightly positive, which leads to the assumption that the a-PVA-SPION are located in lysosomes. Rich in digestive enzymes, the lysosomes are responsible for the dissolution of pathogens. In general, monocytes are patrolling cells within the bloodstream that recognize pathogens or respond to inflammatory stimuli.⁶⁹ Once activated, a differentiation process is initiated that leads to the development of macrophages. These cells primarily ingest the pathogens for rapid killing and elimination and further regulate inflammatory reactions by producing pro-inflammatory or anti-inflammatory cytokines.^{74–76} Incubation with nanoparticles clearly triggers an activation process, whereas monocyte survival is not influenced. But when considering macrophages, it became obvious that the number of differentiated, viable cells was increased by a-PVA-SPION contact. Our results indicate that more monocytes are triggered to become macrophages, thereby assimilating and removing the foreign invader and enhancing immunity. This may facilitate further uses of nanoparticles. Monocytes and macrophages have recently been identified as interesting targets for both diagnosis and treatment of inflammatory diseases such as RA.⁷⁷ A directed uptake of specific nanoparticle formulations by monocytes and macrophages, which migrate into the inflamed tissue, is desirable on the one hand for imaging as shown by the NanoDiarA project results, thereby enabling early diagnosis of RA.⁷⁸ On the other hand, conjugation of nanoparticles that invade inflamed tissue with disease modifying drugs may optimize treatment strategies in the future.

Conclusion

Our study clearly demonstrates that there are mixed effects of a-PVA-SPION on human immune cells. Thus they do not appear to have any observable toxic effects on the survival of the immune cells investigated. However, they promote

differentiation to and increased survival of MDMs, which are important human professional phagocytes and thus favor the expression of immunity. The analysis of cytokine secretion revealed that a-PVA-SPION trigger dose-dependent release of some cytokines that is comparable to an acute inflammatory process of unknown duration. Moreover, the concentration of a-PVA-SPION that would be used in imaging remains to be determined. It may be similar or lower than the dose range used here. Therefore, we have obtained a nuanced conclusion, where the expected cytokine secretion as an immune response from professional phagocytes to a foreign substance was present but to a lesser extent than anticipated. SPION particles of this kind, which have already been generated with different surface modifications, will require examination in studies of this kind to determine the nature of the human immune responses they might generate. That these *in vitro* studies are of importance is apparent, especially once the local *in vivo* concentrations of SPION have been determined.

Outlook

In future, the use of nanomaterials is likely to increase since these substances open up previously unforeseen opportunities due to their special properties. In particular, nanoparticles represent a promising tool in the field of human medicine, for example as MRI contrast agents. It has been recently demonstrated that these particles are taken up by human MSCs and therefore could probably be used for cell-based therapies.⁵¹ Interestingly, we were able to demonstrate that these particles are also taken up by human macrophages, which are known to migrate into inflamed tissue. This might facilitate very early visualization of tissue inflammation by MRI, thereby enabling an early therapeutic intervention prior to onset of significant joint damage. Some studies using these nanoparticles proved their efficiency to target and monitor *in vivo* the inflammation of knee joints treated with dexamethasone using a-PVA-SPION labeled macrophages.⁷⁹

However, as with any novel technology, a number of critical issues must be resolved before using these a-PVA-SPION in human medicine. Specifically, these particles have first to be modified to a certain degree to be usable *in vivo* in the human system. One possibility would be to use the PVA coating properties to cross-link the polymer on the surface of the SPION or to functionalize them with reactive groups to target specific diseases.⁸⁰ Of course, these new modifications will need further *in vitro* studies. However, once a suitable formulation has been established, it may facilitate prevention of joint damage arising from RA.

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Supplementary materials

Table S1 Characteristics of patients with RA at screening visit

Characteristics	RA patients (n=19)
Female:male	16:3
Age (years)*	56.42 (24/75)
Duration of RA (years)*	10.3 (0.2/37.6)
RF*	231 (1/1,600)
ACPA*	241.7 (0/1,000)
DAS28*	4.81 (2.72/6.86)
ESR (mm/h)*	22.68 (2/60)
CRP (mg/dL)*	0.98 (0.03/6.56)
Glucocorticoid treatment	13 (68%)
DMARDs	19 (100%)
NSAIDs	11 (58%)

Note: Data are given as *mean (minimum/maximum) or number (%).

Abbreviations: RF, rheumatoid factor; ACPA, anti-citrullinated peptide antibodies; DAS28, disease activity score of 28 joints; ESR, erythrocyte sedimentation rate; CRP, c-reactive protein; DMARDs, disease modifying antirheumatic drugs; NSAIDs, non-steroidal anti-inflammatory drugs; RA, rheumatoid arthritis.

Table S2 Influence of a-PVA-SPION on the concentration of secreted cytokines in whole blood samples obtained from RA patients

Cytokine	Ctrl	LPS	PHA	a-PVA-SPION 1 µg/mL	a-PVA-SPION 10 µg/mL	a-PVA-SPION 100 µg/mL	a-PVA-SPION 1,000 µg/mL	Norm-serum- level*
IL1ra [pg/mL]	146	1,164	740	106	546	588	277	133
IL1β [pg/mL]	13	706	125	15	57	1,661	1,303	3
IL2 [pg/mL]	nd	nd	nd	nd	nd	nd	nd	1
IL4 [pg/mL]	8	18	18	7	12	20	15	3
IL5 [pg/mL]	nd	nd	nd	nd	nd	nd	nd	4
IL6 [pg/mL]	16	22,147	5,242	46	474	18,818	20,102	5
IL7 [pg/mL]	nd	nd	nd	nd	nd	nd	nd	21
IL8 [pg/mL]	81	8,025	7,460	508	5,762	17,279	20,050	4
IL9 [pg/mL]	52	129	129	39	68	118	139	12
IL10 [pg/mL]	3	290	220	3	9	70	125	4
IL12 [pg/mL]	31	67	66	36	37	68	82	16
IL13 [pg/mL]	nd	nd	nd	nd	nd	nd	nd	4
IL15 [pg/mL]	nd	nd	nd	nd	nd	nd	nd	1
IL17 [pg/mL]	17	101	182	17	74	128	120	21
Eotaxin [pg/mL]	91	115	87	103	86	101	86	31
FGF basic [pg/mL]	17	24	24	13	17	26	20	7
G-CSF [pg/mL]	35	189	82	29	75	627	1,040	52
GM-CSF [pg/mL]	158	118	121	114	99	145	147	6
IFN-γ [pg/mL]	380	1,249	901	357	487	1,110	1,055	77
MCP1 [pg/mL]	131	1,924	7,455	553	8,911	11,732	10,834	16
MIP1a [pg/mL]	21	1,708	1,381	24	115	1,886	1,845	7
MIP1b [pg/mL]	178	9,369	9,639	467	2,472	11,227	7,897	35
IP10 [pg/mL]	1,267	8,238	11,175	913	1,524	1,372	682	592
PDGF [pg/mL]	1,445	2,496	3,135	1,518	1,963	4,096	2,824	1,571
RANTES [pg/mL]	4,732	8,659	7,048	4,892	6,878	8,491	8,788	204*
TNFα [pg/mL]	40	1,273	661	30	118	348	113	35
VEGF [pg/mL]	48	145	136	70	45	77	203	11

Notes: Supernatants were collected from whole blood survival assays (blood samples obtained from patients with RA) after 20 hours with or without SPION treatment and analyzed by multiplex suspension array. Supernatants from nine RA patients were analyzed, median values are shown for the controls (LPS, PHA) and the different SPION concentrations (1–1,000 µg/mL). *Adapted from Up-regulation of cytokines and chemokines predates the onset of rheumatoid arthritis, *Arthritis & Rheumatism*, John Wiley and Sons. Copyright © 2010 by the American College of Rheumatology. ¹ & Copyright © 2010, Bio-Rad Laboratories, Inc. Adapted from Chapman P, Reyes C, Gupta V. Normal Physiological Levels of Human Cytokines Using Bio-Plex Pro™ Cytokine Assays. Bio-Rad Tech Note. 2010;6029. Available from: http://www.bio-rad.com/webroot/web/pdf/lsl/literature/Bulletin_6029.pdf. Accessed January 29, 2015.²

Abbreviations: nd, not detectable; Ctrl, control; RA, rheumatoid arthritis; LPS, lipopolysaccharide; PHA, *Phaseolus vulgaris*; a-PVA-SPION, amino-polyvinyl alcohol coated superparamagnetic iron oxide nanoparticles.

Table S3 Influence of a-PVA-SPION on the concentration of secreted cytokines in whole blood samples obtained from HD

Cytokine	Ctrl	LPS	PHA	a-PVA-SPION 1 µg/mL	a-PVA-SPION 10 µg/mL	a-PVA-SPION 100 µg/mL	a-PVA-SPION 1,000 µg/mL	Norm-serum- level*
IL1ra [pg/mL]	48	971	1,095	241	333	603	371	133
IL1β [pg/mL]	2	816	528	6	881	2,117	3,233	3
IL2 [pg/mL]	nd	nd	nd	nd	nd	nd	nd	1
IL4 [pg/mL]	3	15	20	4	19	21	20	3
IL5 [pg/mL]	nd	nd	nd	nd	nd	nd	nd	4
IL6 [pg/mL]	5	30,184	23,901	165	14,755	30,615	22,398	5
IL7 [pg/mL]	nd	nd	nd	nd	nd	nd	nd	21
IL8 [pg/mL]	110	15,261	14,485	731	19,613	18,145	15,709	4
IL9 [pg/mL]	161	225	327	198	237	290	317	12
IL10 [pg/mL]	5	600	497	6	62	80	203	4
IL12 [pg/mL]	24	84	79	30	57	47	84	16
IL13 [pg/mL]	nd	nd	nd	nd	nd	nd	nd	4
IL15 [pg/mL]	nd	nd	nd	nd	nd	nd	nd	1
IL17 [pg/mL]	53	144	299	51	148	143	156	21
Eotaxin [pg/mL]	66	91	93	80	85	98	82	31
FGF basic [pg/mL]	6	36	28	27	29	23	30	7
G-CSF [pg/mL]	47	371	256	27	386	538	1,672	52
GM-CSF [pg/mL]	137	198	199	103	134	159	112	6
IFN-γ [pg/mL]	546	3,291	2,710	1,165	799	2,487	1,729	77
MCPI [pg/mL]	111	2,305	4,509	409	6,419	7,188	6,121	16
MIPIa [pg/mL]	17	7,241	11,755	51	2,257	7,059	5,808	7
MIPIb [pg/mL]	214	14,073	16,550	786	11,718	16,665	12,236	35
IPI0 [pg/mL]	551	24,929	26,849	5,744	890	742	689	1,388
PGDF [pg/mL]	1,098	2,310	2,828	1,178	3,100	3,335	2,319	1,571
RANTES [pg/mL]	6,267	10,547	11,419	8,913	10,289	10,928	9,433	204*
TNFα [pg/mL]	61	4,863	3,186	75	1,083	1,315	220	35
VEGF [pg/mL]	32	136	191	59	76	64	125	11

Notes: Supernatants were collected from whole blood survival assays (blood samples obtained from healthy donors) after 20 hours with or without SPION treatment and analyzed by multiplex suspension array. Supernatants from four HD were analyzed: median values are shown for the controls (LPS, PHA) and the different SPION concentrations (1–1,000 µg/mL). *Adapted from Up-regulation of cytokines and chemokines predates the onset of rheumatoid arthritis, *Arthritis & Rheumatism*, John Wiley and Sons. Copyright © 2010 by the American College of Rheumatology. ¹ Copyright © 2010. Bio-Rad Laboratories, Inc. Adapted from Chapman P, Reyes C, Gupta V. Normal Physiological Levels of Human Cytokines Using Bio-Plex Pro™ Cytokine Assays. Bio-Rad Tech Note. 2010;6029. Available from: http://www.bio-rad.com/webroot/web/pdf/literature/Bulletin_6029.pdf. Accessed January 29, 2015.²

Abbreviations: nd, not detectable; Ctrl, control; HD, healthy donors; LPS, lipopolysaccharide; PHA, Phaseolus vulgaris; a-PVA-SPION, amino-polyvinyl alcohol coated superparamagnetic iron oxide nanoparticles.

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