Silver nanoparticles promote procoagulant activity of red blood cells: a potential risk of thrombosis in susceptible population

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Abstract

Background: Silver nanoparticles (AgNP) are widely used in medical practices owing to their distinct antibacterial, antiviral and anticancer activities. However, with increasing use of AgNP, concerns over its potential toxicity are also escalating. Here, we demonstrated the potential thrombotic effect of AgNP which was mediated by the procoagulant activity of red blood cells (RBCs).

Results: In freshly isolated human RBCs, AgNP, but not silver microparticles (AgMP), elicited morphological changes, phosphatidylserine (PS) exposure and microvesicles (MV) generation, the key indicators of procoagulant activity in RBCs at concentration ranges (≤ 100 μg/mL) that were free of significant hemolysis. In line with this, AgNP potentiated thrombin generation and adherence of RBCs to endothelial cells, while AgMP did not. Oxidative stress, intracellular calcium increase and ATP depletion were found to underlie the procoagulant effects of AgNP, which led to altered activity of membrane aminophospholipid translocases. These in vitro findings were well reproduced in rat in vivo, where intravenously exposure to AgNP promoted venous thrombosis significantly. Of note, RBCs isolated from cancer patients, who inherently convey the risk of thrombogenesis, were more sensitive to the procoagulant effects of AgNP. In addition, AgNP significantly potentiated the procoagulant effects of a chemotherapeutic drug, paclitaxel.

Conclusion: Collectively, these results suggest that AgNP may have prothrombotic risks by promoting procoagulant activity of RBCs and caution shall be taken for its use in the population sensitive to thrombosis like cancer patients.

Keywords: Silver nanoparticles (AgNP), Toxicity, Thrombosis, Red blood cells (RBCs), Cancer

Background

Silver nanoparticles (AgNP) are widely used for therapeutic interventions and diagnosis in medical practices and have gained increasing popularity as drug carriers, nanoprobes, bio-imaging and labeling agents [1, 2]. Along with increasing application of AgNP in nanomedicine, however, concerns are also escalating over its potential toxicity against human health [3–5]. Potential adverse effects of AgNP against human health have been first illuminated at a concentration range of 25 ~ 500 μg/mL for 10 ~ 72 h duration in vitro, with respect to its cytotoxicity on human cell lines, reactive oxygen species (ROS) generation, oxidative stress, cell-cycle arrest, pro-apoptotic effects and genotoxicity [5–7].

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this process, which is primed through intracellular calcium increase, ATP- and thiol-depletion [18]. Oxidative stress has been well-established to amplify PS exposure and MV generation, further promoting the procoagulant activity of RBCs [19]. Importantly, association of prothrombotic complications with procoagulant activity of RBCs has been established for cancer patients [18], suggesting that AgNP may present additional thrombotic risk on the cancer patients.

Here, we investigated whether AgNP could affect the procoagulant activity of RBCs, with respect to PS exposure, ROS generation and intracellular calcium increase. The relevance of AgNP-induced procoagulant activity of RBCs to in vivo thrombosis was examined using a rat venous thrombosis model. In an effort to address the thrombosis risk of AgNP in the population susceptible to thrombosis, we examined on the procoagulant effects of AgNP on freshly isolated RBCs from leukemia patients and the synergy with a chemotherapeutic drug (paclitaxel) with thrombotic effects of its own was assessed.

Results
Characterization of AgNP and its cellular uptake by RBCs
The size distribution of silver nanoparticles (AgNP) employed in our study was characterized using scanning electron microscopy (SEM). The majority of AgNP were less than 100 nm in size, and the calculated average particle diameter was approximately 49.3 nm (Fig. 1a). The peak size by intensity frequency of AgNP was 125.4 nm (Fig. 1b). And the zeta potential of AgNP was −32.07 mV at pH 7.4. Ionic silver was separated and quantified using inductively coupled plasma mass spectrometry (ICP-MS) after ultracentrifugation (13,000 rpm for 10 min) and dialysis (24 h), and negligible (<0.01%) ionic silver was detected. Furthermore, such content of ionic silver was examined with no ability of PS exposure after 4 or 24 h treatment (Additional file 1: Figure S1). The physiochemical properties of AgNP as well as AgMP were summarized in Additional file 2: Table S1. To examine whether AgNP may cross cell membranes of RBCs, we determined the cellular uptake of AgNP by RBCs using transmission electron microscopy (TEM). TEM images showed the cellular uptake of AgNP by RBCs accompanied by morphological alterations, which was in a concentration dependent fashion. Compared to control, AgNP-treated cells showed clear structural alteration at 100 μg/mL and at higher concentration of 500 μg/mL severely damaged cell membranes were observed (Fig. 1c), indicating a potential toxic effect of AgNP on RBCs.

A pilot study for the following study in vitro
A preliminary experiment to find the concentrations with no hemolysis was done prior to the main study. Hemolysis was detected spectrophotometrically at various concentrations of AgNP (Fig. 2a). AgNP induced approximately 11.9 and 43.1% hemolysis at the high concentrations (for 250 and 500 μg/mL, respectively), which was in line with the TEM images (Fig. 1b), indicating that AgNP can induce hemolysis at high concentrations of ≥250 μg/mL. As opposed to AgNP, silver microparticles (AgMP) did not induce hemolysis even at high concentrations up to 500 μg/mL, reflecting that size did matter for hemolytic effects of Ag particles.

Phosphatidylserine (PS) exposure of RBCs and microvesicles (MV) generation at non-hemolytic concentrations
While lower concentrations of AgNP (≤100 μg/mL) did not induce hemolysis, TEM observation indicated the cellular uptake of AgNP, suggesting that non-hemolytic effects might be induced in RBCs by AgNP at low doses. Firstly, we investigated the procoagulant activity of RBCs, which is critical to the development of venous thrombosis. Confocal microscopy and SEM observation revealed that the exposure to AgNP induced the occurrence of spiny RBCs, namely echinocytes at 100 μg/mL (4 h) while AgMP was without effects (Fig. 2b, c). A lower effective level could be observed after a longer exposure time compared to the effective level of 50 μg/mL for 4 h incubation with AgNP. As shown in Additional file 3: Figure S2, at 24 h, the effective level of AgNP was 1 μg/mL with around 30% PS exposure, which was 100 times less than that observed at 4 h. It is well known that loss of phospholipid asymmetry is attributable to morphological change in RBCs. During this process, PS can be externalized to outer membrane and membrane surface is lost as MV is generated, which are closely related to procoagulant activity of RBCs [20]. Indeed, in flow cytometry analysis, PS exposure increased in concentration- and time-dependent manners after the exposure to AgNP (Fig. 2d, e). MV generation was also observed upon the exposure to AgNP (Fig. 2f). In contrast to AgNP, even high concentrations of AgMP did not induce any effects on RBCs including hemolysis, shape change, PS exposure or MV generation (Fig. 2).

Effects of AgNP on oxidative stress and intracellular calcium in human RBCs
Oxidative stress can induce PS exposure and MV generation in RBCs, which can promote the procoagulant activity of RBCs [18]. AgNP-induced ROS generation in RBCs, as examined by DCF-DA-enhanced fluorescence using flow cytometry, increased in a concentration dependent manner, while no effects were observed in AgMP-exposed RBCs (Fig. 3a), which was in a good accordance with PS exposure and MV generation. Consistently with ROS generation, the representative intracellular antioxidant,
glutathione (GSH) [21] and ATP were decreased significantly following the exposure to AgNP (Fig. 3b, c).

It is well established that oxidative stress can cause intracellular calcium increase, which is referred as a key stimulus to induce both PS exposure and MV generation [22]. Notably, intracellular calcium level was increased considerably by AgNP (Fig. 3d), which was abolished by the pretreatment with N-acetylcysteine (NAC), a well-known antioxidant (Fig. 3e). Increased intracellular calcium level can activate scramblase, leading to a bidirectional exchange (scrambling) of membrane phospholipids [23] and outward translocation of PS residing on inner membrane leaflet [24]. Indeed, the activity of scramblase was significantly increased as determined by C6-NBD-PC externalization (Fig. 3f). Flippase is another aminophospholipid translocase mediating the inward translocation of PS exposed on outer membrane leaflet [20]. The significant decrease in flippase activity at 10 μg/mL indicated an effect of AgNP on the inner ward phospholipid translocation. Compared with AgNP-exposed groups, no significant effects in ROS generation, GSH depletion, ATP depletion, intracellular calcium level and scramblase activity were observed in AgMP-exposed groups (Fig. 3).

To confirm the role of oxidative stress and intracellular calcium increase in AgNP-induced PS exposure, the effects of antioxidants and a calcium chelator, EGTA, were examined. Pretreatment of antioxidants such as NAC, vitamin C and trolox, and EGTA significantly attenuated PS exposure induced by AgNP (Fig. 4a, b), which could be further confirmed by confocal microscopy and SEM (Fig. 4c, d), supporting that oxidative stress and intracellular calcium increase may be key mechanisms underlying AgNP-induced PS exposure.

**Prothrombotic effects of AgNP on human RBCs**

Prothrombotic effects of AgNP on RBCs were evaluated using examining thrombin generation and RBCs adhesion to human umbilical vein endothelial cells (HUVECs). Procoagulant activity of RBCs results in accelerated thrombin generation, a key step for blood coagulation cascade, and
the adhesion of RBCs to vascular wall, which could ultimately promote thrombosis [18]. To investigate whether AgNP-induced PS exposure could promote blood coagulation indeed, thrombin generation by AgNP-exposed RBCs was evaluated. A concentration-dependent increase in thrombin generation was induced by AgNP-exposed RBCs (Fig. 5a), which was significantly inhibited by the pretreatment with antioxidants (Fig. 5b) and EGTA (Fig. 5c). Additionally, adhesion of RBCs to HUVECs was also enhanced by the exposure to AgNP (Fig. 5d). Increased adhesion of RBCs was completely abolished by NAC and EGTA pretreatment (Fig. 5e), which was further confirmed in fluorescence microscopy (Fig. 5f), supporting that exposure of RBCs to AgNP could accelerate coagulation and increase adherence of RBCs to blood vessel, which may ultimately promote thrombosis. These events were all attenuated by antioxidants and a calcium chelator, suggesting the association with oxidative stress and intracellular calcium increase as similar with PS exposure and MV generation. Compared with AgNP-exposed groups, no significant effects on thrombin generation or adherence of RBCs to HUVECs were observed in AgMP-exposed groups (Fig. 5a, d).
procoagulant activity of RBCs and eventually led to increased venous thrombosis.

Effects of AgNP on procoagulant activity of RBCs isolated from cancer patients and the synergy with chemotherapeutic agents

AgNP have been applied to the management of cancer for therapeutic inventions and diagnosis, but cancer itself, is a well-known risk factor of thrombosis [14, 15, 25, 26]. RBCs isolated from the patients with leukemia were examined for AgNP-induced procoagulant activity. As a result, the exposure to AgNP induced ROS generation and PS exposure in the RBCs isolated from cancer patients in a concentration dependent manner, which was higher in the degree and lower in the effective concentrations when compared with RBCs from healthy controls (Fig. 7a, b).

Chemotherapeutic drugs to treat cancers also have procoagulant effects [27, 28] and have been previously reported to contribute to increased thrombosis in cancer patients [29]. We investigated whether AgNP may have synergy effects on the procoagulant effects of paclitaxel (PTX), a well-known chemotherapeutic drug which can induce procoagulant activity of RBCs [27]. As shown in Figure 7c and d, the combined exposure to PTX and AgNP significantly increased ROS generation and PS exposure in RBCs compared with those treated with AgNP alone or PTX alone.

Discussion

Here, we demonstrated that silver nanoparticles (AgNP) (of sizes less than 100 nm), but not silver microparticles (AgMP), can promote phosphatidyserine (PS) exposure and microvesicles (MV) generation in freshly isolated human red blood cells (RBCs), mainly through reactive oxygen species (ROS) generation and intracellular calcium increase. AgNP-induced procoagulant activity of RBCs was observed at non-hemolytic concentrations and more importantly, confirmed in rat venous thrombosis model in vivo where the exposure to AgNP led to
increased PS exposure and promoted thrombosis supporting the relevance of our findings to real in vivo states (Fig 8). Of note, RBCs of cancer patients were more sensitive to the procoagulant effects of AgNP and AgNP manifested the synergy with a chemotherapeutic agent, paclitaxel in promoting procoagulant activity of RBCs, suggesting that a caution shall be taken for the application of AgNP to the management of cancer patients.

Toxicity of AgNP has been investigated in many studies, including cytoxicity of AgNP in various cells in vitro [4, 6, 30], and assessment of bio-distribution, elimination, and accumulation of AgNP in organs and potential organ toxicity associated with AgNP exposure [7, 31]. Considering the direct contact of blood cells with AgNP entering the body during circulation, potential cardiovascular toxicity of AgNP shall not be ruled out. However, few studies have addressed the mechanism underlying AgNP-induced toxicity on RBCs in relation to pathophysiological effects. Previous studies have focused on the direct toxicity of AgNP on the structural integrity of RBCs or early events such as hemolysis, oxidative stress and membrane damage [11, 32, 33] without extending them into the mechanisms leading to diseases. Here, we demonstrated that AgNP may promote thrombosis through procoagulant activity of RBCs providing an important piece of clue for the potential cardiovascular toxicity of AgNP.

In RBCs, toxicological effects of AgNP have been investigated regarding the hemolysis employing extremely high concentration ranges (up to 500 μg/mL) of AgNP [11, 32]. We could also observe a significant hemolysis at high concentrations of AgNP (from 250 μg/mL to 500 μg/mL), which matched well with previous studies of hemolysis. Actually, this range is too high and raises questions on its clinical relevance. Of note, we observed that AgNP can induce PS exposure and procoagulant activity at non-hemolytic concentrations as low as 50 μg/mL after 4 h exposure, demonstrating that AgNP can cause adverse effects on cardiovascular system at non-hemolytic ranges. This concentration range of AgNP is comparable with those (25 ~ 400 μg/mL) used in the previous literatures on AgNP toxicity against human peripheral blood cells or lung cells [6, 7].
Furthermore, we also observed that a prolonged exposure (24 hr) could significantly lower the effective level to 1 μg/mL as shown in Additional file 3: Figure S2, indicating a more potent toxicity could be expected after a longer exposure. Currently, reports on the toxicity of AgNP in vivo are often controversial and inconsistent. In an attempt to demonstrate AgNP toxicity in vivo, diverse exposure routes, such as oral exposure, inhalation, intraperitoneal and intravenous, in the settings of single or multiple exposure and with various AgNP differed in size, surface charge and surface-coating substances have been explored in many experimental animal species. While oral administration of AgNP unanomously showed no obvious toxicity in acute or subacute studies [34–36], intravenous (i.v.) exposure to AgNPs was not without toxicity. Previous studies observed spleen and liver toxicity of AgNP in mice after 10 mg/kg i.v. administration, and liver and hematological toxicity in rats after i.v. injection of 4–40 mg/kg AgNP [37, 38]. Consistently, in our study, AgNP induced thrombosis at ≥ 10 mg/kg by promoting procoagulant activation of RBCs. After i.v. injection of 10 mg/kg AgNP in rats, peak concentration of 5.2 μg/mL was observed at 8 h, which was basically maintained at around 5 μg/mL up to 96 h [39] of which range was similar to our studies in vitro and in vivo.

It has been still controversial that whether the toxic effect is particle-specific or mediated by silver ions dissolved from AgNP. A previous literature suggested released silver ions were the primary cause of AgNP toxicity on Sphaerium corneum [40], and the effect of silver ion on RBCs was reported by Sopjani et al. [41]. However, Recordati et al. showed that silver ion and AgNP exhibit distinct patterns of tissue distribution and toxicity, and toxicity of AgNP was attributable to the nanoparticle form [37]. Consistently, we could observe negligible release of silver ion from AgNP and more importantly, found that silver ion did not induce PS exposure shown in Additional file 1: Figure S1, although the dissolution may differ depending on physiochemical characteristics (e.g. size, shape, surface charge, pH etc.) [42].

![Fig. 5](image_url) Effects of AgNP on thrombin generation and adhesion of human RBCs to human umbilical vein endothelial cells (HUVECs). Thrombin generation was determined by a prothrombinase assay after RBCs were treated (a) with distilled water (DW; control), various concentrations of AgNP and AgMP for 4 h (n = 8); (b) with 100 μg/mL of AgNP along with preincubation with various antioxidants (n = 4) and (c) with 0, 50 and 100 μg/mL of AgNP along with preincubation with EGTA (n = 4). AgNP- or AgMP-treated RBCs were incubated with HUVECs for 1 h, and then RBCs adhered to HUVECs were counted by microscopic observation (n = 4). e RBCs were pretreated with NAC or EGTA before AgNP treatment, and RBCs were further incubated with HUVECs for 1 h. And then RBCs adhered to HUVECs were counted by microscopic observation (n = 4) or (f) examined using fluorescence microscope as described in methods. Red: glycophorin-A-PE for RBC staining; Green: calcein green for HUVECs staining. Values are mean ± S.E. of 4–8 independent experiments. *represents significant differences from control group (p < 0.05), # represents significant differences from AgNP-treated group (p < 0.05). Scale bars: 100 μm.
ROS and oxidative stress are involved in regulating normal physiology as well as pathophysiological processes. Excessive production of ROS has been well established to orchestrate a series of pathological events such as genotoxicity, inflammation, fibrosis, and carcinogenesis [43, 44]. ROS generation induced by AgNP has been reported to provoke cytotoxicity and genotoxicity in leukemia cells and lung cells [6, 7], indicating that ROS generation may explain the toxicity of AgNP. Indeed, ROS generation played a central role in the procoagulant and prothrombotic effects of AgNP on RBCs. Interestingly, other nanomaterials such as carbon, titanium dioxide (TiO₂) and manganese oxide (MnO₂) have been shown to induce oxidative stress triggering signaling cascades that eventually result in increased expression of proinflammatory, fibrotic cytokines and cell damage [21, 43, 45]. In this context, in addition to AgNP, potential effects of other nanomaterials on procoagulant activity of RBCs warrant further attention as well [46].

Recently, nanotechnology has been incorporated into various fields of medical practice. Nanomedicine has emerged as the mainstream in cancer management. Notably, cancer patients are at a hypercoagulable state as for such reasons as advanced stage of diseases, intravascular metastasis, and increased tumor volume [14, 25, 29]. Clinically detectable thrombotic events were found in 15–20% of all cancer patients [14, 15, 25]. Cancer, itself, is a major thrombotic risk factor increasing thrombosis by 4.1 fold and moreover, chemotherapeutic agents heighten it even further by 6.5 fold [15, 26], reflecting the risk of AgNP-use in cancer management. Of critical, exposure to AgNP in combination of chemotherapeutic agent may aggravate thrombotic risks when combined with the hypercoagulable state of cancer patients, which needs further studies in the future.
Conclusion
Our study demonstrated that AgNP can promote prothrombotic risks by increasing procoagulant activity of RBCs both in vitro and in vivo. The mechanism underlying the effect appears to involve oxidative stress and reactive oxygen species generation. Of note, procoagulant effects of AgNP were enhanced in RBCs of cancer patients and a significant synergy was observed with a chemotherapeutic agent, paclitaxel in promoting procoagulant activity of RBCs, suggesting that caution should be taken for its use in cancer patients.

Methods
Materials
AgNP (silver nanoparticles, < 100 nm particle size, contains 0.2% PVP as dispersant, 99.5% trace metals basis.), AgMP (silver microparticles, 5-8 micro, ≥ 99.9% metals basis), mercury orange, trolox, L-ascorbic acid, glutaldehyde solution and osmium tetroxide, the ATP bioluminescent assay kit, and purified human thrombin were purchased from Sigma Chemical Co. (St. Louis, MO). Phycoerythrin-labeled monoclonal mouse anti-human CD235a and fluorescein-isothiocyanate (FITC)-labeled
annexin V (annexin V-FITC) were from BD Pharmingen (San Diego, CA). Fluo-4 acetoxyethyl ester (fluo-4 AM) and the chloromethyl derivative of 2', 7'-dichlorodihydrofluorescein diacetate (CM-H2DCF-DA) were obtained from Life Technologies. 1-Palmitoyl-2-[6-[[7-nitro-2,1,3-benzoxadiazol-4-yl]amino]hexanoyl]-sn-glycero-3-phospho-L-serine (C6-NBD-PS) and 1-oleoyl-2-[6-[[7-nitro-2,1,3-benzoxadiazol-4-yl]amino]hexanoyl]-sn-glycéreo-3-phosphocholine (C6-NBD-PC) were from Avanti Polar Lipids (Alabaster, AL). Purified human prothrombin, factor Xa and factor Va were from Hematologic Technologies, Inc. (Essex Junction, VT), and S2238 was from Chromogenix (Milano, Italy). Human umbilical vein endothelial cells (HUVECs) and the endothelial cell growth media (EGM) kit were purchased from Lonza. Human recombinant tissue factor (Recombiplastin) was obtained from Instrumentation Laboratory (Lexington, MA) and thromboplastin (Simplastin Excel) was from Biomerieux (Durham, NC).

Assessment of characterization of AgNP
Both AgNP and AgMP were purchased from Sigma Aldrich. Preparation of AgNP suspension was carried out according to the methods previously described [12]. AgNP or AgMP powder were dispersed in distilled water as 100 X stock solution (1–50 mg/mL) and sonicated with a probe type sonicator with a maximum output power, 200 Watts (Branson Sonifier, Danbury, CT) for 2 min to prevent particles self-assembly (agglomeration). In addition, prior to every experiment, AgNP suspensions were vigorously vortexed for 30 sec. AgNP we used in the present study were with 0.2% PVP as dispersant (provided by Sigma Aldrich), while no PVP was in AgMP (provided by Sigma Aldrich). Through mathematical calculation, there might be 0.2 μg/mL PVP in 10 μg/mL AgNP-treated group, while 2 μg/mL PVP in 100 μg/mL AgNP-treated group. In order to estimate the effect of PVP in AgNP on PS exposure, we determined the effect of 2 μg/mL PVP alone on PS exposure of RBCs. Results are as follows: Control (1.52 ± 0.90% PS exposure); 2 μg/mL PVP (1.38 ± 1.46% PS exposure). Since PVP had minimal effects, DW (distilled water) was used as a vehicle control.

For the characterization of AgNP, AgNP suspension was dried and was observed with scanning electron microscope (SEM) (ZEISS, MERLIN Compact) to examine the size frequency. A detailed statistical analysis of AgNP was performed by randomly measuring 118 nanoparticles and the procedure was operated by manually

![Fig. 8](image_url)
outlining the particles from several images taken by SEM. Nanoparticles were used at a concentration of 100 μg/mL in the following determination. The hydrodynamic diameter and the zeta potential of the nanoparticles were measured by dynamic light scattering (DLS-7000) and electrophoretic light scattering (ELS Z-100), respectively. The polydispersity index (PDI) was below 0.3 in all cases. For determination of dissolved silver, ultracentrifuge (13,000 rpm for 10 min) and dialysis (24 h) were prior to the quantification of Ag. Dialysis experiments were carried out in dialysis bags filled with 5 mL silver nanoparticle dispersion (100 μg/mL) and immersed in 495 mL water. The concentration of silver was < 0.1 μg/L determined by inductively coupled plasma mass spectrometry (ICP-MS) with the limit of detection of 0.1 μg/L. The maximum possible percentage of ionic Ag in silver nanoparticle dispersion was 0.01%.

Preparation of RBCs
With the approval from the Ethics Committee of Health Service Center at Seoul National University, human blood was obtained from healthy male donors (20-30 years old) using a vacutainer with acid citrate dextrose (ACD) and a 21 gauge needle (Becton Dickinson, U.S.A.) on the day of each experiments. Platelet rich plasma and buffy coat were removed by aspiration after centrifugation at 200 g for 15 min. Packed RBCs were washed 2 times with phosphate buffered saline (PBS: 1.06 mM KH₂PO₄, 154 mM NaCl and 2.96 mM Na₂HPO₄ at pH 7.4) and once more with Ringer’s solution (125 mM NaCl, 5 mM KCl, 1 mM MgSO₄, 32 mM HEPES, 5 mM glucose, pH 7.4). Washed RBCs were resuspended in TBS buffer to a cell concentration of 5 x 10⁷ cells/mL, and the final CaCl₂ concentration was adjusted to 1 mM prior to use.

Observation of cellular uptake of AgNP by RBCs
For detection of cellular uptake of AgNP by RBCs, isolated RBCs were incubated with AgNP for 4 h, and fixed with 2% glutaraldehyde solution in the refrigerator overnight. On the second day, after washing three times with 0.05 M sodium cacodylate buffer every 10 min, cells were post-fixed with 2% osmium tetroxide for 2 h. Before en-bloc staining with 0.5% uranyl acetate for 30 minutes, briefly washing with DW were done twice, and serially dehydrated once with 30, 50, 70, 80, 90% and three time with 100% ethanol. Next, propylene oxide was used for transition every 10 min for 2 times, and infiltration were done once with propylene oxide and spurr’s resin (1:1) for 2 h, then with only spurr’s resin in the desiccator overnight. On the third day, infiltration was completed with newly spurr’s resin for 2 h in the desiccator, then samples were kept in the 70°C oven overnight for polymerization. Finally, samples were observed under TEM (JEOL, JEM 1010).

Flow cytometric analysis of phosphatidylserine (PS) exposure and microvesicles (MV) generation
Annexin V-FITC was used as a marker for PS detection, whereas anti-glycophrin A-PE was used as an identifier of RBCs. Negative controls for annexin V binding were stained with annexin V-FITC in the presence of 2.5 mM EDTA instead of 2.5 mM CaCl₂. Samples were analyzed on the flow cytometer FACS Calibur (Becton Dickinson, U.S.A.) equipped with an argon-ion laser emitting at 488 nm. The light scatter and fluorescence channels were set on a log scale. Data from 5,000 events were collected and analyzed using Cell Quest Pro software. MV were identified on the basis of forward scatter characteristics after calibration by 1 μM standard beads. The extent of hemolysis was determined spectrophotometrically at 540 nm.

Microscopic observation using confocal and scanning electron microscopy (SEM)
For confocal microscopy, 500 μL RBCs suspension was added and attached for 1 h to a four-chambered coverslip (Lab-Tek; Nalge Nunc Inc., Naperville, IL). After washing the coverslip once with Ringer’s solution containing 2% BSA, RBCs were then incubated with a vehicle control (DW in Ringer’s solution), AgNP or AgMP (dissolved in Ringer’s solution). After incubation, the coverslips were washed once again with Ringer’s solution and then, RBCs were stained with Ringer’s solution containing anti–glycophrin-A–PE for 30 min. Ultimately, samples were washed once and resuspended to be observed using confocal microscopy equipped with an argon laser (Leica, Wetzlar, Germany). Excitation and emission filters were set at 488 nm and 550–600 nm, respectively. For SEM observation, after fixation with 2% glutaraldehyde solution for 1 h at 4°C, the RBCs were centrifuged and washed three times with PBS, followed by post-fixation with 1% osmium tetroxide for 30 min at room temperature in the hood. After washing with PBS twice, the samples were dehydrated serially with 50, 70, 80, 90, and 100% ethanol. After drying and coating with gold, the images were observed on a SEM.

Measurement of reactive oxygen species (ROS) and intracellular glutathion (GSH) level
To detect ROS generation, RBCs were preincubated with 5 μM CM-H₂DCF-DA for 30 min at 37°C. After washing with Ringer’s solution containing 1 mM CaCl₂ to remove excess CM-H₂DCF-DA, RBCs were incubated with AgNP or AgMP. The aliquots were diluted and the fluorescence of intracellular DCF was measured using flow cytometry. GSH level in RBCs was measured according to the method described by Jose Enrique...
O’Connor et al [47]. After incubated with vehicles (DW), AgNP or AgMP, RBCs were treated with 40 μM mercury orange (dissolved in acetone) and then incubated in ice for 5 min. After that, cells were washed and resuspended with cold Ringer’s solution, and then diluted with cold Ringer’s solution to be analyzed with an FACS Calibur flow cytometer (BD Biosciences).

**Measurement of intracellular ATP and calcium levels**

Samples were washed and resuspended in Ringer’s solution containing 1 mM CaCl₂. The aliquot was mixed vigorously with TAE buffer (100 mM Tris-acetate, 2 mM EDTA, pH 7.8) containing 10% trichloroacetic acid, and then cooled in ice for 20 min. The sample was centrifuged and the aliquot of resultant supernatant was mixed with cold TAE buffer. The intracellular ATP level was measured by luciferin/luciferase assay in Luminoskan (Labsystems, Franklin, Massachusetts) using an ATP assay kit. We calculated the ATP concentrations based on the ATP standard curve. To detect intracellular calcium increase, RBCs were loaded with 3 μM Fluo-4 AM (Molecular Probes, Eugene, OR, USA) for 1 h at 37°C in the dark. After excess Fluo-4 AM was removed, AgNP or AgMP were treated to RBCs. The aliquots were diluted and analyzed in flow cytometer. Data from 5,000 events were collected and analyzed using Cell Quest Pro software (Becton Dickinson).

**Phospholipid translocation measurement**

RBCs (5 x 10⁷ cells/mL) were incubated with AgNP or AgMP for 30 min at 37°C and then loaded with 0.5 μM C₆-NBD-PC and C₆-NBD-PS. Aliquots from the cell suspension were removed at the indicated time intervals and placed on ice for 20 min in the presence or absence of 1% bovine serum albumin (BSA). Pellets obtained after 1 min of centrifugation at 10,000 g were lysed in 1% Triton X-100, and the fluorescence intensities were measured (ex 485 nm, em 535 nm). The amount of internalized probe was determined by comparing the fluorescence intensity associated with the cells before and after back-extraction.

**Prothrombinase assay**

After incubation with AgNP or AgMP for 4 h, samples were incubated with 5 nM factor Xa and 10 nM factor Va in Tyrode buffer (134 mM NaCl, 10 mM HEPES, 5 mM glucose, 2.9 mM KCl, 1 mM MgCl₂, 12 mM NaHCO₃, 0.34 mM Na₂HPO₄, 0.3% BSA, and 2 mM CaCl₂ at pH 7.4) for 3 min at 37°C. Thrombin formation was initiated by addition of 2 μM prothrombin. Exactly 3 min after the addition of prothrombin, an aliquot of the suspension was transferred to a tube containing stop buffer (50 mM Tris-HCl, 120 mM NaCl, and 2 mM EDTA at pH 7.9). Thrombin activity was determined using the chromogenic substrate S2238 (chromogenic substrate for thrombin; Chromogenix, Milano, Italy). We calculated the rate of thrombin formation from the change in absorbance at 405 nm using a calibration curve generated with active-site–titrated thrombin.

**Adherence of RBCs to human umbilical vein endothelial cells (HUVECs)**

The HUVECs (6-8 passages) were maintained in the endothelial cell growth media kit at 37°C in a 95% air/5% CO₂ incubator. Endothelial cells (1x10⁵ cells) were seeded into 25T flask for 5 days, AgNP or AgMP-treated RBCs were washed once and resuspended in EBM-2 to a cell concentration of 5x10⁶ cells/mL. After HUVEC was washed twice with EBEm-2, AgNP or AgMP-exposed RBCs were layered onto confluent HUVEC monolayer and incubated for 60 min at 37°C. After the incubation, the flask was rinsed 3 times with EBEm-2 to remove non-adherent RBCs. The number of adherent RBCs was counted on light microscope. The experiments were performed in triplicate and 25 fields were selected randomly to count RBCs number.

**Fluorescence observation of adhered RBCs to HUVECs**

Endothelial cells (2 x 10⁴ cells) were seeded into 4-well-chamber for 2 days and stained with calcein green for 20 min. AgNP or AgMP-treated RBCs were washed once and resuspended in EBM-2 to a cell concentration of 5x10⁶ cells/mL. After HUVECs were washed twice with EBEm-2, AgNP or AgMP-exposed RBCs were layered onto confluent HUVEC monolayer and incubated for 60 min at 37°C. After the incubation, the chambers were rinsed once with EBEm-2 to remove nonadherent RBCs, and glycoporphin A-PE were added for staining RBCs. Adhered RBCs to HUVECs were observed using fluorescent microscopy.

**In vitro assessment of PS exposure and procoagulant activity in rat RBCs**

Male Sprague-Dawley rats (300-400g) were anesthetized with urethane (1.25 g/kg, i.p.). Briefly, citrated whole blood (3.8% sodium citrate) was withdrawn from abdominal aorta and RBCs were isolated as human RBCs preparation. Isolated RBCs were further incubated with AgNP for 4 h, then PS exposure and procoagulant activity were determined mentioned above.

**In vivo experiments**

Male Sprague-Dawley rats (300-400 g) were anesthetized with urethane (1.25 g/kg, i.p.). 2 h after intravenous injection of AgNP (10 or 50 mg/kg), whole blood (3.8% sodium citrate) was collected from abdominal aorta, and PS exposure of RBCs was determined using flow cytometry. Thrombus formation was induced by stasis combined with...
Experiments using RBCs from cancer patients
With an approval from the Ethics Committee of Health Service Center at Seoul National University Hospital, human blood was obtained from cancer patient male donors with leukemia and healthy control male donors using a vacutainer with acid citrate dextrose and a 21-gauge needle on the day of each experiment. RBCs preparation, measurement of ROS generation and PS exposure were done as described earlier.

Statistical analysis
The means and standard errors of means were calculated for all treatment groups. The data were subjected to two-way analysis of variance followed by Duncan’s multiple range test or student t test to determine which means were significantly different from the control. In all cases, a p value of < .05 was used to determine significant differences.

Additional files

- **Additional file 1**: Figure S1. PS exposure of human red blood cells induced by ionic silver (0.1 µM). (TIF 208 kb)
- **Additional file 2**: Table S1. Summary of physicochemical properties of AgNP and AgMP. (TIF 314 kb)
- **Additional file 3**: Figure S2. PS exposure of human red blood cells induced by low concentrations of AgNP after a prolonged exposure. (TIF 232 kb)

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Availability of data and materials
Yes

Authors’ contribution
YB and KWL designed and performed the experiments; KK and TN analyzed the data and performed platelet experiments; IR designed the RBC experiments isolated from cancer patients; CNB performed an additional experiment for the revision; JHC supervised the study. All authors read and approved the final manuscript.

Ethics approval
With the approval from the Ethics Committee of Health Service Center at Seoul National University, human blood was obtained from healthy male donors or cancer patients. All the animal protocols used in vivo experiments were approved by the Ethics Committee of Animal Service Center at Seoul National University.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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