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Polydopamine nanoparticles for treatment of acute inflammation-induced injury

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Abstract

Nanotechnology-mediated anti-inflammatory therapy is emerging as a novel strategy for treatment of inflammation-induced injury. However, one of the main hurdles for these anti-inflammatory nano-drugs is their potential toxic side effects in vivo. Herein, we uncovered that polydopamine (PDA) nanoparticles with structure and chemical properties similar to melanin, a natural bio-polymer, displayed significant anti-inflammation therapeutic effect on acute inflammation-induced injury. PDA with enriched phenol groups functioned as a radical scavenger to eliminate reactive oxygen species (ROS) generated during inflammatory responses. As revealed by in vivo photoacoustic imaging with a H₂O₂-specific nanoprobe, PDA nanoparticles remarkably reduced intracellular ROS levels in murine macrophages challenged with either H₂O₂ or

lipopolysaccharide (LPS). The anti-inflammatory capacity of PDA nanoparticles was further demonstrated in murine models of both acute peritonitis and acute lung injury (ALI), where diminished ROS generation, reduced proinflammatory cytokines, attenuated neutrophil infiltration, and alleviated lung tissue damage were observed in PDA-treated mice after a single dose of PDA treatment. Our work therefore presents the great promise of PDA nanoparticles as a biocompatible nano-drug for anti-inflammation therapy to treat acute inflammation-induced injury.

Keywords: Polydopamine; Acute inflammation injury; Reactive oxygen species; Anti-inflammation therapy.

Introduction

Acute inflammatory response characterized by circulating neutrophils influx into the tissues and organs is a frequent event in the clinic, which causes tissue infection and damage in a short time period.¹⁻³ Among different types of acute inflammation, acute lung injury (ALI) and peritonitis are the simplest form of acute inflammatory responses. ALI is induced by the damage of alveolar epithelial cells and capillary endothelial cells and characterized with the dysregulated neutrophils sequestrated in the lung.^{1, 4-8} These dysregulated neutrophils are capable of releasing proteases, cationic peptides and reactive oxygen species (ROS) to aggravate lung injury.^{4, 9-12} Excessive ROS formation breaks the balance of endogenous anti-oxidative capacity and induces tissue damage and dysfunction. Therefore, prevention of the tissue damage by scavenging ROS could serve as a significant therapeutic option for treatment of acute inflammation-induced injury in the clinic.¹³⁻¹⁵

Polydopamine (PDA) derived from self-polymerization of dopamine shows many unique chemical properties and has attracted much attention in the area of nanomedicine.¹⁶⁻²² PDA, a biocompatible and biodegradable polymer, has similar structure and chemical properties to that of melanin. With the versatile molecular adsorption ability, it has been found that PDA nanoparticles can be used for drug delivery, molecular imaging and cancer theranostics, as demonstrated in many previous studies.^{17-19, 23-27} On the other hand, with a large amount of phenol groups on their surface, PDA nanoparticles show an excellent chelating ability to capture different types of metal ions such as radioisotope ions for nuclear imaging.^{17, 23, 28-30} Moreover, those phenol groups also make PDA an excellent free radical scavenging agent.^{16, 31} However, the applications of PDA nanoparticles as an anti-inflammatory agent to treat acute inflammation-induced injury including acute peritonitis and ALI have not yet been demonstrated to our best knowledge.

In this work, bare PDA nanoparticles were developed as anti-inflammatory nano-drug for treatment of acute inflammation-induced injury. At the in vitro level, it was discovered that PDA nanoparticles efficiently scavenged either H₂O₂- or LPS-induced cellular ROS. By utilizing our recently developed H₂O₂-responsive liposomal photoacoustic imaging nano-probe, we found that PDA nanoparticles were able to effectively suppress the in vivo inflammation process in an acute peritonitis model induced by LPS, as evidenced by the down-regulated ROS (e.g. H₂O₂) and pro-inflammatory cytokine levels after PDA treatment. Furthermore, using a murine model of ALI, we demonstrated that treatment with PDA nanoparticles ameliorated lung injury, with substantially diminished retention of neutrophils and lymphocytes in the lung and down-regulated levels of proinflammatory cytokines. Our results present an interesting anti-inflammation therapy platform by

utilizing the inherent biochemical properties of nanoparticles and suggest the potential application prospect of PDA nanoparticles as an anti-inflammatory nano-drug.

Results and Discussion

PDA nanoparticles were synthesized following a literature method (Figure 1A).^{16, 18} Transmission electron microscopy (TEM) image (Figure 1B) of the PDA sample showed that PDA nanoparticles were successfully formed with a uniform and mono-dispersed spherical structure. As shown in the TEM image, the average diameter of PDA nanoparticles was ~80 nm. Those as-made PDA nanoparticles were stable in water without any obvious aggregation, as revealed by dynamic light scattering measurement (Figure 1C).

ROS are one of main reasons to cause various acute and chronic inflammatory diseases.^{15, 32-36} Previous studies have demonstrated that H_2O_2 is one of the most common ROS types.³⁷⁻³⁹ As such, we firstly examined the ROS scavenging activity of PDA nanoparticles toward H_2O_2 . In our recent work, we fabricated a liposomal nanoprobe by encapsulating horseradish peroxidase (HRP) and its substrate, 2, 2'-azino-bis(3-ethylbenzothiazoline-6-sulfonic acid) (ABTS), into polyethylene glycol modified liposomes, and used such Lipo@HRP&ABTS as a H_2O_2 -specific nanoprobe for in vivo photoacoustic (PA) imaging via the HRP-induced chromogenic reaction.³⁷ Therefore, using Lipo@HRP&ABTS as an indicator, the capability of PDA nanoparticles to scavenge H_2O_2 was evaluated. Firstly, PDA (0.01 mg/ml) was added into H_2O_2 (20 μ M) solutions and incubated for various time periods. Afterwards, the Lipo@HRP&ABTS nanoprobe was added into those mixture solutions and incubated for 1 min under room temperature. As HRP in the presence of H_2O_2 would

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convert colorless ABTS into its oxidized form with strong greenish color and the absorbance peak at \sim 730 nm, the absorbance spectrum changes of the reaction solution could be employed to determine the changes of H_2O_2 concentrations. Interestingly, we found that addition of PDA rapidly reduced the H_2O_2 concentration in the solution, as revealed by a decrease of absorbance from the Lipo@HRP&ABTS nanoprobe (Figure 1D). Furthermore, an increase in PDA concentrations also resulted in a further decrease of H_2O_2 concentrations (Figure 1E). Notably, the H_2O_2 scavenging activity of PDA increased as the solution pH increased from acidic to neutral values (Supporting **Information Figure S1B & 1C).** Such a PDA-concentration-dependent H_2O_2 scavenging phenomenon could be vividly visualized by the obvious color change of PDA-treated H₂O₂ solutions in the presence of Lipo@HRP&ABTS nanoprobe (Figure 1F). Besides, PA signals, which were originated from the 730 nm absorbance of Lipo@HRP&ABTS in response to H_2O_2 , of the reaction solutions with different PDA concentrations were also detected (Figure 1G). As the increase of PDA concentrations, the H₂O₂-specific PA signals from Lipo@HRP&ABTS showed dramatic decreases (Figure 1G). It was noteworthy that even a rather low PDA concentration (0.01 mg/ml) was capable of inducing a significant decrease of the H_2O_2 concentration in the aqueous solution.

To understand the mechanism underlying the ability of PDA to scavenge H_2O_2 , we examined whether PDA caused the decomposition of H_2O_2 by measuring O_2 production in the H_2O_2 solution after adding PDA nanoparticles. Without PDA, the rate of O_2 generation was quite slow as a result of self-decomposition of H_2O_2 (Supporting Information Figure S1D). Interestingly, when PDA nanoparticles were added into the H_2O_2 solution, the O_2 production rate was greatly increased (Supporting Information Figure S1D & 1E), suggesting the 'catalase-like' catalytic performance of PDA to trigger the decomposition of H_2O_2 (Supporting Information Figure S1F). Therefore, we propose that PDA nanoparticles could act as a H_2O_2 scavenger not only as a reducing agent to directly react with such ROS by redox reactions, but also as a catalyst to trigger decomposition of H_2O_2 .

Next, the interactions between PDA nanoparticles and cells were studied at the in vitro level. After incubation with Raw 264.7 cells, PDA nanoparticles could be taken up into the cells (Supporting Information Figure S2). The potential antioxidant and anti-inflammatory activities of PDA were then studied in vitro by a dye of 2,7-Dichloro-dihydrofluorescein diacetate (DCFH-DA), which is specifically responsive to intracellular ROS.⁴⁰ In our experiments, Raw 264.7 cells were pre-incubated with PDA for 6 h and then stimulated by H_2O_2 or LPS, the latter is a component from Gram-negative bacterial cell wall commonly used to stimulate oxidative stresses and induce inflammatory responses. As expected, the cellular ROS levels showed drastic increases upon exposure to either H₂O₂ or LPS. Interestingly, the ROS levels of cells pre-treated with PDA (PDA + H₂O₂ and PDA + LPS groups) were found to be significantly reduced, as revealed by both FACScan analysis and confocal fluorescence images of DCFH-DA stained cells (Figure 2A-C&E and Supporting Information Figure S3A). Moreover, PDA treatment also effectively prevented H_2O_2 induced cytotoxicity to those cells (Figure 2B-D). No obvious toxicity was observed after Raw 264.7 cells were incubated with different concentrations (up to 160 µg/ml) of PDA for 24 h (Supporting Information Figure S3B). On the other hand, PDA nanoparticles could down-regulate the LPS-induced excretion of pro-inflammatory cytokine TNF- α by Raw 264.7 cells (Supporting Information Figure S3C). Therefore, PDA nanoparticles showed significant antioxidant and

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anti-inflammatory activities to reduce ROS production and protect cells from ROS-induced damage at the in vitro level.

Encouraged by the effective anti-inflammation ability of PDA in vitro, we then studied whether PDA nanoparticles could alleviate LPS-induced acute peritonitis in mice. Acute peritonitis was induced by intraperitoneal (i.p.) injection of LPS (Figure 3A) and mice were randomly divided into four groups (group 1-4): phosphate buffered solution (PBS) control group, PDA group, LPS group, and LPS + PDA group. PDA nanoparticles (10 mg/kg) were i.p. injected into the mice at 0.5 h after LPS injection in group 2 and group 4 (Figure 3A). At different time points (24 h, 36 h and 48 h) post PDA injection, the mice were i.p. injected with Lipo@HRP&ABTS and imaged by a LAZR PA Imaging System.³⁷ LPS treatment resulted in strong photoacoustic signals in mouse abdomen owing to the inflammation-induced production of H₂O₂ (Figure 3B&3C). Notably, administration of PDA remarkably reduced H₂O₂-associated photoacoustic signals in the mouse abdomen, suggesting largely decreased H₂O₂ levels in mice treated with PDA (Figure 3B&3C). Furthermore, blood samples were collected at different time points for measurement of inflammation-related cytokines including TNF- α and IL-1 β . It was found that serum levels of TNF- α in 'LPS + PDA' treatment group were significantly reduced at 24 h, 36 h, and 48 h post PDA treatment compared to LPS group (Figure 3D), whereas serum levels of IL-1 β also showed the same decreasing trend at 36 h post PDA treatment (Supporting Information Figure S4). Therefore, both PA imaging and serum cytokine data indicate that the inflammatory response initiated during LPS-induced acute peritonitis is greatly attenuated by treatment with PDA nanoparticles.

ALI is even more common than acute peritonitis in the clinic. In our experiments, ALI model

was established by intranasal instillation of LPS into mice (Figure 4A). Thereafter, these mice were intravenously (i.v.) injected or intranasally (n.a.) dropped with PDA nanoparticles (Figure 4A) at the different time points post LPS stimulation. The biodistribution analysis based on ¹³¹I-labeled PDA nanoparticles revealed that PDA nanoparticles predominantly accumulated in the lung at 24 h post PDA injection or dripping (Figure 4B & C). In comparison with i.v. injection, the n.a. dripping method led to a much higher accumulation of PDA nanoparticles in the lung with lower retention in other organs such as liver and spleen (Figure 4B & C).

Based on the biodistribution analysis, in order to seek the best method and timing of PDA administration to treat ALI, we further studied the therapeutic effect of PDA nanoparticles under either n.a. dripping or i.v. injection. Firstly, the administration time of PDA nanoparticles was assessed. It was uncovered that the earlier PDA nanoparticles were administrated for ALI treatment, the better therapeutic effect could be, no matter n.a. dripping or i.v. injection (Figure 4D - G). Notably, when PDA nanoparticles were administrated within 0.5 h after ALI was initiated, lung myeloperoxidase (MPO) activity, total leukocyte numbers, total protein concentrations, and total neutrophil counts were all significantly down-regulated, suggesting the remarkable therapeutic effect of PDA nanoparticles to treat ALI (Figure 4D - G). Thereafter, the therapeutic efficacy between n.a. dripping and i.v. injection was compared. It is well documented that dysregulated leukocytes, especially neutrophils, play an important role in aggravating ALI.^{4,9} Interestingly, compared to i.v. injection, n.a. dripping of PDA nanoparticles resulted in a more markedly decrease in total protein concentrations, especially the total leukocyte and neutrophil numbers (Figure 4E - G). Hematoxylin and eosin (H&E) staining of lung tissues collected at 24 h also demonstrated that n.a. administration

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of PDA offered better therapeutic outcome compared to that achieved by the i.v. injection route for ALI treatment (**Supporting Information Figure S5**). Therefore, we chose intranasal dripping at 0.5 h post initiation of ALI to evaluate the therapeutic effect of PDA nanoparticles in details.

Based on above findings, the time-dependent efficacy of PDA nanoparticles to ameliorate LPS-induced ALI was further evaluated. ALI model was induced by LPS (Figure 5A) and mice were randomly divided into four groups: PBS group, PDA group, LPS group, and LPS + PDA group. Mice were n.a. dropped by PDA (10 mg/kg) at 0.5 h post LPS inhalation. Administration of PDA markedly attenuated proinflammatory cytokines IL-6, TNF- α , and CXCL-2 in BALF of LPS-induced ALI mice at all tested time points, but did not affect these cytokine levels in normal healthy mice (Figure 5B, 5C & 5D). Meanwhile, PDA significantly reduced total leukocytes and neutrophils counts, lung MPO activity, and total protein levels in BALF of LPS-induced ALI mice (Figure 5E - 5H).

Histological examination of the lung tissues was further conducted to directly evaluate the therapeutic outcome. In the lung of the 'LPS group', epithelioid cell proliferation and infiltration of inflammatory cells (mainly neutrophils) were observed (Figure 6). By contrast, administration of PDA could effectively ameliorate lung morphological changes and significantly reduce neutrophil infiltration as observed in the 'LPS + PDA group' (Figure 6). These results indicated that PDA could observably alleviate LPS-induced ALI in vivo, promising the ALI therapy.

Conclusion

In summary, PDA nanoparticles were developed for treatment of acute inflammation-induced injury, as demonstrated in both murine acute peritonitis and ALI models. The characteristics of PDA

nanoparticles in eliminating ROS were carefully studied for understanding the antioxidation mechanisms. We uncover that PDA nanoparticles could act as a ROS scavenger not only as a reducing agent to directly react with ROS, but also as a catalyst to trigger decomposition of H_2O_2 . At the cellular level, PDA nanoparticles strongly down-regulated intracellular ROS levels in response to inflammatory stimulation. Consistent with the anti-oxidative and anti-inflammatory properties of PDA nanoparticles, in vivo ROS generation during acute peritonitis was substantially relieved in mice treated with PDA, as confirmed by photoacoustic imaging using a H_2O_2 -specific nanoprobe. The anti-inflammation activity of PDA nanoparticles was further evaluated in the ALI model. After a single dose of PDA treatment, PDA nanoparticles markedly improved ALI, with significantly reduced neutrophil infiltration, decreased BALF protein concentrations, and improved lung morphological alterations. Therefore, an interesting therapeutic nano-drug based on PDA nanoparticles is developed in this work, useful for not only protecting the tissue from ROS injury, but also improving the behavior of dysregulated neutrophils to ameliorate acute inflammation. Such a strategy may significantly improve the clinical outcomes for not only ALI, but also other inflammatory diseases with dysregulated neutrophils, such as acute respiratory distress syndrome, acute gout of liver encephalopathy and epilepsy. This study thus presents the great promising of applying nanomedicine for anti-inflammation therapy.

Materials and methods

Materials. Dopamine hydrochloride, 2,2'-Azino-bis(3-ethylbenzthiazoline-6-sulfonic acid) (ABTS), horseradish peroxidase (HRP), lipopolysaccharide (LPS) and 2',7'-Dichlorofluorescin

diacetate (DCFH-DA) were purchased from Sigma-Aldrich and used as received. Hydrogen peroxide (H₂O₂) was purchased from China National Pharmaceutical Group Corporation. Raw 264.7 macrophage cells were obtained from American Type Culture Collection (ATCC). DMEM culture medium was obtained from Thermo Fisher Scientific Inc. Female Balb/c mice were purchased from Nanjing Peng Sheng Biological Technology Co. Ltd.

Preparation of PDA nanoparticles. PDA nanoparticles were synthesized according to a method previously reported.^{16, 18} Briefly, 90 mg dopamine hydrochloride was dissolved in 45 ml deionized water, into which 0.38 ml of 1 M NaOH solution was added. After self- polymerization for 2 h, PDA nanoparticles were obtained by centrifugation (14800 rpm) and washed with deionized water for several times.

Characterization. The morphology of PDA was characterized with FEI Tecnai F20 transmission electron microscopy. UV–Vis–NIR absorption spectra were measured by a GENESYS 10S UV–Vis spectrophotometer. The hydrodynamic light scattering (DLS) was acquired by a Zetasizer Nano Z (Malvern).

Catalysis of H₂O₂ by PDA nanoparticles. The O₂ release in the H₂O₂ solution with or without PDA (0.01 mg/ml) was measured by a dissolved oxygen meter (Rex, JPBJ-608, China). The decomposition of H₂O₂ (100 μ M) was determined by an ABTS assay. A liposomal H₂O₂-specific nanoprobe, Lipo@HRP&ABTS, was synthesized according to our recently reported protocol.³⁷ PDA

solutions with different concentrations (0.01, 0.02, 0.03 mg/ml) were mixed with H₂O₂ for 5-10 minutes, and then incubated with Lipo@HRP&ABTS solution. The absorbance of the mixture after reaction was measured using a UV–Vis spectrophotometer.

Cell uptake of PDA nanoparticles. To test the cell uptake of nanoparticles, Raw 264.7 cells, a murine macrophage cell line, were incubated with FITC labeled PDA nanoparticles (PDA-FITC, 80 μ g/ml) for various periods of time. Afterwards, the cells were washed by PBS three times and imaged under a confocal fluorescence microscope (OLYMPUS, IX73).

Intracellular ROS scavenging by PDA nanoparticles. To evaluate the ROS scavenging in vitro, Raw 264.7 cells were seeded in 24-well plates and incubated with PDA ($80 \mu g/ml$) for 6 hours before adding H₂O₂ (100 and 200 μ M). After 30 minutes of exposure to H₂O₂, the cells were washed by PBS twice and then incubated with DCFH-DA. Flow cytometry (ESP Elite, Beckman Coulter, Fullerton, CA, USA) was used to test the levels of ROS in the cells by detecting the mean fluorescence intensity (MFI) of DCFH-DA fluorescence. For cell viability assay, Raw 264.7 cells were seeded in a 96-well plate and mixed with PDA ($80 \mu g/ml$), with or without adding H₂O₂. After 24 hours incubation, the cells were washed and the cell viability was determined by the cell counting kit (CCK-8) assay. For the anti-inflammatory study of PDA in vitro, PDA nanoparticles ($80 \mu g/ml$) were incubated with Raw 264.7 cells for 6 hours. Then, the cells were treated with LPS. Intracellular ROS levels were assessed using DCFH-DA staining by flow cytometry. Additionally, DCFH-DA fluorescence in the cells was visualized by a confocal fluorescence microscope (OLYMPUS, IX73).

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Peritoneal inflammation amelioration for PA imaging. All animal procedures were performed in accordance with the Guidelines for Care and Use of Laboratory Animals of Soochow University and approved by the Animal Ethics Committee of Soochow University. Female BALB/c mice (8-10 weeks) were purchased from Nanjing Peng Sheng Biological Technology Co Ltd and randomly divided into four groups: phosphate buffered solution (PBS) control, PDA control, LPS-treated group, and PDA + LPS-treated group (4 mice per group), which were intraperitoneally (i.p.) injected with PBS (0.1 ml), PDA (10 mg/kg), LPS (5 mg/kg), and PDA (10 mg/kg) + LPS (5 mg/kg), respectively. At different time points (24 hours, 36 hours, 48 hours) after injection, the mice were i.p. injected with H₂O₂-responsive probe Lipo@HRP&ABTS for photoacoustic (PA) imaging of peritoneal inflammation by in vivo detection of H₂O₂.³⁷ All images on the abdomen were assessed using a Vevo LAZR Imaging System (FujiFilm VisualSonics Inc.). Besides, serum samples in all groups were collected from the mouse eyes. And serum cytokines (TNF- α , IL-1 β) were analyzed with ELISA kits (eBioscience, San Diego, CA, USA), according to the manufacturer's instructions. All animal studies were approved by Soochow University Laboratory Animal Center.

Biodistribution of PDA nanoparticles. ¹³¹I-labeled PDA was prepared according to the previous reports.²³ For intravenous route, female BALB/c mice were intravenously (i.v.) injected with ¹³¹I-PDA (PDA: 10 mg/kg, 200 μ Ci of ¹³¹I). After 24 hours post injection, major organs including liver, spleen, kidney, heart, lung, skin, muscle, bone, stomach and small intestine were collected and measured by the gamma counter (Science and Technology Institute of China in Jia Branch Innovation

Co., Ltd.).

LPS-induced ALI. Female BALB/c mice (8-10 weeks) were used for those experiments. The ALI model was induced with n.a. instillation of LPS, according to the standard method previously reported.⁴¹⁻⁴² At 24, 36 and 48 hours after LPS inhalation, the lung tissues and bronchoalveolar lavage fluid (BALF) were collected after three times lavages and stored at – 80 °C. BALF samples were directly used for analyses without any additional treatment.

Measurement of lung MPO activity. The groups were the same as before: PBS group, PDA control, LPS-treated group, and PDA + LPS-treated group (4 mice per group). Right lung tissues of the mice were homogenized and centrifuged. The supernatant solution was used to analyze and quantify lung MPO activity with the MPO assay kit (Nanjing Jiancheng Bioengineering Institute) according to the vendors' instructions.

Assessment of leukocyte number, total protein concentration, inflammatory cytokines and chemokine in BALF. After lung lavage for three times, BALF was obtained.⁴¹ A standard haemocytometer and Wright-Giemsa staining assay was used to assess the counts of total and differentiated leukocytes. A BCA protein assay kit (Piece, Rockford, IL, USA) was used to measure the total protein concentration in BALF. The levels of inflammatory cytokines (TNF- α , IL-6 and IL-1 β) and chemokine (CXCL2) in BALF were determined by ELISA (eBioscience, San Diego, CA, USA), according to the supplier' instructions.

Lung histological examination. Left lung tissues of the mouse were harvested and fixation in 4 % paraformaldehyde for hematoxylin and eosin (H&E) staining at 24 hours, 36 hours and 48 hours after LPS instillation. Lung slices were observed for evaluating morphological changes by a digital microscope (Leica QWin).

Statistical analysis. The statistical analyses were performed by using Student's t-test for the p value (* p < 0.05, ** p < 0.01, *** p < 0.001). The values were expressed as mean \pm standard deviation (SD) for control and experimental samples.

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Conflict of interest

The authors declare that they have no competing interests.

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Figure 1. Preparation and ROS elimination characterization of PDA. (A) A schematic illustration for the synthesis of PDA and effect of PDA for protecting the tissue from oxidative and inflammatory injury. (B) TEM image of PDA nanoparticles. (C) Hydrodynamic diameters of PDA nanoparticles in water. (D) UV-Vis-NIR absorbance spectra changes of the reaction solutions measured at different time point (i.e. 1 min, 10 min, 30 min) after H₂O₂ (20 μ M) was incubated with PDA (0.01 mg/ml). The absorbance was originated from the Lipo@HRP&ABTS probe in the presence of H₂O₂. (E&F) UV-Vis-NIR absorbance spectra (E) and the photo (F) of the mixture solutions, including free Lipo@HRP&ABTS, free PDA, and Lipo@HRP&ABTS mixed with H₂O₂ (25 μ M) pre-incubated with different concentrations of PDA for 10 min. (G) PA images (up) and PA signal intensities at 780 nm (bottom) of the mixture solutions. 1: Lipo@HRP&ABTS solution. 2: PDA (0.03 mg/ml) solution. 3 - 6: the mixture solutions of Lipo@HRP&ABTS, H₂O₂ (25 μ M) and PDA at different concentrations (0.03, 0.02, 0.01 and 0.00 mg/ml).



Figure 2. In vitro antioxidant and anti-inflammatory experiments. (A&B) Intracellular ROS scavenging by PDA in LPS-treated (A) or H_2O_2 -treated (B) Raw 264.7 cells as detected by the flow cytometer using DCFH-DA as the ROS-specific probe. (C) Mean fluorescence intensity (MFI) calculated based on the flow cytometer data in (A&B). (D) The relative viabilities of Raw 264.7 cells incubated with H_2O_2 (200 µM and 100 µM) with or without PDA (80 µg/ml). ** p < 0.01, *** p < 0.001. (E) Confocal fluorescence images of ROS levels in the LPS-treated cells with or without PDA treatment using DCFH-DA as the ROS probe. Scale bar = 25 µm.



Figure 3. In vivo PA imaging to monitor relief of acute peritonitis by PDA treatment. (A) Schematic illustration showing the experimental design. LPS was i.p. injected into mice to induce acute peritonitis, which was ameliorated by PDA. The therapeutic effect could be monitored by PA imaging for in vivo H₂O₂ detection. (B) In vivo PA images of mouse abdomen at different time points (24 h, 36 h and 48 h) post injection of LPS with or without i.p. injection of PDA. (C) PA signals at 780 nm for mouse abdomen of all groups based on PA imaging data in (B). (D) Serum cytokine TNF- α from mice evaluated at 24 h, 36 h and 48 h post injection of LPS. * p < 0.05 and ** p < 0.01.



Figure 4. PDA injection with different administration routes for in vivo ALI treatment. (A) Schematic illustration showing the establishing process of the ALI model and the two treatment routes. (B&C) The biodistribution of PDA by i.v. injection (B) or n.a. dripping (C) at 24 h post treatment. He: heart; Li: liver; Sp: spleen; Lu: lung; Ki: kidney; St: stomach; Si: small intestine; Sk: skin; Bo: bone; Mu: muscle. (D-G) Comparison of therapeutic responses between i.v. injection and n.a. administration of PDA evaluated at different intervention time points (0.5 h, 2 h, 4 h and 6 h) post model establishment, including the lung MPO activity (D), the total protein concentration (E), the total leukocyte number (F), and the total neutrophil number (G). P values were calculated by the Student's t-test. $\neq p < 0.05$, $\neq \neq p < 0.01$ means n.a. administration group versus i.v. injection group. ** p < 0.01, ** p < 0.001 versus LPS group.



Figure 5. In vivo therapy with PDA for ALI treatment. (A) Schematic illustration showing the ameliorative process of ALI induced by PDA. (B - D) BALF cytokines including IL-6, TNF- α , and CXCL2 from mice measured at 24 h, 36 h and 48 h post dripping of LPS. (E-H) The assessment in lung MPO activity (E), the BALF total protein concentration (F), the BALF total leukocyte number (G), and the BALF total neutrophil number (H) at different time point (24 h, 36 h and 48 h) after LPS inhalation. * p < 0.05, ** p < 0.01 and *** p < 0.001.



Figure 6. In vivo lung histological examination. H&E stained images of the lung tissues collected from all groups. Scale bar (red line) = $500 \ \mu m$; Scale bar (black line) = $50 \ \mu m$.