Acute exposure to silica nanoparticles aggravate airway inflammation: different effects according to surface characteristics

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Silica nanoparticles (SNPs) are widely used in many scientific and industrial fields despite the lack of proper evaluation of their potential toxicity. This study examined the effects of acute exposure to SNPs, either alone or in conjunction with ovalbumin (OVA), by studying the respiratory systems in exposed mouse models. Three types of SNPs were used: spherical SNPs (S-SNPs), mesoporous SNPs (M-SNPs), and PEGylated SNPs (P-SNPs). In the acute SNP exposure model performed, 6-week-old BALB/c female mice were intranasally inoculated with SNPs for 3 consecutive days. In the OVA/SNPs asthma model, the mice were sensitized two times via the peritoneal route with OVA. Additionally, the mice endured OVA with or without SNP challenges intranasally. Acute SNP exposure induced significant airway inflammation and airway hyper-responsiveness, particularly in the S-SNP group. In OVA/SNPs asthma models, OVA with SNP-treated group showed significant airway inflammation, more than those treated with only OVA and without SNPs. In these models, the P-SNP group induced lower levels of inflammation on airways than both the S-SNP or M-SNP groups. Interleukin (IL)-5, IL-13, IL-1β and interferon-γ levels correlated with airway inflammation in the tested models, without statistical significance. In the mouse models studied, increased airway inflammation was associated with acute SNPs exposure, whether exposed solely to SNPs or SNPs in conjunction with OVA. P-SNPs appear to be relatively safer for clinical use than S-SNPs and M-SNPs, as determined by lower observed toxicity and airway system inflammation.

INTRODUCTION

Nanotechnology is a recently developing field that allows for the production of nanoparticles (NPs) at supramolecular levels, typically <100 nm. Nanotechnology has been applied to many industrial fields, including textiles, cosmetics, food packaging and medicines. In particular, the application of nanotechnology in the medical field has improved various biotechnological processes such as drug synthesis and their targeted delivery. Nanotechnology is expected to progress and expand in the future.1–4 The increasing applications of nanotechnology warrant an increased study of the toxicity of NPs, with emphasis on their potential safety issues. Studies thus far have revealed that NPs can alter immune function and cause inflammation in various organs and systems. The toxicity of NPs, particularly associated with the liver, kidney, lung and immune system, has been clearly established.5–10 These toxicity studies and their results have been helpful for determining the concentration, size and surface pattern of NPs that should be selected for safe application.

Of the various clinical routes of NP administration, inhalation is the main route of exposure due to the small size of NPs and their ease of airborne dispersion. Humans can be easily exposed to NPs unintentionally via aerosolization. Recent clinical studies have shown that occupational exposure during the manufacturing of NPs can cause respiratory diseases such as asthma and bronchitis.11–13 Therefore, airway toxicity related to NPs has garnered considerable recent attention. Asthma, a chronic inflammatory airway disease characterized by variable bronchial obstruction, is a common, highly prevalent and significant airway disease. There are

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many risk factors for developing or aggravating asthma and asthma-like airway diseases such as air pollutants, cigarette smoking and viral infection. These factors, as well as fine materials like NPs, may induce and aggravate asthma. Many studies seeking to understand and reduce the toxicity associated with NPs, have sought to examine them in more detail, as well as their characteristics, such as surface charge and surface pattern, which contribute to airway inflammation.

Silica, or silicon oxide (SiO₂), is naturally found as sand or quartz. Silica is a main component of glass and concrete. Owing to its optical, thermodynamic characteristics and chemical durability, silica has been used as a heat-resistant glass and semiconductor; it is commonly used in optical communications systems and ceramic technology. However, silica is a well-known cause of pneumoconiosis, often presenting as chronic bronchitis, emphysema and lung fibrosis. In addition, lung cancer has been associated with silicosis, cigarette smoking and viral infection. These factors, as well as fine materials like NPs, may induce and aggravate asthma. Many studies seeking to understand and reduce the toxicity associated with NPs, have sought to examine them in more detail, as well as their characteristics, such as surface charge and surface pattern, which contribute to airway inflammation. Due to these potential health risks, the use of silica should be carefully regulated and handled with proper safety precautions.

Silica NPs (SNPs) have recently been used in biosciences, medicines and daily life, after studies have revealed that SNPs are relatively safe. Unfortunately, some studies have since revealed inhalant-associated toxicity of SNPs in chronic exposure conditions; however, the acute effects of SNPs on airway systems has not yet been fully evaluated in vivo. Especially, aggravating effects of SNPs on airway systems in asthma mice model induced by ovalbumin (OVA) sensitization and challenges has not been well known. Moreover, there are few studies examining a counterplan for reducing toxicity related to SNPs. In this study, exposure to SNPs alone as well as in conjunction with OVA was induced to determine the toxicity of these NPs on airway systems and examine any associated aggravation of airway inflammation. Furthermore, various surface patterns were studied to determine a possible, relatively safe SNP with reduced toxicity.

MATERIALS AND METHODS
Preparation and characterization of SNPs
This study examined three types of SNPs: spherical (S-SNPs), mesoporous (M-SNPs) and PEGylated (P-SNPs; PEG, polyethylene glycol). To prepare the S-SNPs, 11.87 ml of ethanol (Sigma-Aldrich, St. Louis, MO, USA), 2.1 ml of distilled (DI) water, 0.67 ml of tetraethoxysilane (Sigma-Aldrich) and 0.36 ml of 5 M NH₄OH (Sigma-Aldrich) were mixed in a vial and stirred for 24 h. M-SNPs were synthesized using 53.4 g of DI water, 6.24 g of cetyltrimethylammonium chloride (Sigma-Aldrich) and 0.3 g of sodium acetate (Sigma-Aldrich), were mixed well, and then heated in a 65 °C oil bath. After heating for 1 h, 3.9 ml of tetraethoxysilane was slowly added to the mixture and then stirred for 24 h. The resulting suspensions were purified by centrifugation, re-dispersed in ethanol and DI water, and then finally re-dispersed in DI water.

P-SNPs were prepared by surface functionalization of S-SNP. Briefly, thiol-functionalized S-SNP was prepared by adding (3-mercaptopropyl) trimethoxysilane (10% of tetraethoxysilane) 5 h after the addition of NH₄OH during synthesis of S-SNP. After purification, these particles were conjugated with maleimide-functionalized PEG (molecular weight: 2000) in DI water for 24 h (weight ratio: ~0.51). Excess amount of PEG was removed by centrifugation and dispersed in DI water to obtain P-SNP.

Visualization and surface area calculation of SNPs
To characterize the SNPs, transmission electron microscopy was performed using a Tecnai 20 (FEI Co., Hillsboro, OR, USA). The SNPs were diluted to a concentration of 1 mg ml⁻¹ in a phosphate-buffered saline solution. Transmission electron microscopy specimens of the diluted SNPs were prepared by placing one drop on a carbon-coated copper grid, which was allowed to dry at 60 °C for 15 min. Transmission electron microscopy analyses were performed at 25 ± 2 °C; each SNP measured ~100 nm (Figure 1). These analyses indicated that S-SNPs and M-SNPs had spherical and mesoporous morphologies, respectively. The P-SNPs also demonstrated spherical morphology.

Nitrogen sorption experiments were performed at 77 K with the Micromeritics ASAP 2020 system (Micrometerics, Norcross, GA, USA). Specific surface area and pore size of the particles were calculated using the Brunauer-Emmett-Teller equation and the Barrett-Joyner-Halenda method, based on the desorption branch, respectively.

![Figure 1](image-url) Transmission electron microscopy images of (a) S-SNPs, (b) M-SNPs and (c) P-SNPs. M-SNP, mesoporous silica nanoparticle; P-SNP, PEGylated silica nanoparticle; S-SNP, spherical silica nanoparticle.
Animals
Female BALB/c mice, between 5 and 6 weeks old, were used in this study (Orient, Daegeon, Korea). All mice were maintained at conventional animal facilities under specific, pathogen-free conditions (room temperature of 21–24 °C, relative humidity between 45% and 70%, with a 12-h light/dark cycle). All experimental protocols were approved by the Animal Research Ethics Board of Yonsei University (Seoul, Korea).

Direct acute exposure to SNPs
To establish the effects of SNPs in the airway system, the mice were intranasally inoculated with SNPs (S-SNPs, M-SNPs and P-SNPs in S-SNP group, M-SNP group and P-SNP group, respectively) or saline on days 0, 1, and 2 (Figure 2a). All mice were killed 48 h after the last intranasal inoculation.

Establishment of OVA-induced acute asthma model with SNP exposure (OVA/SNPs asthma model)
To make acute asthma mice model, mice were sensitized with an intraperitoneal injection of either 200 μl of 10 μg OVA (Sigma-Aldrich) suspended in 1% aluminum hydroxide (Resorptar, Indergen, NY, USA) or 200 μl of saline on days 1 and 14. On days 21, 22 and 23, the OVA-sensitized mice were challenged intranasally with 30 μl of OVA (1 mg ml⁻¹) in saline solution. SNPs (S-SNPs, M-SNPs and P-SNPs in S-SNP group, M-SNP group and P-SNP group, respectively) or saline were co-administered by intranasal inoculation of 200 μg (Figure 2b). All mice were killed 48 h after the last intranasal inoculation.

Measurement of airway hyper-responsiveness
Airway hyper-responsiveness (AHR) in response to inhaled aerosolized methacholine was measured using a forced oscillation technique (FlexiVent; SCIREQ, Montreal, QC, Canada) 48 h after the last intranasal inoculation. Cannulas were inserted into anesthetized mice via tracheostomy, after which the mice were connected to a ventilator. Aerosolized phosphate-buffered saline or methacholine at varying concentrations, 3.125, 6.25, 12.5, 25.0 and 50.0 mg ml⁻¹, was diffused to the mice for 10 s via a nebulizer. AHR was measured and continuously recorded for up to 2 min.

Bronchoalveolar lavage and inflammatory cell counting
To collect bronchoalveolar lavage fluid (BALF), lung lavage was performed three times by passing 1 ml of Hank’s Balanced Salt Solution through the tracheal tube. The recovered BALF was centrifuged at 4 °C for 3 min at 1500 g. The cellular pellet was resuspended in 300 μl Hank’s Balanced Salt Solution. One part of the resuspended BALF cells (250 μl) were centrifuged by cytocentrifugation (Cytospin 3; Thermo, Waltham, MA, USA) at 4 °C for 5 min at 1000 r.p.m. and were pelleted to cytospin slides. The slides were stained with hematoxylin and eosin (Hemacolor, Merck, Darmstadt, Germany) and counted the differential inflammatory cells including neutrophils, eosinophils, lymphocytes and macrophages (200 cells per cytospin). The remaining resuspended BALF cells (50 μl) were used for counting the total number of cells using a hemocytometer and trypan blue staining.

Measurement of cytokine levels
Concentrations of interleukin (IL)-5, IL-13, IL-1β and interferon-γ in the lung tissues were assessed by ELISA (R&D Systems, San Diego, CA, USA) according to the manufacturer’s instructions. The ELISA detection ranges for IL-5, IL-13, IL-1β and interferon-γ are 31.3–2000, 62.5–4000, 15.6–1000 and 31.3–2000 pg ml⁻¹, respectively.

Histological analysis
After the use of one lung in the collection of BALF, the other lung was fixed in 4% formalin and embedded in paraffin. Lung sections were cut into 3- to 4-μm thick segments and stained with hematoxylin and eosin and periodic acid-Schiff for histological analysis. The slides were observed under light microscope.

Statistical analysis
All results are expressed as the mean ± s.e. One-way analysis of variance was performed using SPSS statistical software version 12.0 (SPSS Inc., Chicago, IL, USA). The AHR data were analyzed with repeated analysis of variance measures, followed by a post hoc Bonferroni test. The remaining data were analyzed with one-way analysis of variance followed by a post hoc Bonferroni test. P-values <0.05 were considered statistically significant.

RESULTS
Difference and chemical characteristics of three kinds of SNPs
The surface area calculated using the Brunauer-Emmett-Teller equation of S-SNPs and M-SNPs was 12.7 and 70.6 m² g⁻¹, respectively. The surface area of P-SNPs was similar to those of S-SNPs. The average pore diameter of the M-SNPs was calculated to be 3.0 nm using the Barrett-Joyner-Halenda method.

Effects of acute SNP exposure
Significant airway inflammation was observed for each type of SNP treatment; however, the level of inflammation varied according to the surface characteristics of the SNP used. The results of BALF examination revealed statistically significant variation in the differential cell counts performed on the S-SNP group as compared with the control group, including the total cell count, macrophage count and neutrophil count. Lymphocytes were significantly higher in the S-SNP group than in the other groups. The M-SNPs and P-SNP groups also demonstrated higher total cell and macrophage counts than did the control group, but without statistical significance. The M-SNP...
and P-SNP groups induced a smaller increase in cell count than did the S-SNP group (Figure 3a).

Significantly higher AHR was observed in the S-SNP group than in the control group. The M-SNP and P-SNP groups also demonstrated higher AHR than did the control group, but lower than the AHR observed in the S-SNP group. No statistically significant differences were observed (Figure 3b).

In each of the SNP groups, histologic examination revealed higher peribronchial and perivascular inflammation than that observed in the control group. The S-SNP group demonstrated marked increase in bronchial epithelial thickness with abundant inflammatory cell infiltration. The histologic findings between M-SNP and P-SNP groups could not be visually discriminated (Figure 4).

Higher levels were obtained for the S-SNP group than the control group for each cytokine measured, including IL-5, IL-13, IL-1β and INF-γ. The levels of IL-5 and IL-13, which are known as helper T cell type 2 (Th2) cytokines, were lower in the P-SNP group than in the S-SNP group; however, none of the cytokine measurements demonstrated a statistically significant difference (Figure 5).

**Effects of acute SNP exposure in the OVA-induced asthma model**

SNPs, inoculated thrice in conjunction with OVA, induced asthma in the mice. Upon examination of the BALF, the total cell counts in the S-SNP and M-SNP groups were significantly higher than that in the control group. The number of macrophages was significantly higher in the S-SNP and M-SNP groups than in all of the other groups—the control group, OVA-only asthma group and P-SNP group. Eosinophils were significantly elevated in the S-SNP and M-SNP groups than in the control group. The total number of cells and eosinophils were higher in the P-SNP group than in the control group, but lower compared with those of the S-SNP and M-SNP groups (Figure 6a).

AHR measurements in the OVA asthma group were greater than in the control group, yet without statistically significant meaning. Overall, no significant difference in AHR was observed among the various SNP groups (Figure 6b).

Significantly more cellular infiltration in peribronchial and perivascular tissues was observed in the OVA asthma group, as well as in the three SNP groups, as compared with the control group. No statistical difference was observed among the OVA asthma group and each of the SNP groups (Figure 7).

IL-5 and IL-13 levels were statistically significantly greater in the OVA asthma group and SNP groups than in the control group. No significant statistical difference was noted between the OVA asthma group and each of the SNP groups. Similarly, the levels of IL-1β and INF-γ were higher in the OVA asthma group and SNP groups than in the control group; however, no statistically significant difference was observed. The P-SNP group demonstrated lower levels of IL-5, IL-13, IL-1β and INF-γ than the OVA asthma and other SNP groups, except the control group (Figure 8).

**DISCUSSION**

This study demonstrated that acute exposure, simulated just once per day for 3 days, can harm the respiratory system. Acute SNP exposure, as demonstrated here, can induce significant airway inflammation and aggravate allergic airway inflammation in the OVA-induced asthma models.

In this study, short-term exposure to SNPs induced respiratory inflammation with associated AHR in each of the S-SNP, M-SNP and P-SNP groups. Both BALF and histologic examination showed significantly higher cell counts in the S-SNP group than in the control group. Acute exposure to SNPs in conjunction with OVA (OVA/SNPs asthma model) also induced severe airway inflammation when compared with those treated with OVA alone. No significant difference in AHR was observed among all the groups, including mice in the OVA asthma group and OVA/SNPs asthma mice. Both models showed increased levels of Th1 cytokines (IL-1β and interferon-γ) and Th2 cytokines (IL-5, and IL-13), without significant statistical difference. Some studies have shown that Th2 cytokines have a key role in airway toxicity.10,29,30 However, in this study, increased macrophage counts in BALF samples and increased levels of IL-1β and interferon-γ were
observed, particularly in the SNP models. It could be assumed that this airway inflammation was induced not only via the Th2 pathway but also via the Th1 pathway.

Th1- and Th2-related immunity maintain a delicate balance. If one side increases, the other side naturally decreases. Several Th1 enhancers reduce Th2-related immunity. OVA induces Th2-related immunity, whereas SNPs are hypothesized to induce Th1-related immunity. Therefore, OVA/SNP groups will induce a Th2-related response to a lesser extent than that in only OVA-treated groups. Indeed, the AHR and the IL-5 and IL-13 levels were lower in the OVA/SNP group compared to the OVA group. In contrast, the IL-10 and IFN-γ levels were higher in the OVA/SNP group compared to the OVA group. This suggests that the Th1-Th2 balance is perturbed by silica nanoparticles.
IL-13 levels in OVA/SNP groups decreased compared with that in the OVA group (Figures 6b and 8). SNP, a Th1 immunity enhancer, may reduce the Th2-related immunity by lowering AHR and by lowering the levels of IL-5 and IL-13. In contrast, OVA, a Th2 enhancer, may reduce Th1-related immunity. The decrease in macrophage counts in BALF and pathologic inflammation revealed that the OVA/SNP model induced severe inflammation. Although not statistically significant, we believe that SNP slightly reduced Th2-related immunity induced by OVA by inducing Th1-related immunity. Therefore, overall total inflammation induced by OVA/SNP is similar to or more severe than that in the OVA- or SNP-treated groups.

We selected a specific dose of SNPs (200 μg per mouse) for this study. Previous studies have revealed that there is a dose-dependent effect of SNP-induced airway inflammation. Thus, most current studies use a specific dose of SNP (12.5–200 μg) to evaluate its toxicity. The number of SNP in ambient air range from $2 \times 10^4$ to $2 \times 10^5$ cm$^{-3}$, with mass concentrations of > 50 μg m$^{-3}$ near major highways. Average breathing volume per day is approximately 15–25 m$^3$. An individual working near a major highway might therefore inhale anywhere between 750 and 1250 μg of SNP in a day. In mouse experiments of toxicity to a specific material, 10- to 1000-fold of what is considered a toxic dose of the study compound to be investigated is typically used. We selected an SNP dose of ~400-fold, calculated based on the mass of SNP typically inhaled relative to body mass (human: 0.02 μg g$^{-1}$; mouse (in this study): 8 μg g$^{-1}$). We believe that this SNP dose of 200 μg per mouse is clinically feasible. Indeed, these experiments showed significantly increased AHR, which is of clinical importance, inducing clinical signs such as cough and dyspnea under the stimulation of less irritable materials.

Many studies have demonstrated that the toxicity of SNPs depends on several factors, including size, surface area, electric charge and function. Some studies have revealed correlation between the surface pattern of SNPs and the level of toxicity; however, results vary according to experimental setting and designs. In this study, the toxicity of acute SNP exposure varied owing to the surface pattern of the SNP. In acute SNP exposure models, the P-SNP and M-SNP groups induced less airway inflammation than the S-SNP group, without statistical significance. Among OVA/SNP asthma mice, those in the P-SNP group induced lesser signs of airway inflammation than those in the S-SNP and M-SNP groups, and the difference was statistically significant. The contact area is a critical factor for inducing toxicity during the interaction between SNPs and tissues. The contact and surface area of M-SNPs is largest among the three kinds of SNPs (M-SNPs, 70.6 m$^2$ g$^{-1}$; S-SNPs, 12.7 m$^2$ g$^{-1}$). Smaller surface area has lesser opportunity to interact with tissue to induce inflammation. Therefore, the larger surface area of M-SNPs could induce the most severe inflammation compared with the other SNPs in this study. In P-SNPs, the surfaces of S-SNPs are functionalized with the biocompatible PEG ligands. Therefore, P-SNPs exhibited the lowest toxicity among three different types of SNPs.

In summary, the toxicity of SNPs was lower in the P-SNP group than in the S-SNP or M-SNP group. P-SNPs did induce airway inflammation but lesser than did S-SNPs and M-SNPs induce. P-SNPs induce lesser inflammation on the airway, with reduced inflammation in OVA/SNPs asthma models. P-SNPs may be more suitable for use in various medical/pharmaceutical fields, including drug delivery systems, diagnostic carriers and other medical applications. The harmful effects associated with SNPs should be carefully considered before implementation, particularly when exposure of the airway system is possible.

Although various studies have evaluated the toxicity of SNPs, this study cannot be compared with the previous work owing to significant differences in its experimental design. This study focused on examining respiratory system response after short-term exposure to SNPs, with or without OVA sensitization, as well as the different effects of SNPs according to their surface patterns. These results contribute to the existing knowledge base for SNP usage in scientific and industrial fields.
However, the present study did not completely examine the effect of SNPs in conjunction with OVA in the OVA/SNPs asthma models to determine its use as an adjuvant. The OVA-only group demonstrated lower total cell counts and macrophage counts in BALF than the S-SNP and M-SNP groups. As S-SNPs and M-SNPs can induce airway inflammation on their own, it could not be determined whether the increased effects observed in the OVA/SNP asthma models are due to actions of SNPs alone, or because of an adjuvant effect with OVA. To evaluate more accurate pro-inflammatory effects and adjuvant effects of OVA/SNP models, further study is required. Moreover, the present study revealed that Th2 and Th1 cytokines are...
associated with toxic effects of SNPs; however, the mechanisms by which SNPs induce inflammation on the respiratory system have not yet been fully evaluated. A more accurate evaluation of this mechanism, as well as further cytokine and gene studies, is required.

In conclusion, this study demonstrated the toxic effects of acute SNP exposure. Acute SNP exposure can induce significant airway inflammation, and further aggravate airway inflammation, as demonstrated in the OVA asthma model. In terms of surface characteristics, P-SNPs appear to be safer for clinical use than S-SNPs or M-SNPs.

CONFLICT OF INTEREST

The authors declare no conflict of interest.

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