

Published in final edited form as:

Environ Sci Nano. 2014 April ; 1(2): 161–171. doi:10.1039/C3EN00053B.

Quantitative assessment of inhalation exposure and deposited dose of aerosol from nanotechnology-based consumer sprays†

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Abstract

This study provides a quantitative assessment of inhalation exposure and deposited aerosol dose in the 14 nm to 20 μm particle size range based on the aerosol measurements conducted during realistic usage simulation of five nanotechnology-based and five regular spray products matching the nano-products by purpose of application. The products were also examined using transmission electron microscopy. In seven out of ten sprays, the highest inhalation exposure was observed for the coarse (2.5–10 μm) particles while being minimal or below the detection limit for the remaining three sprays. Nanosized aerosol particles (14–100 nm) were released, which resulted in low but measurable inhalation exposures from all of the investigated consumer sprays. Eight out of ten products produced high total deposited aerosol doses on the order of 10¹–10³ ng kg⁻¹ bw per application, ~85–88% of which were in the head airways, only <10% in the alveolar region and <8% in the tracheobronchial region. One nano and one regular spray produced substantially lower total deposited doses (by 2–4 orders of magnitude less), only ~52–64% of which were in the head while ~29–40% in the alveolar region. The electron microscopy data showed nanosized objects in some products not labeled as nanotechnology-based and conversely did not find nano-objects in some nano-sprays. We found no correlation between nano-object presence and abundance as per the electron microscopy data and the determined inhalation exposures and deposited doses. The findings of this study and the reported quantitative exposure data will be valuable for the manufacturers of nanotechnology-based consumer sprays to minimize inhalation exposure from their products, as well as for the regulators focusing on protecting the public health.

[†]Electronic supplementary information (ESI) available. See DOI: 10.1039/c3en00053b

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Declaration of interest

The authors declare no conflict of interest.

Introduction

The use of engineered nanomaterials in consumer products is gradually becoming pervasive. Depending on the geopolitical area, nanomaterials that are engineered but do not have a novel molecular identity and/or those manufactured in small to moderate quantities are not regulated by government agencies.¹⁻⁵ This provides for an increasing share of common consumer products becoming nanotechnology-based, due to introduction of engineered nanomaterials into such products. The types of products where engineered nanomaterials may be or are used include cosmetics and personal care products, nutritional supplements and drugs, household and industrial use chemicals and antiseptics, all of which may be in the form of liquids or powders and can be easily dispersed into the air.⁶⁻¹² Another category of nanotechnology-based products is solid products which cannot be dispersed, for example, electronics and equipment, structural materials, apparel, *etc.*

When nanotechnology-based consumer products are manufactured, used and disposed, engineered nanomaterials may be released, which can lead to human and environmental exposure.¹³⁻¹⁸ Human exposure can occur through ingestion, inhalation or the cutaneous route.¹⁹ The inhalation exposure route has been identified as potentially the likeliest one for those consumer products which are or can easily be dispersed as aerosol during their normal use, such as sprays or powders.^{13,16,17,20-22}

In our earlier study of quantitative exposures from consumer products, we investigated inhalation exposure from several cosmetic powders.²³ Here, we report the results of a quantitative exposure assessment for selected consumer sprays, which are supposed to be used as antiseptics, cosmetics and personal care products. Ten spray products initially identified by Nazarenko *et al.*¹⁶ were investigated. The selection comprised five pairs of nanotechnology-based and non-nanotechnology-based (regular) sprays with each pair consisting of two products with the same purpose of application. Categorization was based on product labeling and/or marketing as nanotechnology-based or non-nanotechnology-based (regular). When the liquid products are sprayed (used) close to the personal breathing zone, they will generate aerosol particles containing engineered nanomaterials if they are present in the original products. Engineered nanomaterials may be distributed in the generated aerosol in complex ways because of agglomeration.²⁴⁻²⁸ At the same time, we found that wide aerosol size distributions were formed due to spraying of both the nanotechnology-based and regular consumer spray products, including production of nanosized particles as well as super-micron agglomerates by all products.¹⁶ These findings confirmed the potential for nanomaterial exposure from particles within a wide range of sizes. The engineered nanomaterials are likely distributed across both the nanosized aerosol fraction (<100 nm) and larger particles in an agglomerated form, sometimes as large as 20 μm , which was the upper limit of our measuring equipment. However, two things remain unknown: (1) the exact quantities of aerosol particles of different sizes that are inhaled and (2) how much aerosol is deposited in various regions of the human respiratory system, thereby potentially delivering engineered nanomaterials into the body *via* the inhalation route. Such quantitative exposure information is crucial for risk assessment and studies investigating health effects of nanomaterial exposure.

The quantitative investigation of inhalation exposure presented here will be valuable for the manufacturers of nanotechnology-based sprays because it provides insights into how such an exposure occurs and what factors influence its magnitude, which may help formulate the products and design sprayers in a way that will minimize generation of aerosol particles of unwanted sizes. Lastly, this paper provides quantitative exposure data for real consumer spray products acquired from the market which we hope will help in risk assessment and development of any consumer-oriented regulations and/or guidelines.

Materials and methods

Summary

All nanotechnology-based and regular consumer spray products were investigated using transmission electron microscopy (TEM) in order to determine the size, shape, state of agglomeration, and electron beam sensitivity of the particles in them. Where particulate matter was observed, we took micrographs at various magnifications. We then reprocessed the original aerosol size distribution measurement data collected in an earlier study¹⁶ for use in the exposure calculations presented here. In that 2011 study, aerosol size distributions were measured when the sprayers containing the original product (when available) were activated manually in the immediate vicinity of a human mannequin head to simulate use by a consumer. The released particles were sampled through stainless steel tubes inserted in the nostrils of the mannequin head to simulate potential inhalation exposure. For this study, the measurement data from Nazarenko *et al.*¹⁶ were exported from the aerosol instrument manager software (AIM Manager, TSI, Inc., Shoreview, MN, USA) as aerosol particle mass distributions, which were then used in a mathematical model to calculate the masses of “inhaled” and “deposited” particulate matter.

Investigated products

The consumer spray products, for which exposure was assessed quantitatively and reported herein, had been described in an earlier study.¹⁶ The selected sprays, for which aerosol size distributions were measured, included 5 products marketed as nanotechnology-based and 5 regular products. These products, along with their compositions reproduced verbatim from the product labels, are listed in Table 1. The brand names of the products were substituted with descriptive names according to the purpose of application. The five regular products matched the five nanoproducts by their purpose of application and included a pair of topical antimicrobial silver sprays, a pair of facial cosmetic sprays, a pair of hair sprays, a pair of surface disinfectant sprays, and a pair of skin hydrating sprays.

TEM characterization of consumer sprays

A small quantity of each liquid consumer spray was spread on an HC300-Cu TEM grid (Electron Microscopy Sciences, Hatfield, PA, USA) using a glass stick and allowed to dry at room temperature (22–23 °C) and humidity (15–35% RH) for at least 24 hours. A transmission electron microscope (2010F, JEOL Ltd, Tokyo, Japan) in the TEM mode was used to examine these specimens at different magnifications. Digital micrographs with automatically inserted scale bars were taken. For the silver particles where the atomic grid

could be observed, the corresponding micrographs were amended with small image insets showing it.

Simulated use of consumer sprays

The methodology for simulated use of the consumer sprays and aerosol sampling is described in detail elsewhere.¹⁶ Briefly, we placed a human mannequin head inside a level II biosafety cabinet (NUAIRE, Inc., Plymouth, MN, USA), which is equipped with a high-efficiency particulate air (HEPA) filtration system. The biosafety cabinet was furnished with a polyethylene curtain covering the front opening of the cabinet, in which gloves for manual activation of the sprays were fitted. Products were sprayed in close proximity of the mannequin head, but the spray cone was directed towards the back wall of the cabinet, in the same direction where the mannequin head was facing. Thus, the products were not sprayed directly into the face of the mannequin but in close proximity. The mannequin head had two stainless steel tubes inserted into its nostrils. The two stainless steel tubes exited the head at the nape where they were joined by a stain-less steel Y-connector, and the combined aerosol stream passed through conductive tubing into a stainless steel flow splitter and then into the aerosol measurement instruments: a Scanning Mobility Particle Sizer (SMPS) (module combination 3080/3786, TSI, Inc.) and an Aerodynamic Particle Sizer (APS) (model 3321, TSI, Inc.). These instruments measured aerosol size distributions in the size range ~14 nm to 20 μm . Three replicates were carried out for each consumer spray. We exported the measurement data using the AIM software (TSI, Inc.) as aerosol particle mass concentrations assuming a spherical shape of particles and a particle density of 1 g cm^{-3} . The justification for using this particle density has been provided previously.¹⁶ The results of the dose assessment reported herein can be adjusted for a different particle density, if such data become available.

This configuration of the experimental setup allowed measurement of released aerosol particles from the simulated personal breathing cloud. Therefore, we assumed that the concentrations and size distributions of the measured particles are representative of those inhaled during the actual spray application by consumers.

Quantitative exposure assessment

The mass-based aerosol concentrations across the measurement size range ~14 nm to 20 μm were used as input in the mathematical model that had been used earlier for the quantitative exposure assessment of cosmetic powders.²³ Similar to that earlier study, the “inhalation exposure” and “deposited dose” were calculated. “Inhalation exposure” is the aerosol mass entering the human respiratory system during an exposure event. It was calculated separately for several aerosol particle size ranges indicated as subscripts in μm : $\text{PM}_{0.1-0.014}$ (ultrafine aerosol fraction), $\text{PM}_{1-0.1}$ (submicron fraction of fine particles), $\text{PM}_{2.5-1}$ (micron fraction of fine particles), $\text{PM}_{10-2.5}$ (coarse particles), and PM_{20-10} supercoarse particles²⁹). “Deposited dose” is the aerosol mass of all measured particle sizes deposited in the human respiratory system during an exposure event. The deposited dose was calculated for the entire respiratory system and individually for each region of the respiratory system: the head airways (HA), the tracheobronchial region (TB) and the alveolar region (AL).

The detailed description of the employed mathematical model and its development was published earlier.²³ We are also providing the equations and definitions of this model in the ESI (Supplementary Methods).[†] The same user profile reported by Nazarenko *et al.*²³ was used for the consumer sprays: inhalation flow rate (Q_{inh}) = 11.0 L min⁻¹ and body weight (bw) = 60 kg, which correspond to the body weight and breathing rate suggested for assessing short-term exposures of 18–60 year old females performing light activities.³⁰ The justification of this choice of user and activity profile was discussed earlier.²³

The duration of each exposure event was assumed to be $T_{\text{contact}} = 1$ min; however, a direct adjustment of this duration may be done to calculate the inhalation exposure of different durations as required by any other exposure scenario.

Similar to the case with cosmetic powders in our previous study,²³ here we also could not obtain information about the amount or fraction of nanomaterials in the investigated consumer sprays in order to determine f_{nano} . As in the cosmetic powder study, we assumed $f_{\text{nano}} = 1$ indicating that aerosolized particles produced from the nanotechnology-based products consist completely (100%) of nanomaterials (the worst case scenario). However, the dose calculations can be easily adjusted should nanomaterial content become known. We assumed aerosol particle losses in the sampling lines as negligible.

The deposited dose was determined as aerosol mass deposited in each of the three regions of the respiratory system and in the entire respiratory system during a 1-minute exposure event per 1 kg of body weight in the same way as in the previous study for cosmetic powders.²³

As in the previous study,²³ we also determined the deposited dose for the HA, TB and AL regions of the respiratory system as percentage of the total deposited dose. This presentation allowed us to illustrate the respiratory system region with the highest deposited dose relative to the other regions.

Results

TEM characterization of consumer sprays

Fig. 1 and 2 demonstrate two selected TEM micrographs for each of the consumer sprays, for which the TEM investigation showed presence of particles or electron-contrast structures. These include two nanotechnology-based silver nanospray and disinfectant nanospray and five non-nanotechnology-based regular silver spray, regular disinfectant spray, regular hair spray, regular skin hydrating mist, and regular facial spray. In three nanotechnology-based products (facial nanospray, hair nanospray and skin hydrating nanomist), no such particles or structures were observed.

The phenomenon of radiolysis (alteration of a material by the energy of the electron beam) above a certain magnification was observed for two regular products: regular skin hydrating mist and regular facial spray. This phenomenon has been described earlier.^{16,31} It can be seen that the particles with clear boundaries as seen in Fig. 2h and j at low magnifications,

[†]Electronic supplementary information (ESI) available. See DOI: 10.1039/c3en00053b

when the diameter of the electron beam is larger, “melted” when viewed at higher magnifications (Fig. 2g and i), when we focused the electron beam in a smaller area of the sample. This is likely an indication of the organic chemical nature of the particles in these two products.

In the silver nanospray (Fig. 1a and b) and the regular silver spray (Fig. 2a and b), we found silver nanoparticles with sizes from ~3 nm to several dozen nanometers and larger. Most particles were agglomerated, especially in the silver nanospray. The silver nanospray contained on average smaller particles than its regular counterpart. In the regular silver spray, we also observed particles and agglomerates larger than 100 nm (up to almost 0.5 μm).

The disinfectant nanospray (Fig. 1c and d) presented only a few low-contrast particles with a very low level of agglomeration. The smallest particles were in the nanosize range (about 70 nm), while the largest particles were slightly larger than 200 nm. The low electron contrast of these particles may be an indication of their organic nature or that they are composed of lighter chemical elements.

The regular disinfectant spray presented a very interesting particle structure as shown in Fig. 2c and d: spherical particles, most of which were below 100 nm in size and looked either nanostructured or as agglomerates of very small nanoparticles (1–2 nm). The electron contrast of these 1–2 nm nanoparticles or nanostructure elements varied from low to high. There is also an additional level of agglomeration of the larger nanostructured or agglomerated particles. They were observed attached to each other in pairs and up to several particles.

The regular hair spray (Fig. 2e and f) had two different kinds of particles, some of which were nanosized (as small as ~16 nm). The particles of the first kind were small spherical and of varying sizes (~16 to above 100 nm). The size of agglomerates reached almost 700 nm. There were also areas containing loosely associated agglomerates extending to 2–3 μm in size.

The TEM micrographs of the regular skin hydrating mist (Fig. 2g and h) and regular facial spray (Fig. 2i and j) looked very similar. However, the smallest visible particles in the regular facial spray were in the nanosize range (as small as ~80 nm), while the smallest particles in the regular skin hydrating mist were ~150 nm. The largest particles in these two sprays were very large compared with the other investigated sprays: >2.5 μm in the regular skin hydrating mist and >6 μm in the regular facial spray. As described above, the particles in these two products were altered due to radiolysis at higher magnifications of the TEM.

Quantitative exposure assessment

Inhalation exposure—Fig. 3 shows the inhalation exposure resulting from a 1-minute use of each of the investigated consumer sprays, expressed as aerosol mass in different particle size fractions per 1 kg of body weight.

It can be seen that the inhalation exposure of the ultrafine aerosol fraction ($PM_{0.1-0.014}$) was in a wide range between ~ 0.002 (disinfectant nanospray) and ~ 0.05 (disinfectant spray) $ng\ kg^{-1}$ bw per application. Disinfectant nanospray, facial spray, silver nanospray, skin hydrating mist, silver spray, and facial nanospray produced the lowest inhalation exposures ($\sim 0.002-0.007\ ng\ kg^{-1}$ bw per application). Hair spray, hair nanospray, skin hydrating nanomist, and disinfectant spray produced inhalation exposures that were approximately an order of magnitude higher ($\sim 0.01-0.05\ ng\ kg^{-1}$ bw per application).

In the submicron fraction of fine particles ($PM_{1-0.1}$), a relatively wide range of inhalation exposure was observed for different sprays: from ~ 0.3 (disinfectant nanospray) to ~ 31 (disinfectant spray) $ng\ kg^{-1}$ bw per application. The use of four products (disinfectant nanospray, silver spray, silver nanospray, and skin hydrating mist) resulted in a very low $PM_{1-0.1}$ below $1\ ng\ kg^{-1}$ bw per application. Three products (facial spray, facial nanospray and skin hydrating nanomist) produced inhalation doses of $PM_{1-0.1}$ around $\sim 4-5\ ng\ kg^{-1}$ bw per application. Three products (hair nanospray, hair spray and disinfectant spray) produced relatively very high $PM_{1-0.1}$ inhalation exposures of $\sim 18-31\ ng\ kg^{-1}$ bw per application.

The inhalation exposure of the micron fraction of fine particles ($PM_{2.5-1}$) varied greatly as well: from ~ 0.06 (silver spray) to ~ 340 (disinfectant spray) $ng\ kg^{-1}$ bw per application. Silver spray and silver nanospray produced $PM_{2.5-1}$ inhalation exposure below $1\ ng\ kg^{-1}$ bw per application: ~ 0.06 and ~ 0.3 , respectively. Higher $PM_{2.5-1}$ inhalation exposures ranging between 1 and $100\ ng\ kg^{-1}$ bw per application were produced by disinfectant nanospray (~ 4), skin hydrating mist (~ 20), skin hydrating nanomist (~ 53), and facial nanospray (~ 78). The highest $PM_{2.5-1}$ inhalation exposures exceeding $100\ ng\ kg^{-1}$ bw per application were produced by facial spray (~ 103), hair nanospray (~ 126), hair spray (~ 205), and disinfectant spray (~ 340).

No coarse particles (the $PM_{10-2.5}$ fraction) and, hence, no corresponding inhalation exposures to coarse particles were determined in two out of ten products: silver nanospray and silver spray. The $PM_{10-2.5}$ inhalation exposure was the lowest for disinfectant nanospray ($\sim 13\ ng\ kg^{-1}$ bw per application) and highest for hair spray ($1200\ ng\ kg^{-1}$ bw per application). For the other consumer sprays, the $PM_{10-2.5}$ inhalation exposure ranged between approximately 200 and $700\ ng\ kg^{-1}$ bw per application.

Particles above $10\ \mu m$ and, accordingly, inhalation exposure of PM_{20-10} (supercoarse particles)²⁹ were detected in only three products: $\sim 1\ ng\ kg^{-1}$ bw per application for hair nanospray, $\sim 15\ ng\ kg^{-1}$ bw per application for skin hydrating mist and $\sim 26\ ng\ kg^{-1}$ bw per application for hair spray.

Based on the two-tailed Student's *t* test assuming equal variance, the inhalation exposures were statistically different: (1) in all aerosol particle size fractions for the nano and regular hair sprays and for the nano and regular disinfectant sprays, (2) in all but the $PM_{10-2.5}$ size fraction for the nano and regular skin hydrating mists, and (3) in the $PM_{10-2.5}$ fraction only for the nano and regular facial sprays.

Deposited dose—Fig. 4 shows the deposited dose in the head airways (HA), the tracheobronchial region (TB), and the alveolar region (AL), as well as the total respiratory system resulting from a 1-minute use of every investigated consumer spray, expressed as aerosol mass in different particle size fractions per 1 kg of body weight. Proportional distribution of the deposited doses in different regions of the respiratory system is shown in Fig. 5. Eight sprays (facial nanospray, hair nanospray, disinfectant nanospray, skin hydrating nanomist, facial spray, hair spray, disinfectant spray, and skin hydrating mist) had very similar looking deposition profiles (Fig. 4). Their lowest and highest deposited doses for HA were $\sim 13 \text{ ng kg}^{-1} \text{ bw}$ per application (disinfectant nanospray) and $\sim 1171 \text{ ng kg}^{-1} \text{ bw}$ per application (hair spray), while the HA deposited doses from the remaining six products were in the range ~ 205 to $\sim 785 \text{ ng kg}^{-1} \text{ bw}$ per application. The TB deposited doses for those eight sprays were between ~ 1 and $\sim 63 \text{ ng kg}^{-1} \text{ bw}$ per application, while those for AL were between ~ 1.4 and $\sim 101 \text{ ng kg}^{-1} \text{ bw}$ per application. The total deposited dose spanned a wide range: from the lowest of $\sim 16 \text{ ng kg}^{-1} \text{ bw}$ per application (disinfectant nanospray) to the highest of $\sim 1335 \text{ ng kg}^{-1} \text{ bw}$ per application (hair spray). The deposited doses from the other six products were in a range between ~ 232 and $\sim 920 \text{ ng kg}^{-1} \text{ bw}$ per application. As can be seen in Fig. 5, these eight sprays have similar proportional distributions of deposited doses in different regions of the respiratory system: ~ 85 – 88% of the total respiratory system deposition occurred in the head airways, ~ 4.6 – 5.2% in the tracheobronchial region, and ~ 7.0 – 9.5% in the alveolar region. Compared to these eight products, silver nanospray and silver spray looked rather differently with substantially lower deposited doses, respectively: in HA – ~ 0.17 and $\sim 0.06 \text{ ng kg}^{-1} \text{ bw}$ per application, in TB – ~ 0.02 and $\sim 0.01 \text{ ng kg}^{-1} \text{ bw}$ per application, and in AL – ~ 0.08 and $\sim 0.05 \text{ ng kg}^{-1} \text{ bw}$ per application. The total deposited dose for silver nanospray and silver spray was ~ 0.3 and $\sim 0.1 \text{ ng kg}^{-1} \text{ bw}$ per application, respectively (Fig. 4) – lower than for the other products. The proportional deposition in different areas in the respiratory system was also different from the other eight products: $\sim 29\%$ (silver nanospray) and $\sim 40\%$ (silver spray) of the aerosol mass deposited in the alveolar region, $\sim 7\%$ for both products in the tracheobronchial region, and $\sim 64\%$ and $\sim 52\%$ deposited in the head airways (Fig. 5).

Discussion

This study found that the release of aerosol particles in various size fractions from different consumer spray products varied greatly from product to product. This is in contrast to the results of our previous study focusing on cosmetic powders, where relatively similar proportions of the concentrations in different particle size fractions were observed for various powders.²³ For easier comparison, the ranges of inhalation exposure and deposited doses for the sprays and the powders are summarized in Table 2. The high variability of aerosol size distributions among consumer spray products suggests substantially different exposure levels to different particle size fractions depending on the product.

In addition, for the consumer sprays, inhalation exposure of the $\text{PM}_{10-2.5}$ fraction was dominant in seven out of ten products (facial nanospray, hair nanospray, skin hydrating nanomist, facial spray, hair spray, disinfectant spray, and skin hydrating mist). For the remaining three products, the $\text{PM}_{10-2.5}$ inhalation exposure was either below the detection limit (silver nanospray and silver spray) or close to the detection limit (disinfectant

nanospray). For powders investigated in that earlier study,²³ inhalation exposure of PM_{10-2.5} was also the highest among the considered size fractions, but PM₂₀₋₁₀ reached levels similar to PM_{10-2.5} in three out of seven powders. In addition, PM_{10-2.5} inhalation exposures for all powders were above the detection limit.

Another notable difference between the consumer sprays and the cosmetic powders is the release of particles in the nanosized aerosol fraction (PM_{0.1-0.014}). This fraction was released above the detection limit levels from *all* tested consumer spray products. In contrast, only two out of seven cosmetic powders produced PM_{0.1-0.014} inhalation exposure that was detected.²³ Additionally, for those products where the release of the nanosized fraction was detected, a substantially higher concentration for consumer sprays was observed compared with cosmetic powders.

Another revealing finding is that those consumer sprays that showed high abundance of nanosized particles in the TEM micrographs (silver nanospray, disinfectant nanospray and silver spray) produced the lowest inhalation total and nanosized aerosol exposures. At the same time, the two other sprays with TEM micrographs showing noticeable presence of nanostructures or nanoparticles (disinfectant spray and hair spray) produced high PM_{0.1-0.014} inhalation exposures. In the case of these two products, however, the TEM micrographs showed high amounts of residue, in which nanosized particles were embedded. This residue was possibly formed by organic compounds present in the products and was visible as electron-contrast plumes of undefined shape. The presence of dissolved substances that likely formed this residue may have led to generation of additional particles during product application and may have facilitated easier dispersion of primary particles. Overall, the presence of nanosized particles and structures in the original liquid products as detected by TEM did not seem to correlate with the inhalation exposure of the nanosized *aerosol* fraction or the larger aerosol fractions. Some spray products that showed a high number of nanosized objects in the TEM micrographs actually produced the lowest total inhalation aerosol exposures, while some produced high inhalation exposures in the nanosized fraction. Since the presence of nanosized objects in products does not seem to correlate with the concentration of airborne nanoparticles, it suggests that simply measuring the size distributions of aerosols created during the use of nanotechnology-based consumer products is not sufficient to accurately predict or assess nanomaterial exposure. A more accurate approach would be to determine the exact and relative quantities of nanomaterial(s) in a given product and then determine the masses of inhaled and deposited aerosol to determine nanomaterial inhalation exposure. At the same time, the mass fractions of nanomaterial(s) in the original product and the inhaled or deposited aerosol may differ due to 1) non-homogeneity of the product, 2) non-uniform distribution of nanomaterial(s) across different size fractions of the produced aerosol and 3) aerosol dynamics after aerosolization, particularly when a spray is not used immediately within the personal breathing zone.

As suggested earlier,^{16,17,23} a substantial fraction of nano-materials is likely found in larger aerosol size fractions – albeit in an agglomerated form. The observation of a large number of nano-objects using TEM in those sprays that produced very low inhalation exposure of nanosized aerosol particles seems to support this supposition.

Another notable phenomenon observed for the consumer sprays is the absence of measurable inhalation exposure of the largest supercoarse particles (PM₂₀₋₁₀) in all but three consumer sprays: hair nanospray, skin hydrating nanomist and hair spray. Moreover, for these three products, the measured inhalation exposures of PM₂₀₋₁₀ were substantially lower than those of the adjacent size fraction, *e.g.*, PM_{10-2.5}. In the case of the cosmetic powders investigated earlier,²³ the simulated use of *all* of the products resulted in relatively high inhalation exposures of PM₂₀₋₁₀. One possible explanation for this phenomenon could be the difference in the mode of application of consumer sprays vs. cosmetic powders. Whereas the cosmetic powders are applied directly using a brush onto the face including the immediate vicinity of the nostrils, the consumer sprays are atomized at a distance from the nose, which leaves more time for the largest particles to settle before they could be inhaled. In addition, and probably more important, liquid atomization is a more energetic dispersion technique compared to powder application by a brush or a pad and disaggregates the material more effectively.

Looking at the deposited doses (Fig. 4 and 5), one can see that between ~52% and ~88% of all particles by mass deposited in the head airways (HA). This range indicates comparatively lower average HA deposition for the consumer sprays than for the cosmetic powders, for which the HA deposition was ~85–93%. However, among all of the investigated consumer sprays, two products with the lowest total inhalation exposure dose (silver spray and silver nanospray) also produced the lowest HA deposition fractions: 52% (regular silver) and ~64% (nano silver). For the other eight consumer sprays, the deposited dose fraction for the HA was ~85–88%. As a result of relatively low HA deposition fractions, for both the regular and nano silver sprays, high alveolar (AL) deposition fractions were computed: ~29% (nano silver) and ~40% (regular silver). In contrast, the highest AL deposited dose fraction for any of the previously investigated cosmetic powders did not exceed 10%. Hence, in the case of these two consumer sprays, both of which were observed to contain nanomaterials, deposition in the alveolar region may be more important compared to depositions in the other regions of the respiratory tract. At the same time, the total inhalation aerosol exposures for these two products were 3–5 orders of magnitude lower compared to those for other sprays, and thus one cannot conclude that the exposure of the alveolar region to particulate matter, potentially containing engineered nanomaterials, would be higher than from other sprays.

Similar to the cosmetic powders,²³ the deposited dose for the tracheobronchial region was the lowest among all regions of the respiratory system for all consumer sprays, specifically ~1.5–5.6 times lower than for the alveolar region, a greater difference than for cosmetic powders (~1.5–2 times). Again, silver nanospray and silver spray had the highest difference between the TB and the AL deposited doses: a factor of ~4 and ~5.6. The factor for the other eight sprays was below 1.8 – not different from that for the cosmetic powders.

Our choice of mass as the metric for aerosol exposure when using consumer products was presented in the earlier study.²³ Briefly, the number-based or surface area-based metrics would not allow adequate representation of the nanomaterial content in the total aerosols where most nano-objects exist in the form of agglomerates.²³ Since all consumer sprays investigated here were in non-pressurized containers and were sprayed using a pump-based

mechanism, the deagglomeration processes are not expected to have been extensive, contrary to what has been shown for the propellant-based spray products.^{21,32,33}

The effect of particle agglomeration leads to the potential nanomaterial inhalation exposure through aerosol particle size fractions larger than 100 nm, and for some spray products, up to supercoarse particles. In addition, major differences between consumer sprays and cosmetic powders include evaporation of solvents from liquid particles after spraying and a longer residence time of aerosol particles between aerosolization and inhalation. These two phenomena provide for a possibility of a more complicated aerosol dynamics. This size distribution may greatly depend on the spray composition, e.g., water and organic solvent content, the way a product is used and the environmental conditions including humidity and temperature. The presence of solvents with molecularly dissolved substances can cause particle formation due to crystallization/solidification of these chemicals. Additionally, the dissolved chemicals can precipitate on the surface and within caverns in other particles and agglomerates influencing their size.³⁴⁻³⁸ After deposition in the respiratory system, certain chemicals can dissolve away from the deposited particles thereby altering their size and state of agglomeration,³⁹⁻⁴⁴ which can greatly influence nanomaterial fate in the live tissue and the resulting potential biological effects.

Compared to pure nanomaterials, the multi-ingredient nature of most nanotechnology-based consumer products (Table 1) is likely to lead to altered nanomaterial properties, including biological effects as well as different aerosol properties. These differences of nanotechnology-based products from pure nanomaterials mean that to accurately estimate potential exposure and health effects, investigation of nanotechnology-based products themselves is necessary. Investigation of pure nanomaterials that are added to nanotechnology-based consumer products should also be performed. Among the challenges warranting analysis of the actual products in addition to pure nanomaterial analysis are: (1) the difficulties in determining the presence and concentrations of engineered nanomaterials in the products, (2) the differences in aerosol production, the resulting size distributions, and its subsequent dynamics for the multi-ingredient consumer products, and (3) the effect of product matrix on surface chemistry and fate of nanomaterials deposited in the respiratory system, such as penetration of engineered nano-objects into the live tissue and the effects there, as well as their possible translocation in the body.

As with certain cosmetic powders,²³ we observed nanosized particles or nanostructures in the TEM micrographs of some consumer sprays, not identified by their manufacturers as nanomaterial-containing, particularly regular silver spray and regular disinfectant spray. This again supports our argument made earlier^{16,17,23} that product identification and labeling as a basis for determining a consumer product's nanotechnology-based status may not adequately represent the actual nature of any given consumer product with respect to its "engineered nano status". As our own experience of trying to analyze consumer products with respect to their content of engineered nanomaterials has shown, it may be difficult or impossible to determine the engineered nano status of any given product using current analytical techniques. For example, we could not obtain clear TEM micrographs of three out of five nanotechnology-based consumer sprays (facial nanospray, hair nanospray and skin hydrating nanomist). This was mostly due to two factors: (1) the presence of multiple

ingredients in the products that obscured particulate matter and/or underwent radiolysis with volatilization under the electron beam, which may cause microscope contamination and (2) low electron contrast of particles. Due to these issues, we emphasize again the importance of accurately reporting the content of engineered nanomaterials in consumer products, so that accurate exposure assessment and health risk analysis could be performed.

Conclusions

For the consumer sprays, we observed a greater variability in the levels of inhalation exposure and deposited doses of aerosols compared with cosmetic powders, which were investigated in an earlier study. We conclude that aerosol exposure would be markedly different depending on the spray product used, which was not the case with certain cosmetic powders explored earlier.

We also found that consumers would receive a measurable inhalation exposure in the nanosized aerosol fraction ($PM_{0.1-0.014}$) from *all* consumer sprays. This is in contrast to the cosmetic powders, only two of which released detectable nanosized aerosol particles.²³ This indicates that exposure to airborne nanosized particulate matter would occur from all investigated products, even from those that are not labeled as containing engineered nanomaterials. This is a very important finding showing that the release of aerosol particles <100 nm alone cannot serve as an indication of engineered nanomaterial exposure.

The inhalation exposure by mass was highest in the $PM_{10-2.5}$ aerosol size fraction for the majority of the consumer sprays. At the same time, it was minimal or below the detection limit for three sprays. This high $PM_{10-2.5}$ variability presented by consumer sprays was a major difference from the cosmetic powders investigated previously. If engineered nanomaterials are present in a product, they are likely distributed in *all* aerosol size fractions. Thus, for those products where particles in the PM_{10-2} size range were observed, most nanomaterials would likely be inhaled with the $PM_{10-2.5}$ aerosol fraction.

We found the head airways to be the primary site of aerosol deposition with ~52–88% of all aerosol particle masses. Hence, as with the cosmetic powders, toxicological research should also focus on head airways along with the other regions of the respiratory system currently receiving more attention.

Accurate engineered nanomaterial exposure assessment can only be conducted when the quantitative content of nanomaterial(s) in the original nanotechnology-based products is known. A mandate requiring that such information from the manufacturers of nanotechnology-based consumer products be provided should be considered as a possible solution to the challenge of quantitative exposure assessment.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgements

This research was supported in part by the National Institute of Environmental Health Sciences (NIEHS) sponsored by the University of Medicine and Dentistry of New Jersey (UMDNJ) Center for Environmental Exposures and Disease (grant no. P30ES005022), the NSF (grant NSF-CBET-1236508), and the New Jersey Agriculture and Experiment Station (NJAES) at Rutgers University. The views expressed in this paper are solely those of the authors and do not necessarily reflect the views of the funding agencies.

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Nano impact

This study provides a quantitative assessment of human inhalation exposure to nanomaterials due to the use of nanotechnology-based consumer sprays. To the best of our knowledge, it is the first quantitative exposure assessment of a wide selection of consumer spray products in a realistic exposure scenario. We expect this study to generate a substantial impact due to high public interest and attention of the governmental and non-governmental agencies to the safety of nanotechnology-based consumer products. This study is published just as the Regulation of the European Parliament and of the Council on Cosmetic Products, PE-CONS 3623/09 (2009) came into legal effect in 2013 mandating reporting of “the reasonably foreseeable exposure conditions” for all of the nanomaterial-containing cosmetic products on the EU market.

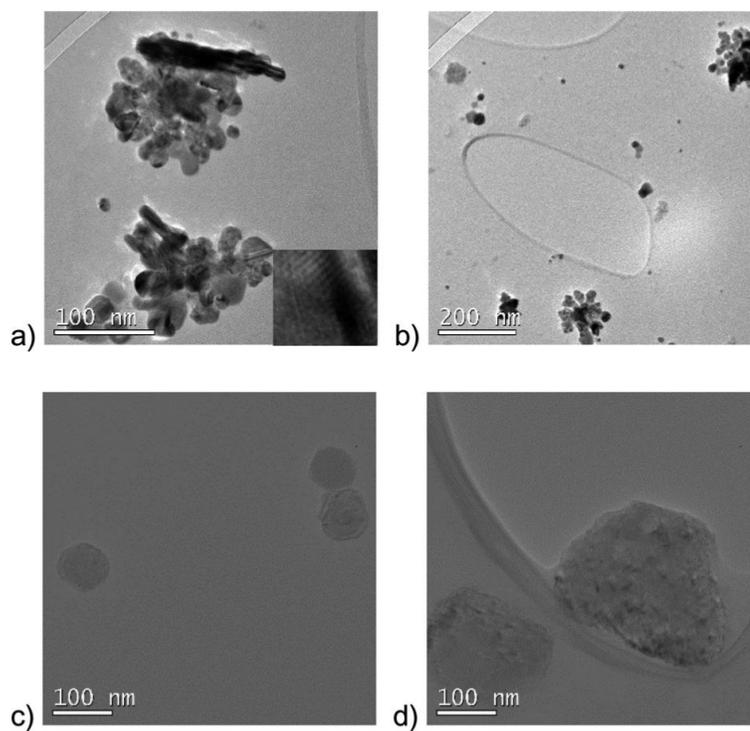


Fig. 1. TEM micrographs of silver nanospray (a, b) and disinfectant nanospray (c, d).

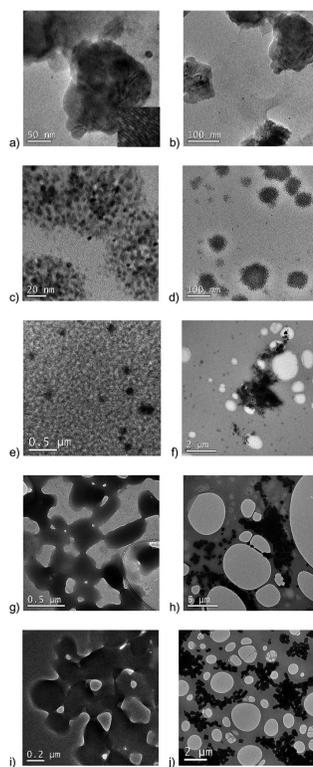


Fig. 2. TEM micrographs of regular silver spray (a, b), regular disinfectant spray (c, d), regular hair spray (e, f), regular skin hydrating mist (g, h), and regular facial spray (i, j).

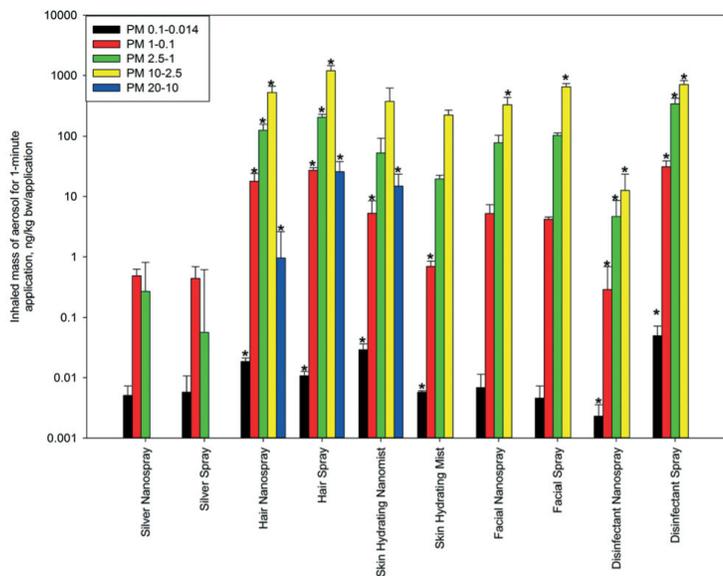


Fig. 3. Inhalation exposure of PM from consumer sprays based on mass concentration of particulate matter in different aerosol particle size fractions sampled with the mannequin head sampler during simulated product application. The data represent averages of three repeats. Error bars represent one standard deviation. * denotes a significant difference ($p < 0.05$) for a given particle size fraction between a nano and a regular product, based on the two-tailed Student's t test. Numerical data used to produce the figure are provided in the ESI, [†] Table S1.

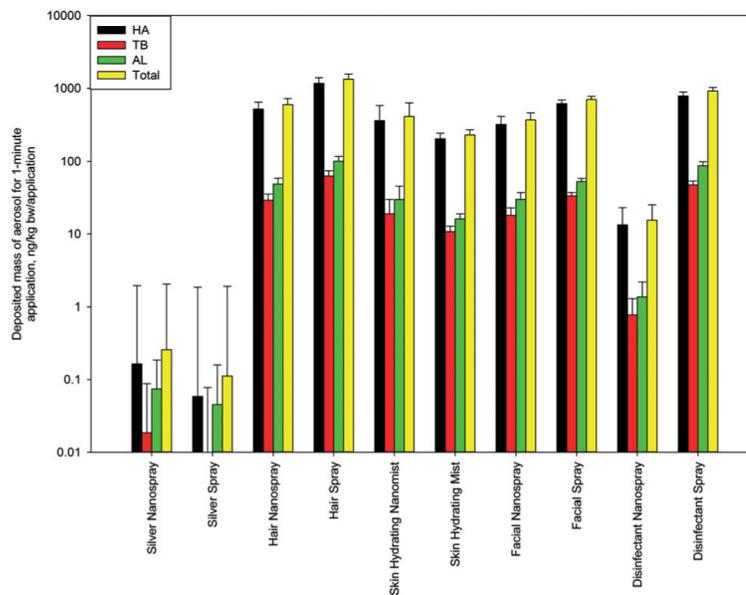


Fig. 4. Deposited dose of PM from consumer sprays deposited in different regions of the respiratory system (the head airways (HA), the tracheobronchial (TB) and the alveolar (AL) regions). The data represent averages of three repeats. Error bars represent one standard deviation and illustrate uncertainty of model results propagating from known uncertainty of experimental data. Numerical data used to produce the figure are provided in the ESI,[†] Table S2.

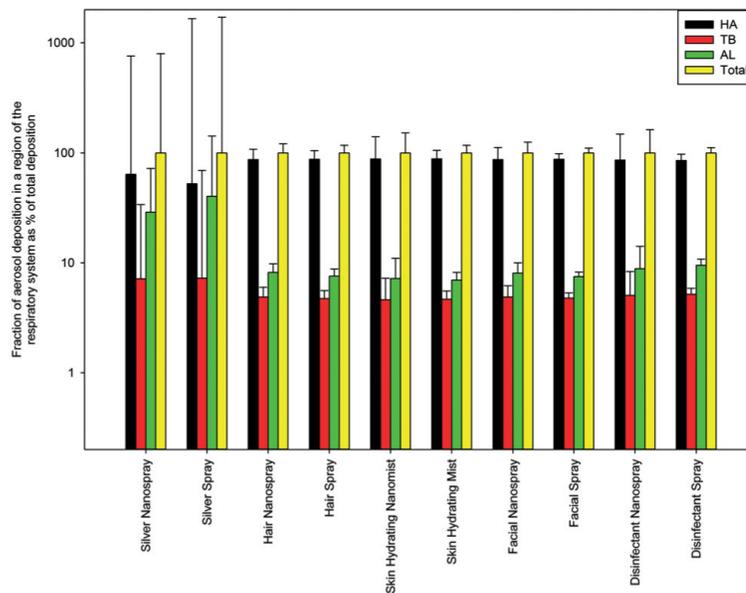


Fig. 5. Percent distribution of PM from consumer sprays deposited in different regions of the respiratory system (the head airways (HA), the tracheobronchial (TB) and the alveolar (AL) regions). The data represent averages of three repeats. Error bars represent one standard deviation and illustrate uncertainty of model results propagating from known uncertainty of experimental data. Numerical data used to produce the figure are provided in the ESI,[†] Table S3.

Table 1

Investigated consumer sprays

Product	Composition ^a
Silver nanospray	Silver nanoparticles, purified water
Regular silver spray	99.99% Pure silver suspended in demineralized water
Facial nanospray	Distilled water, vitamin C, nanosize particles of copper, calcium, magnesium and zinc
Regular facial spray	Water, butylene glycol, glycerin, panthenol, tocopheryl acetate, phenoxyethanol, alcohol denat., methylparaben, lecithin, <i>Rosa centifolia</i> (rose) water, butylparaben, ethylparaben, isobutylparaben, propylparaben
Hair nanospray	Alcohol denat., aqua, PVP/VA co-polymer, isopropyl alcohol, myrtrimonium bromide, parfum
Regular hair spray	SDA alcohol 40-B, water, VA/crotonates/vinyl neodecanoate co-polymer, octylacrylamide/acrylates/butylaminoethyl methacrylate co-polymer, aminomethanol propanol, lauryl pyrrolidone, PEG-75 lanolin, cyclopentasiloxane, fragrance
Disinfectant nanospray	Parachlorometaxylenol – 0.20%, other ingredients – 99.80%
Regular disinfectant spray	<i>o</i> -Phenylphenol – 0.22%, diisobutylphenoxyethoxy ethyl dimethyl benzyl ammonium chloride monohydrate – 0.70%, inert ingredients – 99.08%
Skin hydrating nanomist	Purified water, dimethicone, copolyol, algae extract, mugwort (<i>Artemisia vulgaris</i>) extract, <i>Aloe barbadensis</i> gel, fucogel, plankton extract, lavender (<i>Lavendula angustifolia</i>) oil, calcium PCA, zinc PCA, phenoxyethanol, methylparaben, propylparaben
Regular skin hydrating mist	Water, glycerin, hyaluronic acid, diazolidinyl urea, polysorbate 80, ergothioneine, <i>Aloe barbadensis</i> leaf juice, sodium carboxymethyl <i>b</i> -glucan, <i>Camellia sinensis</i> leaf extract, tetrasodium EDTA, allantoin, citrus <i>Aurantium bergamia</i> (bergamot) fruit oil, citric acid, kinetin, iodopropynyl butylcarbamate

^aReproduced from the product labels exactly.

Table 2

Comparative summary of exposure data for consumer sprays vs. powders

Property	Sprays (ng kg ⁻¹ bw per application)	Powders ^a (ng kg ⁻¹ bw per application)
Range of total inhalation exposure	5.0×10^{-1} –1452.6	38.7–33 317.7
Range of inhalation exposure to PM _{0.1–0.014}	2.3×10^{-3} – 5.0×10^{-2}	0– 5.7×10^{-3}
Range of inhalation exposure to PM _{1–0.1}	2.9×10^{-1} –31.2	7.2×10^{-2} –358.9
Range of inhalation exposure to PM _{2.5–1}	5.7×10^{-2} –338.8	5.4–1011.9
Range of inhalation exposure to PM _{10–2.5}	0–1194.2	18.7–29 874.3
Range of inhalation exposure to PM _{20–10}	0–25.8	14.5–2072.6
Range of total deposited dose	1.1×10^{-1} –1334.5	36.5–32 043.4
Range of HA deposited dose	5.9×10^{-2} –1171.1	33.3–28 650.9
Range of TB deposited dose	8.1×10^{-3} –62.7	1.2–1385.3
Range of AL deposited dose	4.5×10^{-2} –100.7	2.0–2007.2
Range of percent of HA deposited dose relative to total deposited dose	52.4–88.4%	84.5–93.1%
Range of percent of TB deposited dose relative to total deposited dose	4.6–7.3%	2.9–5.2%
Range of percent of AL deposited dose relative to total deposited dose	7.0–40.4%	4.1–10.3%

^aNazarenko *et al.*²³