Molecular Mechanisms of Toxicity of Silver Nanoparticles in Zebrafish Embryos

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ABSTRACT: Silver nanoparticles cause toxicity in exposed organisms and are an environmental health concern. The mechanisms of silver nanoparticle toxicity, however, remain unclear. We examined the effects of exposure to silver in nano-, bulk-, and ionic forms on zebrafish embryos (Danio rerio) using a Next Generation Sequencing approach in an Illumina platform (High-Throughput SuperSAGE). Significant alterations in gene expression were found for all treatments and many of the gene pathways affected, most notably those associated with oxidative phosphorylation and protein synthesis, overlapped strongly between the three treatments indicating similar mechanisms of toxicity for the three forms of silver studied. Changes in oxidative phosphorylation indicated a down-regulation of this pathway at 24 h of exposure, but with a recovery at 48 h. This finding was consistent with a dose-dependent decrease in oxygen consumption at 24 h, but not at 48 h, following exposure to silver ions. Overall, our data provide support for the hypothesis that the toxicity caused by silver nanoparticles is principally associated with bioavailable silver ions in exposed zebrafish embryos. These findings are important in the evaluation of the risk that silver particles may pose to exposed vertebrate organisms.

INTRODUCTION

Nanoparticles are being introduced, in some cases rapidly, into the consumer market but there is a fundamental lack of knowledge on their potential consequences for human and environmental health (reviewed in refs 1 and 2). The use of silver nanoparticles (AgNP) has increased very significantly in recent years, with a current total global production estimated at 500 tonnes per year (reviewed in ref 1). Very little empirical data are available on environmental levels of AgNP but given recent years, with a current total global production estimated at 500 tonnes per year (reviewed in ref 1). Very little empirical data are available on environmental levels of AgNP but given very high concentrations and the particles had been stabilized with starch or BSA to prevent agglomeration. The conditions adopted were likely to enhance uptake in those laboratory exposures, but they may not reflect environmental reality. In juvenile or adult fish, one of the main routes of uptake and toxicity for AgNP appears to be via the gills, and adverse effects include on osmoregulation, similar to that reported for ionic silver.8,9 For AgNP, the effect on Na+/K+-ATPase activity in gills has been attributed to the dissolution of silver ions from the particles, and/or to the presence of silver ions at the surface.
of the AgNP. The mechanisms by which this specific effect of exposure to AgNP occurs are unclear and may be due to damage of the gill surface caused by the presence of particulate matter and/or local dissociation of Ag⁺ from the AgNP trapped in the mucus surrounding the gill membrane. The biological processes affected by exposure to AgNP, reported across a variety of organs, include induction of oxidative damage, alterations to the regulation of enzymes responsible for free radical scavenging, altered regulation of gene expression pathways involved in apoptosis, and disrupted regulation of the cellular machinery involved in storing, detoxification and metabolism of metals (such as metallothionein 2).12–16

The objective of this study was to investigate the molecular mechanisms of toxicity of AgNP (10 nm) in comparison with a micrometer sized counterpart (Ag Bulk, 0.6–1.6 μm) and with Ag⁺, using a recently developed High-throughput (HT-)SuperSAGE approach on an Illumina GA II platform and zebrafish (embryos) as a model organism. Our data demonstrate that most transcriptomic alterations caused by exposure to AgNP and its ionic and bulk counterparts were common across all treatments, supporting the hypothesis that the toxicity of AgNP is principally associated with the toxicity of Ag⁺ at the surface of the particles or dissolved in the water. We also identified some gene changes unique to each treatment, suggesting that there may also be some particle-specific effects.

MATERIALS AND METHODS

Characterization of the Silver Particles and Their Dissolution Rates. Silver particles (designated as AgNP [10 nm] and Ag Bulk [0.6–1.6 μm]) were purchased from Nanostructured and Amorphous Materials Inc., Houston, TX and silver with 2% HNO₃ was obtained from PerkinElmer Life and Analytical Sciences, Shelton, CT. Further details on the characterization of these particles and their dissolution in exposure water, and the biological effects of exposure to AgNP can be found in ref 13 and in the Supporting Information (SI).

Fish Source, Culture, and Husbandry. Adult zebrafish (wild-type WIK strain, originally from the Max Planck Institute, Tubingen, Germany) for the provision of embryos were kept in aquaria at the University of Exeter in 140 L mixed-sex stock tanks, as described in ref 18. Fish were allowed to breed naturally and eggs were collected in glass egg chambers, approximately 1 h postfertilization (hpf). Eggs were then cleaned and unfertilized embryos were removed prior to the exposures.

Embryo Exposures to AgNP, Ag Bulk and Ag⁺ for Global Gene Expression Analysis. Stock solutions for AgNP, Ag Bulk and silver nitrate were made up in ultrapure water, and sonicated for 1 h to ensure dispersal of the particles. Exposures were conducted in glass chambers, at 28 ± 1 °C with a 12 h light:dark photoperiod. Immediately prior to the start of the exposures, glass chambers received 400 mL artificial water, prepared according to the ISO 7346:3:1996 guideline, containing 5 μg/L of 10 nm AgNP, 5 μg/L of Ag Bulk or 0.25 μg/L of silver nitrate. A control chamber was set up containing water alone. These exposure concentrations were well below those causing overt toxicity, based on the mortality curves for the various forms of silver obtained under the same experimental conditions in our laboratory (see SI Figure S1). The dosing of Ag Bulk was chosen to be the same as that for the AgNP (in terms of mass), to allow us to test whether the size of the particles was a determinant of the effects seen. An exposure to silver nitrate at 0.25 μg/L was included to provide a likely worst case scenario for the highest dissolution of Ag⁺ for the AgNP and Ag Bulk exposures in the adopted exposure conditions, based on data from the literature suggesting that dissolution of AgNP is variable but below 5% (Osborne et al 2012 reported this to vary between 0.1 and 2% in zebrafish embryo culture water). Our own experiments (see SI) suggest that dissolution rates for AgNP and Ag Bulk under our experimental conditions were 0.27% ± 0.15% and 0.27% ± 0.07%, respectively. Exposures were initiated at 4 hpf and 585 ± 99 eggs were deployed into each treatment chamber. Unfertilized eggs were identified by visual inspection using light microscopy (Kyowa Optical SDZ PL, Kyowa Optical, Japan) and removed from the exposure chamber (22.8% of the total number of eggs were unfertilized). Solutions were replaced every 12 h during the exposure period, and dead embryos were removed at the times of replacement of the exposure water. At 24 h and 48 h, three pools of 50 embryos were removed from each of the exposure tanks, immediately frozen in liquid nitrogen and stored at −80 °C for analysis of gene expression. The experiment was terminated at 48 hpf.

Embryo Exposures for Investigations into the Effects of Ag⁺ on Oxygen Consumption in Zebrafish Embryos. In order to verify the biological significance of the effect of all treatments on the oxidative phosphorylation pathway, we designed a study to evaluate the effects of exposure to a series of doses of Ag⁺ on embryo oxygen consumption. Zebrafish embryos were exposed to a range of concentrations of Ag⁺ to include those expected to be present in the water following exposure to AgNP, Ag Bulk and Ag⁺ (0, 0.031, 0.062, 0.125, 0.250, 0.500, 1, and 5 μg Ag⁺/L). Exposures were initiated at 4 hpf and maintained for 48 h, and the average oxygen consumption per embryo was determined for two 24 h periods (0–24 h and 24–48 h from the onset of the exposure). The full description of the methodology adopted for these procedures is given in the SI.

RNA Isolation and HT-SuperSAGE Library Construction. RNA extraction was conducted on each pool of 50 embryos using the RNAeasy kit, according to the manufacturer’s instructions (Qiagen, UK). The quality and concentration of the resulting RNA were determined using a NanoDrop ND-1000 Spectrophotometer (NanoDrop Technologies, Wilmington, DE), and an Agilent Bioanalyser 2100 (Agilent Technologies, Inc., Santa Clara, CA). HT-SuperSAGE was conducted according to previously described methods and libraries were sequenced using an Illumina Genome Analyzer (GA) II platform. Further details are given in the SI.

Sequence Tag Preprocessing. In total, over 16.9 million 36-bp sequences were obtained and processed, using the SuperSAGE data analysis procedure presented in Figure 1. A selection of tools from the FAST-Toolkit (http://hannonlab.cshl.edu/fastx_toolkit/) and our own custom Perl scripts were used, running under the NEBC Bio-Linux environment. The FASTQ/A Barcode splitter (FASTX-Toolkit) was used to separate the samples from each lane based on the 4 base barcodes. The barcodes were removed from the sequences using the FASTX/A Trimmer (FASTX-Toolkit), before converting the sequences to FASTA format using FASTQ-to-Fasta (FASTX-Toolkit). A Perl script (SI, File S1) was used to remove all remaining adapter sequences after the last
occurrence of CATG (NruIII restriction site, used for sequence tag preparation) in each of the sequences. FASTX_collapser (FASTX-Toolkit) was used to collapse the sequences and calculate the frequency of unique sequence tags (unitags) in each library.

**Sequence Tag Mapping and Statistical Analyses.** Unitags were mapped to all available zebrafish genes (28 491 genes) in Ensembl Zv9 (release 60) using Bowtie, with no mismatches allowed. Counts of unitags that mapped to the same gene were summed and unitags that aligned with more than one gene were removed from the data set using the Perl scripts provided in SI File S2. For each treatment, the counts of unitags representing each (unique) gene were normalized to 1 million tags. Fold changes (FC) were determined for each gene by dividing the number of tags in the normalized silver treatment libraries by the number of tags in the normalized control libraries. FC values less than 1 (down-regulated genes) were transformed using the following formula: $-1/\text{FC}$, in order to center the fold change values around 0. Differences in gene expression between the silver treatments and the controls were determined using DiscoverySpace 4.0, which implements the Audic-Claverie significance test to account for differences in sample size and P values were adjusted using a false discovery rate correction. Genes that were found to be significantly altered as a result of the silver treatments (adjusted $P < 0.05$) were further investigated. In order to visualize differences and similarities in gene expression between each of the silver treatments and the controls, scatter plots were produced using DiscoverySpace and correlation analysis (Pearson’s) was performed using SigmaPlot 11 (Systat Software UK Ltd.).

**Functional Annotation of Differentially Expressed Genes.** Functional annotation analysis was performed on the lists of differentially expressed genes (adjusted $P < 0.05$) for each treatment using the Database for Annotation, Visualization and Integrated Discovery (DAVID v6.7) and using all genes that were found to be expressed in zebrafish embryos at 24 hpf (12 939 genes) or 48 hpf (13 584 genes), respectively, as background. Gene Ontologies (GO) for Biological Processes, Cellular Components and Molecular Function were considered significantly enriched when $P < 0.05$.

**Pathway Analysis.** Pathway analysis was conducted through the use of Ingenuity Pathways Analysis (IPA; Ingenuity Systems, http://www.ingenuity.com), based on the lists of differentially expressed genes (adjusted $P < 0.05$) for each of the treatments. Enrichment of Canonical pathways was determined using the IPA library of canonical pathways, with a P value cutoff of 0.05 (adjusted for multiple testing using the Fisher’s Exact Test). Similarly, enriched KEGG pathways were determined using the lists of differentially expressed genes (adjusted $P < 0.05$) and all expressed genes at 24 hpf and 48 hpf, respectively, as background using DAVID.

**Other Statistical Analysis.** Statistical analysis of the data for embryo survival and oxygen consumption were conducted within SigmaPlot 11 (Systat Software UK Ltd.). Changes in embryo survival caused by exposure to AgNP, Ag Bulk or Ag+ during the exposure experiments for molecular analysis were analyzed using Chi-square tests. Relationships between the concentration of Ag+ and oxygen consumption were investigated using regression analysis. Differences between treatment groups were determined using one way analysis of variance (ANOVA), followed by an all pair-wise multiple comparison procedure (Student–Newman–Keuls Method). For these tests, data were considered to be significant when $P < 0.05$.

**Data Deposition.** The raw reads and processed unitag frequency tables for each of the HT-SuperSAGE libraries have been deposited in NCBI’s Gene Expression Omnibus and are accessible through GEO Series accession number GSE38125.

**RESULTS AND DISCUSSION**

**Embryo Mortality.** Embryo mortality during the course of the exposures for generation of embryo samples for molecular analysis was 4.6 ± 2.5%, which falls within that expected for zebrafish and within the guidelines for OECD standardized testing guidelines. There were no differences in embryo mortality rates between treatment groups ($P = 0.213$), confirming that all exposures were conducted at concentrations below those causing any overt toxicity.

**HT-SuperSAGE Data Analysis.** The number of sequences obtained for each library ranged from 1.2 million (48 hpf AgNP) to 2.2 million (24 hpf AgNP) and the abundance distributions of these sequence tags were generally consistent between all libraries (SI Table S1). In total, 13 680 994 sequences were taken further for analysis and these were represented by 606 259 unique sequence tags (unitags). In total 99 683 unitags mapped to zebrafish genes, with 77 821 of these mapping to unique genes only (7 690 269 counts in total) and representing 14 793 genes (Figure 1). This is consistent with previous studies that reported a similar number of expressed genes in embryos at similar stages of development using zebrafish Affymetrix microarrays (ArrayExpress database [http://www.ebi.ac.uk/arrayexpress/]; Experiment ID: E-TABM-33) or RNA-Seq. Approximately 80% of the genes were represented by more than one unitag (see SI Figure S2).
consistent with other studies that are based on SAGE methods, which is likely the result of incomplete NlaIII digestion during the library preparations, alternative polyadenylation and/or alternative splicing sites.32−34

Statistical analysis of the gene expression data identified genes differentially expressed between the different treatments and the control, for both 24 hpf and 48 hpf (summarized in SI File S3). A visual representation of the expression profiles of each treatment group compared to the respective control is presented in SI Figure S3. There was a strong correlation between the levels of expression of genes in the controls compared with those in each treatment group (r > 0.937). The number of differentially expressed genes per treatment group, and the overlap between the differentially expressed genes for each treatment were consistent across all treatments, at both 24 h and 48 h of exposure (with the exception of the treatment 24 h Ag+; which showed a greater proportion of differentially expressed genes; Figures 2a, b, and c). A heatmap of the differentially expressed genes across the various silver treatment libraries is shown in SI Figure S4, and demonstrates the consistency in gene expression across all treatments. The condition tree highlights the differences in gene expression
Figure 4. Effects of the silver treatments on transcription of genes belonging to the mitochondrial dysfunction pathway. Shades of green and red represent down-regulation and up-regulation of target genes, respectively. (a) Effects of exposure to AgNP for 24 h on the mitochondrial dysfunction pathway. (b) Summary of the effects of exposure to all treatments on the mitochondrial dysfunction pathway for 24 h and 48 h. Numbers in the table represent fold-change compared to the time-matched control. Illustrations of the mitochondrial dysfunction and oxidative phosphorylation pathways for all six treatments are presented in SI, File S6. Pathways were generated through the use of Ingenuity Pathways Analysis (Ingenuity Systems, www.ingenuity.com).
profiles for the differentially expressed genes between the 24 h Ag⁺ compared to all other treatments and between embryos collected at 24 hpf and 48 hpf.

Some of the differentially expressed genes were found to be specific to each of the different treatments (AgNP = 13 genes; Ag bulk = 1 gene; Ag⁺ = 32 genes), reflecting gene responses distinct to the different treatment (SI Table S2). Distinct genes were found to be differentially expressed at both time points for any one treatment and not present in the list of differentially expressed genes for the other treatments at any time point.

**Commonalities in the Effects of Exposure to AgNP, Ag Bulk and Ionic Ag on Embryonic Gene Expression Profiles.** We explored the biological significance of the changes in gene expression caused by exposure to AgNP, Ag Bulk and Ag⁺ in 24 hpf and 48 hpf embryos by determining the enrichment in GO terms and canonical pathways using DAVID and IPA. Analyses were conducted based on differentially expressed genes, including all up- and down-regulated genes, to obtain a global overview of the biological effects of the three distinct treatments (Figure 3, SI Figure S5 and File S5). The most significantly over-represented GO terms and KEGG pathways (ribosome and oxidative phosphorylation) were common to all three treatments within each time of sampling, demonstrating a very significant overlap in the mechanisms of toxicity of the three forms or silver considered in this study. Based on this overlap, it seems likely that the toxicity of AgNP and Ag Bulk to zebrafish embryos is predominantly associated with the toxicity of free Ag⁺.

The dissolution rates of AgNP and Ag Bulk in our experiments were 0.27% ± 0.15% and 0.27% ± 0.07%, respectively. Parallel studies in our laboratory, investigating the dissolution rates for AgNP and bulk silver in embryo culture water, have found them to vary between 0.1 and 2% over a 48 h period. Furthermore, very recently published data suggest that some silver particles (41.6 ± 9.1 nm AgNPs at a concentration of 0.2 nM) can penetrate the chorion through diffusion via the chorion canals and that once inside the chorion, the dissolution rate can be much higher compared to that observed in the exposure water. Consequently, the amount of bioavailable Ag⁺ in the AgNP and Ag Bulk treatments would likely be sufficient to cause most of the changes in gene expression observed. For many of the genes significantly affected in all the exposures, the changes were more pronounced (in terms of fold-change) for the exposure to Ag⁺ compared with those observed for AgNP and Ag Bulk, and this is likely to be associated, at least in part, with the bioavailability of silver ions in each treatment. However, the pathways affected (as assessed by the analysis of GO and KEGG pathway enrichment (Figure 3, SI Figure S5 and File S5)) remained common across all treatments. These changes are consistent with previous literature reporting effects of exposure to Ag (ions or nanoparticles) on several biological processes across a series of model organisms.

The most significantly enriched GO terms for all treatments, both at 24 and 48 hpf were ribosome, translation and structural constituents of ribosome, for cellular components, molecular functions and biological processes, respectively. Concurrently, the most enriched KEGG pathway was ribosome (Figure 3 and SI File S5). These results align well with previous literature that report disruption to protein biosynthesis pathways in a range of organisms following exposure to AgNP and Ag⁺. Some of the most pronounced changes were observed for ribosomal proteins. For example, rpl10a was down-regulated by 5.6, 3.4 and 5.6-fold (24 h), and rpl18a down-regulated by 12.2, 1.2 and 9.7-fold (48 h), following exposure to AgNP, Ag Bulk, and Ag⁺, respectively. Expression of several genes encoding components of the ubiquitin proteasome system were also altered across all treatments, (at 24 and 48 h), indicating that protein degradation pathways were also altered by the treatments (SI File S3).

Oxidative phosphorylation was affected consistently by the exposure to AgNP, Ag Bulk, and Ag⁺ at both time points analyzed (Figure 3). This is a complex pathway that plays a key role in ATP generation in aerobic organisms, and is comprised of a series of five complexes and other associated molecules that act in concert within the mitochondrial inner membrane. The gene expression profiles following 24 h of exposure to all forms of Ag showed a down-regulation of many of the genes encoding for complex I, II, III and IV (Figure 4 and SI File S6). Within the complexes some genes were up-regulated whereas others

![Figure 5. Oxygen consumption rates of zebrafish embryos exposed to silver ions. Groups of 5 embryos were exposed to seven concentrations of silver ions (0–5 μg/L) in isolated glass vials, and the oxygen consumption was measured for 0–24h (A) and 24–48 h (B) exposure periods (n = six groups of five embryos per treatment). Average oxygen consumptions for individual embryos are presented. There was a significant effect of the exposure to silver ions for the 0–24 h, but not for the 24–48 h exposure periods (One Way ANOVA, P < 0.001, followed by an all pairwise comparison between the groups using the Student–Newman–Keuls Method, P < 0.05). Letters above the box plots represent statistical significance, with treatments with different letters being significantly different from each other.](dx.doi.org/10.1021/es401758d)
were down-regulated, in particular for complex I (for Ag⁺ and Ag Bulk). This complex is comprised of a large number of genes that form 14 central and 32 accessory subunits, encoded by both the mitochondrial and the nuclear genome, and that associate into three functional modules, that oxidize NADH, reduce quinone and pump protons across the membrane (reviewed in ref 39). Our data may reflect a differential regulation of the various subunits and/or a differential regulation of the various functions of complex I. Overall, following 24 h of exposure to all Ag treatments, the gene expression data indicated a down-regulation of all complexes involved in oxidative phosphorylation, possibly with the exception of complex I for Ag Bulk (Figure 4 and SI File S6). This effect is consistent with previously reported observations reporting a down-regulation of this pathway following Ag exposure in a human cell line, using biochemical methods. The degree to which these genes were regulated between the various treatments was consistent with predicted differences in the concentrations of bioavailable Ag⁺, which were expected to be highest for Ag⁺, followed by AgNP and lowest for Ag Bulk.

Data for embryos analyzed following 48 h of exposure to all Ag treatments similarly showed significant alterations in the expression of genes encoding for proteins belonging to all complexes of the oxidative phosphorylation pathway. When taken together, the changes in gene expression suggest an up-regulation of these complexes, consistent with a recovery of the activity in this pathway, during the 24–48 h period, in exposed embryos (Figure 4 and SI File S5).

Quantifying Effects of Ag⁺ on Embryo Metabolism. The changes in gene expression for constituents of the oxidative phosphorylation pathway suggested that Ag may cause a down-regulation in metabolic rates and ATP synthesis during the first 24 h of exposure, with a subsequent recovery from this effect at 48 h for all treatments. To further investigate the feasibility of this hypothesis, we conducted an experiment to investigate the effects of Ag⁺ on the metabolic activity, measuring oxygen consumption as a proxy for aerobic respiration of exposed embryos. Our data demonstrated that Ag⁺ caused a decrease in oxygen consumption during the first 24 h of exposure ($r = −0.469; P < 0.001; Figure S$), with significant decreases in oxygen consumption in embryos exposed to 0.125, 0.5, and 5 μg Ag⁺/L compared with those exposed to 0.031 μg Ag⁺/L, and 5 μg Ag⁺/L compared to the controls during the first 24 h of exposure (One Way ANOVA followed by Student–Newman–Keuls Method; Figure S5a). This effect of Ag⁺ on embryo oxygen consumption was no longer apparent for the second 24 h period (between 24 and 48 h of exposure ($r = −0.114; P = 0.467; Figure S5b$), supporting the inhibitory effects of Ag⁺ on oxidative phosphorylation (and associated decrease in oxygen consumption) during the first 24 h of exposure and subsequent recovery by 48 h. Together, these data support previous studies that have reported inhibition of oxidative phosphorylation following exposure to silver nanoparticles and silver ions in a range of mammalian model systems. Importantly, our data suggest that zebrafish embryos have the capacity to deploy compensatory mechanisms at concentrations of Ag below those causing overt toxicity.

The mechanisms by which exposure to Ag caused the observed changes in the oxidative phosphorylation pathway are difficult to ascertain. Recent evidence from mammalian studies has shown that pyruvate kinase, M2 (PKM2) that catalyzes the rate limiting ATP producing step of glycolysis, plays a key role in maintaining cellular redox homeostasis in cancer cells. This enzyme is specifically oxidized by H₂O₂, resulting in a decrease of its activity, decreased pyruvate formation and increased flux of glycolytic metabolites into the pentose phosphate pathway. This pathway produces reduced NADPH, a crucial source of reducing equivalents for fatty acid synthesis and for the glutathione, peroxiredoxin and thioredoxin systems, which play a key role in the detoxification of reactive oxygen species (ROS), providing cancer cells with a protective mechanism from intracellular ROS (reviewed in ref 45). In contrast, normal mammalian cells express another pyruvate kinase isomerase (PKM1), which is not susceptible to oxidation by hydrogen peroxide. Interestingly, we did not see expression of this later isomerase in either 24 h or 48 h embryos. Instead, zebrafish embryos expressed two forms of the H₂O₂ sensitive isomerase (pkm2a and pkm2b). The principal form of Pkm measured was pkm2a (the average counts per million in our data set were 226.0 ± 14.0 for embryos analyzed at 24 h and 565.5 ± 25.4 for embryos analyzed at 48 h). pkm2b was also expressed, but at much lower levels (0.9 ± 0.6 for 24 h embryos and 40.4 ± 8.4 for 48 h embryos). The fact that both the expressed forms of pkm in zebrafish are homologues to the human H₂O₂ sensitive form suggest that if the intracellular levels of H₂O₂ increase, zebrafish embryos may have the capacity to partially switch to the pentose phosphate pathway, as a mechanism of response to ROS. Increase in ROS is likely to occur during exposure to Ag, as demonstrated by the increase of gene markers for oxidative stress in our study, and in a number of other studies. One of the most significant changes in gene expression across the treatments was for alcohol dehydrogenase class-3 (adhS), which was 70.5, 7.1, and 125.0 fold up-regulated following exposure to AgNP, Ag Bulk, and Ag⁺, respectively, at 24 h. Similarly, after 48 h of exposure, adhS was up-regulated by 151.0, 22.8, and 50.5, as a result of exposure to AgNP, Ag Bulk, and Ag⁺, respectively. Alcohol dehydrogenases (ADH) metabolize ethanol and other alcohols and aldehydes. In mammals, there are several classes of ADH, and class I ADH are responsible for the metabolism of ethanol and other small chain alcohols. Class II ADH enzymes preferentially metabolize larger alcohols and aldehydes (reviewed in ref 47). In the zebrafish, enzymes structurally and functionally similar to the mammalian ADHs have been identified. However, the affinity of these enzymes to alcohols appears to be less specialized than that for mammalian enzymes but the affinity of each enzyme for the various alcohols has not been studied in detail. In addition to their role in alcohol metabolism, ADHs also convert NAD+ to NADH, and this may act as a compensatory mechanism for potential reduction in NADH resulting from the inhibition of the Pkm. The significance of the very pronounced increase in adhS following exposure to all three forms of Ag is, therefore, difficult to ascertain. We hypothesize that this could be associated with the effects of all treatments on the oxidative phosphorylation pathway that may have resulted in a shift toward anaerobic metabolism, with increased accumulation of alcohol within the cells, and detoxification by AdhS.

Specific Effects of Exposure to AgNP, Ag Bulk and Ionic Ag on Embryonic Gene Expression Profiles. Analyses for differentially expressed genes across the different treatments and consistent over time (to investigate for possible particle related effects) identified 1 gene for Ag Bulk, 13 genes for the AgNP treatment and 32 genes for the Ag⁺ treatment, respectively (SI Table S2). These differences, however, would
appear to be relative to the proportion of biologically available Ag⁺ in the different treatments and may not only indicate unique mechanisms associated with each treatment but also be a function of the level of bioavailable Ag⁺.

Considering the uniquely expressed genes in the different treatments, for Ag Bulk, ETS translocation variant S (etv5a) was 1.4 and 1.5-fold down-regulated at 24 h and 48 h, respectively. This gene is involved in the cellular response to oxidative stress (GO:00034599) and in the regulation of transcription, DNA-dependent (GO:00063550).

We identified 13 genes that were specific to AgNP. Among these genes, cryptochrome 1a (cry1a) was strongly down-regulated both at 24 and 48 h of exposure (6.3 and 4.3 fold down-regulated, respectively). This gene acts as a potent repressor of clock function and mimics the effect of constant light to “stop” the circadian oscillator, and is induced by light. This finding suggests that the light availability to embryos exposed to AgNP differed from that available to embryos in other treatments. Despite the fact that we tested relatively low concentrations of AgNP, it is possible that the suspended nanoparticles, and/or nanoparticles at the surface of the embryo, were acting as a barrier to light, and may have resulted in a difference in light availability to the embryos incubated in the AgNP treatment. Other genes responsive to AgNP alone included the proteasome subunit beta type-1 (psnb1), a gene involved in protein catabolism (1.2-fold down-regulated following 24 h of exposure and 1.3-fold up-regulated at 48 h of exposure) and ribosomal protein S6 modification-like protein B (zgc:92166; 1.7-fold and 1.5-fold up-regulated at 24 h and 48 h, respectively). Together, these changes, although small in magnitude, indicate that the protein turnover in the AgNP treatment may be increased compared to the other treatments (in particular to Ag Bulk), suggesting that the potential damage caused by ROS on cellular proteins is compensated through degradation and synthesis of new proteins.

We identified 32 genes that were specific to the Ag⁺ treatment. The genes present in this list belong to gene ontologies that overlap with those over-represented among the genes affected by the other treatments (including oxidative phosphorylation and generation of precursor metabolites and energy). Genes regulated included Cyclic AMP-dependent transcription factor ATF-4 (atf4b1; 2.3 and 2-fold down-regulated at 24 h and 48 h, respectively); a gene involved in gluconeogenesis and regulation of transcription and probable ATP-dependent RNA helicase DDX56 (ddx56; 4.3 and 4.6-fold down-regulated at 24 h and 48 h, respectively), a gene involved in ribosome biogenesis. This adds to the evidence that the genes unique to this treatment reflect responses to higher levels of biologically available Ag⁺ associated with this treatment.

To summarize, we have applied sequencing-based transcription-profiling to investigate the mechanisms of toxicity of AgNP, compared to its ionic and bulk counterparts, in exposed zebrafish embryos, and provide mechanistic evidence for the hypothesis that the toxicity of AgNP is principally associated with the presence of bioavailable Ag⁺. All Ag treatments resulted in an inhibition of the oxidative phosphorylation pathway at 24 h of exposure, potentially due to the increase in ROS within mitochondria. Unique gene responses in the different treatments also supported that differences for responses in the transcriptome could be caused by the differences in bioavailability in Ag⁺ between the treatments. However, a gene response unique to AgNP was identified that may have resulted from alterations in light availability to the developing embryos, demonstrating that not all changes in gene expression observed can be explained by the effects of Ag⁺ alone, and that particle-associated effects may also contribute to the toxicity of AgNP.

Our work suggests that the predominant adverse effects of AgNP appear to be associated with the toxicity of Ag⁺. We would suggest that the risk assessments for AgNP should have a strong focus on understanding the bioavailability of Ag⁺ originating from AgNP, and the toxicology of Ag⁺, to establish what the levels of exposure and health consequences are likely to be in vulnerable environmental compartments.

### ASSOCIATED CONTENT

#### 3 Supporting information

Further details on the methodology and results for the characterization of the silver particles used for the exposures, mortality curves, sequencing analysis, and a number of supporting figures and tables. This material is available free of charge via the Internet at http://pubs.acs.org.

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**Notes**

The authors declare no competing financial interest.

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