The role of chorion on toxicity of silver nanoparticles in the embryonic zebrafish assay

Ki-Tae Kim¹, Robert L. Tanguay²

¹Department of Environmental Engineering, Seoul National University of Science and Technology, Seoul, Korea; ²Department of Environmental and Molecular Toxicology, Oregon State University, Corvallis, OR, USA

Objectives This study was designed to investigate how the size- and surface coating-dependent toxicity of silver nanoparticles (AgNPs) is influenced by the presence and absence of the chorion in an embryonic zebrafish assay.

Methods Normal and dechorinated embryos were exposed to four different AgNPs, 20 or 110 nm in size, with polypyrrolidone (PVP) or citrate surface coatings in a standard zebrafish embryo medium (EM). This was then compared to a 62.5 µM calcium chloride (CaCl₂) solution where agglomeration was controlled.

Results Embryonic toxicity in the absence of the chorion was greater than in its presence. The smaller 20 nm AgNPs were more toxic than the larger 110 nm AgNPs, regardless of the chorion and test media. However, surface coating affected toxicity, since PVP-coated AgNPs were more toxic than citrate-coated AgNPs; this was strongly affected by the presence of the chorion in both EM and CaCl₂.

Conclusions Our results demonstrate the permeability function of the chorion on the size- and surface coating-dependent toxicity of AgNPs. Thereafter, careful experiment should be conducted to assess nanoparticle toxicity in zebrafish embryos.

Keywords Chorion, Silver nanoparticles, Zebrafish
fluences, including chemical exposure. Although the removal of the chorion is helpful in elucidating the mechanisms of toxicity of ENMs, questions concerning interference of the chorion in AgNP toxicity remain, because ecological toxicity testing has, in general, been performed in the presence of the chorion.

Although no comprehensive investigation of chorion permeability or sensitivity to chemicals has yet been reported, the chorion is suspected to be a barrier to the entry of NPs into zebrafish embryos [7,8]. The chorion possesses canals, the pore size of which is approximately 0.6-0.7 μm, larger than the size of the NP. Lee et al. [7] first visualized AgNP penetration through chorion canals. However, although the pore size is known to be larger than the NP size, in particular, the effect of the chorion on NP transport and subsequent biological toxicity may be complicated when NPs are agglomerated. The presence of the chorion is a critical concern on varied developmental toxicity of zebrafish embryos exposed to various formulations of AgNPs to understand the effect of size and surface coating [3,9]. Consequently, the systematic examination of the changes in the biological responses of zebrafish embryos exposed to NPs in the presence or absence of the chorion is necessary.

The objective of this study was to investigate how the size- and surface coating-dependent toxicity of AgNPs varied according to the presence or absence of the chorion in an embryonic zebrafish assay. We used 20 and 110 nm sized AgNPs, and AgNPs were coated with either polyvinylpyrrolidone (PVP) or citrate. The citrate and PVP are the most widespread and most commonly used stabilizer in AgNP synthesis. In order to determine the permeability differential of the AgNPs, we exposed normal and dechorionated embryos to four different AgNPs in a standard zebrafish embryo medium (EM) and 62.5 μM calcium chloride (CaCl₂) solution. We observed that AgNPs were monodispersed in CaCl₂ without agglomeration, and the exposure of zebrafish embryos in CaCl₂ to AgNPs was useful for the observation of the variations in biological toxicity in the presence and absence of the chorion. This can be compared further to EM in which AgNPs were agglomerated.

Materials and Methods

Silver Nanoparticles and Characterization in Test Media

AgNPs synthesized by NanoComposix (San Diego, CA, USA) were used. The AgNPs were 20 nm PVP-coated AgNPs (20AgNPs-P), 20 nm citrate-coated AgNPs (20AgNPs-C), 110 nm PVP-coated AgNPs (110AgNPs-P) and 110 nm citrate-coated AgNPs (110AgNPs-C). This set of AgNPs was one of the ENMs selected by the Nanotechnology Health Implications Research Consortium, an interdisciplinary program of the National Institute of Environmental Health Sciences, which addresses the health and safety concerns of AgNPs. We followed the guidelines for sample handling and storage recommended by the Nanotechnology Characterization Laboratory (NCL).

The NCL provided the physicochemical properties of AgNPs at a concentration of 100 mg/L suspended with 2 mM NaCl. The z-average and zeta potentials of four AgNPs were measured using dynamic light scattering (Malvern Zetasizer Nano ZS Instrument, Southborough, MA, USA) with a back scattering detector (173°). The core diameter was measured by transmission electron microscopy, also provided by the NCL (NCL-NIEHS201111A). In this study, we analyzed and compared the z-average of AgNPs in EM and CaCl₂ by using a ZetaPALS (Brookhaven Instruments, Holtsville, NY, USA) after the preparation of 10-fold diluted aliquot from 1,000 mg/L of stock solution.

Optimization of 62.5 μM Calcium Chloride

Because zebrafish embryos can develop normally in low ionic strength media [10], we tested media of various ionic strengths and found that incubation of embryos in 1 mM CaCl₂ did not induce developmental morbidity or mortality at 120 hours post fertilization (hpf) [6]. To ascertain the minimum concentration of CaCl₂, we carried out a titration, and determined that 62.5 μM CaCl₂ was low sufficient for normal development until 120 hpf and nonagglomeration of AgNPs.

Zebrafish Maintenance and Embryonic Zebrafish Assay

Adult zebrafish (Danio rerio) were raised at Oregon State University’s Sinnhuber Aquatic Research Laboratory in a water flow-through system that maintained a constant temperature of 28°C under a 14:10 hours light-dark photoperiod. Embryos collected and staged [11] were from the tropical SD zebrafish line and were cultured in buffered EM (pH 7 ± 0.2) using reverse osmosis water. The standard EM consisted of 15 mM NaCl, 0.5 mM KCl, 1 mM CaCl₂, 1 mM MgSO₄, 0.15 mM KH₂PO₄, 0.05 mM Na₂HPO₄ and 0.7 mM NaHCO₃ [12].

Healthy staged embryos were loaded into 96-well plates filled with 90 μL of test medium and 10 μL of the AgNP suspension. To achieve the final concentrations of 0.8, 4, 20, 10, and 50 mg/L, after a 10× concentrated, serial diluted concentration of AgNPs was prepared, 10 μL of the AgNP suspension was mixed into the respective wells. Exposure to AgNPs was initialized at 6 hpf. Eight embryos were exposed to each treatment with four replicates. We evaluated a total of 22 endpoints including mor-
tality and malformation at 24 and 120 hpf. The percentage of malformation was calculated by the number of embryos affected per living embryos at each treatment. Screened malformations were: yolk sac edema; axis, eye, snout, jaw, and otic (ear) defects; pericardial edema; brain, somite, and pectoral and caudal fin defects; pigmentation abnormality; circulation and trunk defects; swim bladder abnormality; notochord defects, and touch response abnormality.

Using pronase (Sigma-Aldrich, St Louis, MO, USA), we removed the chorion in an automated dechorinator at 4 hpf. Briefly, after introducing pronase, the plate was shaken for 5 minutes, washed and then the shaking procedure was repeated. While dechorionated embryos were placed into each well using an automated embryo placement systems, normal embryos (i.e., undechorionated embryos) were transferred into a 96-well plate by hand. The ability to produce healthy embryos was confirmed in a previous study [13]. No detectable mortality or morbidity of dechorionated embryos in the control group was observed over 5 days in this study.

Results

Characterization of Silver Nanoparticles

We determined the z-average of each AgNPs in EM and CaCl₂, and compared them to those provided by the NCL (Figure 1). We noted that the stock solution of 1,000 mg/L AgNPs provided by NanoComposix was suspended in 2 mM NaCl, where the NCL measured the reference of the z-average. Depending on the AgNPs, the z-average increased dramatically up to micro-size in EM. The z-average of 20AgNPs-P, 20AgNPs-C, 110AgNPs-P, and 110AgNPs-C increased to 69.7 ± 2.7, 3078.4 ± 400.7, 691.8 ± 40.3, and 1400.5 ± 91.2 nm (mean ± standard deviation), respectively. Although it appeared that the citrate surface coating caused more agglomeration, it is not clear why the diameter of the 20AgNPs-C was larger than that of the 110AgNPs-C. Further, the z-average of AgNPs in CaCl₂ were totally different from those in EM. All AgNPs were stably dispersed (i.e., no increas in z-average), the z-average measured did not differ significantly from those provided by the NCL (Figure 1). The z-average of 20AgNPs-P, 20AgNPs-C, 110AgNPs-P, and 110AgNPs-C were determined to be 30.5 ± 1.2, 26.1 ± 0.8, 123.2 ± 0.9, and 100.4 ± 2.1 nm, respectively. The suspension of 20AgNPs and 110AgNPs exhibited a bright and slightly darker yellow color, respectively, but their color turned to black when transferred into EM, indicating agglomeration, whereas the color did not change in CaCl₂. These z-average corroborate UV-vis spectra analysis (data not shown). The absorbance of AgNPs was maintained over 5 days in CaCl₂ while a drop in absorbance was observed in EM. After 5 days, the absorbance of 20AgNPs-P was attenuated and that of 20AgNPs-C, 110AgNPs-P, and 110AgNPs-C disappeared completely in EM, indicating NP agglomeration. The NCL report stated that all AgNPs exhibited negative zeta-potentials (NCL-NIEHS201111A). Because there is no consistent correlation between zeta-potential and either z-average or UV-vis absorbance spectra in 2 mM NaCl where agglomeration is controlled, zeta-potentials were not measured in EM and CaCl₂ used in this study. The electrostatic repulsion might not be dominant in maintaining AgNP stability in suspension in comparison with the interfacial forces [14]. Taken together, AgNPs were dispersed stably in CaCl₂. We hypothesized that the exposure of nonagglomerated AgNPs helped us to investigate the effect of the chorion on AgNPs with variable size and surface coating.

Embryonic Toxicity of Silver Nanoparticles

The Figures 2 and 3 show the total percentage of mortality and malformation of embryos exposed to four AgNPs in the presence and absence of the chorion in EM and CaCl₂. A total of 18 endpoints evaluated at 120 hpf was used for the toxicity differential of each treatment. The black color indicates the mortality at 120 hpf over the range of AgNP exposure concentrations (0, 0.08, 0.4, 2, 1.0 and 50 mg/L). The percent inci-
Figure 2. Embryonic toxicity of four silver nanoparticles (AgNPs) suspended in (A) embryo medium and (B) 62.5 uM calcium chloride in the presence of the chorion. Eight embryos were exposed and four replicates were used for each treatment. No detectable mortality and malformation were observed at 0, 0.08, 0.4, 2 mg/L at each treatment. 110AgNPs-C, 110 nm citrate-coated AgNPs; 110AgNPs-P, 110 nm polypyrrolidone (PVP)-coated AgNPs; 20AgNPs-C, 20 nm citrate-coated AgNPs; 20AgNPs-P, 20 nm PVP-coated AgNPs.

Figure 3. Embryonic toxicity of four silver nanoparticles (AgNPs) suspended in (A) embryo medium and (B) 62.5 uM calcium chloride in the absence of the chorion. Eight embryos were exposed and four replicates were used for each treatment. No detectable mortality or malformation was observed at 0, 0.08, 0.4, 2 mg/L for each treatment. 110AgNPs-C, 110 nm citrate-coated AgNPs; 110AgNPs-P, 110 nm polypyrrolidone (PVP)-coated AgNPs; 20AgNPs-C, 20 nm citrate-coated AgNPs; 20AgNPs-P, 20 nm PVP-coated AgNPs.

Figure 4. Representative images of embryos exposed to four silver nanoparticles (AgNPs) at 120 hpf: (A) control, (B) unhatched embryos, mainly observed at 50 mg/L of polypyrrolidone (PVP)-coated AgNPs (AgNPs-P) and citrate-coated AgNPs (AgNPs-C) in the presence of the chorion, (C) malformed embryos.

Figure 4 illustrates examples of embryos exposed to each Ag-
As demonstrated, the size- and surface coating-dependent toxic comparison of ENM toxicity with respect to chorion permeability was needed. A further study comparing the effect of physical size effect in the embryonic toxicity of AgNPs, regarding the surface coating in the presence of the chorion. Consequently, presence of the chorion related to the hatching process in the presence of the chorion [13] is lacking in nanotoxicology. Therefore, further experimental investigations of the effect of the chorion should be carefully considered in assessing the toxicity of ENMs and other chemicals in the embryonic zebrafish assay.

**Discussion**

Based on the percentage of mortality and malformation, we ranked the embryonic toxicity of AgNPs in EM in the presence of the chorion as: 20AgNPs-P > 20AgNPs-C > 110AgNPs-C ≈ 110AgNPs-P and in the absence of the chorion as 20AgNPs-C > 20AgNPs-P > 110AgNPs-C > 110AgNPs-P. The incidence of mortality was considered to be more toxic than malformation. The presence of the chorion changed the order of toxicity with respect to the surface coating rather than the size. It is not clear, however, that 20AgNPs-C with a larger z-average in EM were more toxic than 110 nm AgNPs, regardless of the presence of the chorion. In CaCl₂, the embryonic toxicity of AgNPs was ranked as 20AgNPs-P > 20AgNPs-C > 110AgNPs-P ≈ 110AgNPs-C and 20AgNPs-P > 20AgNPs-C > 110AgNPs-P > 110AgNPs-C in the presence and absence of the chorion, respectively. We noted that in the environment in which particle agglomeration (i.e., in CaCl₂) and chorion interference (i.e., in the absence of the chorion) were controlled, PVP was more toxic than the citrate coating, regardless of the particle size, as shown above. In CaCl₂, where agglomeration was controlled, a change in the order of toxicity also occurred, predominantly regarding the surface coating in the presence of the chorion. Consequently, the presence of the chorion did influence the interaction with the chemical surface coating material, rather than the physical size effect in the embryonic toxicity of AgNPs, regardless of the test medium. A further study comparing the effect of chorion on AgNPs with sizes between 20 and 110 nm AgNPs is needed.

To our knowledge, this is the first report of a systematic comparison of ENM toxicity with respect to chorion permeability. As demonstrated, the size- and surface coating-dependent toxicity of AgNPs was controlled by the function of embryonic permeability. Embryonic permeability was affected by particle agglomeration, and primarily caused a differential in toxicity dependent on the surface coating. The embryonic zebrafish assay is a well-established model not only for rapid and cost-efficient toxicity screening [15,16], but also for the identification of molecular mechanisms [17] in assessing ENM toxicity. However, knowledge with regard to the permeability limitations and secondary phenotypic responses to the chorion related to the hatching process in the presence of the chorion [13] is lacking in nanotoxicology. Therefore, further experimental investigations of the effect of the chorion should be carefully considered in assessing the toxicity of ENMs and other chemicals in the embryonic zebrafish assay.

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**Conflict of Interest**

The authors have no conflicts of interest with material presented in this paper.

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