

Engineered nanoparticles interact with nutrients to intensify eutrophication in a wetland ecosystem experiment

MARIE SIMONIN,^{1,2,9} BENJAMIN P. COLMAN,^{1,3} STEVEN M. ANDERSON,^{1,2} RYAN S. KING,^{1,4} MATTHEW T. RUIS,^{1,2}
ASTRID AVELLAN,^{1,5} CHRISTINA M. BERGEMANN,^{1,2} BRITTANY G. PERROTTA,^{1,4} NICHOLAS K. GEITNER,^{1,6}
MENGCHI HO,^{1,7} BELEN DE LA BARRERA,^{1,7} JASON M. UNRINE,^{1,8} GREGORY V. LOWRY,^{1,5} CURTIS J. RICHARDSON,^{1,7}
MARK R. WIESNER,^{1,6} AND EMILY S. BERNHARDT^{1,2}

¹Center for the Environmental Implications of Nanotechnology (CEINT), Duke University, Durham, North Carolina 27708 USA

²Department of Biology, Duke University, Durham, North Carolina 27708 USA

³Department of Ecosystem and Conservation Sciences, University of Montana, Missoula, Montana 59812 USA

⁴Department of Biology, Baylor University, Waco, Texas 76798 USA

⁵Department of Civil and Environmental Engineering, Carnegie Mellon University, Pittsburgh, Pennsylvania 15289 USA

⁶Department of Civil and Environmental Engineering, Duke University, Durham, North Carolina 27708 USA

⁷Duke University Wetland Center, Nicholas School of the Environment, Duke University, Durham, North Carolina 27708 USA

⁸Department of Plant and Soil Sciences, University of Kentucky, Lexington, Kentucky 40526 USA

Abstract. Despite the rapid rise in diversity and quantities of engineered nanomaterials produced, the impacts of these emerging contaminants on the structure and function of ecosystems have received little attention from ecologists. Moreover, little is known about how manufactured nanomaterials may interact with nutrient pollution in altering ecosystem productivity, despite the recognition that eutrophication is the primary water quality issue in freshwater ecosystems worldwide. In this study, we asked two main questions: (1) To what extent do manufactured nanoparticles affect the biomass and productivity of primary producers in wetland ecosystems? (2) How are these impacts mediated by nutrient pollution? To address these questions, we examined the impacts of a citrate-coated gold nanoparticle (AuNPs) and of a commercial pesticide containing Cu(OH)₂ nanoparticles (CuNPs) on aquatic primary producers under both ambient and enriched nutrient conditions. Wetland mesocosms were exposed repeatedly with low concentrations of nanoparticles and nutrients over the course of a 9-month experiment in an effort to replicate realistic field exposure scenarios. In the absence of nutrient enrichment, there were no persistent effects of AuNPs or CuNPs on primary producers or ecosystem productivity. However, when combined with nutrient enrichment, both NPs intensified eutrophication. When either of these NPs were added in combination with nutrients, algal blooms persisted for >50 d longer than in the nutrient-only treatment. In the AuNP treatment, this shift from clear waters to turbid waters led to large declines in both macrophyte growth and rates of ecosystem gross primary productivity (average reduction of 52% ± 6% and 92% ± 5%, respectively) during the summer. Our results suggest that nutrient status greatly influences the ecosystem-scale impact of two emerging contaminants and that synthetic chemicals may be playing an under-appreciated role in the global trends of increasing eutrophication. We provide evidence here that chronic exposure to Au and Cu(OH)₂ nanoparticles at low concentrations can intensify eutrophication of wetlands and promote the occurrence of algal blooms.

Key words: algae; algal bloom; aquatic ecosystems; copper nanoparticles; Egeria; eutrophication; gold nanoparticles; macrophytes; multiple stressors; nanomaterial; nutrients.

INTRODUCTION

The rising concentrations and diversity of synthetic chemicals in the environment is an important marker of the Anthropocene (Lewis and Maslin 2015). However, our understanding of the ecological consequences of emerging contaminants lags far behind that of other global change drivers, such as elevated atmospheric CO₂, habitat loss, or climate change (Bernhardt et al. 2017). Although many emerging contaminants, like metal nanomaterials, are becoming globally ubiquitous (Jambeck et al. 2015, Stehle and Schulz 2015, González-Alonso et al. 2017), they are found at both the highest concentrations and in the greatest diversity in

wastewaters and agricultural runoff, where they co-occur with high nutrient loads and historically problematic contaminants (King et al. 2016). The nutrient pollution side of this issue is well studied. Eutrophication resulting from excess phosphorus (P) and nitrogen (N) inputs to water bodies stimulates algal blooms and low oxygen conditions that often result in a loss of biodiversity (Smith et al. 1999). Though eutrophication is recognized as the primary water quality issue in freshwater ecosystems worldwide (Smith and Schindler 2009), little attention has been paid to how emerging synthetic chemicals, like nanomaterials, may interact with nutrient pollution and alter ecosystem productivity and biodiversity.

There is growing evidence that our failure to examine the interactions between nutrient and other synthetic chemical pollutants may be compromising our understanding of impaired aquatic ecosystems. For instance, two studies found that phosphorus availability modulated the response

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⁹E-mail: simonin.marie@gmail.com

of phytoplankton and periphyton communities to silver nanoparticles in lakes, leading to a reduction of the contaminant toxicity for pelagic primary producers and an increased toxicity for the benthic communities (Das et al. 2014, Norman et al. 2015). The deleterious effects of synthetic chemicals may be amplified under high nutrient loads due to increased uptake rates and faster transfer of the contaminants through exposed food webs (Berglund 2003). Or, on the contrary, the uptake of toxic compounds in freshwater food webs has been shown to decrease in the presence of high algal biomass, especially during algal blooms in eutrophic systems (Pickhardt et al. 2002). Because eutrophication often shifts ecosystem pH and alters organic matter composition and concentrations, nutrient enrichment can thus alter the fate, transport and reactivity of pollutants (Skei et al. 2000, Valenti et al. 2011, Kong et al. 2017). While the ecological consequences of both nutrient pollution and synthetic chemical exposures are often mediated through complex top-down or bottom-up effects, such complicated trophic interactions are still rarely included in ecological or ecotoxicological studies (Rohr et al. 2006, Gessner and Tlili 2016), at the exception of some well-studied conventional pesticides (Van den Brink et al. 2000, 2009, Lin et al. 2012, Yin et al. 2018).

One important class of emerging contaminants that represents a rapidly growing sector of the synthetic chemical industry are engineered nanoparticles (NPs) that are being produced in increasing quantity to perform a wide range of applications in medicine, food, cosmetics, electronics, or as agrochemicals (Kah 2015, Mitrano et al. 2015, Servin et al. 2015, Sun et al. 2015). Nanomaterials are now recognized as emerging contaminants of terrestrial and aquatic ecosystems where they join other emerging and established contaminants (e.g., nutrient pollution, metals, pharmaceuticals). While many studies have examined the effects of engineered nanomaterials on organisms, they have typically examined them in isolation from other likely co-occurring contaminants that may modulate their ecological effects (Simonin and Richaume 2015, Bundschuh et al. 2016).

Hence, in this study, we asked two main questions: To what extent do engineered nanomaterials affect the biomass and productivity of primary producers in wetland ecosystems? How are these impacts modulated by nutrient enrichment? To answer these questions, we assessed the impact of repeated exposures of wetland ecosystems to two different nanomaterials: citrate-coated gold nanoparticles and a nanopesticide containing $\text{Cu}(\text{OH})_2$ nanoparticles (later called AuNPs and CuNPs, respectively). With the growing use of nanomaterials in agrochemical products and the application of biosolids containing high concentrations of NPs, wetlands receiving waters from agriculture fields will likely be a major environmental sink for these emerging contaminants (Dale et al. 2015, Kah 2015). We added CuNPs in the form of a commercial pesticide (Kocide 3000, DuPont, Wilmington, Delaware, USA), whereas the AuNPs were custom synthesized. Our intention in the AuNP enrichment was to use these particles as a tracer for nanoparticle behavior because AuNPs typically have low aqueous solubility and, in contrast with Cu, there are low natural background concentrations of Au in the environment (Ferry et al. 2009, Keller et al. 2017). We conducted our experiment in large wetland mesocosms that were

assigned to one of two nutrient conditions (ambient vs. nutrient enriched). Over the course of a nine-month experiment, we examined the interactive effects of continuous, low dose additions of AuNPs or CuNPs (low $\mu\text{g/L}$ range) with nutrient enrichment (inorganic N and P additions) on aquatic primary producers within a set of outdoor replicated wetland mesocosms. The submerged portion of our mesocosms was dominated by the aquatic macrophyte *Egeria densa* and supported complex aquatic food webs including numerous macroinvertebrate taxa and large populations of *Gambusia holbrooki*. In this paper, we specifically assessed the individual and interactive effects of nanoparticles and nutrient enrichment treatments on water chemistry, biomass, metal accumulation, and activity of aquatic primary producers.

We hypothesized that CuNPs would have significant ecological impacts on primary producers due to the well-described toxicity and antimicrobial activity of this fungicide, even in the $\mu\text{g/L}$ range in aquatic ecosystems (Wang et al. 2011b, Thwala et al. 2016, Keller et al. 2017). The direction of this effect was difficult to predict a priori, as CuNP fungicides could directly affect algal communities and the photosynthetic potential of macrophytes (Regier et al. 2015), or they might indirectly benefit autotrophs because of their toxicity to competing microbes. We expected to observe significant effects of AuNP on wetland autotrophs, as AuNPs have been found to induce toxicity and to be bioavailable for different model organisms (Barena et al. 2009, Sabo-Attwood et al. 2012, Tsyusko et al. 2012, Glenn and Klaine 2013). We anticipated significant treatment interactions between nutrient enrichment and both NPs, as previous studies have reported interactive effects of nutrient enrichment and pollutants on algal and periphyton stoichiometry and on macrophyte growth (Fulton et al. 2009, Das et al. 2014, Norman et al. 2015). However, the direction of this interaction was difficult to predict as nutrient availability may either compensate for or enhance the impact of metal pollution on aquatic primary producers (Gessner and Tlili 2016).

MATERIALS AND METHODS

Wetland mesocosm setup and experimental design

Our outdoor wetland mesocosm facility is located in the Duke Forest, a research forest adjacent to Duke University in Durham County, North Carolina, USA. Each mesocosm is a large box built from weather-treated lumber (dimensions $3.66 \times 1.22 \times 0.81$ m) that was partially filled with sand. This sand was graded to create a flat, deep segment 0.8 m in length adjacent to a 2.8 m hillslope rising at a 13° slope and then lined with a food grade liner. This setup allowed us to create three different hydrologic zones within each mesocosm: a permanently flooded zone (aquatic zone), a periodically flooded zone (transition zone), and a rarely flooded zone (upland zone). More details about the construction, set-up and monitoring of the mesocosms are further described in Lowry et al. (2012) and Colman et al. (2014). In early June 2015, we filled the mesocosms with a sandy-loam soil (Soils and Sand, Durham, North Carolina, USA) with a texture of 56.1% sand, 24.3% loam, 19.6% clay, total organic matter content of 3.05%, and a pH of 6.3. Initially,

250 L of groundwater from the experimental site in the Duke Forest was used to fill the mesocosms. The organisms were introduced sequentially in the system between June and September 2015. In the aquatic zone, we added the macrophyte *Egeria densa*; the aquatic snails *Physella acuta* and *Lymnaea* sp.; and the fish *Gambusia holbrooki* (eastern mosquitofish). Algae and zooplankton inoculum was added in 250 mL of water from a local wetland biweekly to avoid strong divergences between mesocosms over time because of limited dispersal (Hall et al. 2004). To homogenize the water chemistry and algal composition between mesocosms, the water was circulated between all the mesocosms throughout July 2015 using submersible pond pumps. In the transition zone, the following plant species were planted: *Juncus effusus*, *Carex lurida*, and *Lobelia cardinalis*. In the upland zone, *Panicum virgatum* and *Chasmanthium latifolium* were planted, *Lolium multiflorum* and *Andropogon gerardii* were seeded. These mesocosms were open to colonization by other organisms and populations of insects and spiders were also monitored during the experiment.

Mesocosms were designated as either nutrient enriched or ambient nutrient, with nutrient additions to the nutrient enriched mesocosms beginning on 28 September 2015, over three months before the beginning of the nanomaterial additions. The ambient nutrient mesocosms received 1 L of mesocosm water each week without any nutrients added to mimic dosing disturbance, while the Nutrient enriched mesocosms received 1 L of mesocosm water each week that had been supplemented with 88 mg of N as KNO_3 and 35 mg of P as KH_2PO_4 . While our addition rate had an N:P molar ratio of 5.5, the measured nutrient N:P ratios in the water column averaged at 16.8 ± 1.6 in the nutrient enriched mesocosms (23 ± 3.0 in the ambient nutrient mesocosms). Our goal was to push the mesocosms towards eutrophic conditions without attaining hypereutrophic conditions. We achieved this with an annual addition rate of 2.5 g N/m^2 of and 1.0 g P/m^2 , which falls between thresholds for eutrophication in wetland ecosystems receiving water from agroecosystems (Verhoeven et al. 2006).

On 18 January 2016, we randomly assigned the NP treatments to cross the nutrient treatments, resulting in a full-factorial experiment with three replicate mesocosms for each of six treatment/nutrient-status combinations: control-ambient nutrient, control-nutrient enriched, AuNP-ambient nutrient, AuNP-nutrient enriched, CuNP-ambient nutrient, CuNP-nutrient enriched. Nanoparticle additions were done weekly over the 270-d (9-month) experiment, with NP additions mixed with 1 L of mesocosm water, which was then distributed throughout the entire aquatic zone of each mesocosm directly below surface water. Control treatments received the same volume of mesocosm water without any NPs.

The citrate-stabilized AuNPs were synthesized by the Center for the Environmental Implications of Nanotechnology (CEINT) at Duke University and had an average primary particle diameter of $11.9 \pm 1.2 \text{ nm}$ (transmission electron microscopy measurement) in the stock suspension and an average hydrodynamic diameter of $10.9 \pm 1.4 \text{ nm}$ and an apparent zeta potential of $-13 \pm 6 \text{ mV}$ in mesocosm water without nutrient addition at pH 7.5 (Dynamic Light Scattering measurement based on particle number; Zetasizer Nano ZS, Malvern, UK). In the nutrient enriched mesocosm water,

the hydrodynamic diameter was $12.0 \pm 0.2 \text{ nm}$ and the zeta potential was $-14 \pm 1 \text{ mV}$, indicating overall a high stability of the AuNPs in mesocosm water. The mesocosms exposed to AuNPs received a weekly dose of 19 mg Au uniformly applied to the aquatic zone resulting in a total dose of 750 mg Au added over the 9 months of the experiment.

The copper hydroxide NPs (CuNPs) were a commercial nanopesticide (KOCIDE 3000; DuPont, Wilmington, Delaware, USA). As a commercial product, this pesticide does not contain only $\text{Cu}(\text{OH})_2$ NPs and present some trace amounts of other elements including a polymer matrix that could influence its toxicity and bioavailability (Keller et al. 2017). The CuNPs had an average primary particle diameter of $38.7 \pm 8.2 \text{ nm}$ in the stock powder and an average hydrodynamic diameter of $120 \pm 30 \text{ nm}$ and zeta potential of $-32.0 \pm 5.5 \text{ mV}$ in the mesocosm water without nutrient addition at pH 7.5. In the nutrient enriched mesocosm water, the hydrodynamic diameter was $106 \pm 16 \text{ nm}$ and the zeta potential was $-32.8 \pm 1.6 \text{ mV}$. The mesocosms exposed to CuNPs received a weekly dose of 35 mg of Kocide, except for the first week of the treatment when they received an initial pulse of 347 mg that resulted in a total dose of 1.664 g Kocide per mesocosm over the 9 months of the study. The measured Cu content in the Kocide product is 27% (dry mass). Thus, the total dose of Cu-based NPs added to the mesocosms was then 450 mg of Cu over the course of the experiment. The dissolution rates of the CuNPs were assessed in situ using Float-ALyzer G2 membrane dialysis devices (Spectrum Laboratories, Rancho Dominguez, California, USA) with a molecular weight cutoff of 8–10 kDa. It was observed that after 48 h, $60\% \pm 8\%$ of the CuNPs were dissolved in absence of nutrient addition and $70\% \pm 3\%$ in nutrient enriched conditions (Vencalek et al. 2016).

The concentrations of CuNP and AuNP applied to the mesocosms were different because they were added for different goals. The concentration for CuNP was designed to simulate field rates. We assumed a conservative ratio between contributing land surface to wetland area of 10:1 for the aquatic compartment of our wetland (1.83 m^2 at initiation of treatments). Based on the application rates from the label of Kocide 3000, we assumed an application rate of 20 kg/ha (intermediate value between mean rates for tree crops and field crops). We then used a literature value measured for a Kocide $\text{Cu}(\text{OH})_2$ fungicide for the rate of loss of agrochemicals from surface soils of 6% (Rice et al. 2001). When scaled to the 9-month window of our experiment, this gave an addition rate of 450 mg of Cu per mesocosm. AuNPs were used as a particle tracer (low dissolution) that would behave more like a particle than CuNPs (high dissolution). As such, we needed to ensure that it would be readily detectable across time and in various compartments of the mesocosms. Based on past experiments, we chose 750 mg of Au addition per mesocosm. As a whole, the amount of both CuNPs and AuNPs was designed to yield water column concentrations in the low $\mu\text{g/L}$ range.

Au and Cu concentrations in unfiltered surface water were measured before and after dosing every week. Samples were acid digested (HNO_3 and HCl , 3:1 ratio) and measured using ICP-MS (7900 ICP-MS, Agilent, Santa Clara, California, USA). On average, the Au concentration in surface water 2 h after dosing was $74.1 \pm 47.9 \mu\text{g/L}$ and 7 d after

dosing (right before the new weekly dosing) was $4.4 \pm 3.7 \mu\text{g/L}$. On average, Cu concentration in water after dosing was $50.8 \pm 30.0 \mu\text{g/L}$ and 7 d after dosing was $13.1 \pm 10.9 \mu\text{g/L}$. In the control mesocosms, the Au and Cu concentrations were below the method detection limit ($0.15 \mu\text{g/L}$) for most of the measurements.

Water chemistry and ecosystem metabolism rates

Over the course of the experiment, we extensively monitored the mesocosm water volume, temperature, turbidity, pH, conductivity (Manta 2, Eureka, Austin, Texas, USA), anion concentrations: NO_3^- , SO_4^{2-} , Cl^- , Br^- (ICS 2000 Ion Chromatograph, Dionex, Sunnyvale, California, USA) and ortho- PO_4 (DU-64 Spectrophotometer, Beckman Coulter, Atlanta, Georgia, USA) on a daily or weekly basis. The weather conditions were also monitored continuously during the experiment for air temperature, precipitations, wind, relative humidity, and barometric pressure (Weather station, Campbell Scientific, Logan Utah, USA). Some selected parameters are presented in Appendix S1.

On a monthly basis, the concentrations of total N and dissolved organic carbon (DOC) were measured on a TOC-VCPH Analyzer with a TNM-1 module (Shimadzu, Kyoto, Japan) and total P on a Beckman DU-64 Spectrophotometer.

Dissolved oxygen (DO) concentrations (at 15 cm deep) were measured weekly before dawn using a YSI DO200 or YSI 556 (YSI, Yellow Springs, Ohio, USA). Continuous DO measurements (15-min intervals) for one to four consecutive days were conducted monthly using a EXO1 data sondes probe (YSI) in order to calculate the ecosystem metabolism rates: ecosystem gross primary productivity (GPP) and ecosystem respiration (ER) that were estimated from the day–night dynamics of DO in surface waters. We performed metabolism rate calculations using the LakeMetabolizer package in R (Winslow et al. 2016) with the *k.vachon* function to estimate the coefficient of gas exchange across the air–water interface (*k*) and the *metab.ols* function using the ordinary least squares model to estimate the GPP and ER rates.

Algae

Knowing that our treatments could cause modifications in the abundance of planktonic and epiphytic algae, we collected both quantitative data on phytoplankton and qualitative data on epiphytic algae. For phytoplankton, chlorophyll *a* (chl *a*) concentrations were measured weekly in surface waters at 15 cm deep (Submersible fluorometer, Turner Designs, San Jose, California, USA and CR1000 data logger, Campbell Scientific). Epiphytic algal abundance was recorded in weekly photographs of each mesocosm. Based on published correlations between planktonic and epiphytic algal abundance under low to moderate nutrient enrichment (Sand-Jensen and Søndergaard 1981, Sand-Jensen and Borum 1991), an algal bloom threshold was established based on chl *a* concentrations in the water column that was consistent with the literature for phytoplankton (Carlson 1977, Pace et al. 2017), and consistent with evidence of the existence and persistence of a mat of epiphytic algae as documented in mesocosm photographs. The algal bloom threshold was estimated at $3.37 \mu\text{g/L}$ chl *a* in this experimental system.

Macrophytes

To measure stem-level rates of photosynthesis and respiration and the accumulation of the NPs in new plant growth, three stems from the macrophyte *Egeria densa* were collected (7 cm clippings shoots) at the water surface every three months (Days 90, 193, and 269). These stems were thoroughly rinsed to remove periphyton before measuring photosynthesis and respiration rates in a bottle assay. The stems were incubated in 40 mL vials in filter-sterilized mesocosm water (from where the stem originated) for 3–4 h under light and for 3–4 h under dark conditions in a growth chamber (25°C , photosynthetically active radiation [PAR] $250\text{--}450 \mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$). The CO_2 concentration was measured in the headspace using a LI-COR LI 6200 and 6250 (LI-COR, Lincoln, Nebraska, USA) and the slope of accumulation or decline of CO_2 over time was used to calculate the photosynthesis and respiration rates. To express the photosynthesis and respiration rates per unit of surface area, the leaf area was measured by scanning each individual leaf and using the WinFolia software (Regent Instruments, Quebec, Canada). The dried *Egeria densa* stems were later digested using either $\text{HNO}_3\text{:H}_2\text{O}_2\text{:HCl}$ (6:3:22) for control or Au-exposed tissues, or $\text{HNO}_3\text{:H}_2\text{O}_2$ (2:1) for Cu-exposed tissues. Total metal concentration was then measured using ICP-MS (Agilent 7700; Agilent, Santa Clara, California, USA). The absence of drift during the analysis was checked by measuring two standards (5 and 10 ppb, Cole-Parmer, Vernon Hills, Illinois, USA) every 10 samples. The calculated method detection limit was $0.5 \mu\text{g}$ of Au or Cu per g of dry plants.

At the same three-month intervals, *Egeria densa* growth rates were measured by harvesting all the plant biomass that had colonized mesh columns (Tenax Cintoflex-M Aquaculture and Hatchery Netting, 2 cm mesh size, column diameter 20 cm, height 50 cm) located in the aquatic zone of the mesocosms. Three mesh columns per mesocosm were positioned in the aquatic zone along a depth gradient. At each harvest, the biomass collected in the three mesh columns was combined and dried at 60°C for at least 72 h and the water volume inside each column was measured to determine the *Egeria densa* growth rates per unit volume.

Statistical analyses

Analysis of chl *a*, dissolved oxygen, GPP, ER, *Egeria densa* photosynthesis, respiration rates, growth rates, metal concentrations, total N, total P, and DOC concentrations were conducted using generalized linear mixed-effects modeling to determine the effects of NP exposure (Control, AuNPs, CuNPs), nutrient status (ambient nutrient, nutrient enriched), and their interactions. In model selection and post hoc tests, we chose $P < 0.05$ to discriminate significant effects. Main effects and interactions were nested by day and mesocosm was treated as a random effect in the models to account for serial correlation among observations from the same mesocosms over time (Zuur et al. 2009). The models were fit following a framework similar to the one described in King et al. (2016) using the *glmer* function of the *lme4* package in R 3.2.3 (R Core Team 2015). For non-repeated measurements, such as the number of algal bloom days, a generalized linear model was used to test the effects of

nanoparticle exposure, nutrient enrichment, and their interaction by using the glm function and a similar framework to fit the model to the data. The most appropriate probability distribution (family) for each analyte was initially constrained by the type of response variable (e.g., for counts, Poisson or negative binomial; for continuous, Gaussian or Gamma; for proportions, binomial). Model fitting was then done using a tiered approach. First, to test and correct for disparities between the assumed and actual distribution of analytes, multiple models were compared using different link functions (e.g., identity, inverse, log) with residual plots and Aikake information criterion (AIC) values used to select the most parsimonious model (Zuur et al. 2009). Second, to test for and correct for the presence or absence of interactions, the full model with all interactions was compared with models with nonsignificant interactions being removed and

residual plots and AIC values again being used to select the most parsimonious model. To determine at what levels the differences arose due to treatment, nutrient status, and/or time, post hoc comparisons were performed using the lsmeans function in the lsmeans package in R. Results were graphed using the ggplot2 package in R.

RESULTS

Exacerbated eutrophication in nutrient enriched AuNP and CuNP mesocosms

The water column total N concentration was more than 30% higher in both AuNP-nutrient enriched and CuNP-nutrient enriched treatments compared with the control-nutrient enriched mesocosms on Days 27 and 56 (Fig. 1a, b; NP

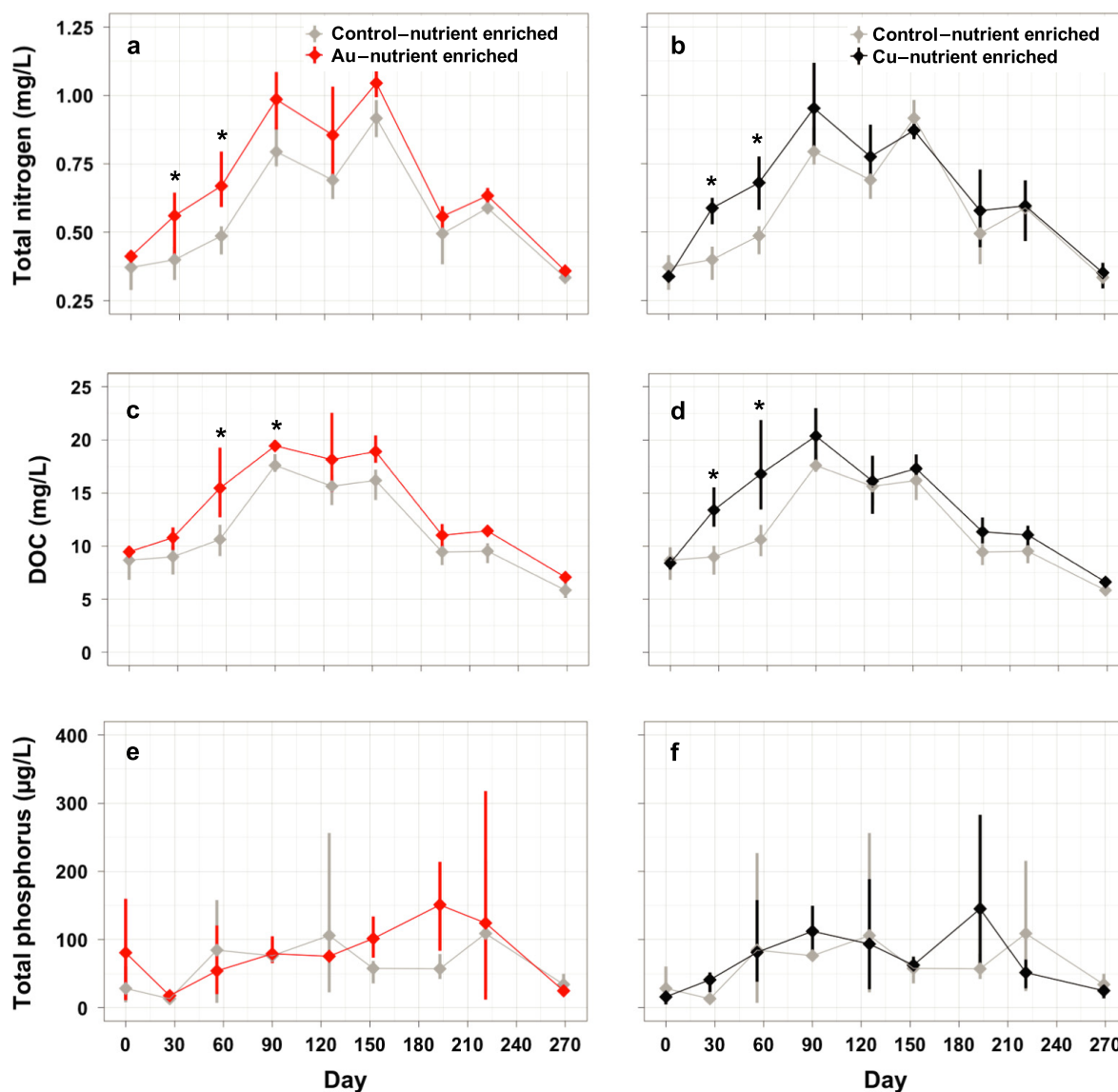


FIG. 1. (a, b) Total nitrogen, (c, d) dissolved organic carbon (DOC), and (e, f) total phosphorus in the water column in the nutrient enriched treatments only. The citrate-coated gold nanoparticle (AuNP; a, c, e) and $\text{Cu}(\text{OH})_2$ nanoparticles (CuNP; b, d, f) nutrient enriched treatments are displayed in different panels along with their respective control treatment means and 95% confidence intervals are presented and significant effects ($P \leq 0.05$) are indicated by asterisks.

effect, $P = 0.004$). Similarly, DOC concentrations were more than 30% higher than controls in the AuNP-nutrient enriched on days 56 and 90 (Fig. 1c) and more than 49% higher in CuNP-nutrient enriched mesocosms than controls on days 27 and 56 (Fig. 1d, NP effect [$P < 0.001$], nutrient effect [$P = 0.005$]). In the AuNP-ambient nutrient and CuNP-ambient nutrient mesocosms, there was no detectable change in total N or DOC concentrations compared to control-ambient nutrient mesocosms (data not shown). Unlike total N or DOC, total P concentration were not significantly altered by any experimental treatment on any date (Fig. 1e, f).

Repeated nanomaterial exposure in nutrient enriched conditions increased algal bloom frequency and magnitude

On multiple dates, phytoplankton biomass (measured as water column chl *a*) was significantly higher in the AuNP-nutrient enriched (Fig. 2b) and CuNP-nutrient enriched treatments (Fig. 2d) compared to the control-nutrient enriched treatment. In contrast, we never observed a significant chl *a* response to nutrients added without NPs (significant NP $\times$ nutrient interaction, $P = 0.006$). In the absence of nutrient enrichment, the chl *a* concentration was significantly higher or lower compared to the control-ambient nutrient treatment at several dates in both CuNP (higher at two dates, lower at one date, Fig. 2c) and AuNP (higher at three dates, lower at three dates, Fig. 2a; overall NP effect,

$P < 0.001$). The highest chl *a* concentrations recorded in this experiment were observed in the AuNP-nutrient enriched treatment ($>30 \mu\text{g/L}$) during major algal bloom events (between Day 150 and 210, summer season). Over the course of the experiment, the number of days when a large algal bloom was observed was 3.5 times higher in the nutrient enriched AuNPs and CuNPs exposed mesocosms (77–79 d) than in the control-nutrient enriched mesocosms (23 d, Fig. 3). In the absence of nutrient enrichment, the number of algal bloom days was not significantly different in the mesocosms exposed to NPs and control conditions (Fig. 3).

Consequences to dissolved oxygen and metabolism rates

Under nutrient enriched conditions, pre-dawn dissolved oxygen (DO) levels were consistently lower than nutrient enriched controls in both the AuNP treatment between Day 20 and 116 (Fig. 4b) and in the CuNP treatment between Day 20 and 63 (Fig. 4d). NPs alone had no consistent effect on DO concentrations in the absence of nutrient enrichment (Fig. 4a, c). All mesocosms had pre-dawn DO concentrations consistently below 10% throughout the summer (Day 132 and 234) and were predominantly hypoxic.

Gross primary productivity (GPP) and ecosystem respiration (ER) varied widely over the course of the experiment and the highest ecosystem metabolism rates were observed in the spring (Day 86, Fig. 5). The metabolism rates were

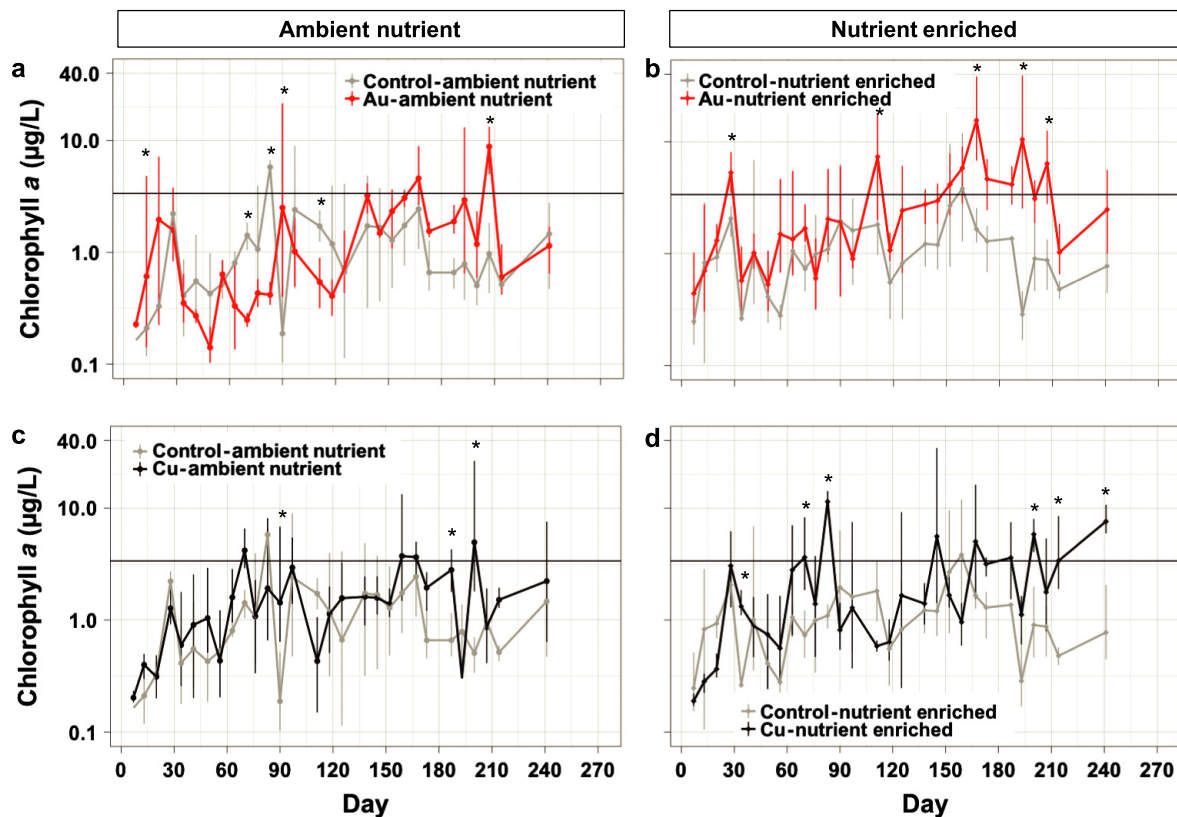


FIG. 2. Chlorophyll *a* concentration on a log scale in the surface water over the 9 months of the experiment. The different treatments are displayed in different panels along with their respective control treatment: (a) AuNP-ambient nutrient, (b) AuNP-nutrient enriched, (c) CuNP-ambient nutrient, (d) CuNP-nutrient enriched. Significant effects ($P \leq 0.05$) are indicated by an asterisk. The horizontal line represents the threshold at which a major algal bloom event was observed. Values are mean $\pm$ 95% confidence intervals.

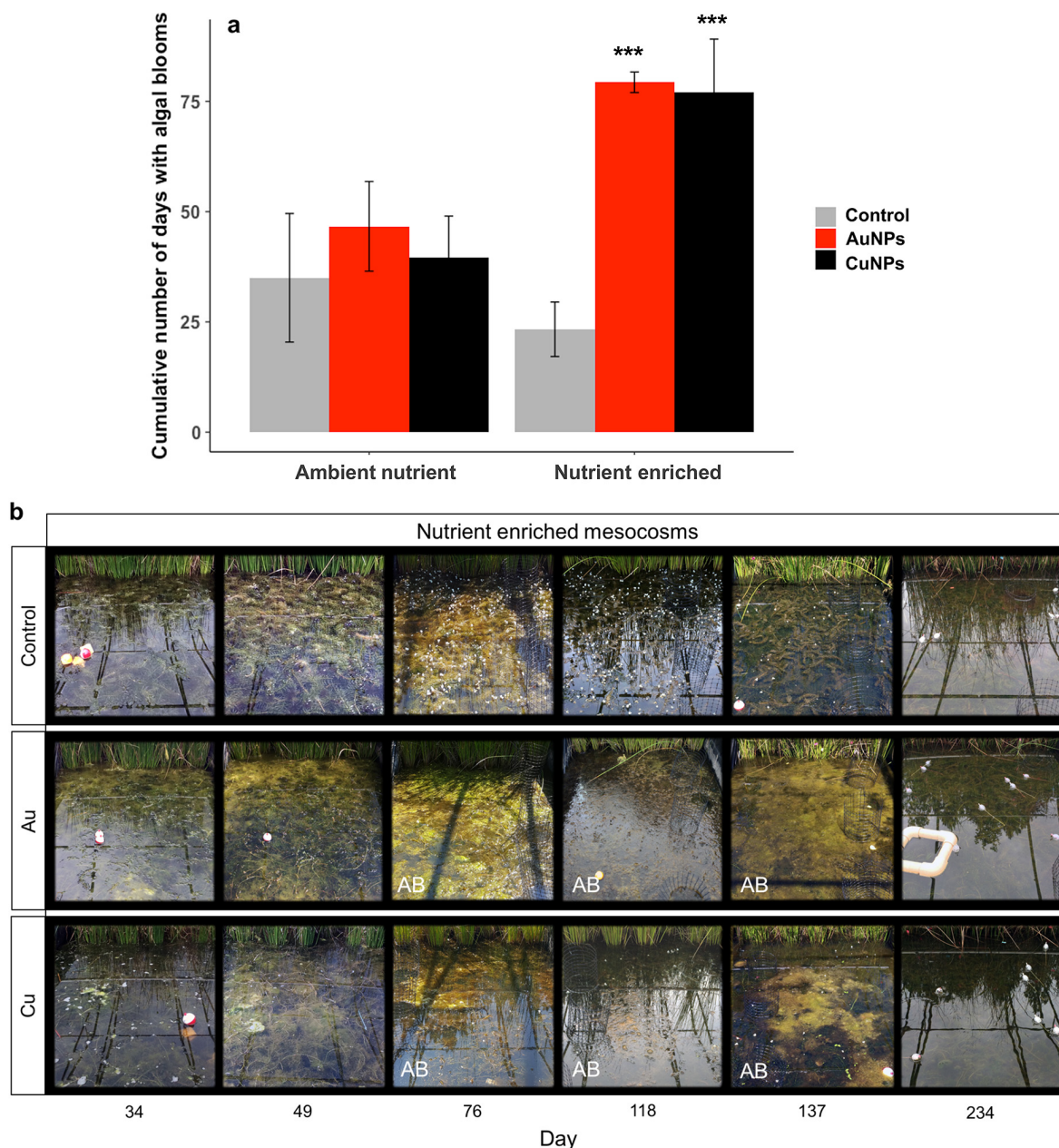


FIG. 3. (a) Cumulative number of algal bloom days in each treatment. Three asterisks denote significant ($P \leq 0.001$) difference between each NP treatment and the control at the same nutrient level. Values are mean $\pm$ 95% confidence intervals. (b) Photographs of the aquatic zone of three representative mesocosms in the nutrient enriched control, AuNP, and CuNP treatments at six time points during the experiment. Photographs of mesocosms experiencing a major algal bloom are indicated with the letters "AB." The small white forms that we can observe on Day 76, 117, and 138 in the control mesocosm are *Egeria densa* flowers.

only significantly affected by the AuNP-nutrient enriched treatment (Fig. 5). In this treatment, GPP was suppressed to near zero on 4 measurement dates (Day 155, 156, 191, 192; -91 to -94%), while ER was significantly reduced on one date (Day 155, -94%).

Physiological effects of Au and CuNPs and bioaccumulation in Egeria densa

In the AuNP-ambient nutrient treatment, photosynthesis rates (measured on new stems on Days 90, 193, and 269) were significantly reduced at all dates (-22% to -39% ,

Fig. 6), while in the AuNP-nutrient enriched treatment, photosynthesis was decreased only on Day 90 (-42% , Fig. 6). On Day 90, a reduction of the respiration rates was found in the nutrient enriched AuNP mesocosms (-63% , Fig. 6) and on Day 193 in the ambient nutrient AuNPs mesocosms (-63% , Fig. 6). In the mesocosms exposed to CuNPs, no effects were observed on *Egeria* photosynthesis, but respiration rates were significantly decreased on Day 90 (-80% , Fig. 6) in the nutrient enriched conditions.

On all destructive sampling dates, Au and Cu concentrations in *Egeria* stems (new growth) were significantly higher in plants collected from NP treatments than those from

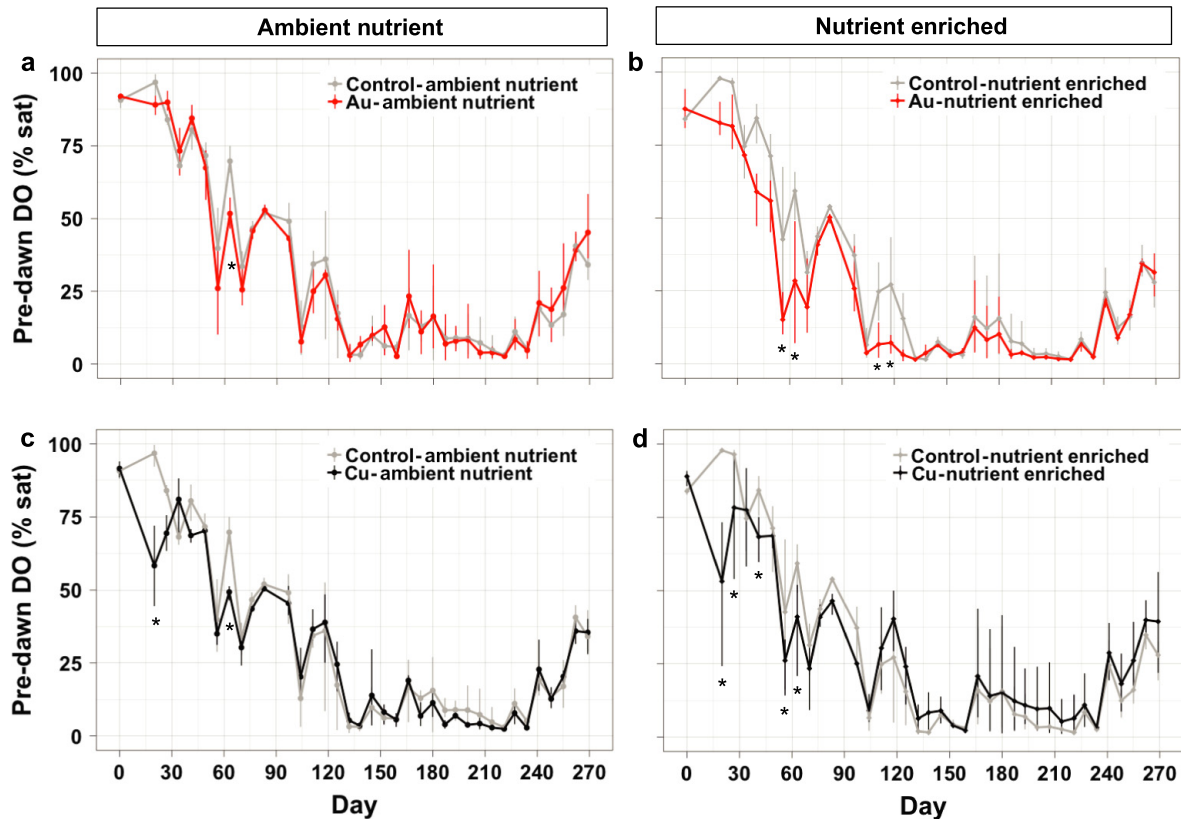


FIG. 4. Dissolved oxygen (DO) saturation (sat) at the surface water before dawn over the 9 months of the experiment. The different treatments are displayed in different panels along with their respective control treatment: (a) AuNP-ambient nutrient, (b) AuNP-nutrient enriched, (c) CuNP-ambient nutrient, (d) CuNP-nutrient enriched. Asterisks denote significant ($P \leq 0.05$) difference between NP treatment and control of each sampling date. Values are mean $\pm$ 95% confidence intervals.

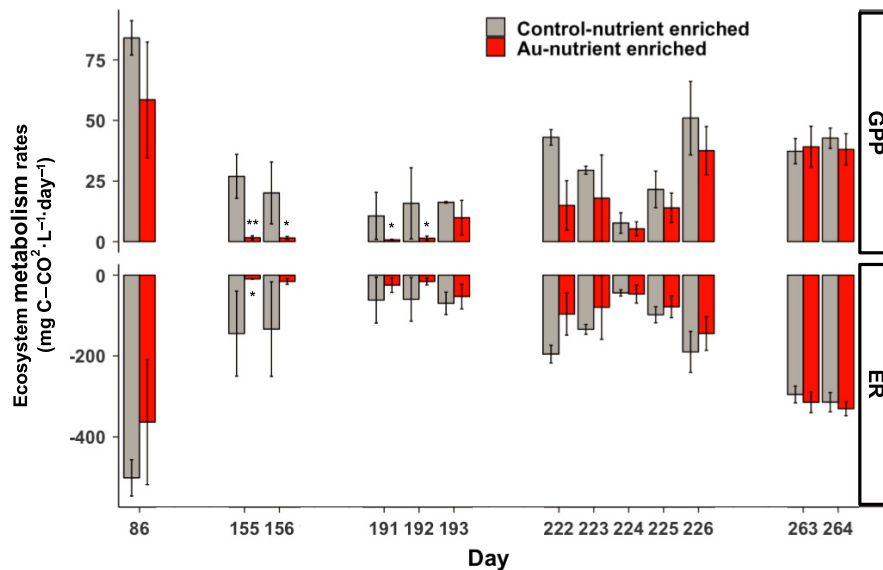


FIG. 5. Ecosystem metabolism: gross primary productivity (GPP, top) and ecosystem respiration (ER, bottom) in the AuNP and control-nutrient enriched treatments measured at discrete times between Day 86 and the end of the experiment. Asterisks denote significant ($*P \leq 0.05$; $**P \leq 0.01$) difference between NP treatment and control of each sampling date. Values are mean $\pm$ 95% confidence intervals.

controls ($P < 0.001$, Fig. 7). The highest concentration of Au was observed in *Egeria densa* stems on Days 90 and 193 with concentrations ranging from 137 to 298 mg/kg on

average (Fig. 7a), while the Cu concentrations averaged between 37 and 61 mg/kg during the experiment (Fig. 7b). The Au concentrations decreased significantly over time in

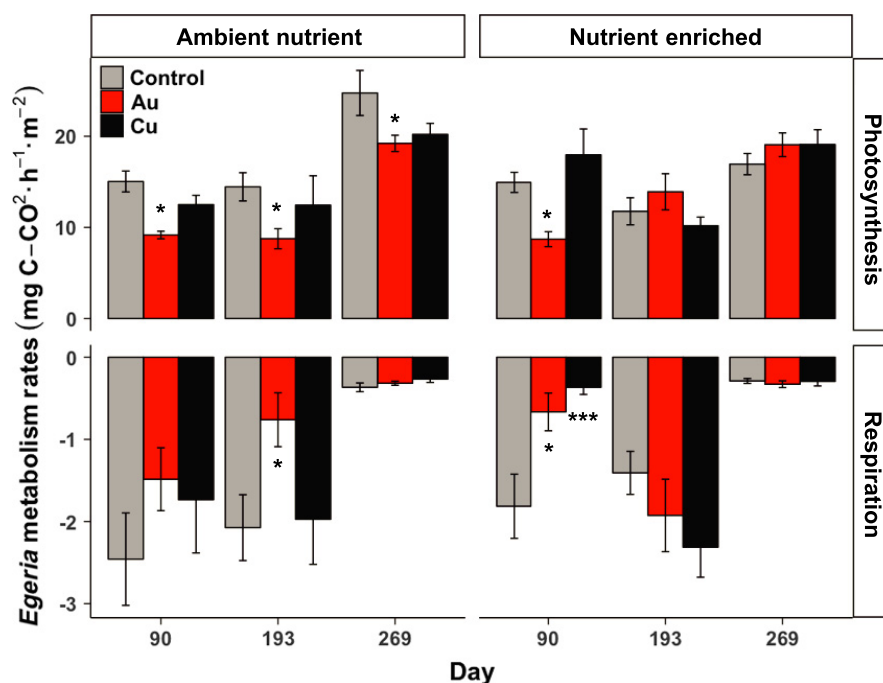


FIG. 6. *Egeria densa* stem photosynthesis (top) and respiration (bottom) rates in the ambient nutrient treatments (right) and in the nutrient enriched treatments (left). Asterisks denote significant (* $P \leq 0.05$; *** $P \leq 0.001$) difference between NP treatment and control at the same nutrient level of each sampling date. Values are mean $\pm$ 95% confidence intervals.

Egeria densa stems, especially between day 193 and 269. In both Ambient Nutrient and Nutrient Enriched conditions, Cu concentrations increased between Day 90 and 193 and then remained stable to Day 269.

During the same months, *Egeria densa* growth rates were measured in mesh columns placed in the water column and the highest growth rates in the controls were observed during the summer (Day 193, Fig. 8). The nutrient enrichment treatment did not significantly alter *Egeria* growth rates ($P = 0.82$). The only significant treatment effect observed was a 52% decline in *Egeria* growth rates in the AuNP-nutrient enriched on Day 193 (Fig. 8).

DISCUSSION

We found that when added together with nutrients, AuNPs enhanced the frequency and duration of algal blooms and this was accompanied by a reduction in macrophyte growth and photosynthesis, leading to extended periods of water column hypoxia and reduced ecosystem productivity. Adding CuNPs in combination with nutrient enrichment caused similar increases in algal blooms and reductions in dissolved oxygen, but these changes were not accompanied by a decline in ecosystem productivity. In the absence of nutrient enrichment, there were no consistent effects of AuNPs or CuNPs on primary producers and whole ecosystem behavior.

Larger impacts of gold nanoparticles than copper-based nanopesticide

We were surprised to discover that our “nanoparticle-tracers,” AuNPs, caused larger and more frequent ecosystem

responses than the Cu-based nanopesticide. The macrophyte response to AuNPs could result from a direct effect of the rapid Au accumulation into *Egeria densa* in the first 6 months of the experiment, while CuNPs showed both less bioaccumulation and non-significant effects on leaf-level *Egeria densa* physiology. These patterns in bioaccumulation and impact may be related to the contrasting behavior of CuNPs and AuNPs in the water column. The CuNPs used in this experiment had a larger initial size than the AuNPs, showed evidence of aggregation, but also rapidly dissolved in situ in mesocosm water (dissolution half-life time, $t_{1/2} \sim 8$ h) and dissolved Cu²⁺ was likely complexed by dissolved organic matter (Vencalek et al. 2016). In contrast, AuNPs were less aggregated in the water column (~ 10 nm) and are assumed to have low solubility under environmental conditions (Lee and Ranville 2012). The two metal NPs were dosed at different concentrations in the mesocosms, but our results still suggest that under these environmentally realistic conditions, the metal bioaccumulation rate in an aquatic plant was higher when exposed to the small and stable AuNPs than the much more soluble CuNPs.

We expected *Egeria* tissue metal concentrations to be similar at each harvest since the stems collected were about the same age (new growth) and were exposed to a similar water column NP concentrations. This expectation was met for CuNPs but not for AuNPs. Tissue Cu concentrations in *Egeria* was stable across the three harvest dates. However, in the AuNP treated mesocosms, the highest Au accumulation was measured on Day 90 and then tissue Au concentration in *Egeria densa* stems declined over the remaining six months of the experiment. This result raises a number of questions about the environmental variables that mediate AuNP uptake in plants. It is unlikely that these large decreases were

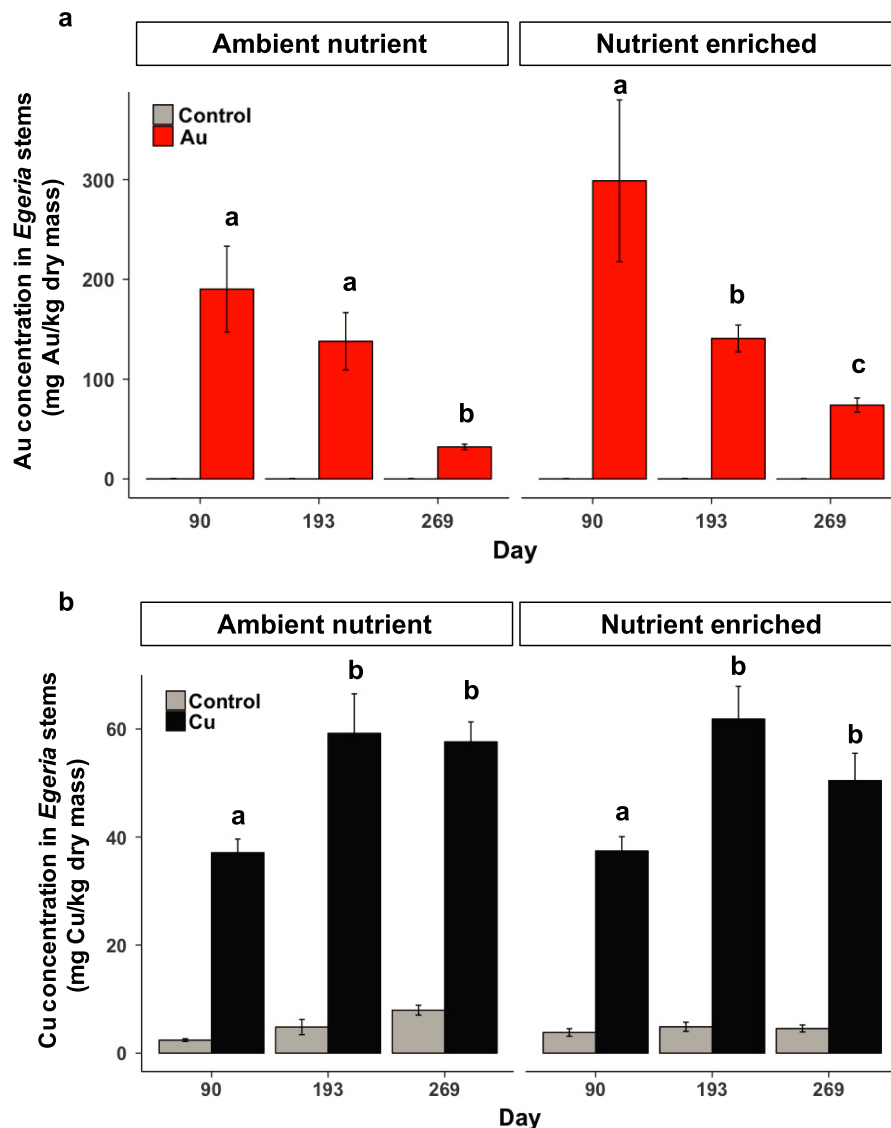


FIG. 7. Gold (a) and copper (b) concentrations (mg of metal per kg of plant, dry weight) in *Egeria densa* stems. Note that the y-axis scales are different between the two graphs. Different letters indicate significantly different dates in the same treatment. Within each nutrient level, treatment means of the three sampling dates annotated by the same letters are not discernible at $\alpha = 0.05$. Values are mean $\pm$ 95% confidence intervals.

explained by higher plant turnover or regulation of Au concentrations in tissue but they may have been controlled by an external factor sensitive to seasonal variations and/or NP dosing. We speculate that changes in the composition and capacity of the periphyton growing on *Egeria* stems could be responsible for the differences in bioaccumulation over time that we observed. Periphyton can act as a barrier to prevent contaminant uptake by the macrophytes (e.g., NPs strongly adsorbed to the biofilm) or could be involved in the facilitation of the uptake of NPs by mediating their dissolution or modifying NP aggregation and surface properties (Schwab et al. 2016). It is also possible that *E. densa* exudes a sulfhydryl rich metal binding protein, which could modulate the bioavailability of Au as it has been previously shown for AgNPs (Bone et al. 2012, Unrine et al. 2012).

We cannot resolve whether the higher tissue concentrations or a higher toxicity of AuNPs led to more severe effects of AuNPs than CuNPs in this experiment. In previous lab

based assays, metal NPs have been reported to reduce both algal and macrophyte growth and photosynthesis (Hoecke et al. 2013, Thwala et al. 2016). In these studies, researchers typically attribute these negative impacts on increases in oxidative stress or membrane damage caused by the NPs but the literature is scarce on the specific toxicity mechanisms of AuNPs in plants and algae. However, one study in *Caenorhabditis elegans* demonstrated that citrate coated AuNPs can cause unfolded protein response and endoplasmic reticulum stress by denaturing proteins that they come into contact with (Tsyusko et al. 2012).

Despite the known toxicity of Cu to aquatic organisms, Cu from CuNPs was less bioavailable and had less impact on autotrophs than AuNPs. The study of the toxicity and ecological impacts of AuNP has received less attention than other NPs, such as Ag-NPs and TiO₂-NPs, because of their lower industrial production rates. The use of AuNPs in pharmaceuticals is growing (bioimaging, gene/drug delivery,

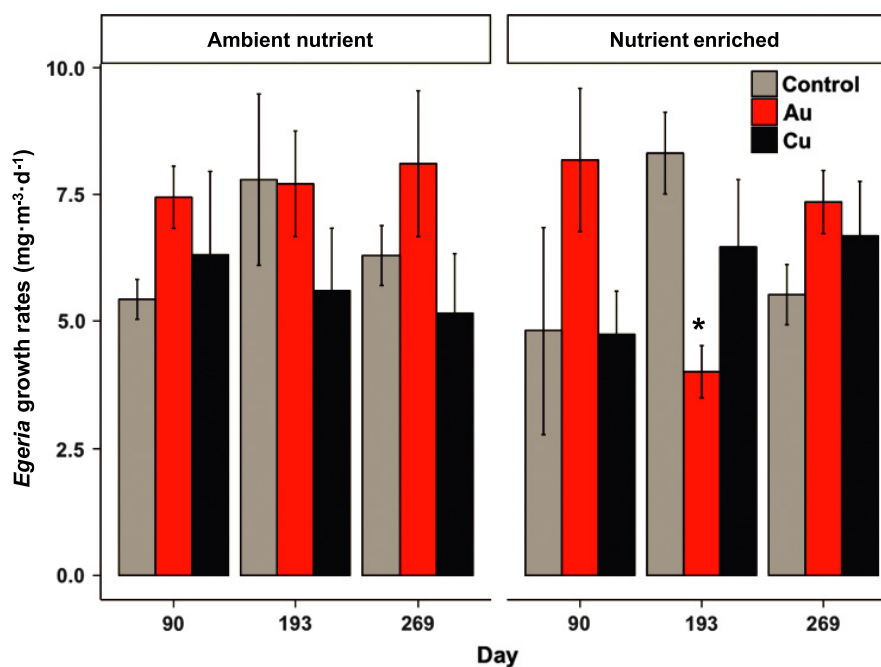


FIG. 8. *Egeria densa* growth rates in the Ambient Nutrient treatments (right) and in the Nutrient Enriched treatments (left). Asterisk denotes significant ($P \leq 0.05$) difference between NP treatment and control at the same nutrient level of each sampling date. Values are mean $\pm$ 95% confidence intervals.

phototherapy, home diagnostics; Ghosh et al. 2008, Rossi et al. 2016) but widespread environmental exposure to this contaminant is unlikely. The important implications of this study for future ecological research on contaminant impacts are the recognition that lab based toxicity studies can have limited power to predict ecological impact. What is needed is research that uncovers the mechanisms through which AuNPs can disrupt and alter ecosystem processes so that we can prevent environmental exposures of contaminants possessing similar traits.

Metal nanoparticle contaminants can intensify eutrophication in wetland ecosystems

We hypothesized that nutrient enrichment would alter the ecological impact of NPs and we indeed found multiple lines of evidence showing interactive effects between nutrient enrichment and NPs. We observed very limited effects of both AuNPs and CuNPs in the absence of nutrient enrichment, but when added together with nutrients, both NPs exacerbated the adverse effects of eutrophication in wetlands (algal blooms, hypoxia, and decreased productivity). This experiment suggests that nutrient status can greatly influence the ecosystem-scale impact of emerging contaminants, and that metal-based synthetic chemicals may be playing an underappreciated role in the global trends of increasing eutrophication (Heisler et al. 2008, Smith and Schindler 2009).

We found little evidence that nutrient enrichment directly affect NP properties, with neither NP aggregation nor surface charge being affected by nutrient enrichment. The rate of AuNP and CuNP assimilation into macrophyte tissues was unaffected by nutrient enrichment, suggesting that other unmeasured properties of the NPs (e.g., organic coatings,

surfacial chemistry) were not significantly altered by the nutrient regime. Together these results suggest that there are limited direct interactions between nutrient status and the bioavailability or surficial reactivity of these NPs. However, our results demonstrate that nutrient enrichment can alter organismal responses to NPs. Photosynthetic rates of individual *Egeria* plants were more suppressed by AuNPs in the absence of nutrient enrichment than under high nutrient conditions. Under low nutrient conditions, the energetic costs to take up nutrients might be more important and thus limiting the energy available to be invested to offset the effects of external stressors. This result supports the nutrient rescue hypothesis, that high nutrient availability allows individuals to invest in detoxification responses or compensate for the energetic costs of contaminant stress more easily (Leflaive et al. 2015, Aristi et al. 2016). Despite this finding at the individual plant scale, this “rescue” did not propagate to ecosystem scales where we consistently observed ecological responses to NP exposure only in the nutrient enriched treatments.

Under nutrient enriched conditions, AuNPs and CuNPs were associated with an increased frequency and magnitude of algal blooms in these experimental systems. These algal blooms led to decreases in dissolved oxygen in the water column and lowered primary productivity after 5 and 6 months of AuNP exposures (late spring–summer season). The extended algal bloom in the AuNPs-nutrient enriched treatment lead to 91%–94% declines in ecosystem GPP. We hypothesize that this pronounced response arises from relatively small disparities in the effects of our contaminant stressor on algae vs. macrophytes (Fig. 9). In the presence of the NPs, floating algae were better able to capitalize on the nutrients added in our nutrient enrichment treatments and thus

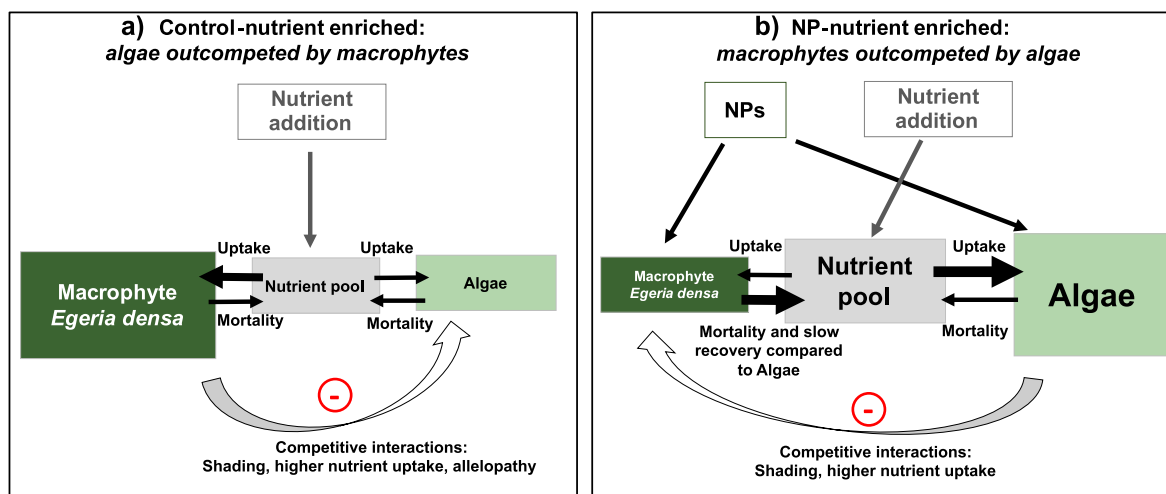


FIG. 9. Conceptual diagram representing our hypothesis of how nutrient availability in the water column and the competitive interactions between macrophytes and algae are modified under (a) nutrient enrichment alone compared to (b) nutrient enrichment combined with NPs additions.

could shade out their benthic competitors. We hypothesize that the 52% of the reduction we observed in macrophyte growth rates in the AuNPs-enriched nutrient treatment resulted from altered competitive interactions (in particular shading) rather than from direct AuNP toxicity.

There are many direct and indirect mechanisms that may explain how NPs and nutrients together caused the observed increased magnitude and duration of bloom conditions in the summer months of the experiment. In the first three months of the experiment, total N and DOC concentrations increased by 30–60% in the water columns of mesocosms in the NPs-nutrient enriched treatments compared to the nutrient only controls. These increases could be explained by mortality of the algae and macrophytes immediately after we initiated the contaminant exposure (previously observed by Roussel et al. 2007, Colman et al. 2014) or by an increase in consumer excretion rates in response to contaminant exposure (Taylor et al. 2016). Either mechanism could explain the internal eutrophication we observed in the presence of NPs, and could explain the observed intensification of the effects of our nutrient enrichment treatment. This compounding of internal mineralization with nutrient enrichment likely generated the appropriate conditions for the cyclical algal blooms observed in the NPs-nutrient enriched treatments.

An alternative explanation for the observed algal blooms is that the NP exposures selected for tolerant or resistant algal taxa that were then competitively superior to macrophytes under NP exposure scenarios. Since our mesocosms contained only a single macrophyte species, such community adaptation was not possible for these benthic plants. We speculate that the slower growth and recovery rates of macrophytes relative to algae would disadvantage these benthic autotrophs. Adding to the complexity of this competitive interaction, previous studies have documented that increased nutrient availability or NP stress can decrease the production of allelopathic secondary compounds (e.g., phenolics, flavonoids) by macrophytes (Richardson et al. 1999, Gross et al. 2007, Wang et al. 2011a). Such a shift in

resource allocation by macrophytes would further advantage their planktonic competitors. Altogether, these results suggest that early in the experiment, mesocosms exposed to nutrient enrichment and AuNPs saw a combination of an initial increase in nutrient availability and modifications in competitive interactions between algae and macrophytes (Fig. 9). This combination of direct and indirect effects of NPs likely contributed to the cascading effects that caused a suppression of ecosystem productivity four to five months later during algal blooms.

Our results provide an example of emerging contaminants having large ecosystem-level impacts on aquatic ecosystems, with both AuNPs and CuNPs causing higher frequency and magnitude of algal blooms. Our experiment also demonstrated that such impacts are highly context dependent, as we only observed these ecosystem impacts when NPs were superimposed with nutrient enrichment. Studying the interactive effects of multiple stressors in realistic biological systems is crucial, as this work again confirms that their consequences on ecosystem structure and function are very difficult to predict. While many questions about the mechanisms responsible for these effects remain unanswered, our findings provide evidence that synthetic chemicals such as metal NPs can exacerbate aquatic ecosystem eutrophication.

CONCLUSIONS

Perhaps the most fascinating outcome of this experiment was the discovery that the addition of small amounts of a single synthetic nanoparticle could in fact catalyze the transformation of our wetland ecosystems from clear waters to turbid waters with large floating algal mats. The increasing frequency and magnitude of algal blooms is expected to increase with climate change but the role of synthetic chemicals, like metal pollutants or pharmaceuticals in these events is not currently considered (Heisler et al. 2008, O'Neil et al. 2012). The effects observed here might be specific to these metal NP contaminants but we postulate that any chemical contaminant that causes differential stress for algae relative

to macrophytes or modifies nutrient availability has the potential to intensify eutrophication (Fig. 9). This experiment shows that large declines in the rates of ecosystem productivity and changes in the relative dominance of different autotrophs can result from sub-lethal effects at very low concentrations. We cannot determine from the current experiment whether AuNPs reduced the ability of macrophytes to compete for nutrients and produce allelochemicals or if AuNPs selected for bloom-forming algal species. Further work should explore how each of these competitive interactions may be affected by contaminant exposure.

This study also shows that the ability to detect these ecosystem level effects is highly sensitive to the timing of experimentation and sampling. The most marked treatment responses were observed during the warmest months of the year and dissipated in the subsequent autumn, even as we continued to add both nutrients and NPs to the mesocosms. This suggests that the effect of these materials, and perhaps many conventional contaminants, may be most pronounced and most measurable during periods of intense competition, which in many aquatic ecosystems are characterized by warm temperatures, high rates of grazing pressure, and high productivity by both macrophytes and algae.

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SUPPORTING INFORMATION

Additional supporting information may be found online at: <http://onlinelibrary.wiley.com/doi/10.1002/eap.1742/full>

DATA AVAILABILITY

Data available from Figshare: <https://doi.org/10.6084/m9.figshare.6118553>