Phytoremediation potential of switchgrass (*Panicum virgatum*), two United States native varieties, to remove bisphenol-A (BPA) from aqueous media

Jacob C. Phouthavong-Murphy1,2, Alyssa K. Merrill1,2, Stephanie Zamule3, David Giacherio3, Beverly Brown1, Carol Roote1 & Padmini Das1*

Plastic wastes burdening Earth’s water and accumulating on land, releasing toxic leachates, are one of the greatest global threats of our time. Bisphenol-A (BPA), a potent endocrine disrupting compound, is a synthetic ingredient of the polycarbonate plastics and epoxy resins used in food containers, cans, and water bottles. Bisphenol-A’s rising concentrations in the environment require a sustainable alternative to current removal practices, which are expensive and/or ecologically unsafe. Switchgrass offers a safe alternative. To investigate its potential for BPA removal, two United States native switchgrass varieties were tested in hydroponic media. Results show minimal hydrolysis and photolysis of BPA over 55 days, confirming its persistence. Both generic and heavy metal switchgrass exhibited statistically significant (*p* < 0.0001) BPA removal (40% and 46%, respectively) over approximately 3 months, underscoring switchgrass’s effectiveness for BPA removal. Significantly higher (*p* < 0.005) BPA accumulation in roots than shoots and nonsignificant variances in biomass, chlorophyll (*p* > 0.19), and peroxidase between BPA-treated and untreated plants indicates substantial BPA tolerance in both varieties. Kinetic parameters of BPA removal and translocation factors were also determined, which will inform the design of BPA removal phytotechnologies for a variety of soil conditions, including landfills where BPA accumulation is greatest.

Discovered in the 1890s, as a potential drug, 4,4′-(propane-2,2-diyl)diphenol (bisphenol-a, BPA) was later found to be useful in plastic production1. Manufacturing companies use large quantities of BPA in polycrylamide, which is distributed to the public in food containers, baby bottles, and a variety of other consumer products2. In 2012, worldwide BPA production reached an estimated 6.5 million tons per year, with predicted increases of 4.6% annually until 20193,4. In 2015, an estimated 7.7 million tons of BPA were produced, representing an even greater increase (6% per year) than initially predicted5. By the end of 2022, production is estimated to be 10.6 million tons5. Products containing BPA do not remain in use indefinitely, it is estimated over 450 tons of BPA are released into the environment annually6.

In 2012, the United States (U.S.) Food and Drug Administration banned BPA in infant products due to controversy over developmental toxicity risks7,8. However, BPA is still utilized in the production of a variety of plastics for other consumer uses, including polyvinyl chloride, polycarbonate, and thermal receipt paper9. Despite its widespread use, BPA has been linked to numerous adverse health outcomes, including infertility, tumor growth, and obesity10–12. Bisphenol-A exposure occurs, since the potential exists for BPA to leach out of disposed BPA containing products into the environment through degradation, posing a risk to humans who accidentally consume BPA through food and/or drinking water and organisms who come into contact with polluted areas13–15.

Cellular function disruption due to BPA has been reported at levels as low as 0.23 ng/L, with the solubility of BPA approximately 100 mg/L16. A National Health Survey conducted by the U.S. Center for Disease Control and Prevention found over 95% of human urine had BPA levels above 10 ng/L16,17. At present, no maximum

1Nazareth College of Rochester, 4245 East Ave, Rochester, NY, 14618, USA. 2These authors contributed equally: Jacob C. Phouthavong-Murphy and Alyssa K. Merrill. *email: pdas8@naz.edu
permissible concentration of BPA in water has been established by the U.S. Environmental Protection Agency, though the intake limit is set at 0.05 mg/kg/day. As the production of plastics containing BPA continues to rise, the environmental concentrations of BPA most likely will increase concurrently. With landfill leachate concentrations presently documented as high as 17 mg/L, exposure to dangerous levels of BPA is expected to escalate as more BPA containing products are released into the environment.

While the potential exists for degradation of environmental BPA through hydrolysis or photolysis, the quantity of BPA eliminated through degradation in pure water is not significant. Rather degradation occurs only in the presence of humic substances, due to reactive oxygen species found in conjunction with humic substances. A previous study found the half-life of BPA in surface water ranged from 66 hours to 160 days from photolytic degradation, due to water condition variation. Further, the degradation observed over 160 days did not prevent BPA from traveling through the water system and harming aquatic organisms. Thus, additional methods of BPA removal must be found.

Researchers seeking methods to remove environmental BPA face a multitude of challenges. Widely utilized remediation techniques are expensive and not ecologically viable. Chemical degradation using nanotechnology, TAML (Tetra-amido macroyclic ligand)/H₂O₂ complex and other methods have demonstrated only minimal success, and the potential toxicity of these chemicals on the environment is a concern. Phytoremediation shows promise as a sustainable alternative to remove BPA from contaminated soil and water systems in a cost-effective, ecologically-friendly, and socially-acceptable way. Phytoremediation has been successfully utilized for the removal of a number of other environmental contaminants, including tetracycline, polychlorinated biphenyl, and trinitrotoluene. Organisms previously shown to take up BPA include aquatic plants: Azolla filiculoides (water velvet), Potamogeton illinoensis (Illinois pondweed), Lemna species (duckweed), Spirogyra (algae), terrestrial plants: Portulaca oleracea (common purslane), Dracaena sanderiana (lucky bamboo), Fragaria vesca (wild strawberry), Vicia faba (broad bean), Oryza sativa (Asian rice); and fungi: Fusarium sporotrichioides, Fusarium moniliforme, Aspergillus terreus, and Aspergillus nidulans. Although a number of plants capable of up taking BPA have been identified, few meet the criteria for a landfill phytoengineering technique required in the U.S.

Environmental BPA levels are much higher in landfill leachate (17 mg/L) compared to surface water (12 µg/L), making it essential to develop remediation technologies to extract BPA from landfills, thus preventing further leaching into the surrounding soil and groundwater, and greater exposure to organisms. Phytoremediation using a BPA-tolerant plant offers a promising remediation method, however, selecting an appropriate plant species to be applied in a landfill environment poses a number of challenges. Native varieties are preferable to avoid risks to environmental health caused by invasive species. Edible plants should be bypassed to reduce the risk of further accumulation of BPA in the food chain. The plant should be terrestrial and capable of tolerating other environmental contaminants frequently present in landfill conditions. Prior to implementation, the non-edible, terrestrial plants native to the area of interest should be thoroughly studied to ascertain the plant’s ability to tolerate BPA toxicity and remove BPA from contaminated systems.

Thus a phytoengineering technique using native, non-edible plants to extract BPA from landfills in the U.S. must be developed. Most BPA-tolerant plants currently reported are either aquatic, edible, or not native to the U.S., and therefore do not satisfy all criteria for successful landfill application in the U.S. The search for a suitable plant capable of thriving in landfill conditions in the U.S. led to the study of switchgrass (Panicum virgatum); a terrestrial, perennial, high biomass grass, native to all of the U.S., except California and the Pacific Northwest. Current uses of switchgrass include erosion control, wild life habitat, fodder, ornamental purposes, paper pulp, and biofuel. Switchgrass has been previously shown to remove heavy metals including lead, cadmium, zinc, chromium, as well as organic contaminants like atrazine. The phytotolerance of switchgrass to a variety of toxic contaminants adds promise for switchgrass’ potential survival in contaminated soils and landfills, where a wide range of pollutants are frequently present in variable concentrations. Additionally, switchgrass is hardy and capable of surviving in a wide range of soil types and climates.

The present investigation aimed to compare the relative potential of two switchgrass varieties to extract BPA from aqueous media at a concentration relevant to the maximum BPA concentrations reported in U.S. landfills. This appears to be the first attempt to investigate the potential of BPA removal using switchgrass, whose effectiveness in removing other toxic chemicals from contaminated systems and phytotolerance against them has been documented. Based on previous research, both varieties of switchgrass were expected to remove a significant amount of BPA from an aqueous media and translocate it into the stalk, making switchgrass a suitable solution to environmental BPA pollution. Investigating the relative effectiveness of BPA removal by two varieties of one plant species is also a novel approach in phytoremediation and will contribute valuable insight to implementation, as some varieties of one plant could be native to one geographical area and others may not. The kinetics of BPA toxicity and remove BPA from contaminated systems.

Results

Hydrolysis and photolysis. Hydrolysis and photolysis of an environmentally relevant (5 mg/L) concentration of BPA solution without plants were tested in the absence (hydrolysis) and presence (photolysis) of light, respectively. In minimal BPA degradation (hydrolysis = 0%, photolysis = 14.9%, control = 11.4%). No significant differences (hydrolysis p = 0.1398, photolysis p = 0.7482) were detected between control and hydrolytic/photolytic samples over the experimental duration of 55 days (Fig. 1), demonstrating BPA does not significantly degrade when exposed to hydrolytic and photolytic conditions.
BPA removal from aqueous media by *panicum virgatum*. Two switchgrass varieties were tested for their ability to remove BPA (20 mg/L) from aqueous media in a greenhouse setting. Both generic and heavy metal switchgrass varieties removed a significant amount of BPA in 84 days ($p < 0.0001$, Fig. 2) and 86 days ($p < 0.0001$, Fig. 3), respectively, compared to no-plant controls. Bisphenol-A concentrations were reduced by 40.4% in the presence of generic switchgrass and 46.1% in the presence of heavy metal switchgrass. Controls exhibited an insignificant decline of BPA at 5.4% (Fig. 2) and 10.6% (Fig. 3) for generic and heavy metal varieties, respectively.

**Kinetic modeling.** Both varieties of switchgrass demonstrated pseudo-second-order removal kinetics, exhibiting very strong $R^2$ values, 0.9794 for the generic variety and 0.9927 for the heavy metal variety (Fig. 4). Comparison of the initial rate constants (day 0 to day 46) of the two varieties showed the heavy metal variety rate constant was significantly higher ($p < 0.001$). Although both generic and heavy metal varieties removed a similar fraction of BPA after 84 to 86 days, respectively, the heavy metal variety showed higher initial BPA removal.
kinetics than the generic one (Supplementary Fig. S1). As evident from the $t_{25}$ and $t_{40}$ values, 25% of BPA was removed from the aqueous solution around day 15 for the heavy metal variety, whereas the generic variety did not reach this level until day 46. Then 40% of BPA removal occurred by day 46 in the heavy metal variety, whereas the generic variety required 84 days to achieve the same level of removal.

Biochemical analyses. Phytotoxic effects of BPA on switchgrass were evaluated through chlorophyll content and biomass as a function of BPA exposure. Chlorophyll A, B, and total chlorophyll content were measured after the final day of BPA exposure and compared with unexposed controls. Chlorophyll levels between the controls and BPA-treated plants in both varieties were not statistically significantly different (generic $p = 0.7348$, heavy-metal $p = 0.1984$) (Fig. 5).

Biomass of generic switchgrass was measured in plants on day 0 and day 84 after BPA exposure and compared with unexposed control plants to evaluate the health of the plant post-BPA exposure. The difference between the controls and exposed plants’ biomass before and after the experimental duration is approaching statistical significance ($p = 0.0568$). The mean comparison showed an increase in weight for the unexposed controls and the
The maintenance of a steady weight for BPA-exposed plants over the course of the experiment (Fig. 6). The growth inhibition percentage was 77.9% when generic switchgrass was subjected to 20 mg/L of BPA for 84 days.

Bisphenol-A accumulation in the plant tissues was significantly higher in the roots as compared to the shoots for both the generic \( (p = 0.0021) \) and heavy metal varieties \( (p < 0.0001) \) (Fig. 7). Heavy metal switchgrass accumulated significantly more BPA in roots than the generic variety \( (p < 0.001) \). No difference was observed in shoot accumulation between either variety \( (p = 0.3728) \).

As the mechanism of BPA metabolism might be through the peroxidase pathway, peroxidase activity between BPA-treated and untreated tissues of the generic switchgrass is compared in the roots and shoots. No significant difference \( (p = 0.0677) \) was detected between control and experimental plant samples, suggesting the utilization of a different BPA metabolic pathway in switchgrass (Fig. 8). Peroxidase activity in the heavy metal variety was not tested, as the likelihood peroxidase is the method of degradation in one variety of switchgrass and not the other was determined to be minimal.

**Discussion**

Bisphenol-A's persistence and the lack of photolytic and hydrolytic degradation found in the current study supports previous reports, where BPA was found not to be subject to these effects in pure water systems. Previous studies found BPA degradation under photolytic conditions when humic substances are present, although complete degradation has never been achieved. In the presence of humic substances, the half-life of BPA has been reported to be between 66 hours and 160 days, depending on the water conditions, when subjected to photolytic degradation. Without humic substances, the present experiment, concurrent with Im and Loffler's findings, showed nonsignificant degradation of BPA in photolytic or hydrolytic conditions over 55 days. These findings reaffirm the environmental persistence of BPA and the potential global risk it poses from rising environmental concentrations due to disintegrating plastics in soil, water, and landfills. Studies thus far establish a need to explore sustainable BPA removal technologies from any contaminated media. While phytoremediation shows promise as a sustainable alternative, global application of a BPA-tolerant plant could pose potential harm to the environment if the plant is invasive to the geographical area of implementation. Moreover, using edible plants would increase the risk of exposure owing to bioaccumulation. Thus, native varieties of non-edible plants should be studied on a regional basis to allow for a phytoengineered BPA removal processes to be designed to address the specific needs of the area.

Two varieties of switchgrass, native to the U.S., except for California and the Pacific Northwest, were investigated to evaluate the potential for a phytoengineering technique to remove BPA from disintegrating plastics in soil, water, and landfills to prevent leaching into surrounding environments. Bisphenol-A is still utilized in...
almost every major plastic production facility in the U.S., though exposure is known to lead to adverse health effects in humans\(^2\). To date, BPA is found in food containers, medical supplies, and a variety of other products that humans regularly come into contact with\(^2\). As a result, BPA concentrations in landfill leachate have risen to levels as high as 17 mg/L\(^{18}\). With BPA still currently in use, this level will likely continue to increase. Many plants are capable of removing BPA from aqueous media, but none meet the requirements for a phytoengineering technique in the U.S. The most promising plant tested was the broad bean which removed almost all BPA after 21 days when exposed to 50 mg/L\(^{31}\). Unlike the broad bean, switchgrass is inedible to humans, decreasing the likelihood of further human BPA exposure. Recent studies have shown a number of photosynthetic organisms native to the U.S. are capable of removing aqueous BPA, including: water velvet (80% removal, 20 days), Spriogyra (95% removal, 20 days), and duckweed (96% removal, 20 days), but these aquatic organisms would not survive in soil or landfill environments due to inadequate water supply\(^{32,38}\). As a terrestrial, high-biomass, fast-growing, perennial, native grass, switchgrass provides a broader scope of implementation to remove BPA from soil, shallow lagoons, and landfill leachate by creating phytoengineered constructed wetlands surrounding the landfill area.

When analyzing the phytoremediation potential of a plant species, understanding the kinetics of contaminant removal are crucial for designing a phytoengineered remediation process. Both switchgrass varieties showed pseudo-second-order BPA removal from aqueous media. Although the initial kinetics differed between the varieties, the plants showed similar rates of removal after their respective experimental time periods of 84 and 86 days. While a prior study described a kinetic model of the removal of BPA by duckweed and Spriogyra\(^{32}\), BPA kinetics of removal by a terrestrial plant have not been studied. Establishing kinetic parameters of BPA removal from aquatic media is a positive preliminary step towards the optimization of implementation factors in the phytoengineered BPA removal processes. These parameters will aid in calculating the required plant concentration at

---

**Figure 7.** BPA Concentration in Two Plant Tissues (mg/kg) in Generic (A) and Heavy Metal (B) Switchgrass. (A) Data are expressed as mean (n = 3) ± one standard deviation. Mean comparison was carried out by Tukey-Kramer HSD Test and shown in the indented table. (B) Data are expressed as mean (n = 3) ± one standard deviation. Mean comparison was carried out by Tukey-Kramer HSD Test and shown in the indented table.

**Figure 8.** Peroxidase Activity (mU/mg) in Generic Switchgrass as a Function of BPA Treatment. The graph shows the difference in peroxidase level between the plants not exposed to BPA and the plants with 20 mg/L BPA exposure. Data are expressed as mean (n = 6) ± one standard deviation. Mean comparison was carried out by Tukey-Kramer HSD Test and shown in the indented table.
given contaminant loads and establishing a balance between plant and BPA uptake. Similar kinetic parameters of BPA removal by switchgrass from soil systems need to be determined and compared with the hydroponic study in future. Moreover, additional investigations of other kinetic parameters of BPA translocation, transpiration, and its biotransformation to other metabolites will provide valuable insight to the phytotolerance of BPA in switchgrass varieties.

Switchgrass showed slower BPA removal rates compared to plants in previous studies providing a probable explanation for the higher BPA tolerance compared to other plants at similar or greater concentrations\(^3\)\(^1\),\(^3\)\(^2\),\(^3\)\(^4\),\(^3\)\(^6\),\(^3\)\(^8\),\(^3\)\(^9\),\(^4\)\(^0\),\(^4\)\(^1\). Chlorophyll levels, a marker of plant-health, remained unchanged compared to the controls in the BPA-treated heavy metal plants, as opposed to the generic variety, which showed slight phytotoxic symptoms. Other studies assessing the chlorophyll content of algae exposed to BPA, such as Monoraphidium braunii and Stephanodiscus hantzschii, found these exhibited phytotoxic symptoms\(^4\)\(^5\),\(^4\)\(^6\). Switchgrass withstands the effects of BPA with minimal phytotoxic symptoms, showing promise in a long-term phytoengineering technique.

Previous studies have also compared the change in growth over the experimental duration. Lower concentrations of BPA hinder growth in both Monoraphidium braunii (10 mg/L, 4 days) and Stephanodiscus hantzschii (15 mg/L, 16 days) with a growth inhibition of 89.23% in the latter\(^4\)\(^5\),\(^4\)\(^6\). Higher concentrations previously tested were found to affect growth in Chlorella fusca (Chlorophyta) (18 mg/L and 41 mg/L, 7 days), Loliun perenne (perennial ryegrass) and Raphanus sativus (radish) (46 mg/L, 16 days), and water velvet (50 mg/L)\(^3\)\(^8\),\(^3\)\(^9\),\(^4\)\(^0\),\(^4\)\(^1\). Both perennial ryegrass and radish had approximately 50% less weight than the controls after 16 days\(^4\)\(^2\). In contrast, the broad bean, one of best BPA-extracting plants reported at present, did not exhibit significant weight difference as compared to the unexposed controls with BPA concentration as high as 50 mg/L after 21 days\(^4\)\(^3\). Similar results were obtained in the current study, with switchgrass showing minimal phytotoxic effects and nonsignificant biomass degradation at 20 mg/L BPA over 84 days. These results indicate switchgrass has higher BPA tolerance than most other species of organisms tested, potentially due to the detoxification mechanism(s) resulting in higher efficiency of BPA removal from contaminated media.

Analysis of plant tissues showed much higher BPA accumulation in the roots compared to the shoots. The root cells may have a better coping mechanism not present in the shoot cells or BPA translocation to the shoots may have been prevented to avoid further phytotoxicity. Proximity of the roots to the BPA contaminated solution could also contribute to this difference. When contaminants are translocated into the shoot harvesting the plant for proper disposal is easier, but the likelihood of BPA transfer is higher if the plant were eaten. Since switchgrass is sometimes used as a forage crop, animals may eat the shoots, but if BPA accumulation into the shoots is minimal, the risk is low.

Enzymes including peroxidases and polyphenol oxidases in ring cleavage reactions have been implicated as contributors in the break down of BPA\(^4\). Peroxidase concentration in the roots and shoots of the generic variety switchgrass were measured in an effort to determine a potential mechanism for BPA tolerance. As the peroxidase analysis showed no difference in any tissue type for the controls and BPA-exposed samples, the likelihood this enzyme is responsible for BPA tolerance in switchgrass is low. \(\beta\)-glucosylation has been previously identified as the pathway that metabolizes BPA in Tobacco Bright Yellow-2 cells, which could be an alternative mechanism utilized by switchgrass\(^4\)\(^8\). Studies are underway to identify a pathway for BPA metabolism and to identify BPA-induced changes in the metabolic profile of switchgrass. Alternatively, if a mechanism for degradation cannot be identified within the switchgrass, BPA may be sequestered within the plant tissue. If sequestration is occurring the tissues may be burned to eliminate the BPA. BPA elimination is possible from the sealed burning of plant matter, allowing for the concentrated collection of BPA or complete combustion breakdown of BPA once the plants are removed from the contaminated environment\(^4\)\(^9\).

Two non-edible varieties of switchgrass, native to most of the U.S., were evaluated as phytoremediation agents capable of extracting BPA from contaminated landfills. Switchgrass, a high-biomass, fast-growing, perennial, native grass, which is environmentally stable and BPA stress-tolerant under long-term exposure, offers a promising phytoremediation strategy\(^4\)\(^0\),\(^4\)\(^1\). Removal efficiencies of both switchgrass varieties were not statistically different, indicating either may be utilized for BPA removal based on availability and cost-effectiveness. The slower initial kinetics of BPA removal in the generic variety may help the plant adapt to BPA-induced stress, resulting in the minimal phytotoxicity symptoms observed at continuous BPA exposure of 20 mg/L over approximately three months. Peroxidase did not show a notable increase in activity in the root and shoot tissues of switchgrass, highlighting the need for further investigation to determine the biochemical mechanisms of BPA tolerance in switchgrass.

Current results are highly promising and have paved the way for an ongoing greenhouse study to fully characterize switchgrass’ potential BPA extraction capacity from soil systems. Once the soil system study is completed, a pilot experiment will determine the feasibility of using switchgrass in the landfill environment. Landfill leachate will be collected and filtered through a constructed wetland where switchgrass is used to reduce the BPA concentration before the leachate is released into the environment. Kinetic parameters will be compared to the current aqueous study, where BPA was fully available for uptake by switchgrass, and the ongoing soil study, where BPA uptake is fractional, relying on the retention and release of BPA as a function of soil properties. The comparative data will provide valuable insights into optimization of a phytoengineering technique to ensure maximum removal of BPA from contaminated landfills and to achieve implementation of this technique in the U.S. to reduce the risk of BPA exposure.

**Materials and Methods**

**BPA analysis.** An Agilent Technologies (Santa Clara, CA) 1260 Infinity High Performance Liquid Chromatography (HPLC) was utilized to analyze aqueous samples for BPA concentration. The mobile phase of 30% water, 70% acetonitrile was run through an EC-C18 2.7 um Poroshell 120 column (4.6 mm × 50 mm) (Agilent Technologies). Injection volume was 10μL with a total run time of 4 minutes, analyzed at 228 nm (BPA
was conducted in Summer 2016 (n = 75%) to simulate an average summer in New York state. The generic variety long-term exposure experiment conducted under minimal light exposure with 0.2 g of plant tissue. A solution of 80% acetone was added to the tissue type were ground using a mortar and pestle while adding liquid nitrogen (Jackson Welding, Rochester, NY) until the tissue was sufficiently frozen. The remaining tissue was stored at −80 °C. Chlorophyll analysis was conducted in Summer 2016 (n = 3) and the heavy metal variety long-term exposure experiment was conducted in Summer 2017 (n = 10). Plants were acclimatized to half-strength Hoagland’s solution over seven days. Hoagland stock solution (100 ×) was composed of 13.52 g/L KH₂PO₄, 50.6 g/L KNO₃, 118.0 g/L Ca(NO₃)₂·4H₂O, and 40.0 g/L MgCl₂·6H₂O. Upon completion of acclimatization, 20.0 g of wet plant biomass were exposed to 1 L of 20 mg/L BPA (in 2% acetonitrile with ½ Hoagland’s Solution as the diluting agent), prepared from the same stock as mentioned above, for 84 days (generic variety) and 86 days (heavy metal variety) respectively. Each plant was placed in a one liter Erlenmeyer flask for the duration of the experiment. Controls included no-plant with BPA solution, no-plant without BPA, and plant without BPA. Over the course of the experiment, samples taken were filtered through a 0.45 μm PTFE syringe filter and analyzed by HPLC, as previously described. An analysis for t₂₅ and t₄₀ was conducted by identifying the time at which 25% and 40% BPA uptake were achieved. After the study was completed, the weight was recorded for a biomass analysis using a Fisher Scientific Scale SLF303 (Max: 300 g, d = 0.001 g) (Hampton, NH).

Biochemical analyses. Plants were separated at the base of the shoot, frozen at −80 °C, and pieces of each tissue type were ground using a mortar and pestle while adding liquid nitrogen (Jackson Welding, Rochester, NY) until the tissue was sufficiently frozen. The remaining tissue was stored at −80 °C. Chlorophyll analysis was conducted under minimal light exposure with 0.2 g of plant tissue. A solution of 80% acetone was added to the 0.2 g to obtain a final volume of 2 mL. The solution was vortexed and centrifuged at 10,000 rpm for 15 minutes at 4 °C using an Eppendorf 5424 R (Hamburg, Germany). The supernatant was filtered through a 0.45 μm PTFE syringe filter and absorbance measured at 645 nm and 663 nm on a ThermoFisher Genesys 20 Spectrophotometer (Hampton, NH). Chlorophyll a, chlorophyll b, and total chlorophyll were calculated according to the equations in Sunkar and Peroxidase was analyzed with an Invitrogen Molecular Probes Amplex Red Hydrogen Peroxide Peroxidase Assay Kit (ThermoFisher Scientific, Waltham, MA), according to the manufacturer’s instructions and summarized herein. Ground tissue was combined with assay buffer in a ratio of 0.01 g tissue/500 μL buffer. The solution was vortexed, then centrifuged at 15,000 rpm for 10 minutes using an Eppendorf 5424 R. A volume of 50 μL was removed from the supernatant and analyzed according to the kit instructions. Analysis was performed using a Bio-Rad iMark Microplate Reader at 570 nm.

Kinetic modeling and statistical analysis. Outlier analysis and two-way ANOVA, for statistical analysis of multiple groups, followed by mean comparisons using Tukey-Kramer for post hoc analysis were conducted with JMP Version 13. Letters within all graphs indicate differences between the means of replicates. Points with the same letter are not statistically different (p > 0.05). All tests were conducted with 95% confidence. Kinetic parameters such as reaction rates for BPA removal from aqueous media were determined following Equation 4 as described by Robati.

Data availability The datasets generated during and/or analyzed during the current study are available from the corresponding author on reasonable request.

Received: 15 July 2019; Accepted: 3 December 2019; Published online: 21 January 2020.
References

1. Rochester, J. R. Bisphenol A and human health: a review of the literature. Reproductive Toxicol. 42, 132–155 (2013).
2. Vogel, S. A. The politics of plastics: the making and unmaking of bisphenol a “safety.” American J. Public Health 99(Suppl 3), S559–S566, https://doi.org/10.2105/ajph.2008.159228 (2009).
3. Corrales, J. et al. Global assessment of bisphenol a in the environment. Dose Response 13(3) (2015).
4. Im, J. & Loffler, F. E. Fate of bisphenol A in terrestrial and aquatic environments. Environ. Sci. Technol. 50, 8403–8416 (2016).
5. Industry Experts. Bisphenol A - a global market overview. Retrieved from: http://industry-experts.com/verticals/files/articles/pc021-bisphenol-a-a-global-market-overview.pdf (2016).
6. U.S. Environmental Protection Agency (USEPA). Bisphenol a action plan. CASRN 80-05-7. Retrieved from: https://www.epa.gov/sites/production/files/2015-09/documents/bpa_action_plan.pdf (2010).
7. Golub, M. S. et al. Bisphenol A: Developmental toxicity from early prenatal exposure. Birth Defects Res. B. Dev. Reprod. Toxicol. 89(6), 441–466 (2010).
8. U.S. Food and Drug Administration (USFDA). Questions & answers on bisphenol a (BPA) use in food. Contact Applications. Retrieved from: http://www.fda.gov/Food/IngredientsPackagingLabeling/FoodAdditivesIngredients/ucm355155.htm (2016).
9. Bauer, B. A. What is BPA and what are the concerns about BPA? Nutrient and Healthy Eating. Retrieved from: https://www.mayoclinic.org/healthy-lifestyle/nutrition-and-healthy-eating/expert-answers/bpa/faq-20058331 (2016).
10. Al-Hiyasat, A., Darmani, H. & Elbetieha, A. M. Effects of bisphenol A on adult male mouse fertility. Eur. J. Oral Sci. 110(2), 163–167 (2002).
11. Newbold, R. R. Impact of environmental endocrine disrupting chemicals on the development of obesity. Hormones 9(3), 206–217 (2010).
12. Yang, M., Ryu, J. H., Jeon, R., Kang, D. & Yoo, K. Y. Effects of bisphenol A on breast cancer and its risk factors. Arch. Toxicol. 83, 281–285 (2009).
13. Careghini, A., Mastorgio, A. F., Saponaro, S., Sezzana, E. & Bisphenol A, nonylphenols, benzophenones, and benztiazoles in soils, groundwater, surface water, sediments, and food; A review. Environ. Sci. Pollut. Res. 22, 5711–5741 (2015).
14. Kang, J. H., Aasi, D. & Katayama, Y. Bisphenol A in the aquatic environment and its endocrine-disruptive effects on aquatic organisms. Crit. Rev. Toxicol. 37(7), 607–625 (2008).
15. Wang, S. et al. Effects of bisphenol A, an environmental endocrine disruptor, on the endogenous hormones of plants. Environ. Sci. Pollut. Res. Int. 22(22), 17653–17662 (2015).
16. Welshons, W. V., Nagel, S. C. & Von Saal, F. S. Large effects from small exposures. III. Endocrine mechanisms mediating effects of bisphenol at levels of human exposure. Endocrinol. 147(6), s56–s69 (2006).
17. Calafat, A. M. et al. Urinary concentrations of bisphenol A and 4-nonylphenol in a human reference population. Environmental Health Perspectives 113(4), 391–395 (2005).
18. Canesi, L. & Fabbri, E. Environmental effects of BPA: Focus on aquatic species. Dose Response, https://doi.org/10.1177/1559325815598304 (2015).
19. Yamamoto, T., Yasuhara, A., Shiraiishi, H. & Nakasugi, O. Bisphenol a in hazardous waste landfill leachates. Chemosphere 42, 415–418 (2001).
20. Klecka, G. M., Gonser, S. J., West, R. J., Goodwin, P. A. & Markham, D. A. Biodegradation of bisphenol A in aquatic environments: River die-away. Environ. Toxicol. Chem. 20(12), 2725–2735 (2001).
21. Grosart, C. P., Okkerman, P. C., Pijnenburg, A. M. & Pijnenburg, C. M. Chemical study on bisphenol A. BKH Consulting Eng. Report: RIKZ/2001.027 (2001).
22. Zhan, M., Yang, X., Xian, Q. & Kong, L. Photosensitized degradation of bisphenol A involving reactive oxygen species in the presence of humic substances. Chemosphere 63, 378–386 (2006).
23. Ghosh, M. & Singh, S. P. A review of phytoremediation of heavy metals and utilization of it’s by products. Asian J. Energy and Environ. 6(04), 214–231 (2005).
24. Karn, B., Kulken, T. & Otto, M. Nanotechnology and in situ remediation: A review of the benefits and potential risks. Environ. Health Perspectives 117(12), 1823–1831 (2009).
25. Onundi, Y. et al. A multidisciplinary investigation of the technical and environmental performances of TAML/peroxide elimination of bisphenol a compounds from water. Green Chem. 19, 4234–4262 (2017).
26. Peuke, A. D. & Rennenberg, H. Phytoremediation: Molecular biology, requirements for application, environmental protection, public attention and feasibility. EMBO Reports. 6(6), 497–501 (2005).
27. Aken, B. V., Correa, P. A. & Schnoor, J. L. Phytoremediation of polychlorinated biphenyls: New trends and promises. Environ. Sci. Technol. 44(8), 2767–2776 (2010).
28. Das, P., Datta, R., Makris, K. C. & Sarkar, D. Vetiver grass is capable of removing TNT from soil in the presence of urea. Environ. Pollut. 158, 1980–1983 (2010).
29. Datta, R. et al. Phytoremediation potential of vetiver grass [Chrysopogon zizanioides (L)] for tetracycline. Int. J. Phytoremediation 15(4), 343–351. [Abstract] (2013).
30. Chat, W. et al. Biodegradation of bisphenol A by fungi. Appl. Biochem. Biotechnol. 120, 175–182 (2004).
31. Ferrara, G., Loﬀredo, E. & Senesi, N. Phytotoxic, clastogenic and bioaccumulation effects of the environmental endocrine disruptor bisphenol A in various crops grown hydroponically. Planta 223, 910–916 (2006).
32. García-Rodriguez, A., Matamoros, V., Fontas, C. & Salvado, V. The inﬂuence of Lema sp. And Spirorgya sp. On the removal of pharmaceuticals and endocrine disruptors in treated waters. Int. J. Environ. Sci. Technol., https://doi.org/10.1007/s13762-014-0632-x (2014).
33. Imai, S. et al. Removal of phenolic endocrine disruptors by Portulaca oleracea. J. Biosci. Bioeng. 103(5), 420–426 (2007).
34. Okuhata, H. et al. Floricultural Salvia plant have a high ability to eliminate bisphenol A. J. Biosci. Bioeng. 110(1), 99–101 (2010).
35. Saiyood, A., Vangnai, A. S., Thiravetyan, P. & Inthorn, D. Bisphenol A removal by Dracaena plant and the role of plant-associating bacteria. J. Hazard Mater. 178, 778–785 (2010).
36. Takahashi, M., Tsukamoto, S., Kawaguchi, A., Sakamoto, A. & Morikawa, H. Phytoremediators from abandoned rice field. Plant Biotechnol. 22(2), 167–170 (2005).
37. Trueeman, R. J. & Erber, L. Invasive species may offer advanced phytoremediation of endocrine disrupting chemicals in aquatic ecosystems. Emirates J. Food Agriculture 25(9), 648–656 (2013).
38. Zazouli, M. A., Mahdavi, Y., Bazrafshan, E. & Balarad, D. Phytodegradation potential of bisphenol A from aqueous solution by Azolla filiculoides. J. Environ. Health Sci. Engineering 12, 66 (2014).
39. Noureddin, M. I., Furamoto, T., Ishida, Y. & Fukui, H. Absorption and metabolism of bisphenol A, a possible endocrine disruptor, in the aquatic edible plant, water convolvulus (Ipomoea aquatica). Biosci., Biotechnol., and Biochem. 66(6), 1398–1402 (2004).
40. U.S. Department of Agriculture (USDA). Switchgrass plant fact sheet. Retrieved from: https://plants.usda.gov/factsheet/pdf/fs_pavi2.pdf (2006).
41. NL Agency - Ministry of Economic Affairs. Switchgrass (Panicum virgatum L.). A perennial biomass grass for efficient production of feedstock for the biobased economy. Retrieved from: https://english.rvo.nl/sites/default/files/2013/12/Switchgrass%20report%20AgNl%20June%202013_0.pdf (2013).
42. Arora, K., Sharma, S. & Monti, A. Bio-remediation of Pb and Cd polluted soils by switchgrass: A case study in India. Int. J. Phytoremediation 18(7), 704–709 (2016).
43. Chen, B., Lai, H. & Juang, K. Model evaluation of plant metal content and biomass yield of the phytoextraction of heavy metals by switchgrass. *Ecotoxicol. Environ. Saf.* **80**, 393–400 (2012).
44. Murphy, I. J. & Coats, J. R. The capacity of switchgrass (*Panicum virgatum*) to degrade atrazine in a phytoremediation setting. *Environ. Toxicol. Chem.* **30**(3), 715–722 (2011).
45. Gattullo, C. E., Bahrs, H., Steinberg, C. E. W. & Loffredo, E. Removal of bisphenol A by the freshwater green alga Monoraphidium braunii and the role of natural organic matter. *Sci. Total. Environ.* **416**, 501–506 (2012).
46. Li, R. *et al.* Toxicity of bisphenol A and its bioaccumulation and removal by a marine microalga *Stephanodiscus hantzschii* *Ecotoxicol. Environ. Saf.* **72**, 321–328 (2009).
47. Hirooka, T. *et al.* Biodegradation of bisphenol A and disappearance of its estrogenic activity by the green alga *Chlorella fusca var. Vacioudata.* *Environ. Toxicol. Chem.* **24**(8), 1896–1901 (2005).
48. Loffredo, E., Gattullo, C. E., Traversa, A. & Sensi, N. Potential of various herbaceous species to remove the endocrine disruptor bisphenol A from aqueous media. *Chemosphere* **80**, 1274–1280 (2010).
49. Nakajima, N. *et al.* Processing of bisphenol-A by plant tissues: Glycosylation by cultured BY-2 cells and glycosylation/transformation by plants of Nicotiana tabacum. *Plant Cell Physiol.* **43**(9), 1036–1042 (2002).
50. Berhane, T. M., Levy, J., Krekeler, M. P. S. & Danielson, N. D. Kinetic sorption of contaminants of emerging concern by a palygorskite-montmorillonite filter medium. *Chemosphere* **176**, 231–242 (2017).
51. International Conference on Harmonisation (ICH). Validation of analytical procedures: Text and methodology Q2(R1). Retrieved from: http://www.ich.org/fileadmin/Public_Web_Site/ICH_Products/Guidelines/Quality/Q2_R1/Step4/Q2_R1__Guideline.pdf (2005).
52. Missouri Botanical Garden. *Gardening Help* http://www.missouribotanicalgarden.org/PlantFinder/PlantFinderDetails.aspx?kempercode=c460.
53. Das, P., Sukar, D., Makris, K. & Datta, R. Urea-facilitated uptake and nitroreductase-mediated transformation of 2,4,6-trinitrotoluene in soil using vetiver grass. *J. Environ. Chem. Engineering* **3**, 445–452 (2015).
54. Hoagland, D. R. & Arnon, D. I. The water-culture method for growing plants without soil. *Circular. California Agricultural Experiment Station* **2(347)**, 32.
55. Sunkar, R. *Plant Stress Tolerance: Methods and Protocols.* New York, New York (2010).
56. Robati, D. Pseudo-second-order kinetic equations for modeling adsorption systems for removal of lead ions using multi-walled carbon nanotube. *J. Nanostructure Chem.* **3**(55), https://doi.org/10.1186/2193-8865-3-55 (2013).

Acknowledgements
The authors acknowledge Nazareth College for funding, greenhouse, and laboratory facilities; Jane Shebert, for analytical assistance; Sharon Luxmore for purchasing materials; Annie Pysnik, Ashley Harford, Julia Widmer, Dani Painter, and Elena Peydro for assistance in sample preparation, and Lindsay LaPlant and William Lammela for providing feedback about this manuscript.

Author contributions
P.D. designed the research. J.C.P. and A.K.M. conducted the greenhouse study and performed the BPA analysis, biomass estimation, chlorophyll estimation, and enzyme analysis. P.D., J.C.P. and A.K.M. analyzed and interpreted the data. J.C.P., A.K.M., P.D. and S.Z. wrote the paper with contributions from D.G., C.R. and B.B.

Competing interests
The authors declare no competing interests.

Additional information
Supplementary information is available for this paper at https://doi.org/10.1038/s41598-019-56655-w.

Correspondence and requests for materials should be addressed to P.D.

Reprints and permissions information is available at www.nature.com/reprints.

Publisher’s note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/.

© The Author(s) 2020