Degradation study of BPA-containing plastics in water samples collected from an urban reservoir

Abstract

Water pollution is a significant problem worldwide, particularly in Brazil where the majority of urban populations rely on drinking water from surface water reservoirs. The accumulation of plastic debris, such as polycarbonate blends, in reservoirs and other waterways is a public health issue because of health and environmental concerns associated with their chemical degradation. A compound commonly found in plastics, 2,2-bis(4-hydroxyphenyl)propionate (bisphenol-A; BPA), has become a serious environmental problem due to its release in water and its estrogen-like properties. This paper focuses on understanding the degradation process of two types of plastics containing BPA, polypropylene (PP) and poly(lactic acid) (PLA), in surface waters. The strength of the chemical binding of BPA to PP and PLA was examined using thermogravimetric analysis (TGA) and Fourier transform infrared (FTIR) spectroscopy. Release of BPA from both types of plastics was determined in surface water samples collected from the Billings Reservoir, the largest source of drinking water of Sao Paulo residents. The results show that BPA bound to PP-based plastics is released more rapidly into surface waters than that bound to PLA-based plastics.

Keywords: BPA degradation, PLA, PP, urban water reservoir, biodegradable polymer

Introduction

Contamination of water resources is one of the main environmental problems nowadays, and the concern with micro pollutants, i.e., substances that are present in concentrations in the order of μgL⁻¹ and ngL⁻¹, has increased significantly in recent years.¹⁻³ Despite being detected in low concentrations, micro pollutants are capable of causing serious effects to biota. Among the substances that can be classified as micro pollutants, we highlight 2,2-bis(4-hydroxyphenyl)propionate (BPA), a substance that has estrogenic activity. BPA has become ubiquitous in the environment over the past 80 years because of its presence in a wide variety of products, including food and beverage packaging, flame retardants, adhesives, construction materials, electronic components, and paper coatings.¹ BPA is a non-steroidal xenoestrogen that exhibits approximately 1/10000th of the activity of estradiol.⁴

The estrogenic effects of BPA were initially reported by Dodds & Lawson.⁵ Recent studies indicate that BPA may be as effective as estradiol in triggering certain receptor responses and may act as an androgen receptor antagonist.⁶ There are an impressive number of studies on the effects of BPA exposure on laboratory animals, but few studies are dealing with the effects of BPA on wild species in field or laboratory settings.⁷ BPA is considered a pseudo-persistent compound: although short-lived, it is ubiquitous in the environment because of its continuous release.⁸ The release of BPA into the environment may occur during chemical manufacturing, transportation, or processing as well as via discharge of effluents from municipal wastewater treatment plants, leaching from sanitary landfills, combustion of household waste, and leaching from plastics. Some researchers were unaware that BPA could leach from plastics until 1993,⁹ and subsequent studies have confirmed that BPA can also leach from polycarbonates and epoxy resins. Several studies indicate that BPA concentrations in various aqueous media, including surface water, saltwater, and groundwater, have increased in recent years.¹⁰ However, BPA leaching is much faster in salt water than in freshwater as its bioavailability increases with salinity.¹¹ The half-life of BPA in surface waters is 150 days.¹² However, due to leaching and its inadequate disposal, BPA is considered persistent in the environment.

BPA is present in a variety of products, such as soda cans, water bottles, plastic cups, and food packaging. Polypropylene (PP) is a plastic material widely used in various industrial applications; however, PP has poorer mechanical strength and thermal stability than most engineering polymers. To overcome these limitations, BPA polycarbonate can be incorporated into PP.¹³ Poly(lactic acid) (PLA) also has limitations and the addition of BPA polycarbonate can be beneficial in this regard.¹⁴ PP and PLA are widely used in packaging and can generate BPA and its derivatives during thermal degradation. Metal water bottles, especially those made of aluminum, are coated with epoxy resins that may contain BPA to combat corrosion. Food stored in glassware may not be free of exposure to BPA because the metal lids typically used in this type of storage often have an anti-corrosive epoxy coating that contains BPA. Furthermore, plastic bowls and other containers used for cooking in microwave oven may release BPA during heating.¹⁵ This study evaluates the release of BPA incorporated into two different formulations of plastic materials, PLA, a biodegradable polyester, and PP, a polylefin, as well as the degradation of these two polymers. The two types of plastics were tested in the form of pellets placed in contact with water samples collected from the Billings Reservoir, which is one of the largest and most important water reservoirs in the Metropolitan Region of São Paulo, Brazil. Certain parts of the reservoir are polluted with chemicals and metals by domestic and industrial sewage and, for
Analysis of plastic materials

Thermogravimetric analyses (TGA) were performed on a Perkin Elmer STA 6000 analyzer at a heating rate of 10°C/min from 25 to 600°C under a nitrogen atmosphere. Fourier transform infrared (FTIR) spectra were recorded on a Perkin Elmer Frontier 94942 spectrometer by applying a scanning range of 500 to 4000 cm⁻¹ and 64 scans, using thin polymer films.

Analysis of surface water samples

The surface water samples used in the tests were collected in August 2015 from the Billings Reservoir using Van Dorn bottles. The variables water temperature, dissolved oxygen, electrical conductivity, and pH were measured in the field using an Oakton PCD 650 multiparameter probe. In the laboratory, the primary nutrient contents (nitrogen, phosphorus, and potassium), total Kjeldahl nitrogen, and ammonical nitrogen, and nitrite and nitrate contents were determined.

Determination of BPA

A high performance liquid chromatography (HPLC) method was developed for the determination of BPA according to the validation parameters proposed by Snyder et al. An Agilent 1200 HPLC-diode array detector (DAD) apparatus was used, equipped with a automatic injector, column oven, Open LAB v. A.01.05 data acquisition software, and Phenomenex kinetex C18 column (30mmx2.1mmx5um). The parameters of the method were isocratic mode of separation, mobile phase 50:50 (v/v) water acidified with orthophosphoric acid (PA, LabSynth, final pH = 3.0, 3mol L⁻¹ H₃PO₄) and acetoniitrite (HPLC grade, JT Baker), flow rate of 1mL min⁻¹, injection volume of 5µL, and readings at λ = 228nm. A 100mg L⁻¹ BPA stock solution was prepared with acetoniitrite, and, after subsequent dilutions, a calibration curve was plotted with concentrations between 5 mg L⁻¹ and 100 mg L⁻¹ in order to evaluate the linearity of the method. Limits of quantification and detection were determined by analyzing the standard deviation of the response (σ) and the slope of the curve (S), in a ratio of 3.3σ/S for detection and 10σ/S for quantification. Caffeine (aqueous solution, Agilent) and diethylstilbestrol (99.9%, Sigma-Aldrich) standards were used to verify the method’s selectivity and specificity. The precision of the method (repeatability and reproducibility) was assessed by analyzing the intra-assay and inter-assay variance coefficients. In order to study the degradation process of BPA-containing PP and PLA in surface waters, tests were performed as described below.

The water samples collected from the reservoir had their pH adjusted to 7.5 using 0.1mol L⁻¹ NaOH (Synth). Then, 100mL of the samples was added to glass flasks, which were placed in a water bath at 23°C, 53°C, or 83°C, in triplicate. Subsequently, 0.5g of PLA incorporated with BPA (PLA-BPA) was added to each flask. After 1min, a 1mL aliquot was collected and filtered through a 0.45µm membrane using a glass syringe. Immediately after sample collection, the flasks were capped and returned to the water bath. Additional aliquots were collected from each flask daily for 8 days and analyzed using the HPLC method developed for BPA determination. To evaluate the release of BPA from PP, 50mL of the collected water samples with pH adjusted to 7.5 and approximately 0.1g of PP incorporated with BPA (PP-BPA) were added to glass flasks, in triplicate. The flasks were placed in a water bath under the same temperatures used in the PLA assays. After 1min, 1mL aliquots were withdrawn from each flask and filtered through a 0.45µm membrane using a glass syringe. Additional aliquots were withdrawn after 48h, 72h, and 7 days and analyzed using the HPLC method developed for BPA determination.

Results and discussion

Analysis of plastic materials

Figure 1 shows the FTIR spectra and TGA curves of PLA, PLA-BPA, PP, and PP-BPA samples. Regarding PLA, Figure 1A shows FTIR spectra expanded between 500–4000cm⁻¹. The characteristic peaks of PLA occur between 1180 and 1745cm⁻¹. Peaks at 1033cm⁻¹, 1045cm⁻¹, and 1160cm⁻¹ correspond to the stretching of C-O bonds of aromatic ethers present in the PLA structure. The 1312cm⁻¹ band refers to the methyl group of PLA, and the peak of C-C bond is found at 1541cm⁻¹. Between 1720 and 1740cm⁻¹, a C=O band that corresponds to aldehydes was identified at 1734cm⁻¹. Bands at 2329, 2852, and 2919cm⁻¹ correspond to the C-H aliphatic bonds. The vibrational band at 2360cm⁻¹ refers to the presence of atmospheric CO₂, and the peaks at 2850 and 2921cm⁻¹ probably correspond to asymmetric C-H stretching of residual carbons. Regarding PP, Figure 1A shows FTIR spectra expanded between 1370–1390cm⁻¹. The peak at 1375cm⁻¹ corresponds to the angular deformation caused by the methyl bond. Bands between 1430 and 1470cm⁻¹ refer to the CH₃ functional group present in the PP structure, whereas C-H bonds caused peaks at 2836 and 2925cm⁻¹. The characteristic band of PP occurs between 2700 and 3000cm⁻¹. The characteristic FTIR peaks of PP-BPA occur above 3000cm⁻¹. Our results showed a peak at 3726cm⁻¹, corresponding to the O-H bond of BPA, proving its presence in the material. The peak at 1542cm⁻¹ corresponds to the C=C bond of aromatic rings of the BPA molecular structure. Peaks between 2852 and 2919cm⁻¹ were attributed to the C-H functional group present in the polypropylene structure, similar to the results reported by Nikjé et al.

The incorporation of BPA into PLA caused the disappearance of the 1045 and 1160cm⁻¹ peaks, referring to the C-C and C-O bonds. With respect to PP, the incorporation of BPA caused the disappearance of the peaks 996, 1375, and 1460cm⁻¹, which correspond to the C-C, CH₃, and CH₂ bonds, respectively. These results show the interaction of BPA with the polymers. Figure 1B shows the TGA curves describing the behavior of PLA and PP at different temperatures. For PLA, there was no mass loss up to 300°C. PLA degradation occurred in a single stage at 430°C, contrary to the results reported by Zhang et al. in which PLA degradation occurred at approximately 400°C using a heating rate of 10°C/min in an N₂ atmosphere. The TGA
curve describing the behavior of PLA-BPA is similar to that of pure PLA; the differences are given by the initial mass loss, which occurs at slightly lower temperatures (at approximately 426°C), possibly because of the thermal characteristic of BPA, which degrades at temperatures in the range of 150°C. This result suggests that there is some interaction between BPA and PLA, probably due to the polarity of PLA and the hydroxyl groups of BPA, which prompts a sudden and rapid separation of BPA from PLA at temperatures above 426°C. The PP TGA curve in Figure 1B shows that mass loss started only at 360°C and that polymer degradation occurred at approximately 482°C, differently from the results of Pérez et al. in which degradation of PP occurred at approximately 450°C, demonstrating a better thermal stability than pure PLA, promoted by the same heating rate condition at 10°C min⁻¹ under N₂ atmosphere. The TGA curves of PP-BPA and pure PP showed a similar mass loss due to the good mixing of the components. The degradation of PP-BPA began in the range of 480°C, and PP-BPA and pure PP curves showed that degradation took place until 600°C.

Analysis of surface water samples

The results of the analysis of surface water samples were as follows: electrical conductivity of 307.0μS cm⁻¹, pH ranging from 7.8 to 8.4, and maximum total phosphorus concentrations of 740.0μg L⁻¹. The high nutrient load found in the selected sampling points of the Billings Reservoir, especially the total phosphorus content, allows the classification of these waters as hypereutrophic, according to the criteria proposed by Cunha et al. for tropical and subtropical lake systems. This trophic state was also observed in previous studies performed in different distributaries of the Billings Reservoir. According to Yang et al., this trophic condition is quite common in shallow reservoirs due to the high vulnerability of these ecosystems to anthropogenic interventions. Under these conditions, the salinity of water increases, which can facilitate the leaching of BPA from both PP and PLA.

HPLC Analysis

The HPLC method validated and applied for BPA analyses, presented limits of detection and quantification of 0.360mgL⁻¹ and 1.09mgL⁻¹, respectively. Values calculated based on the signal-to-noise ratio based on the standard curve (\( y = 2.2548 \times 10^{-5} x + 2.19469 \); \( r^2 = 0.99451 \)) suggest that the detection system is suitable for the analyses performed. The figures of merit, repeatability and reproducibility, determined using the concentration values of the analytical curve, returned %CV (coefficient of variance) values of 1.5 to 3.9%, respectively, which are excellent results, as the analysis employed complex environmental samples. In Figure 2, it is possible to observe that PP rapidly released BPA into the medium under the experimental conditions used and that PLA took longer to release BPA into the medium. This occurs because PLA has oxygen in its chemical structure, which makes it more polar and increases the strength of its interaction with BPA through van der Walls interactions. This finding corroborates the decrease in the immediate estrogenic activity that BPA would have if released immediately under normal pH conditions of surface water.

Figure 1 (A) FTIR spectra and (B) TGA curves of PLA, PLA-BPA, PP, and PP-BPA samples.

Figure 2 Chromatogram of (A) PP-BPA at 83°C and 48h, (B) PLA-BPA at 83°C and 48h, (C) PLA-BPA at 83°C and 0h, (D) PLA-BPA at 83°C and 0h using acetonitrile:acidified water (50:50, v/v) as the mobile phase, injection volume of 5μL, detection at 228nm, 40°C, and flow rate of 1mL min⁻¹. Graph showing BPA release from PP and PLA at different temperatures.

Conclusion

The results show that there is greater interaction between BPA and PLA than between BPA and PP. TGA results further confirm this observation. The interaction probably occurs due to the polarity of PLA and the presence of hydroxyl groups in BPA. PP-based plastics release BPA more readily into surface waters than PLA-based plastics. The validated HPLC method is suitable for verifying that polymeric materials containing BPA can be a leaching source of BPA in surface waters. The presence of dissolved BPA in surface waters at sufficient concentrations could cause estrogenic-related impacts on aquatic life as well as humans, exposed through the consumption of drinking water, especially considering that the studied water reservoir receives a large amount of plastic waste and since traditional water treatment cannot eliminate residues of these materials it is necessary that we have methodologies to verify the quantity of these substances in the treated water that will go to the residences of the population.

Acknowledgements

The authors Derval dos Santos Rosa and Elizabete Campos de Lima are grateful to UFABC (grant no. 447180/2014-2), FAPESP (grant no. 306401/2013-4), and CNPq (no. 2013/12559-8) for their financial support.

Conflict of interest

The authors declare there is no conflict of interest.

References
