Rapid determination of tetrabromobisphenol A and its main derivatives in aqueous samples by ultrasound-dispersive liquid–liquid microextraction combined with high-performance liquid chromatography

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ABSTRACT

A method of ultrasound-dispersive liquid–liquid microextraction (US-DLLME) combined with high-performance liquid chromatography/variable wavelength detection (HPLC-VWD) has been developed for rapid measuring tetrabromobisphenol A and its five derivatives in water. Parameters affecting the extraction efficiency including the extraction solvents and dispersive solvents and their volume, ionic strength of the sample, and ultrasound time were optimized, and further validated by orthogonal array design (OAD). The optimized conditions provided enrichment factors for analytes of 74–490. Most analytes had linear responses between 2 and 500 $\mu\text{g L}^{-1}$, with correlation coefficients (r^2) of 0.9923–0.9994. Limits of detection were 0.13–0.63 $\mu\text{g L}^{-1}$. Relative standard deviations (RSDs) for five replicates ranged from 2.6% to 4.5% for all analytes. When applied to spiked samples of real water, the method provided recoveries of 88.6–106.3% for tap water, 87.8–108.5% for Mi River water, 82.7–113.5% for chemical wastewater, 45.5–115.3% for urine, and 46.4–126.2% for fruit juice, with RSDs ($n=5$) less than 4%, 6%, 8%, 10%, and 9% respectively.

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1. Introduction

Environmental and human health problems caused by brominated flame retardants (BFRs) have recently been receiving increased attention [1]. One of the most important BFRs, tetrabromobisphenol A (TBBPA), has the highest production volume, exceeding those of Deca-BDE and HBCD, covering around 60% of the total BFRs market [2,3], and has been confronted with increasing environmental scrutiny [4].

Available evidence indicates that TBBPA and TBBPA-bis (allyl ether) (TBBPA-BAE) lead to potentially toxic endocrine-disruptor effects, immunotoxicity, and neurotoxicity, all because their structures resemble the thyroid hormone thyroxine (T4) and bisphenol A [5–7]. Other studies have shown that TBBPA can be debrominated under anaerobic and aerobic conditions [8,9] to yield bisphenol A, a widely used compound with a disruptive endocrine function [10]. Therefore, it is urgent to evaluate and monitor the trace levels of these compounds in the environment.

TBBPA has been measured in some water samples. Most of those studies used high-performance liquid chromatography (HPLC) coupled with mass spectrometry (MS) [11,12]. But TBBPA derivatives have received less attention. One study recently measured TBBPA, TBBPA-BAE, and TBBPA-bis (2,3-dibromopropyl ether) (TBBPA-BDBPE) in real samples by LC-MS/MS by using accelerated solvent extraction (ASE) as the sample-extraction method, gel-permeation chromatography (GPC), and solid-phase extraction (SPE) as cleanup technique [13]. Another study proposed pressurized fluid extraction (PFE) and fluidized bed extraction (FBE) as extraction procedures and HPLC-DAD/MS for the determination of TBBPA-BDBPE in environmental samples [14]. However, these extraction methods are time-consuming, tedious and often require large amounts of toxic organic solvent. To overcome those disadvantages, time-saving and high-efficiency microextraction techniques have drawn much attention.

DLLME (dispersive liquid–liquid microextraction) is a new sample pretreatment technique, which is based on a ternary solvent system like homogeneous liquid–liquid extraction and cloud point extraction [15]. In this method, a cloudy solution is formed when an appropriate mixture of a few microliters of organic extraction solvent and a small volume of a dispersive

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solvent are injected into an aqueous sample of the analytes. After centrifuging, the analytes in the sedimentary organic phase can be measured by conventional chromatographic techniques. DLLME is simple, rapid and environmentally friendly, and has been successfully measured trace organic pollutants in environment samples [15–20]. The present work was undertaken to determine TBBPA and its five derivatives by using ultrasound-dispersive liquid–liquid microextraction (US-DLLME) coupled with HPLC. To the best of our knowledge, TBBPA and its five derivatives, especially for TBBPA-bis (glycidyl) ether (TBBPA-BGE), have not been measured this way before. This study evaluated the effects of various experimental parameters, and an optimized method was successfully applied to the real water sample analysis.

2. Experimental section

2.1. Materials and chemicals

Bisphenol A (BPA), TBBPA, TBBPA-BAE, TBBPA-bis (2,3-dibromopropyl ether) (TBBPA-BDBPE), TBBPA-BGE, and TBBP-A bis (2-hydroxyethyl ether) (TBBPA-BHEE) were obtained from Sigma-Aldrich (St. Louis, MO, USA). Their structures are given in the Supporting Information (Table S-1, along with the official abbreviation [21]). Stock solution of each was prepared at a concentration of 100 mg L^{-1} in methanol and stored at 4°C . Working solutions were prepared by appropriately diluting the stock solution with ultrapure water.

Analytical-grade dichloromethane (DCM), chloroform, methanol, acetone, and tetrahydrofuran (THF), and HPLC-grade acetonitrile (ACN) were purchased from J.T. Baker (Phillipsburg, NJ, USA). 1-butyl-3-methylimidazolium hexafluoro phosphate ([BMIM][PF₆]) (99%) was purchased from J&K Chemical Ltd. (Beijing, China), and 1-butyl-3-methylimidazolium bis (trifluoro) ([BMIM][SF₃]) (99%) from Adamas Reagent Co. Ltd. (Shanghai, China). Ultrapure water (18.2 M Ω) was generated with a Milli-Q (Millipore, Billerica, MA, USA) water purification system.

Samples of natural water and chemical wastewater were collected from the Mi River and local chemical plant (Shouguang, China). Urine samples were provided by volunteers. Fruit juice was bought from supermarket. All samples were filtrated through a $0.45 \mu\text{m}$ glass fiber membrane (Automatic Science, China) and were stored at 4°C before being used.

2.2. Instrumentation

Chromatographic analysis was performed with an Agilent1100 HPLC system (Agilent Technologies, USA) equipped with a manual injector and variable wavelength detector (VWD). A reverse phase C18 HPLC column (Acclaim120, $5 \mu\text{m}$, $150 \text{ mm} \times 4.6 \text{ mm}$, Dionex) was used for separating of the analytes. The mobile phase consisted of ACN/water, and used at a constant flow of 1.0 mL min^{-1} . The initial ratio (20% ACN) was held for 3.0 min, then decreased gradually to 0% at the end of the 12-min run. The detector wavelength was 214 nm. Microsyringes of 500–1000 μL and 5–50 μL (SGE, Australia) were used to inject the extraction solvent and dispersive solvent into the aqueous sample and transfer the sedimentary organic phase into HPLC injector, respectively.

2.3. Ultrasound-dispersive liquid–liquid microextraction (US-DLLME)

A scheme of the US-DLLME has been depicted in Fig. 1. A volume of 5.0 mL water sample was placed in a 10 mL screw-cap glass tube with conical bottom. To optimize the extraction, a 5.0 mL water sample spiked with each analyte at $100 \mu\text{g L}^{-1}$ was used. Then 0.6 mL of THF (as dispersive solvent) and 25 μL of chloroform (as extraction solvent) were rapidly injected into the sample solution. The mixture was gently shaken for 10 s and ultrasonicated in ultrasonic bath (KQ-300DE, Shanghai, China) for 2 min to form a homogeneous cloudy solution. The phase was separated by a centrifuging at 4000 rpm for 5 min (DT5-2, Beijing, China). The volume of the sedimentary organic phase was about 15 μL , from which 10 μL was removed for injecting into HPLC. All operations were carried out at room temperature (25°C).

2.4. Calculation of enrichment factor (EF)

In order to evaluate the effect of experimental conditions on the performance of DLLME, an enrichment factor (EF) was calculated [15]:

$$\text{EF} = \frac{C_s}{C_i}$$

C_s , and C_i are the analyte concentration in the sedimentary organic phase and the initial aqueous samples, respectively.

Fig. 1. A scheme of the US-DLLME method.

3. Results and discussion

To maximize extraction efficiency, each factor in the extraction process was optimized. The experiments were performed in triplicate, and the means used.

3.1. Selection and volume of the extraction solvent

The choice of extraction solvent is very important for the DLLME process, in order to maximize the extraction efficiency and selectivity for the target compounds. The extraction solvent should be denser than water, behave well during chromatography, extract the target compounds efficiently, has low solubility in water, and be miscible with the dispersive solvent [15]. These considerations led to selecting halogenated alkanes (dichloromethane, chloroform) as a universal extraction solvent and ionic liquids ([BMIM][PF₆], [BMIM][SF₆]) as a “green” extraction solvent. Chloroform was chosen because it gave the highest EFs (Fig. 2a).

This study aimed to determine the minimum volume of extraction solvent necessary to achieve the highest EF. Volumes of chloroform between 10 and 55 μ L were tested. With less than 10 μ L, centrifuging could not properly separate the phases, and a

viscous phase with matrix components remained. When the chloroform volume was increased, an optimum phase separation was observed. The highest EFs were reached with 25 μ L of chloroform (Fig. 2b). With 10 μ L of chloroform, lower EFs were observed because the volume was too small to quantitatively extract the BFRs. Increasing the volume of the extraction phase decreased the EF of the analytes, when C_0 is constant, the sedimentary organic phase volume will increase with the increment of extraction phase volume and the consequent dilution effect of the analyte in sedimentary phase is reduced C_s . Therefore, 25 μ L was selected to separate the phases and provide high enrichment factors.

3.2. Selection and volume of the dispersive solvent

A key property of dispersive solvents is miscibility with the extraction solvent and the aqueous sample. For this purpose, methanol, ACN, acetone and THF were investigated. Aqueous samples were extracted in 25 μ L of chloroform and 0.6 mL of the dispersive solvent. Dispersive solvents affect the dispersion, stability and equilibrium of the ternary-phase system. An appropriate dispersive solvent will disperse the extraction solvent to fine

Fig. 2. Effect of (a) different extraction solvents and (b) volume of the optimized extraction solvent on enrichment factors of TBBPA and its main derivatives. Dispersive solvent was 0.6 mL of THF.

Fig. 3. Effect of (a) different dispersive solvents and (b) volume of the optimized dispersive solvent on enrichment factors of TBBPA and its main derivatives. Extraction solvent was 25 μ L of chloroform.

droplets in water sample and thereby increase the surface area for transferring the target compounds from sample matrix to extraction solvent, and then, the different volumes of sedimentary phase can be obtained after centrifugation. The volumes of sedimentary organic phase obtained with methanol, ACN, acetone and THF were 7, 10, 8, and 15 μL , respectively. THF gave the highest enrichment factor for all the analytes (Fig. 3a), and so was chosen as the dispersive solvent.

The influence of THF volume was tested for 0.4, 0.6, 0.8, and 1.0 mL. With increasing volume, the extraction efficiency first increased for all the analytes, then decreased (Fig. 3b). For smaller volumes of THF, a stable cloudy system could not be well formed, resulting in lower enrichments. At higher volumes, the solubility of BFRs in water increased, and led to lower extraction efficiencies. These results led to choosing 0.6 mL of THF as the optimum.

3.3. Effect of ionic strength

Increasing the ionic strength of the sample solution by adding salt usually improves the extraction efficiency in conventional liquid–liquid microextraction via a salting-out effect. Therefore, the effect of ionic strength on US-DLLME was studied by adding up to 20% (w/v) of NaCl (Fig. S-1, in Supporting Information). The highest EFs for each target analyte were obtained without NaCl, however. This negative result was attributed to the increase in volume of sedimentary organic phase from 15 to 25 μL as a consequence of decreased solubility of the extraction solvent in the aqueous phase after adding the NaCl. Apparently, the dilution dominated the salting-out effect, which was agreement with the results of other researchers [22–24]. Hence, NaCl was not used.

3.4. Effect of ultrasonication time

Ultrasonication can accelerate the homogenization/emulsification step and the formation of the submicrometer droplets that greatly increases the contact surface between liquids and facilitate

the transfer of analyte from aqueous sample to organic solvent [15]. To study the effect of ultrasonication time, 0, 2, 5, and 10 min were tried. The EFs significantly increased up to times of 2 min, then leveled off, suggesting that 2 min was enough to form a stable cloudy solution and complete the extraction. Consequently, 2 min of ultrasonication time was chosen.

3.5. Orthogonal array design (OAD)

Orthogonal array design (OAD) was used to confirm the optimal experimental conditions. OAD arranges experiments and analyzes data via orthogonal arrays. $L_{16}(4^5)$ orthogonal design was used here to find the optimum combination of five influence factors for each target analyte (extraction solvent, dispersive solvent, volume of extraction solvent, volume of dispersive solvent and addition of salt). Each influence factor includes four states (Table S-2 in Supporting Information), which made 16 “Extraction Conditions” in all. Extraction Condition no. 8 was found to be the best (to yield the highest enrichment factors) for all the BFRs except BPA, whose best was Condition no. 4. Thus, Condition no. 8 was adopted for this study, and used in conjunction with sample volumes of 5 mL and ultrasonication times of 2 min.

3.6. Performance of the US-DLLME

In order to validate the developed US-DLLME method, its linearity, correlation coefficients, detection limits, enrichment factors, and repeatability were tested using spiked samples under the optimized conditions. The range of linearity was 2.0–500 $\mu\text{g L}^{-1}$ for most of the target compounds, with the correlation coefficient (r^2) ranging from 0.9923 to 0.9994 (Table 1). The limits of detections (LODs) based on a signal-to-noise ratio (S/N) of 3 ranged from 0.13 to 0.63 $\mu\text{g L}^{-1}$. The limits of quantification (LOQs) based on a signal-to-noise ratio (S/N) of 10 ranged from 0.43 to 2.10 $\mu\text{g L}^{-1}$. The EFs of BFRs were 74 to 490. The relative standard deviations (RSDs) for five

Table 1
Performance of US-DLLME.

Analyte	Linear range ($\mu\text{g L}^{-1}$)	r^2	EF ^a	LOD ^b ($\mu\text{g L}^{-1}$)	LOQ ^c ($\mu\text{g L}^{-1}$)	RSD% ($n=5$)	Repeatability (RSD%, day-to-day)
BPA	10–500	0.9985	74	0.34	1.13	4.5	5.6
TBBPA	5.0–500	0.9951	356	0.13	0.43	4.3	5.3
TBBPA-BAE	2.0–500	0.9923	121	0.63	2.10	3.1	4.7
TBBPA-BDBPE	2.0–500	0.9994	140	0.61	2.03	2.9	6.5
TBBPA-BGE	2.0–500	0.9956	490	0.26	0.87	2.6	3.6
TBBPA-BHEE	10–500	0.9965	215	0.24	0.80	3.9	4.3

^a EF, enrichment factor.

^b LOD, limit of detection for a $S/N=3$.

^c LOQ, limit of quantification for a $S/N=10$.

Table 2
Comparison of US-DLLME with other sample preparation methods for TBBPA and its derivatives in water.

Methods (compounds)	Pre-treatment	Extraction time (min)	LOD	Recovery (%)	RSD (%)	Ref.
LC-ESI-MS (TBBPA)	Filtration, isotope imprinting	> 90	0.01 $\mu\text{g/L}$	> 95.0	1.0–4.0	[11]
LC-ESI-MS/MS (TBBPA)	SPE	60	0.02 $\mu\text{g/L}$	> 85.0	5–16	[12]
LC-APPI-MS/MS (TBBPA and 2 derivatives)	ASE/GPC/SPE	120	0.03–1.28 $\mu\text{g/L}$	39–80	–	[13]
HPLC-DAD/MS (TBBPA-BDBPE)	PFE/FBE	60	10–72 $\mu\text{g/L}$	35–98	< 2%	[14]
NACE-PDAD (TBBPA and phenolic compounds)	SPE	> 60	0.8–1.5	73–106	1.9–3.4	[25]
LC-ESI-MS/MS (TBBPA)	SPE, filtration	< 20	300 ng g^{-1} (dry weight)	> 95.0	–	[26]
HPLC-VWD (TBBPA and 4 derivatives)	SPME	30	0.4–0.9 $\mu\text{g/L}$	86.5–103.6	1.2–5.1	[27]
HPLC-VWD (TBBPA and 5 derivatives)	US-DLLME	2	0.13–0.63 $\mu\text{g/L}$	82.7–113.5	2.6–4.5	This paper

replicate runs varied from 2.6% to 4.5% at $50 \mu\text{g L}^{-1}$. The repeatability of day-to-day was 3.6–6.5% (RSDs). Thus the proposed method has high sensitivity and good repeatability.

The optimized US-DLLME method was compared with other methods for extracting and measuring of TBBPA in water (Table 2).

Almost all papers have described methods that are only capable of detecting and quantifying TBBPA or one to two derivatives [11–14,25,26]. TBBPA and five derivatives were simultaneously determined in this work only. The extraction time of 2 min was much faster than for traditional methods. The LODs, recoveries and RSDs obtained here are acceptable. Thus, the proposed method is simple, rapid, low-cost, environmentally friendly, and high-throughput.

3.7. Real water sample analysis

To investigate the applicability of the optimized US-DLLME method, samples of tap water, the Mi River, wastewater from a chemical plant, urine, and fruit juice were analyzed. To estimate possible matrix influences, aqueous samples were spiked with different concentrations of analytes to determine the recovery of the target analytes. Typical chromatograms for real samples and those spiked with analytes (Fig. 4) showed that several target compounds were found in original aqueous samples. The spiking recoveries were 88.6–106.3% for tap water, 87.8–108.5% for the Mi River, 82.7–113.5% for chemical wastewater, 45.5–115.3% for urine, and 46.4–126.2% for fruit juice. The RSDs ($n=5$) were less than 4%, 6%, 8%, 10%, and 9% for the five types of samples (Table 3). These results showed that the US-DLLME method is suitable for measuring BFRs in various aqueous samples, and is not affected significantly by sample matrices.

4. Conclusions

A sensitive and simple ultrasound-DLLME method coupled with HPLC-VWD has been developed for the determination of six BFRs in water. The method provides good repeatability, high enrichment factor, and good recovery. In comparison with traditional sample pretreatment methods, such as filtration, SPE, ASE/GPC/SPE, PFE/FBE, and SPME, the presented method has some obvious advantages in total extraction time (only 2 min), cost (instruments and chemicals), green techniques (a small amounts organic solvents). Therefore, the proposed method is rapid, efficient, environmentally friendly and feasible. Since only 5 mL of sample is needed, the method may be widely applied to different compounds and other aqueous samples such as biological and food samples, although the possible problems posed by their more complex matrices will have to be checked first.

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Fig. 4. Chromatogram of (a) the original and spiked Mi River samples ($50 \mu\text{g L}^{-1}$ for each analyte); (b) the original and spiked wastewater samples ($100 \mu\text{g L}^{-1}$ for each analyte). (1) BPA, (2) TBBPA-BHEE, (3) TBBPA, (4) TBBPA-BGE, (5) TBBPA-BAE, and (6) TBBPA-BDBPE.

Table 3
Spiking recoveries and RSDs ($n=5$) of the BFRs in real water samples.

Analyte	Tap water ^{a,b}				Mi River ^{a,b}				Chemical waste water ^{a,b}				Urine sample ^{a,b}				Fruit juice sample ^{a,b}			
	Recovery (%)		RSD (%)		Recovery (%)		RSD (%)		Recovery (%)		RSD (%)		Recovery (%)		RSD (%)		Recovery (%)		RSD (%)	
	a	b	a	b	a	b	a	b	a	b	a	b	a	b	a	b	a	b	a	b
BPA	92.5	90.6	3.8	3.5	88.7	90.3	4.1	3.4	82.7	85.5	6.7	5.6	56.7	61.2	7.3	8.1	64.1	70.2	5.4	6.1
TBBPA-BHEE	96.8	91.5	2.3	3.8	103.9	105.5	5.6	4.8	84.5	83.3	7.1	7.5	45.5	50.0	4.2	5.0	64.8	68.7	5.6	6.6
TBBPA	106.3	105.4	3.1	3.0	104.8	108.5	5.1	4.6	110.9	108.7	5.3	6.3	67.1	65.5	7.6	8.5	82.9	114.2	6.5	7.1
TBBPA-BGE	96.2	88.6	2.2	3.5	92.8	93.5	3.6	4.7	113.5	112.2	4.9	4.8	58.6	56.5	6.4	6.2	126.2	124.5	8.8	7.6
TBBPA-BAE	95.4	91.0	1.8	2.6	89.2	90.1	4.5	5.5	90.1	85.8	7.3	6.6	71.1	67.8	9.5	8.8	46.4	50.2	4.7	6.5
TBBPA-BDBPE	102.5	93.5	2.5	2.6	87.8	89.2	3.8	4.8	83.0	86.5	6.9	7.8	115.3	110.1	7.8	8.9	123.0	119.5	9.1	8.2

^a Spiked with $50 \mu\text{g L}^{-1}$.

^b Spiked with $100 \mu\text{g L}^{-1}$.

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Appendix A. Supplementary material

Supplementary data associated with this article can be found in the online version at <http://dx.doi.org/10.1016/j.talanta.2013.08.011>.

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