DETECTION OF BISPHENOL A CONTAMINATION IN CANNED CARBONATED BEVERAGES BY GAS CHROMATOGRAPHY

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ABSTRACT

Objective: The purpose of this study was to develop sensitive, selective, and valid methods for the detection of bisphenol A (BPA) contamination in beverage samples using gas chromatography (GC)-flame ionization.

Methods: The optimized analysis system employed a long HP-1 capillary column (30 m, inner diameter 0.25 mm, film thickness 0.25 µm), gradient column temperature (150°C-260°C at 10°C/min), and nitrogen as a carrier gas (1 mL/min). Samples were prepared for analysis using ethyl acetate as the extraction solvent.

Results: This method yielded a linearity coefficient of 0.9998, while the limit of detection (LOD) and limit of quantitation (LOQ) were 0.287 µg/mL and 0.956 µg/mL, respectively. All validation parameters, including linearity, selectivity, accuracy, precision, LOD, and LOQ, met recognized acceptability criteria. Contamination analysis showed that one of the three beverage brands tested contained 2.4090 µg/mL BPA, and contamination was even higher after heating.

Conclusion: BPA contamination may occur in canned beverages, especially under improper storage conditions. This GC-based BPA detection system may be useful for the detection of BPA contamination in consumer beverages.

Keywords: Bisphenol A, Gas chromatography, Contamination analysis, Validation, Canned beverages.

INTRODUCTION

Bisphenol A (BPA) has been used by polymer industries since 1950 for the production of polycarbonate plastic polymers and epoxy resins. Products containing BPA include infant milk bottles, food and drink containers, thermal papers, compact discs, tape, water pipe, vehicle spare parts, and coatings on metal cans. The widespread use of BPA leads to frequent human exposure through room air, dust, ingestion through food and beverages inside containers, and the water supply [1].

There is a growing concern that this BPA exposure may cause health problems. In fact, BPA has been linked to miscarriage, premature birth, obesity (and associated diseases), immune dysfunction, and disruption of neural and hormonal signaling. Several studies have reported that BPA compounds can be released from containers containing polycarbonate or epoxy resins under certain conditions, thereby contaminating food and beverages [2]. The European Food Safety Authority (EFSA) has set 5 µg/kg as the maximum tolerable limit of consumed BPA (EFSA, 2014), a substantial decrease from the previous limit of 50 µg/kg BW [3], in response to studies demonstrating the risks from consuming BPA.

BPA can be released from polymers through exposure to heat, acidic solutions, and alkaline solutions. Therefore, factory sterilization may cause the release of BPA into canned beverages. In addition, leaving unconsumed beverage in an opened can without refrigeration and exposed to sunlight may also cause contamination.

Due to the potential dangers of trace BPA contamination, a sensitive analytical method is required for product testing [4]. Such analytical methods have been developed based on high-performance liquid chromatography (HPLC) with UV detection [5,6], electron catching detection [7,8], mass spectrometry (MS) [9], FID/MS gas chromatography (GC) [10-13], and enzyme-linked immunosorbent assay [14]. However, GC is more selective and sensitive than HPLC and can detect contaminants with shorter analysis times and greater single compound resolution. GC using a longer column yields even higher separation efficiency (Wang and Burleson, 1999) [15,16]. Therefore, we developed a GC-based method using a long column for the detection of BPA and employed this method to demonstrate the presence of BPA in test samples of canned beverages.

MATERIALS AND METHODS

Materials

In this study, 18 canned beverage samples of three different brands were tested (coded A, B, and C for samples with 6 months until the expiry date and D, E, and F for samples with 2 years until the expiry date). Samples were obtained from Depok City, West Java. Standards and suppliers were as follows: BPA (Sigma-Aldrich), acetate ethyl (Merck), methanol pro-HPLC (Merck), and distilled water (Aqua Bidest pro HPLC).

Optimal analysis conditions were identified as follows. First, the optimal detector temperature was chosen by injecting 3 µL standard BPA solution at 100 ppm into the GC system with gas water flow at 1 mL/min and one of the three column temperature conditions, 250°C [5,17], 280°C [6,18], or controlled gradient increasing from 150°C to 260°C at 10°C/min [7,19]. Next, the optimal analysis volume was chosen by injecting 1, 2, or 3 µL BPA standard at 15 ppm into the GC system using 1 mL/min gas water flow and optimized temperature conditions. The optimal gas water flow was chosen by injecting 3 µL of standard 35 ppm BPA solution into the GC system with gas water flow at 0.8, 1, or 1.2 mL/min. The optimal analysis condition for each parameter was scored based on the separation between the two nearest peaks (or resolution, R), peak sharpness, follow-up factor, peak release resistance time, and column efficiency (theoretical plate number [N] and height equivalent to the theoretical plate [HETP]).
Next, system conformity tests were conducted. A standard 15 ppm BPA solution was prepared and 3.0 µL was injected into the GC system under optimized conditions (above). Results were graded by the repetition score (%CV) for resolution (R), peak sharpness, follow-up factor, peak release resistance time, and column efficiency (N and HETP).

The BPA analysis method was validated by linearity testing and construction of calibration curves. Briefly, 100 µg/mL standard BPA solution was diluted in methanol to 2, 4, 6, 8, 10, and 15 µg/mL (ppm), and 3.0 µL of each solution injected into the GC system. The peak (y) versus analytic concentration (x) was then subjected to regression analysis to obtain a correlation coefficient (r). From the obtained calibration curve, the limit of detection (LOD) and limit of quantification (LOQ) were derived.

For selectivity analysis, a blank solution (solvent). BPA standard at 15 ppm, and spiked BPA solutions extracted as for beverage samples (below) were injected at 3 µL and retention times were compared. The spiked solution was prepared using the standard addition method to sample 50%, 100%, and 150% BPA concentrations. Each solution was injected into the GC system at 3 µL under optimized conditions. Tests were conducted in triplicate for selectivity analysis and 6 times for the evaluation of precision. Acquisition tests were conducted to assess accuracy, and %KV was calculated to assess precision.

Finally, BPA levels in the beverage samples were examined. 3 µL of each extraction solution was injected into the GC system under optimized conditions, and the acquired peak was analyzed. Canned beverages were first sonicated for 30 min to remove carbon dioxide. Samples (30 mL per container) were extracted by mixing with 30 mL 90% acetone ethyl, shaking gently for 30 s, and then leaving the mixture to sit until clear separation of the organic layer and water layer. Acetone ethyl was used because its close polarity to BPA suggests the suitability for extraction. After layer separation, the 30-mL organic layer was transferred to a reaction tube and the water layer transferred to a glass breaker for the evaporation of water. The organic layer was dissolved in 5 mL methanol and filtered using Whatman filter paper with 0.45 µm pore size. Samples of extract (3 µL) were then injected into the GC system under optimized analysis conditions. Each sample was tested in triplicate.

**RESULTS AND DISCUSSION**

**Determination of optimal analysis conditions**

**Optimal detector temperature selection**

Comparison of chromatographic results using different temperatures (two isothermal conditions and a gradient of 10°C/min from 150°C to 260°C) defined the optimal temperature for separation in methanol as 243–250°C, at which the retention time was 9.713 min (Table 1).

**Injection volume selection**

An optimal injection volume of 3 µL was chosen based on highest peak area, follow-up factor, and resolution among 1, 2, and 3 µL samples of 15 ppm BPA (Table 2).

**Flow rate selection**

A flow rate of 1 mL/min was chosen among 0.8, 1.0, and 1.2 mL/min based on highest theoretical plate number, HETP, and resolution as well as shortest retention time (Table 3).

**System comparability test**

The %KV for the GC system under optimized conditions was 1.377% as calculated from six repeated injections of a standard 15 ppm BPA solution (Table 4), which meets the International Council for Harmonization criteria (ICH, 1996).

**Analysis method validation using KCKT**

**Calibration curve construction and linearity testing**

A calibration curve was constructed under these optimized conditions using standard BPA concentrations of 2, 4, 6, 8, 10, and 15 ppm (Table 5). The calibration curve equation obtained was $y = 4.1465x - 75.602$, and the correlation coefficient ($r = 0.9998$) meets the ICH linearity standards (ICH, 1996).

**LOD LOQ determination**

The LOD and LOQ were derived from the calibration curve regression equation as 0.2870 µg/mL and 0.9563 µg/mL, respectively.

**Selectivity test**

In selectivity testing, BPA solutions subjected to the same extraction process as beverage samples showed no difference in retention time compared to BPA standards under optimized conditions (9.714 min), while a matrix solution produced no BPA signal, indicating selectivity for BPA as shown in Fig. 1.

**Accuracy and precision**

Accuracy and precision were assessed from the results of three replicates each at 2 and 4 ppm and 6 replicates at 6 ppm. Data obtained met the smart criteria, with accuracy within 80–110% and %KV precision no more than 5% (Table 6). [8,20].

**Beverage sample BPA analysis**

Samples A, B, and C had expiry dates in 6 months, and samples D, E, and F had expiry dates in 2 years. Samples A, B, and C were stored in a fridge, at room temperature, or in an oven for 30, 60, or 120 min before the extraction. Results are presented in Table 7. One brand (A) exhibited BPA contamination even when refrigerated and contamination increased with heating, while another brand (C) exhibited contamination only after heating.

**DISCUSSION**

We have established a GC-based method for the detection of BPA in canned beverages with high linearity, sensitivity, specificity, and accuracy. This method was able to detect relatively small amounts of BPA in extracted beverage samples. In fact, one sample with an expiry date in 6 months exhibited BPA contamination even when stored under refrigeration. Further, contamination increased when the samples were heated. In general, BPA levels were higher in cans closer to the expiry date (6 months vs. 2 years). These results suggest that BPA contamination of canned beverages is possible and is exacerbated by improper storage, underscoring the importance of this optimized GC-based detection method.

This BPA contamination may be caused by layering differences (type, amount, etc.) or the sterilization process (high temperature) used by many canned beverage manufacturers. Unintentional heat exposure during storage and transportation may also contribute. Xu-Liang et al. measured BPA concentrations of 0.18–0.45 µg/l in 72 canned beverage samples, with 150°C to 260°C at 10°C/min.
Table 2: Optimal temperature results using a 100 ppm BPA standard

| Injection Volume | Area (µV/s) | Retention Time (min) | Theoretical Plates (N) | HETP | Follow-up Factor (Tf) | Resolution (R) |
|------------------|-------------|----------------------|------------------------|------|----------------------|----------------|
| 1 µL             | 205         | 9.698                | 15875.709              | 0.189| 1.248                | 89.234         |
| 2 µL             | 300         | 9.704                | 13656.681              | 0.219| 1.093                | 84.114         |
| 3 µL             | 572         | 9.714                | 36615.439              | 0.081| 1.367                | 90.016         |

Table 3: Optimal flow rate using a 15 ppm BPA standard

| Flow Rate (mL/min) | Area (µV/s) | Retention Time (min) | Theoretical Plates (N) | HETP | Follow-up Factor (Tf) | Resolution (R) |
|--------------------|-------------|----------------------|------------------------|------|----------------------|----------------|
| 0.8                | 226         | 10.553               | 53060.240              | 0.056| 1.073                | 134.953        |
| 1                  | 508         | 9.740                | 36846.159              | 0.031| 1.126                | 71.932         |
| 1.2                | 594         | 77.66.668            | 776.66.668             | 0.000| 3.015                | 31.080         |

Table 4: System comparability test for 15 ppm BPA

| Peak Wide (µV/s) | Retention Time (min) | Standard Deviation (SD) | Variation Coefficient (%KV) | Theoretical Plates (N) | HETP | Follow-up Factor (Tf) | Resolution (R) |
|------------------|----------------------|-------------------------|----------------------------|------------------------|------|----------------------|----------------|
| 572              | 9.714                | 7.98878                 | 1.3771.377                | 36615.44               | 0.081| 1.367                | 90.016         |
| 572.3            | 9.721                | 7.78878                 | 1.3771.377                | 34450.65                | 0.087| 1.183                | 143.478        |
| 564              | 9.721                | 36677.92                | 1.3771.377                | 36902.85                | 0.081| 1.126                | 143.725        |
| 567              | 9.711                | 36425.84                | 1.3771.377                | 36425.84                | 0.081| 1.057                | 169.779        |
| 568              | 9.709                | 41849.312               | 1.3771.377                | 41849.312               | 0.071| 1.148                | 169.690        |

Table 5: Calibration curve, LOD, and LOQ BPA data in chosen condition

| Concentration (ppm) | Peak Wide (µV/s) | Yi | (y−yi)² | S (y/x) | LOD (µg/mL) | LOQ (µg/mL) |
|---------------------|------------------|----|---------|---------|-------------|-------------|
| 2                   | 11               | 7.368 | 13.191 | 3.9675 | 0.287 | 0.956 |
| 4                   | 91.1             | 90.338 | 0.581 | 7.885  | 0.771 | 25.503 |
| 6                   | 170.5            | 173.308 | 7.885 | 25.503 | 169.779 |
| 8                   | 255.4            | 256.278 | 7.885 | 25.503 | 169.779 |
| 10                  | 334.4            | 339.248 | 25.503 | 25.503 | 169.779 |
| 15                  | 550.8            | 546.673 | 17.032 | 25.503 | 169.779 |

LOD: Limit of detection, LOQ: Limit of quantitation

Fig. 1: Chromatogram of selectivity test for sample solution (a) and 100 ng/mL standard BPA solution (b). Conditions were 3 µl injection volume and gradient temperature (10°C increase/min from 150 to 260°C)
samples from local supermarkets in Canada using a GC-MS method with a LOD of 0.0074 µg/L [9,21]. Therefore, the BPA concentrations found in the current study are relatively high.

In this study, samples B and C still meet the safety standards of the BPOM and the US Environmental Protection Agency limit of 0.6 ppm. However, sample A is not safe for consumption as the BPA concentration was above 0.6 ppm. If an adult weighing 60 kg consumed one sample A canned beverage (330 mL) each day, BPA exposure would equal 1.325 µg/kg body weight/day, which greatly exceeds the tolerable limit of 5 µg/kg BW established by the EFSA. Exposure will be even higher if the beverage is not consumed immediately or is stored improperly, such as at high temperature. Thus, storage conditions during transportation and before sale may be critical for the reduction of BPA contamination risk.

CONCLUSION

We have developed and validated an optimized GC-based method for the detection of BPA contamination in canned beverages. The method uses a long HP-1 capillary column (column length 30 m, inside diameter 0.25 mm, and film thickness 0.25 µm), flame ionization detection, a gradient column temperature (150°C to 260°C at 10°C/min), and nitrogen as the gas carrier at a flow rate of 1 mL/min. Total analysis time was 14 min. The method meets the international standards for linearity (r=0.998 linearity) with LOD of 0.287 µg/mL and LOQ of 0.947 µg/mL. Storage temperature and storage time facilitate BPA migration into canned beverages. BPA concentrations appear higher (closer to the expiry date) than newly distributed products.

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