

## Determination of Bisphenol A Using a Supramolecule and Photo-CIDNP

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As a type of endocrine disruptor, bisphenol A (4,4'-isopropylidenediphenol : BPA)<sup>1</sup> has many harmful effects on the human body. However, bisphenol A is still widely used as a dental treatment agent and in the production of polymers.<sup>2-6</sup> Detection methods of small amounts of this compound in environmental samples have been developed by many analytical and environmental scientists.<sup>7-9</sup> Even with these extensive detection methods, we are still confronted with difficulty on how to make a precise and fast determination of bisphenol A. This report details a new detection method using a supramolecule and compares its effectiveness with that of other methods.

### Experimental Section

**Apparatus.** <sup>1</sup>H and <sup>13</sup>C NMR spectra were recorded using a Varian Unity Plus 300FT-NMR Spectrometer and a Bruker AC-200 FT-NMR Spectrometer. Mass spectra were recorded using a Micromass Platform II GC/FAB/MS. UV/vis absorption spectra were recorded at 25 °C, using a Shimadzu UV 3101-PC spectrophotometer.

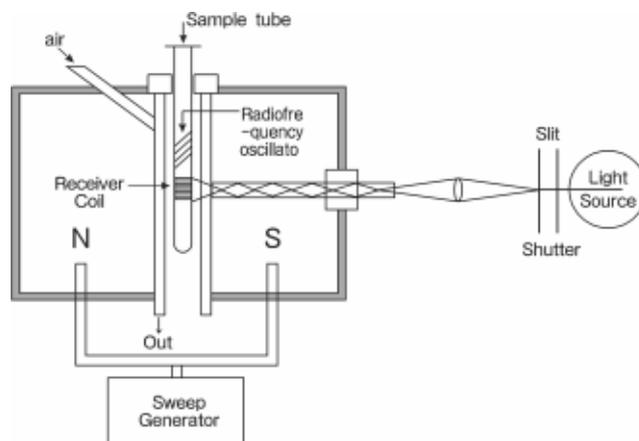
**Preparation of Indicator.** A type of supramolecule, 4,4'-bis(dibenzylammonium)-bipyridyl-tetrahexafluorophosphate (4-H<sub>2</sub>)<sup>4+</sup>(4PF<sub>6</sub>)<sup>-4</sup> was prepared by Stoddart's method.<sup>10</sup> A solution of 4,4'-bipyridine (36 g, 0.23 mol) in 2.5 L of dimethyl sulfoxide was added to a warm solution of HCl salt (260 g, 0.92 mol) of *N*-benzyl-*para*-chloromethylbenzylamine in dimethyl sulfoxide (2.5 L). The reaction mixture was stirred under reflux for 48 hrs. The reaction progress was detected by using UV/vis absorption spectra. A white precipitate formed and was filtered off after the suspension had been cooled to room temperature. The solid was washed with dimethylformamide and diethyl ether before being dissolved in water. A saturated aqueous solution of ammonium hexafluorophosphate was added until no further precipitation occurred. The white solid (4-H<sub>2</sub>)<sup>4+</sup>(4PF<sub>6</sub>)<sup>-4</sup> was filtered off and dried, yield: 165 g, 61%, m.p. 240 °C with decomposition. FAB/MS: *m/z* 1013 [M-PF<sub>6</sub>]<sup>+</sup>; <sup>1</sup>H NMR (CD<sub>3</sub>COCD<sub>3</sub>): δ = 4.63 (s, 4H), 4.70 (s, 4H), 6.21 (s, 4H), 7.40-7.50 (m, 6H), 7.50-7.60 (m, 4H), 7.73 (s, 8H), 8.77 (d, 4H), 9.44 (d, 4H).

**Procedure.** Approximately 50 μL of acetone-*d*<sub>6</sub> were placed into a one mL serum bottle, and the bottle was stoppered with a rubber septum cap. The air in the bottle was

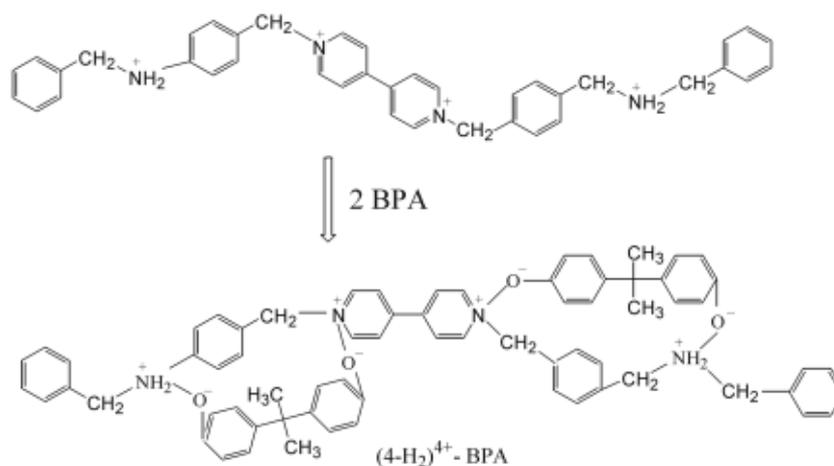
purged with argon gas using a micro syringe. One-half milliliter of (4-H<sub>2</sub>)<sup>4+</sup>(4PF<sub>6</sub>)<sup>-4</sup> reagent in acetonitrile-*d*<sub>6</sub> was then added to the serum bottle with a micro syringe. The derivative, (4-H<sub>2</sub>)<sup>4+</sup>(4PF<sub>6</sub>)<sup>-4</sup> solution was shaken continually until the sample was completely dissolved in the acetone-*d*<sub>6</sub>. A 0.1 μL aliquot was then pipetted and injected into the probe, as adopted from Wans method.<sup>11</sup>

Detection involved using a technique of photochemically induced magnetic polarization (photo-CIDNP) as shown in Figure 1. Irradiation from the side of the sample probe was performed using a high-intensity UV light beam source. A mixture of the (4-H<sub>2</sub>)<sup>4+</sup>-BPA solution was placed in a quartz cuvette, degassed, and sealed. The sample was irradiated in a photo-CIDNP reactor equipped with uncoated low pressure mercury lamp (254 nm) for 1 hr. The formation of the product was observed on an optical bench equipped with a high pressure mercury lamp (Oriol HBO 400 W in a 7340, lamp housing) and an Oriol high intensity monochromator (Model 7240) with a 5 nm band pass at 254 nm. The irradiation beam was focused on the sample solution at a critical level: 2.8 mm above the receiver coil. After irradiation the polarized product, (4-H<sub>2</sub>)<sup>4+</sup>-BPA, was observed inside the receiver coil.

Canned beverage samples were purchased in Busan, South Korea, between March and April 2002. The cans were coated with epoxy over a total area of 2.8 to 3.3 ng/cm<sup>2</sup>. The concentration was converted into the same ratio of coated



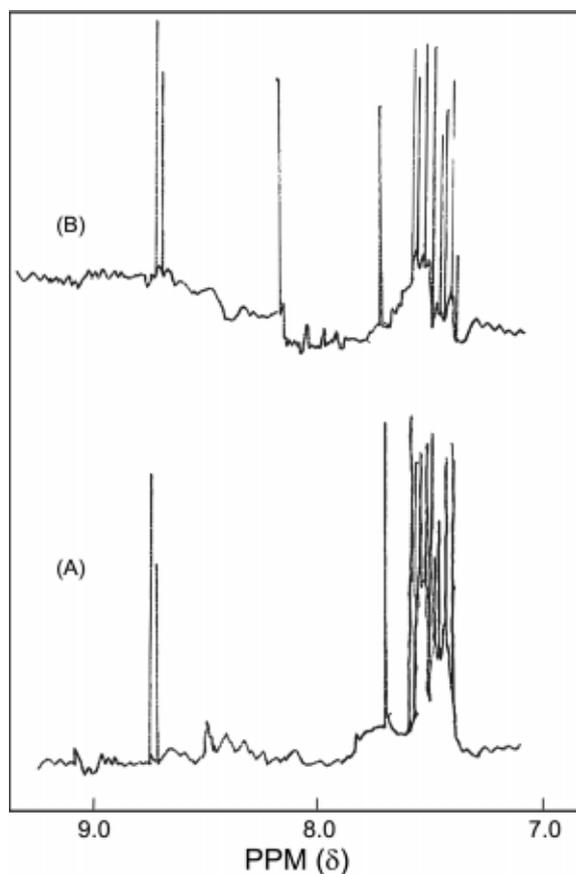
**Figure 1.** Schematic diagram of the irradiation arrangement in photo CIDNP detection.

**Scheme 1**

area after a determination for the same sample. The samples of beverage were concentrated to 7.0-8.0% (w/v), using a rotary vacuum evaporator, then they were extracted in chloroform.

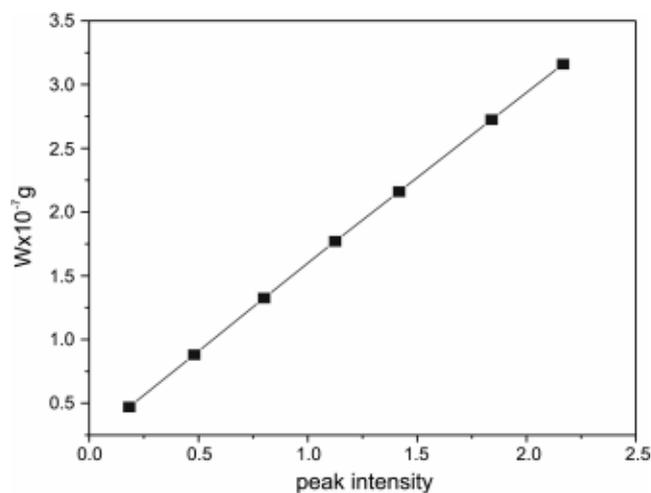
### Results and Discussion

In the determination of bisphenol A, two phenoxide anions combining with two cation sites of (4-H<sub>2</sub>)<sup>4+</sup>(4PF<sub>6</sub>)<sup>-4</sup> were observed, as shown in Scheme 1. The proton NMR



**Figure 2.** Proton-NMR Spectra recorded (A) before and (B) after photo -CIDNP of 4-H<sub>2</sub><sup>4+</sup>-BPA.

peaks of (4-H<sub>2</sub>)<sup>4+</sup>(4PF<sub>6</sub>)<sup>-4</sup> are shown in the range of 7.40-7.50 ppm and 7.50-7.60 ppm with multiplets; 7.73 ppm with singlet and 8.77 and 9.44 ppm with doublets. Figure 2 shows the proton-NMR spectra, which were recorded before and after photo-CIDNP. The <sup>1</sup>H-NMR spectra agree with the expected results corresponding to the loss of PF<sub>6</sub><sup>-</sup> and to the binding of 2 moles of BPA, respectively. The intensity of <sup>1</sup>H-NMR spectra correlated well with the concentrations of BPA. The <sup>1</sup>H-NMR spectrum of (4H<sub>2</sub>)<sup>4+</sup>(4PF<sub>6</sub>)<sup>-4</sup> was not observed in any other peaks in the region of 7.9-8.6 ppm(δ). However, a new peak at 8.20 ppm(δ) appeared soon after the addition of BPA. It is supposed that the new peak corresponds with the product of the binding between pyridinium ions and phenoxide oxygen. The height of the peak is proportional to the concentration of BPA. Figure 3 shows the correlation between the concentration of BPA in the sample solution and the peak intensity in photo-CIDNP. When the concentrations of BPA were changed from 4.98 × 10<sup>-8</sup> g to 32.0 × 10<sup>-8</sup> g the intensities on the <sup>1</sup>H-NMR spectra changed from 0.232 to 2.080 linearly as shown in Figure 2. This means the photo-CIDNP method is a potentially accurate



**Figure 3.** Correlation between the concentration of bisphenol A in solution and the peak intensity in photo-CIDNP.

**Table 1.** A Typical Comparison of Bisphenol A Concentrations (ppb) in Canned Beverages

Brand	Photo-CIDNP <sup>a</sup>	Known Method <sup>b</sup>
1	3.8	3.7
2	4.7	4.9
3	4.2	4.2
4	5.7	5.8
5	8.5	8.3
6	4.8	5.3
7	6.4	6.9

<sup>a</sup>detected by using a supramolecule, (4-H<sub>2</sub>)<sup>4+</sup>-BPA. <sup>b</sup>Detected by using the method of ref.9.

way to detect bisphenol A in the sample. To compare the photo-CIDNP method with the published detection method,<sup>9</sup> the concentrations of bisphenol A in the same sample solutions were determined using both methods, as shown in Table 1. The concentrations of BPA determined by photo-CIDNP agree with those of the previous known detection method,<sup>9</sup> even though they show a little bit of deviation in a few beverage samples.

The photo-CIDNP method is potentially more useful to detect bisphenol A due to the simple preparation of the sample pretreatment and its faster determination compared with the previous known detection methods,<sup>7-9</sup> which require a solid phase extraction, concentration, determination, and finally confirmation using the combined techniques of GC, HPLC and GC/MS.

In summary, we have established a new determination method of bisphenol A, using a supramolecule and an application technique of photochemically induced magnetic polarization (photo-CIDNP). The probe intensities of photo-CIDNP are correlated linearly with the concentrations of bisphenol A in the spiked and canned beverage samples.

This determination is simple and fast compared with other reported methods. The photo-CIDNP method is shown to be a superior procedure to detect bisphenol A in environmental samples.

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