

Basic performance of solid-phase extraction device "Magic Chemisorber[®]"

[Background] Solid-phase micro extraction (SPME) is a simple and rapid sample preparation technique used to extract and concentrate organics in liquid or gaseous matrices. Organics extracted into the solid phase are thermally desorbed and analyzed using GC/MS. The traditional SPME fibers are easily damaged because they are quite fragile and they exhibit a limited extraction capacity due to the small amount of sorbent on the fiber surface. To address these issues, a durable solid-phase extraction device, the "Magic Chemisorber®" (MC), was developed by Frontier Laboratories Ltd. In this report, the basic performance of the Frontier MC is described.

[Experimental] The Frontier MC (Fig. 1) consists of a deactivated titanium capillary tube (L=6 or 30 mm, i.d. =1.2 mm, o.d. =1.6 mm) coated with a thick film (500 µm) of PDMS (polydimethysiloxane). The PDMS is chemically bonded to the titanium tube which stabilizes the PDMS during the thermal desorption of the extracted organics. Before use, the tube is conditioned by heating it (150-280°C at 20 °C/min, 10 min hold) in a helium atmosphere. A pesticide, benfluralin (bethrodine MW 335.3) was used to demonstrate the performance of the MC. A series of aqueous standard solutions ranging from 2.5 to 2.5 x 10⁴ ng/L were prepared. The extracted benfluralin on the MC surface was analyzed using both evolved gas analysis (EGA)-MS and thermal desorption (TD)-GC/MS.

[Results] The EGA-MS thermogram of benfluralin extracted from an aqueous solution (40 mg/L) is shown in Fig. 2. The thermogram was obtained by heating the MC/benfluralin sample: 50 to 280°C at 40 °C/min. The desorbed benfluralin was measured using the MS in the SIM acquisition mode (m/z 292). Analysis of the EGA thermogram indicated that the desorption process is completed when the temperature reaches 280°C. A typical plot of extraction time vs. peak intensity is shown in Fig. 3. The plot indicates that 45 min is required to reach the partition equilibrium of benfluralin at the water-PDMS interface. Analysis of a series of aqueous benfluralin solutions (2.5 to 2.5 x 10⁴ ng/L) shows that the linear dynamic range is ca. 10⁴. Fig. 4 shows a mass chromatogram of 20 mL distilled water spiked with 2.5 ng/L benfluralin. The detection limit calculated from the SN ratio is 0.5 ng/L. These results illustrate the usefulness of the MC as a solid-phase extraction device.



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