Emergent chemical in sewage treatment plant: di-(2-ethylhexyl) phthalate

E. Lovatel¹, L. Calábria² & I. Nascimento³

¹Chemical Engineering Department, University of Caxias do Sul, Brazil
²Materials post-graduate program, University of Caxias do Sul, Brazil
³Center of Exact Sciences and Technology, Institute of Environmental Sanitation, University of Caxias do Sul, Brazil

Abstract

In this work a simple gravimetric method was described for the study of di-(2-ethylhexyl) phthalate (DEHP) in wastewater samples from the inflow and outflow points of the sewage treatment plant (ETE) at the University of Caxias do Sul (UCS). The organic phase of the sample was precipitated by adding concentrated HCl. The precipitate was filtered in glass columns packed with treated cotton. After drying, the columns containing the precipitate were carefully weighted and extracted with n-hexane. After solvent evaporation, the organic extract was re-diluted and analyzed by Gas Chromatography with Flame Ionization Detection (GC/FID). The average rate of DEHP removal was 27.50%. The median DEHP concentration was 6.01 µg mL⁻¹ at the outflow point. This DEHP concentration is well above the maximum concentration level reported in the literature.

Keywords: plasticizers, liquid residues, GC/FID, DEHP.

1 Introduction

Plasticizers are organic esters added to polymers to facilitate processing and to increase flexibility and toughness of the final product by internal modification of the polymer molecule. Some of these compounds are called phthalates and dominate polymer production. DEHP in many cases is the sole general-purpose plasticizer produced.

DEHP can interfere in the functioning of the endocrine system and in the action mechanism of hormones. They are called endocrine deregulators or...
endocrine disruptors and can cause the following problems: reproductive anomalies (morphological and functional gonadal dysfunction, e.g., infertility and decreased libido) and congenital malformations [1-3]. Bornehag and co-workers [4] found associations between dust concentrations of specific phthalate esters in house dust and asthma, rhinitis, and eczema in children. Phthalate esters were also studied in the serum of young Puerto Rican girls with premature breast development [5]. High levels of dimethyl, diethyl, dibutyl, and di-(2-ethylhexyl) phthalates and their main metabolite mono-(2-ethylhexyl) phthalate were as detected in 28 (68%) samples from the larche patients. The long term exposure to high levels of DEHP can produce damages to the liver and testicles in mammals and cause death in aquatic species. Bisphenol A, another important plasticizer, is suspected of acting as an environmental estrogen [6]. Plasticizers can also influence the mobility and bioavailability of toxic substances like polychlorinated biphenyls and metal ions by changes in their water or lipid solubility [7].

As DEHP is used specially in soft polyvinyl chloride production [6], it is widespread in all environments, since it is used in packaging, clothes, films, paints, adhesives, cosmetics, ink printers and many other products. One of the main forms of exposure to phthalates is through food, in addition to medical materials and by occupational contact [8]. On the other hand, DEHP was also detected in mineral, ultra pure and tap waters [9]. Other kinds of samples where DEHP was detected are: municipal solid waste compost [10], sewage and wastewater treatment sludge [11], river sediments [7], landfill leachate [6, 12, 13] and swine slaughterhouse wastewaters [14], among others.

Most of the DEHP concentration that enters the treatment plants remains adsorbed to the waste sludge and suffers biodegradation [15]. On the other hand, the DEHP remaining in the aqueous phase is much less biodegraded. In this way, plasticizers can persist in the environment and contaminate superficial and groundwater.

A number of studies have shown the removal of phthalates in wastewater treatment plants [16-18]. The sample extraction methods in those and several other works were based on liquid-liquid extraction (LLE) or solid-phase extraction (SPE). In the LLE the employment of larges volumes of sample (1 – 2 L) and organic solvents and the recovery grade and detection limit depend on solvent purity. SPE has the advantage of a great reduction of the sample volume (usually 100 mL), but the extraction of water samples containing solids or heavily contaminated by particulate matter can block the cartridges and the previous sample must be filtered. The SPE can also be performed with disks replacing the cartridges but the sample’s volume is drastically enhanced (3 L). In a solventless extraction method called SPME [19, 20], for the extraction of semi-volatile compounds, such as pesticides, PAHs and phthalates from aqueous matrices, only compounds with high octanol-water partition coefficients ($K_{ow} > 4$) can be extracted with high recovery grade. Thus, simple and low cost aqueous samples extraction methods that require minimum amounts of sample and organic solvents are still needed.
The main objective of this work is to investigate the presence of DEHP at inflow and discharge points of a sewage treatment plant, using a simple gravimetric method modified from the one proposed by Thurman and Malcom [21] for sample extraction. The instrumental analysis was performed by Gas Chromatography with Flame Ionization Detection.

2 Materials and methods

The physical-chemical characteristics of the samples are shown in Table 1.

Table 1: Physical-chemical characteristics of the samples (mg L\(^{-1}\)).

<table>
<thead>
<tr>
<th>Extraction</th>
<th>Inflow</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>4</th>
<th>5</th>
<th>Average</th>
</tr>
</thead>
<tbody>
<tr>
<td>QOD</td>
<td></td>
<td>612</td>
<td>632</td>
<td>1879</td>
<td>610</td>
<td>282</td>
<td>803</td>
</tr>
<tr>
<td>BOD</td>
<td></td>
<td>433</td>
<td>407</td>
<td>784</td>
<td>389</td>
<td>137</td>
<td>430</td>
</tr>
<tr>
<td>Suspended Solids</td>
<td></td>
<td>51</td>
<td>71</td>
<td>136</td>
<td>153</td>
<td>45</td>
<td>91.2</td>
</tr>
<tr>
<td>Outflow</td>
<td></td>
<td>1</td>
<td>2</td>
<td>3</td>
<td>4</td>
<td>5</td>
<td>Average</td>
</tr>
<tr>
<td>QOD</td>
<td></td>
<td>72</td>
<td>44</td>
<td>111</td>
<td>99</td>
<td>149</td>
<td>95</td>
</tr>
<tr>
<td>BOD</td>
<td></td>
<td>51</td>
<td>62</td>
<td>82</td>
<td>57</td>
<td>34</td>
<td>57.2</td>
</tr>
<tr>
<td>Suspended Solids</td>
<td></td>
<td>63</td>
<td>44</td>
<td>56</td>
<td>99</td>
<td>92</td>
<td>70.8</td>
</tr>
</tbody>
</table>

2.1 Samples and materials

The ETE-UCS treats, exclusively, 100 m\(^3\)/d of sewage generated at the university (ca. 20,000 people). The treatment system comprises an aeration lagoon (hydraulic residence time: 4 d), a sedimentation lagoon (hydraulic residence time: 2 d) and two maturation lagoons (total hydraulic residence time: 26.9 d). Monthly (from December 2007 to April 2008) wastewater samples (6 L) were collected from the sewage treatment plant of the Caxias do Sul university located in Caxias do Sul, Rio Grande do Sul, southern Brazil, in glass bottles (1 L capacity) with caps protected by aluminum foil to prevent contamination. The sampling period was from 8:00 am to 8:00 pm, every 2 h. All the samples were mixed and 1 L sample was collected from the mixture (composite sample). The samples were filtered at low pressure, maintained at 4 °C and protected from light until the extraction procedures (at maximum 24 h after the collection). All the solvents were distilled twice in glass apparatus. All reagents and solvents were of analytical grade and distilled twice in a glass apparatus, when necessary. All glass material was washed with n-hexane, acetone and dichloromethane and dried at 120 °C for 4 h. Plastic or rubber materials were not used to avoid contact with samples or solvents.

2.2 Sample extraction

The sample extraction was a modified version of the one proposed by Thurman and Malcom [21] for the isolation of aquatic organic substances and utilized by
Calace and Petronio [22] for the characterization of high molecular weight organic compounds in landfill leachate. The extraction procedures were as follows:

a) A volume of 100 mL of the sample was acidified (until pH ~2) with concentrated HCl and magnetically stirred for 15 min;

b) After precipitation of the organic phase (ca 20 min), the sample was filtered in a glass column (7.0 cm long x 1.0 cm i.d.) packed with 1.0 gram of cotton previously extracted (Soxhlet extraction by 4 h with n-hexane);

c) After drying (4 h at 35°C) the organic phase was extracted with 5 mL of n-hexane;

d) After volume reduction to 1.0 mL, the organic extract was analyzed by Gas Chromatography.

2.3 Instrumental analysis

A Perkin Elmer Gas Chromatograph model Autosystem XL, with Flame Ionization Detector was used for identification and quantification of DEHP by the internal normalization method. An Elite-5 fused silica capillary column (30 m x 0.25 mm i.d. crossbond 5% diphenyl – 95% dimethyl polysiloxane, 0.25 µm film thickness) was used for the GC separation using the following oven temperature program: 150°C (5 min hold) heating to 220°C at 3°C/min and heating to 300°C at 10°C/min (5 min hold). The injector temperature was 250°C. The volume of 1.0 µL was injected (triplicate of injection) in the split mode (1:50).

2.4 Recovery grade and detection limit

The recovery grade was measured by the extraction of a real sample spiked with a DEHP standard solution (resulting in a net concentration of DEHP of 20µg mL⁻¹). The detection limit was evaluated by successive dilutions of a DEHP standard solution (100 µg mL⁻¹). The detection limit was 0.5 µg mL⁻¹ and the detector response was linear between 0.5 and 25 µg mL⁻¹ (R² = 0.97; equation: y = 3.726x + 7.384).

The recovery grade was 75.95% (±5.13) and lower than the 95% reported by Ballesteros et al. [23] but those authors employed more complex sample extraction and preparation methods (Solid-phase extraction and silylation) and Gas Chromatography with Mass Spectrometry detection (GC-MS).
At the inflow point, the median concentration of DEHP was 8.29 µg mL\(^{-1}\) and the concentrations ranged from 3.50 (±0.90) to 26.83 (±2.69) µg mL\(^{-1}\). At the outflow point the median concentration was 6.01 µg mL\(^{-1}\) and the concentrations ranged from 3.49 (±0.95) to 20.89 (±1.16) µg mL\(^{-1}\).

The median DEHP concentration found in our work at the outflow was much higher than the 0.72 µg g L\(^{-1}\) found by Furtmann [16] in industrial sewage wastewaters. The DEHP removal grade (27.50%) was lower than the 90%
reported by Paxéus et al. [17]. These results make it quite clear that the treatment system of the ETE-UCS will require much improvement as far as DEHP removal is concerned.

Since DEHP has water solubility as low as $5.10 \times 10^{-4}$ mg L$^{-1}$ [24], a high adsorption of this chemical to the surface of the suspended solid particles and organic matter must be expected. In the studied samples (Table 1), the QOD and BOD removals were relatively high (88.17 and 86.70%, respectively) but the suspended solids remain high (22.37% reduction). These sample characteristics can enhance the DEHP partitioning and explain the high median concentration of DEHP at the exit point. Yamamoto et al. [25] reported a possible relationship between relatively high BOD values of landfill leachate samples and contamination by bisphenol A (a well known plasticizer that is also suspected of endocrine disrupting action).

The PVC materials (like the tubes used in water and wastewater canalization) can be another source of DEHP contamination. According to Tienpoint et al. [26] the PVC concentration in wastewater treatment plant sludge ranged from 18 to 508 µg g$^{-1}$. If plasticizers are contained within a plastic matrix, one would expect that they were not very available for biodegradation [15].

4 Conclusions

The gravimetric method employed in this work is very simple, with a good recovery grade and it can be applied to heavily contaminated water samples, without the limitations normally associated to the traditional extraction methods. The amounts of DEHP at the ETE-UCS outflow are well above the maximum limit allowed by official regulators, as the US-EPA, and this may be due to the physical-chemical characteristics of the sample, mainly suspended solids and organic matter content. The results suggest the need for additional treatment of the wastewater for DEHP removal and replacement of the PVC tubes of the treatment plant to avoid the post-contamination of the samples by plasticizers.

Our next step will be to evaluate how the organic matter and the PVC content can affect DEHP partitioning in the studied samples.

Acknowledgements

To CNPq for the financial support and the LASAN of the Caxias do Sul University for the physical-chemical analysis.

References


