Effect of the Bromine-Based Flame Retardant Plastic Pyrolysis of Hydrotalcite

N. Morita¹, Y. Kawabata¹, T. Wajima², A. T. Saito², and H. Nakagome²

¹Department of Urban Environment System at Chiba University, Japan.
²Chiba University, Japan.

Abstract. In this study, a method is presented to decrease halogen compounds in the product oil from thermolysis of polystyrene and polypropylene mixed plastic spiked with tetrabromobisphenol A. A mixture of hydrotalcite and plastic was pyrolyzed in a glass reactor at 400 °C under a nitrogen atmosphere. Bromine compounds in the residual substances were measured. The yield of product oil increased using hydrotalcite as an additive. The bromine compounds that were the major ingredients in the oil after thermolysis at 400 °C from the mixed plastic, which also included toluene, ethyl benzene, styrene, and 1-methylethyl benzene, were 2-bromohexane, 3-bromo-1-propenyl benzene, 4,5-dibromodecane, 1-bromomethylbenzene, 3-bromophenol, and 4-bromo-2,6-dimethylbenzaniline. However, bromine compounds were not detected in the product oil, residue, or gas when hydrotalcite was added. After the thermolysis of the plastic, bromine compounds in the product oil may decrease because bromine was captured by the added hydrotalcite.

1 Introduction

Plastics originate from fossil fuels and are becoming a problem because of the large quantity of waste. Methods for recycling plastic include thermal, chemical, and material recycling. Chemical recycling by pyrolysis is capable of returning plastics to their raw materials, and has attracted attention for making plastic derived fuel. There is an ever increasing volume of electronic products and associated waste electrical and electronic equipment (WEEE). Moreover, new models of electronic devices are replacing the old at a higher rate than ever before. WEEE includes computers, cellphones, hair dryers, and refrigerators, as well as the cathode-ray tube television [1]. However, the flame retardant materials added in these plastics form a product oil that includes halogen compounds. There are many recyclable materials, e.g., glass and metal; however, the energy consumption of the recycling process is problematic. Protecting the environment is desirable, but there are now more than 15 types of plastic commonly used in electronics [2]. The quantity of WEEE per person in a member nation of the 27 European Union (EU27) countries was estimated as 17 kg per person in 2005 [3]. The EU has introduced laws for the reuse and recycling of plastics to reduce landfill, and control WEEE destruction by thermal decomposition [4]. Plastics can be converted into gas for use as chemical raw materials, and fuel. Oil and residual carbon produced by pyrolysis can also be useful [5]. There has been considerable research on pyrolysis using catalysts and the chemical recycling of plastic [6-17]. For example, the yield of the product oil can be increased using ZSM-5 zeolite red mud mesoporous molecular sieve in the pyrolysis of plastics [18, 19]. However, the toxic brominated flame retardant is a problem for plastic recycling of household electrical appliances [20-24]. The generation of harmful byproducts has been studied to prevent the release of halogen gas, and additives have been used to promote recycling [22]. For example, polybrominated compounds mixed with polypropylene (PP), polyethylene, polystyrene (PS), and polyvinyl chloride have been decomposed safely with a carbon complex of calcium carbonate [25]. Additionally, the quality of the pyrolysis oil has been improved using the commercial hydrogenation catalyst DHC-8 [26]. The yield of product oil was improved to 96% from 93% when the zeolite catalysts, H-USY or H-Beta, were used in a steam contact process for pyrolysis [27]. The content of the bromine compounds in the product oil was decreased by adding hydrotalcite (HT) in the pyrolysis of brominated plastic [28-30]. HT is composed of layers of hydroxides that can form a metal complex hydroxide, containing anionic hosts and Mg²⁺ and Al³⁺ interlamellar guests [31, 32]. HT is an ion exchanger [31-34] and a catalyst [35] for gas removal [36, 37], and these applications have been receiving increasing attention in recent years [38].

The high anion exchange ability of HT, which is similar to organic ion exchange resins, has attracted attention as an adsorbent for the removal of various pollutants from water. The anionic adsorption properties include substitution reactions with chemicals such as...
borate, nitrate, fluoride, phosphate, sulfate, chromate, arsenic acid, and selenic acid. The anionic adsorption mechanism of HT has been proposed as an interlamellar complex formed of anionic exchange reactions [39]. In addition, HT has attracted attention as a catalyst, and reports have shown a yield increase to 77% from 62% in biodiesel fuel generation [40, 41].

In this study, pyrolysis at 400°C of PS and PP plastic mixtures spiked with tetrabromobisphenol A (TBBA) is examined using HT. The effects of additives on the product oil and the behavior of bromine compounds is evaluated.

2 Materials and Methods

2.1 Materials

Plastic samples containing bromine were prepared by mixing 10 g of PS (PS) (Asahi Kasei Chemicals Corporation Co., Ltd.) and 10 g of PP (Teijin Co., Ltd.) with 2 g of TBBA (Tokyo Chemical Industry Co., Ltd., Tokyo, Japan). To reduce bromine compounds in the product oil, synthetic HT, KW-1000, (Kyowa Chemical Industry Co., Ltd.), sea sand (Wako Pure Chemical Industries, Ltd.), molecular sieve 4A (Wako Pure Chemical Industries, Ltd.), and molecular sieve 13X (Wako Pure Chemical Industries, Ltd.) were used as additives. The experimental conditions are listed in Table 1.

Table 1: Experimental conditions used in this study

| Condition | Contents of the sample                                      |
|-----------|-------------------------------------------------------------|
| I         | Polystyrene (PS) + Polypropylene (PP) + Tetrabromobisphenol A (TBBA) |
| II        | PS + PP + TBBA + Sea sand                                   |
| III       | PS + PP + TBBA + Molecular sieves 4A                        |
| IV        | PS + PP + TBBA + Molecular sieves 13X                       |
| V         | PS + PP + TBBA + Hydrotalcite                                |

2.2 Experimental apparatus and procedures

The experimental apparatus used in this study is shown in Figure 1. A 22-g quantity of PS+PP+TBBA, or mixtures with an additive (20 g) was added to a glass reactor. The reactor was purged with nitrogen gas at a flow rate of 50 mL/min for 60 min to remove oxygen. After oxygen substitution, the flow of nitrogen gas was stopped and the pyrolysis experiment was performed. The temperature of the plastic was measured as the initial temperature, and the reactor temperature was increased to the decomposition temperature (400°C) at a heating rate of 5°C/min. The gases produced by pyrolysis were condensed and recovered. Non-condensable gases, which pass through the condenser, were collected in a gas pack via bubbling in an alkaline aqueous solution. Residue remained in the reactor after pyrolysis. The mass balance of the pyrolysis products (oil, gas, and residue) was calculated after the experiment. The pyrolysis products obtained after experiments under each condition were analyzed.

3 Results and Discussion

3.1 Product oil yield

The yields of product oil from the pyrolysis experiments are shown in Figure 2, which varied for different additives. The case using HT showed the highest yield, which was approximately 1.5 times that of the additive-free case. This result indicates the catalytic action of HT [40, 41]. In addition, the gases generated by pyrolysis were all less than 1 L.
3.2 Product oil components

Components of the product oils are given in Table 2. All the oil products contained toluene, ethylbenzene, styrene, and 1-methylethyl benzene. Bromine compounds obtained from the product oil included 2-bromohexane, 3-bromo-1-propenyl benzene, 4,5-dibromodecane, 1-bromomethylbenzene, 3-bromophenol, and 4-bromo-2,6-dimethylbenzaniline.

The results in Table 2 show that bromine compounds were not detected in the oil produced under condition V, which is the plastic containing HT.

Table 2. Components of product oils obtained under each experimental condition.

| Compound                        | Condition |
|---------------------------------|-----------|
|                                 | I  | II  | III | IV  | V  |
| Toluene                         | ○  | ○   | ○   | ○   | ○  |
| 4-Methylheptane                 | ○  | ○   | nd  | nd  | ○  |
| 2-Bromohexane                   | nd | nd  | nd  | nd  | ○  |
| 2,3-Dimethyl-2-heptane          | ○  | nd  | nd  | ○   | nd |
| 2,4-Dimethyl-1-heptene          | nd | nd  | ○   | nd  | nd |
| 2,6-Dimethyl-3-heptene          | nd | nd  | ○   | nd  | nd |
| Ethyl benzene                   | ○  | ○   | ○   | ○   | ○  |
| Styrene                         | ○  | ○   | ○   | ○   | ○  |
| 2,4,6-Trimethyl-3-heptene        | ○  | ○   | nd  | nd  | ○  |
| 4,4,5-Trimethyl-2-hexene         | nd | nd  | nd  | ○   | nd |
| Cumene                          | ○  | nd  | nd  | nd  | nd |
| 1-Methylethylbenzene            | ○  | ○   | ○   | ○   | ○  |
| Phenol                          | ○  | nd  | nd  | nd  | nd |
| 3-Bromo-1-propenyl benzene      | nd | nd  | nd  | ○   | nd |
| 4,5-Dibromodecane               | ○  | nd  | nd  | ○   | nd |
| 1-Bromomethylbenzene            | ○  | nd  | nd  | nd  | nd |
| 3-Bromophenol                   | ○  | nd  | nd  | nd  | nd |
| 4-Bromo-2,6-dimethylbenzaniline | ○  | ○   | nd  | nd  | nd |
| 1,3-Diphenylpropane             | ○  | ○   | ○   | ○   | ○  |

○ : detected, nd : not detected

3.3 Residue analysis

The surface of each sample was observed by EDS after pyrolysis to analyze any bromine remaining in the residue and the element distribution maps are shown in Figure 3.

3.4 Bromine content in the gas

The analysis results of the gas generated by pyrolysis are shown in Figure 4 and Table 3. The gas obtained from thermal decomposition was analyzed for bromine-containing compounds. The bromine included in the plastic from the added TBBBA was 2 g, and the bromine in the generated gas was 1843 mg for all the conditions. Most bromine was distributed in the product oil and the residue. The decomposition of the plastic causes gases, such as H₂ and CH₄, to be expelled, but these were low after the addition of HT. This thermal decomposition reaction does not distribute bromine to the gas because of HT, and they are degraded in the product oil and residue.
Table 3. Components of the product gas

| Condition | H₂ | CH₄ | C₂H₄ | C₂H₆ | C₃H₆ | C₆H₆ |
|-----------|----|-----|------|------|------|------|
| I         | o  | nd  | nd   | nd   | o    | o    |
| II        | o  | nd  | nd   | nd   | o    | o    |
| III       | o  | nd  | nd   | nd   | nd   | o    |
| IV        | o  | o   | o    | o    | o    | o    |
| V         | o  | nd  | nd   | nd   | o    | o    |

○ : detected, nd : not detected

4 Conclusion

In this study, we investigated a method of reducing bromine compounds in the product oil obtained by the thermal decomposition of a TBBA-spiked PS and PP mixture using HT. Plastics containing brominated flame retardants can be thermally decomposed to product oils containing no bromine compounds by adding HT. The yield of product oil after adding HT improved. This indicates that the utility of the HT is high, which may be because of a catalytic effect. The residue after adding HT showed more bromine than without additive, suggesting the HT absorbed the bromine compounds. Thus, the product oil does not contain bromine. The efficacy of plastic thermal decomposition is increased by adding HT, because the bromine compounds are contained in the residue and reduced in product oil.

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