The Use of Methyl Ethyl Ketone in Nitric Acid Leaching Processes for Enhancement of Ag Recovery from Used X-ray Films

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A feasibility study was carried out to improve conventional recycling processes of used X-ray film, where silver was recovered mainly by concentrated nitric acid leaching. In the X-ray film, an emulsion layer containing silver is attached to polyester film base, and the enhancement of Ag leaching was expected by separating the emulsion layer from the polyester film base using methyl ethyl ketone (MEK), because the separation could increase the contact surface between nitric acid and Ag in the emulsion layer. First, the separation behaviors using MEK were investigated and the separation efficiencies of the emulsion layer increased with increasing temperature but decreasing film size and pulp density. The separation efficiency increased to 100% in MEK solution within 3 min under the following conditions; temperature 50°C; setting agitation speed 400 rpm; pulp density 50 g/L; film size 1 × 1 cm. In the followed nitric acid leaching test, the leaching result using the separated emulsion layer showed higher leaching efficiency than that using the unseparated X-ray film. Second, the effect of adding MEK to nitric acid leaching on the leaching efficiency of Ag was investigated by considering the amount of MEK added and temperature as experiment factors. The leaching efficiency of Ag increased with increasing the amount of MEK to 5%, but further addition of MEK to 7% rather reduce the leaching efficiency. With 5% of MEK addition, the leaching efficiency of Ag increased up to 95.9% in 120 min at 50°C whereas it increased and then decreased to 2.7% in the test at 70°C due to the formation of AgCl. These results indicate that the leaching efficiency of Ag could be enhanced using MEK in the nitric acid process for the recovery of Ag from the used X-ray film. [doi:10.2320/matertrans.M2017278]

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1. Introduction

Silver (Ag) has been used in various fields such as photography, radiography, electronics, photonics, electricity, catalysts, batteries, jewelry, silverware, dental materials, biomedicine, medicines and infectious diseases, and its demand has increased consistently.1) About 18% of silver demand has been supplied by recovering silver from photographic wastes.2,3) Silver, especially silver nanoparticle, has been reported to be very harmful to the human body when released into the environment.4,5) Therefore, the recycling of photographic wastes has been required to recover silver and prevent its release in the environment.6–8)

Figure 1 shows that the cross sectional structure of X-ray film. The emulsion layer is mainly composed of gelatin matrix, in which silver components are dispersed as silver halide such as AgBr, AgCl, and AgI6,7) and it has been reported that X-ray films contain around 1.5% of silver.2,3,8) Burning processes had been used to concentrate the silver but the processes have disadvantages such as the loss of film base.9) Hydrometallurgical processes have been recognized as an alternative.

Various leaching agents such as cyanide,9) nitric acid,10,11) and thiosulfate12) were investigated to dissolve silver from used X-ray film. As shown in Fig. 1, because the emulsion layer containing silver components is attached to the film base, only one side of the emulsion layer is exposed to the leaching agents, which could reduce leaching efficiencies of silver. It was expected that the leaching efficiency of silver could be enhanced by separating the emulsion layer.2,3,6,8,13–15) Some separation processes were developed using oxalic acid,6,13) sodium hydroxide,2,14) biologically-produced protease,3,8,15) Methyl ethyl ketone (MEK) has been well known as its relatively low cost and toxicity.16) It has been reported that this organic solvent could swell some resins, where swollen resin behaves like solution or suspension.17) Also MEK has been famous for cleaner or thinner for polyester because MEK could change the surface property of polyester used as film base in X-ray film.18) Thus, it was expected that addition of MEK could separate the emulsion layer from the film base. Therefore, two kinds of experiment sets were investigated; the separating behaviors of the emulsion layer in MEK solvent and the effects of MEK addition on the leaching of X-ray film in nitric acid solution. The effects of various experimental factors such as temperature, agitation speed, film size, and pulp density were examined in the tests.

2. Experiments
2.1 Materials

Used X-ray films were collected from a recycling company in Korea, and they were used for inspection during shipbuilding. The content of Ag was measured to be...
22800 mg/kg, which was analyzed by an analytical company (Yucheon Tech. Co.). The used X-ray films were cut to the proper sizes (1 × 1 cm, 2 × 2 cm, 4 × 4 cm) before experiments. Nitric acid (Junsei Chemical Co., Ltd., Japan) and methyl ethyl ketone (Junsei Chemical Co., Ltd., Japan) used in this study are of reagent grade.

2.2 Separation test of emulsion layer

Separation tests were performed in a 250 cm³ water-jacketed reactor, which was connected to a heating bath circulator (WCB 6, Daihan Scientific Co., Ltd., Korea), to maintain temperature of solution. The reactor was charged with 100 cm³ of MEK and capped with silicone stopper. After increasing the temperature of MEK to 30–60°C, the prepared X-ray films were placed into the reactor. During the test, the solution was stirred with magnetic bar at up to 400 rpm, where magnetic stirring system was chosen because the emulsion layer, after separating, floated and interrupted blade-type stirring. Separation efficiency was calculated at designated time interval with the following equation.

\[
\text{Separation efficiency (\%) = Ns/Nt \times 100}
\]  

where \(N_s\) indicates the number of X-ray film pieces separated and \(N_t\) is the whole number of X-ray film pieces.

2.3 Nitric acid leaching and analysis procedures

Nitric acid leaching was conducted in a 500 cm³ three-necked Pyrex glass reactor using a heating mantle to maintain temperature. The reactor was fitted with a stirrer and a reflux condenser to avoid solution loss at high temperatures. Leach solution was prepared by adding MEK to 1 M HNO₃. A 200 cm³ of leach solution was placed in the reactor and allowed to reach thermal equilibrium at desired temperatures (40–70°C). The used X-ray film pieces were added to the solutions and agitation speeds were set at 400 rpm. During the tests, 2 cm³ of the solution sample was withdrawn periodically at a desired time interval. The solution was filtered with a 0.45-µm membrane filter and then diluted with 5% HNO₃ solution. The concentration of Ag was measured with AA7000 atomic absorption spectrometry (Shimadzu Co. Ltd., Japan), and leach residue was analyzed with MESA 50 X-ray fluorescence (XRF) analyzer (Horiba Ltd., Japan). Leaching efficiency of Ag was obtained from the following equation;

\[
\text{Leaching efficiency (\%) = } \frac{M_s}{M_s + M_r} \times 100
\]  

where \(M_s\) and \(M_r\) represent the mass of silver in solution and in leach residue, respectively. The Ag concentration in the leach residue was measured after digesting with concentrated nitric acid, and, during the leaching test, the Ag contents in the leach residue was obtained as follows; after converting the Ag concentration measured in the next sampling, added the converted Ag content to that in final leach residue.

3. Results and Discussions

The effects of agitation speed (100 rpm to 400 rpm) on the separation of emulsion film from the used X-ray film were examined in MEK solution at 50°C with 20 g/L pulp density and 1 × 1 cm film size. The emulsion layers were separated completely from the film bases within 3 min regardless of the agitation speeds in the range between 100 rpm and 400 rpm (data not shown). As shown in Fig. 2, the used X-ray film piece (Fig. 2(a)) was separated into wrinkled-shape emulsion layer (Fig. 2(b)) and polyester film base (Fig. 2(c)). Since the polyester base is blue, it was easily distinguished from the black X-ray film, and the separation efficiencies could be calculated. It has been found that MEK could swell some resins¹⁷) or change the surface energy of polyester.¹⁸) The swelling of the emulsion layer was not observed in the test. Gelatin is a main component in the adhesive layer as well as in the emulsion layer although minor components are not disclosed. Therefore, since generally gelatin and polyester are not swollen by MEK, the separation of the emulsion layer from the film base would be caused by the change of surface energy of polyester due to MEK. Although the separation was completed rapidly at all agitation speeds used in this study, since the film pieces were observed to occasionally interfere with the agitation, the agitation speed was fixed at 400 rpm to ensure a stable agitation force in all subsequent separation tests.

The effects of temperature on the separation were investigated in MEK solution at 400 rpm with 20 g/L pulp density and 1 × 1 cm film size. The separation tests were conducted in the temperature range of 30°C to 60°C. Since the boiling point of MEK is 79.6°C, the temperature range was set to avoid the evaporation of MEK. As shown in Fig. 3, a faster separation is achieved at higher temperatures. The separation was completed within 3 min at 50 and 60°C, whereas it took 40 min at 30°C. Therefore, the temperature of 50°C was selected in all subsequent separation tests. The effects of pulp density were examined in MEK solution at 400 rpm and 50°C with 20–80 g/L pulp density and 1 × 1 cm film size. As shown in Fig. 4, the separation was achieved completely within 3 min at 20 g/L or 50 g/L pulp density, but, in the test at 80 g/L pulp density, the separation
efficiency was 54.1% at 12 min. In this study, a magnetic stirring system was employed because the separated emulsion layer floats and interrupt the stirring of blade-type stirrer. In the separation test with 80 g/L pulp density, the stirring was interrupted by the film samples, which could cause the low separation efficiency, and the test did not last more than 12 min. Further research will be required on favorable and proper stirring methods.

The polyester film bases are also important valuables, and thus, generally in the conventional recycling processes, used X-ray films are cut into an appropriate size (mostly 2-4 cm length) for subsequent polyester recycling before recovering Ag from X-ray films. Therefore, the effects of film size on the separation were investigated in MEK solution at 400 rpm and 50°C with 20 g/L pulp density. Three sizes (1©1 cm, 2©2 cm, 4©4 cm) of X-ray film were prepared, and the results were given in Fig. 5. The separation was completed within 3 min in the test with 1©1 cm size and within 6 min in the tests with 2©2 cm and 4©4 cm sizes. The solvents penetrate more easily between the layers when the film has the smaller the area.

A comparative leaching test was carried out in 1 M nitric acid solution at 50°C and 400 rpm with 10 g/L pulp density. The leaching behavior of silver during nitric acid leaching was well-established because the effects of agitation speed, pulp density, nitric acid concentration, and temperature were reported in the conventional studies. In this study, it was expected that the leaching efficiency of Ag could be enhanced by separating the emulsion layer from the film base, which could expose the both sides of emulsion layer to nitric acid. Therefore, for comparison, two kinds of sample were prepared; one is the X-ray film piece itself (1©1 cm size) as shown in Fig. 2(a), and the other is the separated emulsion layer as shown in Fig. 2(b). The separated emulsion layer pieces were collected from the preliminary separation test with 10 g/L pulp density and 1©1 cm X-ray film size. Figure 6 shows that the leaching results; the leaching efficiency of Ag increased to almost 100% (232 mg/L) within 120 min using the separated emulsion layer, while, in the leaching test with the X-ray film, the efficiency was 73.0% at 180 min. In the case of the leaching test with the separated emulsion layer, the color of the solution turned black within 30 min and then became a clear solution within 120 min. However, in that with the X-ray film, the emulsion layer was observed like black sludge on the surface of film base as a leach residue. This result indicates that the separation of the emulsion layer in advance is more preferable for the leaching of Ag from X-ray film.

Although the separation process using MEK showed beneficial effects of Ag recovery, it is not easy to add new process to the conventional process. In this study, a feasibility of adding MEK to nitric acid leaching process was investigated. As mentioned above, since the leaching behaviors of Ag in nitric acid solution were well-established, only the effects of MEK addition to nitric acid solution and temperature were investigated in this study.
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4. Conclusion

The use of MEK was investigated to enhance Ag recovery from the used X-ray film in the conventional nitric acid leaching processes. Two types of tests were carried out; nitric acid leaching of Ag after separating the emulsion layer from polyester film base using MEK, and adding MEK to nitric acid leaching of Ag from the used X-ray film.

The leaching efficiency of Ag from used X-ray film was enhanced by separating the emulsion layer from the polyester film base using MEK because the separation process increased the contact surface between nitric acid and silver in the emulsion layer. The separation efficiency increased with increasing temperature but decreasing pulp density and film size. Any significant effect was not observed in the agitation speed test. The feasibility of adding of MEK to nitric acid leaching was investigated. The addition of 5% MEK is most advantageous for recovering Ag from the used X-ray film. When MEK was added to nitric acid leaching, the leaching efficiency of Ag increased at up to 50°C but further increasing to 70°C rather reduce the leaching efficiency due to the formation of AgCl. These results indicate that silver could be recovered by using MEK at 1 M nitric acid concentration. It is also possible to improve the conventional process by adding MEK to the nitric acid leaching process without new plant. Since the boiling points of MEK and nitric acid are 79.6°C and 83°C, respectively, further study will be required to investigate the reuse of MEK and nitric acid employing a boiling process for economic and environmental efficiency.

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