Simplified Mercury Extraction from Coal Fly Ash for Quantification of Total Mercury by ELISA-based Immunoassay

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A simplified two-step mercury extraction procedure enabled the selective and reproducible mercury recovery from actual coal fly ash (CFA). The optimized extraction procedure involving conventional enzyme-linked immunosorbent assay (ELISA)-based immunoassay allowed the ultra-sensitive quantification of total mercury content in CFA. The total mercury content of 41 CFA samples were successfully determined using the above-mentioned method, and the results were in agreement with those obtained by standard instrumental analysis (thermal decomposition atomic absorption spectrometry) within a 15% coefficient of variation. Our method for total mercury quantification is not only simple but suitable for management of the mercury content at coal-fired electric power plants and landfill sites, which deal with large amounts of waste CFA.

Keywords Mercury, coal fly ash, immunoassay, simplified analysis, waste management

(Received September 9, 2019; Accepted December 2, 2019; Advance Publication Released Online by J-STAGE December 13, 2019)

Introduction

Coal-fired power plants, which account for ~41% of global electricity production,1 discharge large quantities of coal combustion residues, such as coal fly ash (CFA). Current environmental regulations require the collection of this ash, which is commonly carried out using a dust collector such as an electric precipitator or a bag filter. The resulting coal fly ash that is not reused in the production of cement and concrete is disposed of safely in a designated landfill or impoundment site.

Although the mercury present in CFA is relatively chemically stable,2,3 it can undergo methylation by environmental bacteria to produce highly neurotoxic organic mercury.4,5 Indeed, in river sediments close to large-scale coal ash spill sites, mercury methylation by sulfate-reducing bacteria has been suggested to take place through use of the sulfate ions present in the ash as a substrate.6,7 Furthermore, recent developments aimed at minimizing gaseous mercury emissions through the use of mercury controlling and capturing technologies,8 such as selective catalytic reduction, sorbent injection systems, and low temperature electronic precipitators, will likely lead to increased mercury content in the CFA.

Although frequent analysis is somewhat complicated and costly, evaluation of the mercury content in CFA and its distribution in waste disposal sites is essential to the management and risk assessment of the site. Conventionally, the total mercury content in environmental samples, such as CFA, have been determined by techniques including thermal decomposition atomic absorption spectrometry (TD-AAS),9 cold vapor atomic absorption spectrometry (CV-AAS),10 atomic fluorescent spectroscopy,11 inductively coupled plasma mass spectrometry,12 or induced coupled plasma atomic emission spectrometry.13 Although these instrumental methods are accurate and sensitive, the development of simpler and more cost-effective techniques is required.

In this context, novel simplified detection principles have been actively studied, with examples including the use of nucleic acids such as DNAzyme and oligonucleotides,14-16 task-specific ionic liquids,17,18 and functionalized nanoparticles.19,20 However, these methods suffer from interference from other chemicals present in the fly ash samples. As such, the excellent specificity and relatively low cost of immunoassay21,22 render them an attractive alternative. Our group previously developed an antibody that is specific to chelated mercury ion, i.e., (Hg2+)2EDTA,23,24 and developed an immunoassay for trace mercury ion analysis in environmental water samples.23 However, although our antibody exhibited good specificity, the assay interference from coexisting metal ions was an issue since some of these ions are present in significantly greater concentrations than mercury itself.

To address this issue, rapid and inexpensive purification of mercury from a sample extract containing multiple other metal species is necessary using either a chelating resin or an ion exchange resin (IER). Although the lower cost and milder elution conditions offered by IERs make them better suited than chelating resins for mercury purification,21 their application to samples with high concentrations of other species, such as extracts of environmental solid samples, has not yet been
examined in detail.

Thus, in this study, we developed a simplified two-step mercury-selective extraction method that can follow mercury using immunoassay to enable high-throughput and inexpensive quantitative determination of the mercury content in CFA. The proof of concept is demonstrated by a comparative analysis of 41 CFA samples from a coal-fired power plant; each of the samples was separately analyzed by TD-AAS, and mercury extraction was carried out by ELISA (enzyme-linked immunosorbent assay)-based quantitative determination for comparison.

Experimental

Reagents and chemicals

Two standard CFA samples, namely JSAC0521 (containing 140 μg Hg/kg) and JSAC0522 (mercury negative, <10 μg Hg/kg), were purchased from The Japan Society for Analytical Chemistry (Japan). Actual CFA samples were collected from the electrostatic precipitator systems of full-scale pulverized coal-fired power plants in Japan, which burn imported coal. The mercury standard solution (HgCl₂, 100 mg/L), potassium permanganate (Kmno₄, 99.5%), hydrochloric acid (HCl, 99.5%), nitric acid (HNO₃, 99.5%), hydroxylamine hydrochloride (HONH₃HCl, 97%), and strongly basic anion exchange resin (DOWEX 1 × 2, 200 - 400 mesh) were purchased from Wako Pure Chemical Industries, Ltd. (Japan). 5Sodium-2,2-dispersal of the ash by vigorous shaking, the mixture was placed in 50 mL plastic bottle (I-boy, ASONE Inc., Japan). Following a slight modification to our previously published extraction procedure, the sample of CFA (0.5 g) was dispersed in distilled, deionized water obtained from an in-house using solutions of 137 mM NaCl, 3 mM KCl, 20 mM phosphate buffer (10 mL; pH 6.5) was added to each well and incubated for 1 d at 4°C. After this time, the excess protein conjugate was removed by washing three times with PBSN (200 μL/well). An aliquot of the sample solution (standard mercury solution, or SPE eluate, 2 mL) was mixed with the EDTA solution (2 μL, 20 mM) and after a few minutes, the anti-Hg²⁺-EDTA antibody was added to wash the coexisting metal adsorbed on the ion exchange resin. The total mercury content in the CFA samples was accurately determined using TD-AAS with gold amalgamation (Nippon Instruments, Co., Ltd., Japan). The mercury ions present in the aqueous samples were determined using ICP-AES (Agilent 720-ES, Agilent Technologies, Inc., CA, USA). The content of all other metal ions present in the aqueous samples was determined using ICP-MS (Agilent 7900, Agilent Technologies, Inc., CA, USA). The mercury ions present in the aqueous samples was determined using ICP-MS (Agilent 7900, Agilent Technologies, Inc., CA, USA). A calibration curve was used for all instrumental determinations carried out herein.

ELISA for mercury determination

The antigen-protein conjugate (metal-EDTA-BSA) for ELISA was prepared (see Supporting Information) as described previously. In this study, Cd³⁺-EDTA, instead of Hg²⁺-EDTA, was employed as the quasi-antigen because of the instability of the latter in a protein-containing aqueous solution. The assay was carried out using a 96-well plate, where Cd³⁺-EDTA-BSA (50 μL/well, 2 μg/mL) diluted with 50 mM MES buffer (pH 6.5) was added to each well and incubated for 15 min at 37°C. After this time, the excess protein conjugate was removed by washing three times with PBSN (200 μL/well). An aliquot of the sample solution (standard mercury solution, or SPE eluate, 2 mL) was mixed with the EDTA solution (2 μL, 20 mM) and after a few minutes, the anti-Hg²⁺-EDTA antibody was added (20 μL, 1 mM final concentration), and the resulting mixture was incubated at 25°C for 15 min. The antibody-containing measurement sample (50 μL) was then added to the well, and incubation continued for 15 min at 25°C. After washing four times with PBSN (200 μL/well), the HLP labelled antibody to mouse IgG(H+L) (1.0 μg/mL, KPL 074-1806) diluted with PBSN (50 μL/well) was added and incubation continued for 30 min at 25°C. After this time, the excess secondary antibody was removed by washing five times with PBSN (200 μL/well). The colorimetric peroxidase substrate ABTS (100 μL/well, KPL 50-62-00) was then added and the mixture incubated for 15 min at 25°C. To stop the coloration, the peroxidase stop solution (50 μL/well, KPL 50-85-01) was added. Finally, optical measurements were carried out for each well using a multiwall plate reader (EL808, BIO-TEK instruments, Inc., VT). The results of the immunoassay were fitted using the four logistic functions as described in a previous report, using the least squares method.

Results and Discussion

Solid-phase purification

Several coexisting metals were found in the extract of standard CFA samples (Table S1, Supporting Information). In particular, the levels of Mn, Al, and Ca in the JSAC0521 extract exceeded that of mercury by a factor of several thousands, and trace Cd
but highly reactive to antibody in use, Table S2, SI) was found in both extracts. To eliminate the above coexisting metals, extract purification was investigated.

Initially, the Hg$^{2+}$ adsorption and elution conditions for the mockup extract were investigated. Thus, to evaluate the volume of IER required and the effects of supplementing the extract with HCl and HAC prior to passage through the column, the mock up extract (5 mL) containing 5 $\mu$g/L Hg$^{2+}$ (0.5 M H$_2$SO$_4$ and 1 g/L KMnO$_4$) supplemented or not supplemented with HAC (0.03 g/L final concentration) and HCl (0.1 M final concentration) was applied to the SPE column packed with different volumes of the ion exchange resin (i.e., 150, 250, and 350 $\mu$L). Here, supplementation with HCl facilitated the formation of a Hg$^{2+}$-chloride anionic complex that was adsorbed on the anion exchanging resin, while supplementation with HAC allowed one to scavenge KMnO$_4$, which would otherwise damage the resin. Following collection of the eluate, the Hg$^{2+}$ concentrations were measured, and as shown in Fig. 1, >250 $\mu$L of the ion exchange resin was required to achieve the optimal adsorption of Hg$^{2+}$ (>95.5%) regardless of HAC supplementation.

This adsorbed Hg$^{2+}$ from the mockup extract was then eluted using a 0.1 M Tris-HNO$_3$ buffer (10 mL, pH 7.5) and the various fractions collected. As shown in Fig. 2, the use of 4 - 5 mL of eluent resulted in the elution of 98.9 - 99.5% of the Hg$^{2+}$ where HAC had been added to the applied sample, whereas 10 mL of eluent was required to elute 82.4% of the Hg$^{2+}$ when HAC was absent from the system. The use of HCl and HAC supplementation of the extraction, in addition to 250 $\mu$L IER resin and 5 mL eluent were therefore applied for the following experiments.

Subsequently, elimination of the coexisting metal species from the CFA extract was examined. Thus, residual metals present in the SPE column after the addition of the extracts (5 mL) prepared from the standard CFA samples (JSAC0521 and JSAC0522) were determined using the collected eluate, as outlined in Table 1. Although the elimination rates of coexisting metal ions were 77 - 100%, non-negligible concentrations of Mn, Mg, Fe, Al, Ca, and Cd were present following extract addition (see Table 1). In particular, the elimination of Cd, which exhibits a strong cross reaction to the antibody, is desirable. Thus, to eliminate any residual metal from the column, washing was carried out using pure water and/or 0.1 M HCl. More specifically, following application of the extract prepared from the standard CFA samples, 5 mL of either or both washing solutions were applied, and finally the Tris–HNO$_3$ buffer (5 mL) was added to recover the Hg$^{2+}$.

As presented in Fig. 3, several metal species, including Cd, coexisted with the recovered Hg$^{2+}$ (76.7% for JSAC0521, see Table 2) when washing was omitted. However, washing was found to result in the removal of residual metal species and elimination of the coexisting metals from the resulting elution buffer (<10 $\mu$g/mL), as shown in Fig. 3. In particular, washing with HCl and H$_2$O gave superior results because of the efficient removal of coexisting metals (Cd, Mn and Al) to negligible levels in terms of cross reactivities of antibody in use$^{23}$ (Table S2, with relatively high mercury recovery (69.9%)). Therefore,
Finally, the reproducibility of the mercury recovery was tested using the developed purification procedure through its application to actual samples (CFA1 – 7). Thus, the extracted solutions obtained from the CFA samples were mixed with HAC and HCl prior to loading onto the column, and after column washing with HCl - H2O, the adsorbed Hg\(^{2+}\) was recovered using the elution buffer. As outlined in Table 3, the recovery rate of mercury ranged from 63.3 to 79.6% (average = 71.5%) and corresponded well with the result obtained using the standard sample (JSAC0521).

### Table 2 Mercury recovery through different washing procedures followed by elution

| Washing procedure | Recovery rate, % | CV, % |
|-------------------|-----------------|------|
| None              | 76.7 ± 3.5      | 4.5  |
| H2O               | 41.0 ± 3.9      | 9.6  |
| HCl               | 70.0 ± 3.4      | 4.9  |
| HCl–H2O           | 69.9 ± 3.8      | 5.4  |

* a. Recovery rate (%) was calculated as \([\text{Hg}^{2+}\ \text{concentration in eluate}/(\text{Hg}^{2+}\ \text{concentration in sample}) \times 100]\).

Actual sample analysis

To demonstrate the quantitative utility of the described analytical procedure, 41 actual CFA samples were pretreated by optimized extraction and then measured by ELISA. For this purpose, calibration curves were first obtained using the CFA samples with known total mercury content through the use of TD-AAS (two standard CFA samples, 35, 139, 181, 232, and 957 μg/kg). Each immunoassay plot was fitted using the sigmoidal regression \((A = 211, B = 1.38)\), as shown in Fig. 4. The detection limit of the total mercury content in the CFA samples calculated using the 3-sigma method was found to be 32.6 μg/kg.

Forty-one actual CFA samples were extracted and purified using the optimized process described above and then analyzed using our developed immunoassay. As presented in Fig. 5, an excellent correlation was found between the determined values and the standard TD-AAS technique (slope = 1.03, \(R^2 = 0.98\)). In addition, all data points presented a <15% deviation from the ideal correlation (slope = 1), which is sufficient for a quantitative method using immunoassay.\(^{27}\)

### Conclusion

We have reported the development of a simplified mercury-selective and reproducible extraction method to enable high throughput and inexpensive quantitative determination of mercury content in CFA using the conventional ELISA-based
immunoassay. The low cost (US$ 11.0/sample, see Table S3, SI) and simplicity of the complete process is believed to render it applicable for the monitoring of mercury in industrially produced CFA, which is a subject of worldwide concern. In the future, we will extend the analytical protocol developed herein to other solid wastes, such as sludge from waste water treatment at coal-fired power plants.

Acknowledgements

We would like to thank Editage (www.editage.com) for English language editing.

Supporting Information

Preparation of the antigen-protein conjugate. Contents of mercury and other metals in the standard CFA samples. Binding of NX2C3 antibody to mercury and other metals. Cost of present total Hg determination in CFA. This material is available free of charge on the Web at http://www.jsac.or.jp/analsci/.

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