

## Development of the analytical method for sulfur isotope in SO<sub>2</sub> gas

\*高橋 雅昭<sup>1</sup>、佐瀬 裕之<sup>1</sup>、大泉 毅<sup>1</sup>

\*Masaaki Takahashi<sup>1</sup>, Hiroyuki Sase<sup>1</sup>, Tsuyoshi Ohizumi<sup>1</sup>

1. アジア大気汚染研究センター

1. Asia Center for Air Pollution Research

### 1. Introduction

Sulfur is a major source of acid deposition. Usually sulfur is emitted as SO<sub>2</sub> and it is oxidized to SO<sub>4</sub><sup>2-</sup> in atmosphere. S isotopic analysis of water and particulate samples were conducted in many studies. But gaseous SO<sub>2</sub> was rarely used for isotopic study. Isotopic ratio of SO<sub>2</sub> is strongly influenced by emission source near the sampling point. Because most of SO<sub>2</sub> emitted far from sampling point is oxidized to SO<sub>4</sub><sup>2-</sup> during transportation. It is possible to distinguish domestic emission from transboundary emission. Moreover, dry deposition processes of SO<sub>2</sub> and particulate SO<sub>4</sub><sup>2-</sup> and their interactions with tree canopy could be discussed more precisely. Isotopic analysis of SO<sub>2</sub> is considered as a powerful method to investigate the behavior of S in the atmosphere and ecosystems. We studied about the analytical method for S isotope in SO<sub>2</sub>.

### 2. Method

#### (1) Equipment

SO<sub>2</sub> concentration in Japan is very low (e.g. annual 0.02ppb at Sado). We need to collect air sample at least 1m<sup>3</sup> that is the amount we can obtain 1.2mg of BaSO<sub>4</sub> precipitation enough for isotopic analysis. High volume air sampler is generally used to collect around 1m<sup>3</sup> of air sample. It uses filter, so we examined filter collection of SO<sub>2</sub> by K<sub>2</sub>CO<sub>3</sub> filter generally used to collect SO<sub>2</sub> in filter-pack method.

#### (2) Filters

We used quartz filter for particulate SO<sub>4</sub><sup>2-</sup> and K<sub>2</sub>CO<sub>3</sub> filter for gaseous SO<sub>2</sub> next to it. K<sub>2</sub>CO<sub>3</sub> filter is made on the basis of EANET manual. But to get enough flow rate for high volume air sampler, the size of K<sub>2</sub>CO<sub>3</sub> filter is larger (20×26 cm, same size with quartz filter) than the manual and already dried before sampling.

#### (3) Flow rate/leak test

Under flow rate 100~400 L/min in this system, we confirmed weather sampling can be carried out stably. And we checked leak using two K<sub>2</sub>CO<sub>3</sub> filters.

#### (4) Sample treatment

On the basis of EANET manual, the sample was extracted with 0.05%  $\text{H}_2\text{O}_2$  aq and filtered. Then 2N HCl was added and concentrated to remove  $\text{CO}_3^{2-}$ . Next 1%(w/w)  $\text{BaCl}_2$  was added. We filtered forming  $\text{BaSO}_4$  precipitation. We examined the method to get  $\text{BaSO}_4$  precipitation and to remove the components from  $\text{K}_2\text{CO}_3$  filter.

### 3. Results

#### (1) Sampling and sample treatment

Sampling could be conducted under flow rate 100~400 L/min. Sampling efficiency was about 80%..

Several conditions to get  $\text{BaSO}_4$  precipitation were examined, it could be gotten by the method as below.

Filter was cut and ultrasonic extracted with 0.05%  $\text{H}_2\text{O}_2$  aq 30 minutes and filtered. 2.5 ml of 2N HCl was added and concentrated to around 10 ml. Then 1 ml of heated 1 %(w/w)  $\text{BaCl}_2$  was added under pH 4~6. Next evaporated all solvent and added 10 ml of water.  $\text{BaSO}_4$  precipitation was observed. After put for one night, the precipitation was separated by centrifuge 3,600 rpm for 10 minutes and we removed the solvent. The precipitation was washed twice by 10 ml of water and filtered. At last the filter was ashed for four hours in muffle furnace in 800 °C .

#### (2) Actual analysis

We used two  $\text{K}_2\text{CO}_3$  filters and collected 2160.2  $\text{m}^3$  of air sample under flow rate 300 L/min. 3.80 mg of particular  $\text{SO}_4^{2-}$  and 0.89 mg (first) and 0.14 mg (second) of  $\text{SO}_4^{2-}$  from  $\text{SO}_2$  were obtained.  $\delta^{34}\text{S}$  of particular  $\text{SO}_4^{2-}$  was 8.64 ‰ and that of  $\text{SO}_2$  from the first filter was -1.54 ‰. The amount of precipitation from second filter was not enough for isotopic analysis. This difference was caused by effect of sea salt mainly. But nss-  $\delta^{34}\text{S}$  of particular  $\text{SO}_4^{2-}$  was 4.78 ‰. It was still higher than  $\text{SO}_2$ . It suggested that nss-  $\text{SO}_4^{2-}$  contains both transboundary  $\text{SO}_4^{2-}$  and domestic  $\text{SO}_4^{2-}$ . According to the relevant literature, domestic  $\text{SO}_2$  may have negative isotopic values because of imported oil from Middle East. Considering that this sampling was carried out in January, possible effect of transboundary  $\text{SO}_2$  with higher isotopic values should also be taken into account for source identification of the collected  $\text{SO}_2$ .

We would like to discuss at the conference with the latest data.

キーワード：硫黄同位体比、 $\text{SO}_2$ ガス

Keywords: sulfur isotope,  $\text{SO}_2$  gas