# Antimony in Urban Roadside Surface Soils: Concentration, Source and Mode of Occurrence

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**Abstract:** Antimony (Sb) is one of the elements of increasing environmental significance. In this study, concentrations of Sb were measured on 21 roadside topsoil samples collected from a medium-size city, Xuzhou (China), in order to assess the magnitude of contamination and to identify the possible contamination sources. The mode of occurrence of Sb and the effect of particle size fractions on Sb concentration distributions were also investigated from two specific soil samples. Median of Sb concentrations of the investigated urban surface soils is 0.96 mg kg<sup>-1</sup>. The Sb in the Xuzhou topsoils is mainly attributed to the inputs of coal combustion and almost independent of the particle size fractions. The most common mode of occurrence of Sb is in association with organically-bound fraction.

Keywords: Antimony, Source, Urban soils.

#### **1. INTRODUCTION**

Antimony is a global environmental contaminant. Compared to Hg, Pb, Cd or As, there have been remarkably few studies with respect to the predominant sources, biogeochemical behavior and ultimate fate of Sb in the environment [1]. Antimony has been determined in rocks, soils, fossils, volcanic gases and waters, as well as in plant and animal tissues. Antimony has low concentrations in the earth's crust  $(0.2-1.0 \text{ mg kg}^{-1})$  [2]. The anthropogenic inputs are significant in many areas in comparison to the natural ones. Previous studies have shown that up to  $3.5 \text{ t yr}^{-1}$  are emitted to the atmosphere from human activities and up to 18 000 and 26 000 t yr<sup>-1</sup> are estimated as inputs to the aquatic ecosystems and soils, respectively [2]. The human contributions of Sb are 100- to 200- fold higher than the atmospheric Sb emissions from natural sources (ca. 2600 t yr<sup>-1</sup>) [3]. Antimony anthropogenic sources mainly include both domestic and industrial coal and fuel combustion, pyrometallurgical nonferrous Cu-Ni-based metal production, incinerators and road traffic [2, 4-8]. In addition, a large number of other industrial processes like the manufacture of paints, semiconductors, computer chips, plastics, compact discs, bactericides, etc., involve the use of Sb compounds. Due to these wastes and emissions into the environment, Sb constitutes a very important subject for environmental and health-related studies [9-11].

Urban roadside topsoils are the "recipients" of large amounts of pollutants from a variety of sources including industrial wastes, vehicle emissions, coal burning waste as well as other activities. Accordingly, roadside topsoils have become an increasingly important environmental sampling medium for assessing anthropogenic pollutant levels in the urban environment [12, 13]. Despite the clear impact of Sb in the environment, so far few studies have focused on measuring Sb in urban soils. Moreover, the toxicity of antimony is not only dependent upon the total concentration but also its chemical form in the environment. Therefore, it is also important to know the mode of occurrence of Sb in urban soils. The concentrations of most heavy metals in soil are usually controlled by the particle size because finer particles, due to their high specific area (surface area per unit volume), may adsorb more heavy metals. Thus, the effects of particle size fractions on concentration distributions of Sb is of importance for transport of soil-bound Sb and pollution control by various remedial options. In this study, concentrations of antimony in urban roadside topsoils were determined and the mode of occurrence of Sb investigated. The predominant anthropogenic sources of Sb and the effects of particle size fractions on concentration distributions were also elucidated.

# 2. MATERIALS AND METHODS

#### 2.1. Studying Area and Sampling

Xuzhou,  $(33^{0}43' - 34^{0}58'N, 116^{0}22' - 118^{0}40'E)$ , is in the northwestern part of Jiangsu, one of the provinces of China. Xuzhou (China) is an important centre of historic and modern day industrial activities where coal provided the fuel for manufacturing. A total of 21 roadside topsoil samples (0 - 10 cm depth) were collected within the city of Xuzhou. At each sampling point, three sub-samples, with a 20×20 cm surface, were taken and then mixed to obtain a bulk sample. Such a sampling strategy was adopted in order to reduce the possibility of random influence of urban waste not clearly visible. All the samples were collected with a stainless steel spatula and kept in PVC packages. The soils samples were air-dried and sieved through a 2-mm nylon sieve.

In addition, two sampling sites (sample # 1 and sample # 11) were selected to investigate the effect of particle size on Sb concentration distribution and the mode of occurrence.

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These two sites were chosen to reflect differences in land use within an urban setting. The first (sample # 1) is in the area close to the Xuzhou Steel-iron Plant and the second site (sample #11) is close to the city centre in an area with mostly automobile traffic.

#### 2.2. Determination of Sb Concentrations

The Sb concentrations in 21 soil samples were dissolved with HF-HNO<sub>3</sub>-HCl acid mixture and determined by Inductively Coupled Plasma Mass Spectrometry (ICP-MS, X-Seris, Thermo Electro Co. USA) [14]. The conditions of analytical determination for Sb are described in Table **1**.

Caution was taken in preparing and analyzing samples to minimize contamination from air, glassware and reagents, which were all of superior quality. Replicated measures of standard reference materials provided by China Environmental Monitoring General Station, reagent blanks and duplicated soil samples randomly selected from the set of available samples were used to assess accuracy and precision.

#### Table 1. The Parameters of ICP-MS Used for Sb Determination

Parameters	Values	
Forward power	1200W	
Nebuliser flow rate	0.80 L/min	
Auxiliary flow rate	0.75 L/min	
Cool flow rate	13.50 L/min	
Internal standards	Rh, Re	

## 2.3. Chemical Fractionation Method for the Determination of the Mode of Occurrence of Sb

Two samples (Sample # 1 and # 11) were characterized by the three-step "BCR" extraction scheme. A detailed description of the BCR-protocol is presented by Quevauviller *et al.* [15]. Briefly, the sequential extraction was performed by sequentially extracting 0.5 grams of the soil samples in a 50 ml centrifuge tube at room temperature using a mechanical shaker with:

Fraction 1: acid extractable species. The sample was extracted with 20 ml 0.11 mol  $I^{-1}$  acetic acid (16 h shaking period);

Fraction 2: bound to Fe-Mn oxide species. The residue was extracted with 20 ml 0.1 mol  $\Gamma^1$  hydroxylamine hydrochloride (16 h shaking period);

Fraction 3: bound to organic species. The residue was extracted with a two-fold 5 ml 8.8 mol  $\Gamma^1$  H<sub>2</sub>O<sub>2</sub> (pH 2-3) digestion at 85 °C and subsequently with 25 ml 1 mol  $\Gamma^1$  ammonium acetate (16 h shaking period);

Fraction 4: bound to silicate species. The residue was extracted with HF-HNO<sub>3</sub>-HCl acid mixture (This step is not a speciation of the BCR protocol).

After shaking, the soil suspension was each time centrifuged at 3000 rpm and the supernatant solution was decanted, filtered through a Millipore filter (pore size  $0.45 \mu$ m) and set aside for analysis. After each of the steps, the soil residue was washed every time with 10 ml  $H_2O$  (15 min). After centrifuging, the washing solution was carefully decanted and discarded. The Sb concentrations in leachate were determined by ICP-MS. Each ICP-MS measurement was repeated twice. Relative standard deviations remained 5%. Accuracy was checked by measuring one of the standard solutions as unknown sample.

# 2.4. Effect of Particle Sizes on Sb Concentration Distributions

The effect of particle sizes on enrichments of Sb was also obtained from the two representative soil samples (#1 and # 11). These two samples were initially sieved through a 1000  $\mu$ m nylon sieve to remove coarse sand and then divided by particle diameter into five physical size fractions, 500-250, 250-125, 125-74, 74-45 and < 45  $\mu$ m. The soil samples in various particle size fractions were dissolved with HF-HNO<sub>3</sub>-HCl acid mixture and the Sb concentrations were measured by ICP-MS.

#### **2.5. Statistical Analysis**

Statistical analysis of data was performed with SPSS Ver. 14.0 for windows software.

#### **3. RESULTS AND DISCUSSION**

# **3.1.** Total Concentrations and Predominant Sources of Sb

The Sb concentrations of surface soil samples ranged from 0.79 to 53 mg kg<sup>-1</sup>, with a median of 0.96 mg kg<sup>-1</sup> (Table **2**). The highest Sb concentration (53 mg kg<sup>-1</sup>) was found in sample # 2 and almost 65 times of the regional background value. The sampling site (Sample # 2) is close to Xuzhou Mechanical Plant. The reason why the Sb concentration is so high is still not clear. The Sb concentrations in Xuzhou urban roadside topsoils are comparable to those reported in Table **3**.

Certain trace metals have been successfully used to identify natural or anthropogenic sources for the urban environment. Ely *et al.* [17] used metals such as Cu, Zn and Pb to evaluate the contribution of traffic to the surrounding urban environment. The correlations between metals (e.g. Ga, Al and Ti) and clays in the soil samples have also been used to indicate the natural origin of these metals [13, 18]. Previous studies have shown that toxic elements Hg and As are principally derived from coal combustion in Xuzhou urban surface soils [13]. The significant positive correlations (P < 0.01) between Sb and Hg or As (Table 4) indicate that the Sb in Xuzhou surface soils is probably mainly ascribed to inputs of coal combustion (though traffic emissions cannot be totally disregarded in any urban environment).

It is well documented that the Cu/Sb ratio is commonly used to identify the Sb contamination source. Sternbeck *et al.* [19] proposed diagnostic criteria for brake wear particles that included a ratio of 4.6  $\pm$ 2.3 for Cu/Sb. Adachik *et al.* [20] reported the ratio of Cu/Sb of 2.5 in street dust, which was within the diagnostic criteria for brake wear particles. Hjortenkrans *et al.* [15] found in roadside soils the Cu/Sb ratio of between 3.6 and 150, with a mean of 20 $\pm$ 19. The wide interval was attributed to the contribution from both Table 2. Antimony Concentrations (mg kg<sup>-1</sup>) in Xuzhou Topsoil Samples

Mean	Median	Standard Deviation	Min.	Max.
3.46	0.96	11.3	0.79	53

Table 3. Antimony Concentrations (mg kg<sup>-1</sup>) in Soils in the World

Material	Range	References
Urban roadside soil in La Coruna, Spain	0.29 - 8.81	[2]
Roadside topsoil in southern Sweden	0.25 – 37	[16]
Urban roadside soil in Xuzhou, China	0.79 – 53	This study

Table 4. Pearson Correlation Coefficients for Antimony, Mercury and Arsenic

	Sb	Hg	As
Sb	1	0.725** <sup>a</sup>	$1.00^{**^{a}}$
Hg		1	0.728** <sup>a</sup>

<sup>\*\*&</sup>lt;sup>a</sup> Correlation is significant at 0.01 levels (two tailed). Data for As and Hg from [13].

traffic and other emissions. In the present study, the mean Cu/Sb ratio is 11 and differs significantly from the results reported for brake wear particles. However, the result obtained in this study is compared to that reported by Hjortenkrans *et al.* [15].

## **3.2. Effect of Particle Sizes on Sb Concentration Distribu**tion

Metal partitioning as a function of grain size is very important for both transport of soil-bound metals and pollution control by various remedial options. In general, most heavy metals (Cu, Zn, Pb, etc.) appear to be accumulated in the finest soil particles because of the increase in specific surface area and concomitant increase in the proportion of reactive substrates. Fig. (1) shows that Sb concentrations in two specific soil samples are almost independent of the particle sizes. This is likely due to the fact that Sb, an oxyanion in natural systems, can adhere to pre-existing particles of any dimension.

# 3.3. Mode of Occurrence of Sb in Two Specific Soil Samples

The mode of occurrence of Sb in two specific soil samples was investigated using BCR sequential extraction scheme and the results are illustrated in Fig. (2). The Sb in the soil sample #1 is strongly associated with organically-bound fraction (45%) with very small amounts in the residual fraction. The percentages of acid-extractable and Fe-Mn oxide phases are almost equal (less than 25%, respectively). The results for soil sample # 11 are similar, but with less association with organically-bound phase and a higher percentage with the residual fraction. The difference in amounts of Sb in these two soil samples may be due to the influence of nearby industrial activities.

#### 4. CONCLUSIONS

The data obtained in this study demonstrate that the Xuzhou urban surface soils are not significantly contaminated by Sb. However, monitoring of Sb should be carried out in a

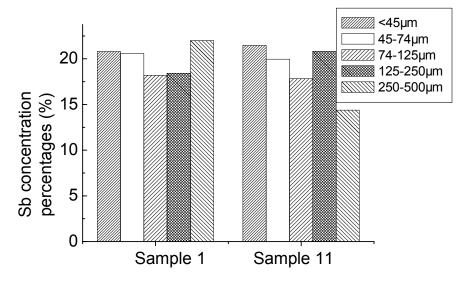


Fig. (1). Antimony distribution in various particle size fractions.

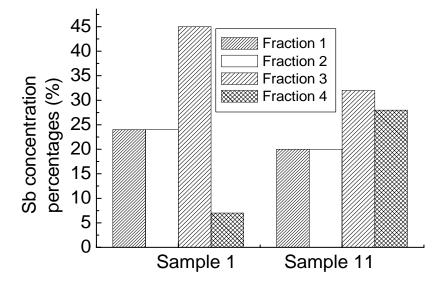


Fig. (2). Mode of occurrence of Antimony following fractionation method (Fraction 1: acid extractable species; Fraction 2: Fe-Mn oxide species; Fraction 3: organic species; Fraction 4: silicate species).

systematic fashion to detect the possibility of increase from the present levels due to the diversity of commercial and industrial activities, different sources of Sb and the potential high toxicity at a particular site. Different from the common heavy metals (such as Pb, Cu, or Zn), the distribution of Sb concentrations is independent of soil particle size fractions. The difference in Sb amounts bound in residue phase in the investigated two topsoil samples is of significance.

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# REFERENCES

- Shotyk W, Krechler M, Chen B. Antimony: global environmental contaminant. J Environ Moint 2005; 7: 1135-6.
- [2] Cal-Prieto MJ, Carlosena A, Andrade JM, et al. Antimony as a tracer of the anthropogenic influence on soils and estuarine sediments. Water Air Soil Pollut 2001; 129: 333-48.
- [3] Nriagu JO. Global metal pollution: posioning the Biosphere. Environment 1999; 32: 7-11.
- [4] Shotyk W, Chen B, Krachler M. Lithogenic, oceanic and anthropogenic sources of atmospheric Sb to a maritime blanket bog, Myrarnar, Faroe Islands. J Environ Monit 2005; 7: 1148-54.
- [5] Furuta N, Iijima A, Kambe A, Sakai K, Sato K. Concentrations, enrichment and predominant sources of Sb and other trace elements in size classified airborne particulate matter collected in Tokyo from 1995 to 2004. J Environ Monit 2005; 7: 1155-61.
- [6] Gomez DR, Gine MF, Bellato ACS, Smichowski P. Antimony: a traffic-related element in the atmosphere of Buenos Aires, Argentina. J Environ Moint 2005; 7: 1162-8.
- [7] Amereih S, Meisel T, Scholger R, Wegscheider W. Antimony speciation in soil samples along two Austrian motorways by HPLC-ID-ICP-MS. J Environ Monit 2005; 7: 1200-6.

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- [8] Månsson NS, Hjortenkrans DST, Bergbäck BG, Sörme L, Häggerud AV. Sources of antimony in an urban area. Environ Chem 2009; 6: 160-9.
- [9] Maher WA. Antimony in the environment-the new global puzzle. Environ Chem 2009; 6: 93-4.
- [10] Filella M, Williams PA, Belzile N. Antimony in the environment: knowns and unknowns. Environ Chem 2009; 6: 95-105.
- [11] Oorts K, Smolders E. Ecological threshold concentrations for antimony in water and soil. Environ Chem 2009; 6: 116-21.
- [12] Manta DS, Angelone M, Bellanca A, Nerir R, Sprovieri M. Heavy metals in urban soils: a case study from the city of Palermo (Sicily), Italy. Sci Total Environ 2002; 300: 229-43.
- [13] Wang XS, Qin Y, Sang SX. Accumulation and sources of heavy metals in urban topsoils: a case study from the city of Xuzhou, China. Environ Geol 2005; 48: 101-7.
- [14] Westerhoff P, Prapaipong P, Shock E, Hillaireau A. Antimony leaching from polyethylene terephthalate (PET) plastic used for bottled drinking water. Water Res 2008; 42: 551-6.
- [15] Quevauviller PH, Rauret G, Muntau H, et al. Evaluation of sequential extraction procedure for the determination of extractable trace metal contents in sediments. Fresen J Anal Chem 1994; 349: 808-14.
- [16] Hjortenkrans D, Bergback B, Haggerud A. New metal emission patterns in road traffic environments. Environ Monit Assess 2006; 117: 85-98.
- [17] Ely JC, Neal CR, Kulpa CF, Schneegurt MA, Seidler JA, Jain JC. Implications of platinum-group element accumulation along U.S. roads from catalytic-converter attrition. Environ Sci Technol 2001; 35: 3816-22.
- [18] Wilcke W, Muller S, Kanchanakool N. Urban soil contamination in Bangkok: heavy metal and aluminium partitioning in topsoils. Geoderma 1998; 86: 211-28.
- [19] Sternbeck J, Sjödin A, Andreasson K. Metal emissions from road traffic and the influence of resuspension- results from two tunnel studies. Atmos Environ 2002; 36: 4735-44.
- [20] Adachi K, Tainosho Y. Characterization of heavy metal particles embedded in tire dust. Envrion Int 2004; 30: 1009-17.

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