Mercury contamination in the Red river watershed: a review of the contaminated sites

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Accepted 27 December 2014

ABSTRACT

This study is based on two sampling campaigns covering the whole Vietnamese Red River watershed in 40 sites at low and high water levels during 2009 in order to determine the seasonal and spatial variations of Hg concentrations in stream sediments. The results showed that the Hg concentrations during low water level were clearly higher than that during high water level, demonstrating an effect of particle (grain) size (i.e. significant contribution of less mercury-contaminated particles in high water level). In addition, in order to assess spatial distribution of the Hg contamination in the Red River watershed and localize geochemical anomalies, multidimensional statistical analyses combined with Hg maps generated by GIS tool were used. We observed that the high mercury anomalies are originated from the important pollutant sources (mining/smelting) located in the upstream and mid-Red River watershed, demonstrating an impact of anthropogenic activities on river Hg concentrations. Finally, the comparison between the Hg concentrations in the stream sediment from the Red River watershed and the consensus-based freshwater sediment quality guidelines (SQGs) indicated that the sediment quality for important areas of the Red River watershed is rather poor and probably represents serious threats to aquatic life.

Keywords: Red River, Hg contamination, anomaly, sediment quality, GIS.

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1. Introduction

Mercury is known as one of the environmental pollutants, enhancing its toxicity for aquatic organisms and bioaccumulation in the food chain (Balogh et al., 1998; Brigham et al., 2009; Maria et al., 1997). Hg contaminated sites persist and many studies have been performed to determine contamination levels, assessment of risks, behavior and fate of this element and its compounds in the different environmental compartments. The both natural (rock weathering, erosion) and anthropogenic sources contribute in variable proportions to mercury concentration in river (Balogh et al., 1998; Brigham et al., 2009; Guangle et al., 2006; Maria et al., 1997; Mlayah et al., 2009). Stream sediment represents one of the ultimate sinks for metal discharged into the aquatic environment and is commonly used in environmental studies to evaluate concentrations of chemical elements in soils over wide areas and to identity possible sources of anomalies such as ore deposits (geochemical exploration) or anthropogenic sources (Guangle et al., 2006; Mlayah et al., 2009).

The Red River (Fig. 1), drainage in Himalaya, is the one of largest rivers in the South-East Asia and contributes significantly to the global sediment budget and water discharge to the ocean (Dang
Much research has been done on the contamination of heavy metal such as As, Cu, Pb and of nutrient salts in the Red River Delta in Vietnam, showing a serious health threat to millions of people living in this area (Berg et al., 2007; Dang T.H. et al., 2010; Ho. et al., 2010; Le T.P.Q. et al, 2007; Nguyen M.P. et al., 2010). However, metal contaminations in the upstream Red River watershed are generally poorly documented, especially mercury element.

In order to determine spatial - seasonal Hg concentration variations and localize Hg anomalies in the Red River watershed, this paper presents the first data of mercury concentrations measured in stream sediments from 40 points throughout the Red River watershed in Vietnamese part in two contrasting hydrological conditions during 2009: high and low water levels. In addition, mercury concentrations of the Red River were compared to the consensus-based freshwater sediment quality guidelines (SQGs) for estimation of potential toxic effects on aquatic environment.

![Figure 1. Description of the Red River watershed and sampling strategy: location of study sites during two spatial sampling campaigns (black points) and the sampling site coordinates. The black dashed line represents the China/Vietnam boundary](image)

**2. Materials and methods**

**2.1 Area descriptions**

The Red River originates from the mountainous area of Yunnan Province in China, flows 1200 km south-eastward and then flows through seven Vietnamese provinces before flowing into the Gulf of Tonkin in the South China Sea, with the total watershed area of 169,000 km² (50.3% of which in Vietnam, 48.8% in China and 0.9% in Laos; Fig. 1). The red laterite soils are abundant in this mountainous area and give the river its characteristic color Schäfer et al.,2006).

The main tributaries of the Red River are the Da River, on the right bank, and the Lo River, on the left bank. The Da River has its source in the Yunnan Province, near to that of the upstream Red River, at an elevation >2000m (Adrienne et al., 2008).

The Red River watershed is characterized by two distinct seasons: the wet season from May to
October and the dry season from November to April, due to the South West monsoon in summer and the North East monsoon in winter, respectively. The summer season is warm and very humid, with mean temperatures ranging from 27°C to 29°C whereas the winter season is cool and dry with mean monthly temperatures ranging from 16°C to 21°C (Fig. 2). Average annual rainfall in the Red River System is 1600 mm, with 85%-95% of this falling during the summer season.

Figure 2. Description of monthly averages of temperature, water discharge and rainfall of the Red River (LW, HW: sampling spatial campaign at low water and high water levels)

2.2. Data and methodology

Sampling frequency: during 2009, sampling campaigns were performed at 40 points throughout the Vietnamese part of the Red River watershed in two contrasting hydrological conditions: low water discharge in Mars - April 2009 and medium to high water discharge in August - September 2009 (Fig. 1, 2). The coordinates of the sampling location were recorded with a differential GPS. At each site, the dissolved, stream sediment samples were measured (Fig. 1). The coordinates of the sampling are listed in Figure 1.

Sampling and sample treatment: stream sediment samples were manually collected using a plastic spatula, consisting of the uppermost 1cm of sediment from several recent depositional pockets within a distance of twenty five meters. Stream sediment samples were sieved (<63µm; nylon sieves) to remove coarse material which was obviously not representative of typical grain size of suspended particulate matter. Thus, sediments were dried in an over at 50°C for 24 hours.

Mercury analyses: Sediment Hg concentration was analyzed from the dry, homogenized material by cold vapor atomic absorption spectrometry after calcination under an O2 stream and amalgamation on a Au trap using automated Hg analyzers (ALTEC-AMA-254 and MILESTONE-DMA-80). The results were expressed in µg/kg (dry weight) and the detection limit (3 times the standard deviation of 5 blank values) varied daily from 1 to 2 µg/kg. The accuracy of the analytical results was checked by analyzing certified reference materials (CRM 320, PACS-1, SL-1) after each set of 4 samples. Precision (coefficient of variation of 5 replicate samples) was better than 5% and results were consistently within the certified concentration intervals. These analyses were performed at the Laboratory of UMR 5805 EPOC (France).

3. Results and discussion

3.1. Variability of total Hg concentrations in stream sediment

The Figure 3A presents the statistical parameters (including the arithmetic average, median, 10- and 90-centiles, minimum and maximum) of Hg concentration in stream sediment from 40 sampling sites in the Vietnamese part of the Red River watershed measured during high and low water levels in 2009. The results showed that the mercury concentration in stream sediments is strong spatial and seasonal variations (Fig. 3A). In fact, in low water level, the Hg concentrations varied between 71 and 475µg/kg in low water condition (i.e. max/min ratio ~ 6.7) and its values varied between 66 and 241µg/kg in high water condition (i.e. max/min ratio ~ 3.7; Fig. 3A). The average Hg concentrations in stream sediment were 126µg/kg and 195µg/kg in high water and low water levels, respectively. We noted that the Hg concentrations in stream sediment of the Red River during high water level were clearly lower than that during low water level. This phenomenon has been observed in many rivers in the world and can explain by the important contribution of less Hg-contaminated sediments (alluvial floor of the channel and bank deposits) and/or a higher percentage of relatively coarse material in rainy season (e.g. 5, 12).
Figure 3. Box plots of Hg concentrations in stream sediment of the Red River in two specific campaigns during low and high waters (LW and HW) in 2009 (A), and Box plots of Hg concentrations (in µg/kg) in stream sediments of the Red River watershed in tailing and non-tailing groups in high water (HW) and low water (LW) conditions (B).

As a comparison, the Hg concentrations measured in the Red River were comparable to those observed in the other rivers like the Garonne River (200µg/kg, (Schäfer et al., 2006) or the Lawrence River (180µg/kg, (Adrienne et al., 2008). They are higher than those measured in the OuedMellègue, Tunisia (40µg/kg, (Mlayah A et al., 2009) or the Resurrection Creek River, Alaska (60µg/kg, (MacFarlane, 2004). As suspected, they are really lower than those of the Lot River (880µg/kg, e.g. Coynel et al., 2009), the Yamuna (tributary of the Ganges River, 5000µg/kg, (Subramanian et al., 2003) or the Guyana River (2400µg/kg, afterSpadini et al., 2003). It’s very interesting to note that the world average value of Hg concentration in stream sediment is 150 µg/kg, established by Viers et al. 2009.

3.2. Spatial distribution analysis assisted by GIS tool

The Hg concentrations measured in stream sediment in the Red River watershed showed strong spatial variations. In order to clearly the spatial distribution and determine Hg sources of Hg concentrations in Vietnamese part of the Red River watershed, Hg concentrations obtained from 40 sites in the Vietnamese part were used to establish the maps assisted by GIS tool. Geographic Information System (GIS; Mapinfo version 7.5) was used to treat Topographical, geological data and metal/metalloid concentrations. According to the coordinates of the sampling locations of 40 sites in the hydrological network during two specific campaigns, the limits and areas of the respective drainage basins have been extracted. Mercury concentrations were considered as generally representative of the watershed area upstream from the sampling point.

For purposes of comparison with the world average concentration of 150µg/kg, the measured concentrations ranges were separated into 4 classes, each class represents concentration ranges <150, 150-250, 250-350 and > 350µg/kg (Fig. 4).

Figure 4. Spatial distribution of Hg concentrations in stream sediment from the Red River watershed at low and high water levels
The Figure 4 shows that the high mercury anomalies are observed in the upstream of Da and Lo River (e.g. points 15, 16, 17, 18, LC, 37, 38 and 39) but also in the mid-catchment (e.g. 7, 11, 13, 14 and 32) in both high and low water levels. The source of this anomaly in the different sub-catchments of the Red River watershed may be due to natural and/or anthropogenic. Based on an inventory of tailings, waste deposits etc. (from the mining map in Vietnam, 2008), we subdivide the sub-catchments into 2 groups: tailing group (visible presence of mining, smelting or urban wastes in the sub-catchment); and non-tailing group (no visible presence of mining, smelting or urban wastes in the sub-catchment). The result obtained is presented in Figure 3B. We observed that the Hg concentrations in the tailing group are clearly higher than those in the non-tailing group, showing the strong impact of anthropogenic activities (mining and smelting) on river Hg concentrations (Figure 3B). However, to obtain more precise results, future researches need to measure Hg concentration in dissolved and particulate phases at the point sources and in the whole Red River watershed with high temporal and spatial resolutions.

### 3.3. Estimation of potential toxic effects

The sediment quality guidelines (SQG) were designed to assess contamination, protect aquatic life, support or maintain designated uses of freshwater, estuarine and marine environments, and to assist the quality of stream sediments (Balogh et al., 1998; Brigham et al., 2009; MacDonald et al., 2000; Maria et al., 1997). In this study, the Hg concentrations in stream sediments from the Red River watershed were compared to the consensus-based freshwater sediment quality guidelines (SQGs) proposed by MacDonald et al. (2008). The so-called Threshold Effect Concentration (TEC) represents concentrations of sediment-associated contaminants below which adverse effects on sediment-dwelling organisms are not expected to occur (<20%). The Probable Effect Concentration (PEC) defines concentrations of sediment-associated contaminants above which adverse effects on sediment-dwelling organisms are likely to be observed (>90%) (MacDonald et al., 2000).

For the mercury metal, the TEC and PEC values were 180µg/kg and 486µg/kg, respectively. The Hg concentrations measured in the Red River were lower than the PEC level in the whole watershed (Figure 3A, Table 1). Thus, Hg concentrations (C) in the sediments from the Red River watershed were subdivided in 3 classes: i) [Hg]<TEC, ii) TEC< [Hg] <2TEC, and iii) 2TEC<[Hg] <PEC and presented in table 1.

#### Table 1. Mercury concentrations in comparison with the consensus-based freshwater sediment quality guidelines (SQGs) proposed by MacDonald et al.(2008)

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C1: class 1 ([Hg]<TEC), C2: class 2 (TEC< [Hg] <2TEC) and C3: class 3 (2TEC<[Hg] <PEC)

The Table 1 showed that during high water condition, in almost all the Red River watershed, the mercury concentrations measured were lower than the TEC level. In contrast, during low water condition, 50% values of Hg concentration measured exceeded the TEC level and 12.5% values varied between 2TEC and the PEC values, suggesting that the sediment quality for important areas of the Red River watershed during low water level is rather poor and the potential effects of sediment-associated contaminants in aquatic life of the Red River are relatively high (~20-90%). These observations are very important for
assessing the different sediment management objectives in the Red River watershed.

4. Concluding remarks

Based on mercury concentrations measured in stream sediments from 40 sampling sites in the Red River watershed, the spatial and seasonal variations of Hg concentrations were determined. The Hg concentrations varied between 71 and 475 µg/kg in low water condition and between 66 and 241 µg/kg in high water condition, showing a strong spatial-seasonal variation of Hg concentrations. The high anomaly of Hg concentrations may be due to anthropogenic sources (mining, smelting…) located in the Red River watershed. Finally, the estimation of potential toxic effects based on the sediment quality guidelines suggested serious effects of sediment-associated contaminants in aquatic environment of the Red River, especially during low water condition.

Acknowledgement

The studies received the supports from University of Bordeaux 1 (France) and funded by the INSU-ST River Song program.

References


