Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

Atmospheric Chemistry and Physics Discussions

MANUSCRIPT SUBMITTED TO ACP: GMOS-A

Title

A SMART NANOFIBROUS MATERIAL FOR ADSORBING AND REAL-TIME DETECTING ELEMENTAL MERCURY IN AIR

Authors

Antonella Macagnano[2], Viviana Perri[1], Emiliano Zampetti[2], Andrea Bearzotti[2], Fabrizio De Cesare[4], Francesca Sprovieri[3], Nicola Pirrone[2]

[1]University of Calabria, via Pietro Bucci, Arcavacata, Rende 87036 (CS), Italy; [2]Institute of Atmospheric Pollution Research-CNR, Via Salaria km 29,300 Montelibretti 00016 (RM), Italy; [3]Institute of Atmospheric Pollution Research-CNR, Division of Rende, c/o UNICAL-Polifunzionale Arcavacata, Rende 87036 (CS) Italy; [4]DIBAF-University of Tuscia, Via S. Camillo de Lellis, 01100 Viterbo, Italy

Correspondence to: *Antonella Macagnano, a.macagnano@iia.cnr.it; antonella.macagnano@cnr.it*

Abstract

The combination of gold affinity for mercury with nanosized frameworks has allowed to design and fabricate novel kinds of sensors with promising sensing features for environmental applications. Specifically, conductive sensors based on composite nanofibrous electrospun layers of titania easily decorated with gold nanoparticles were developed to obtain nanostructured hybrid materials, capable of entrapping and revealing GEM traces from environment. The electrical properties of the resulting chemosensors were measured. Few minutes of air sampling were sufficient to detect the concentration of mercury in the air, in the range between 20-100 ppb, without using traps or gas carriers (LOD ~ 1.5 ppb). Longer measurements allowed the sensor to detect lower concentrations of GEM. The resulting chemosensors are expected to be low-cost, very stable (due to the peculiar structure), and requiring low power, low maintenance and simple equipment to work.

1 Introduction

Mercury (Hg) is released into the atmosphere both by human's activities, predominantly fossil fuel combustion, and naturally, for example, from soil out-gassing, volcanoes and evasion from the sea (Pirrone et al., 2010; Pacyna et al., 2010). One of the more troublesome questions in recent years

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

has been to quantify not only the strength of emission sources but also the effects of re-emission of previously deposited Hg on the overall distribution, concentration and speciation of Hg in the atmosphere (Hedgecock et al., 2003). The deposition of atmospheric Hg depends on its chemical speciation, where the term speciation is used to distinguish between the gaseous elemental (GEM) and gaseous oxidized forms of Hg [GOM and Particle bound mercury (PBM)] and their chemical-physical characteristics (Lyman et al., 2010; Sprovieri et al., 2016a,b). To be precise, Total gaseous mercury (TGM) mainly comprises GEM with minor fractions of other volatile species (e.g., HgO, $HgCl_2$, $HgBr_2$, CH_3HgCl, or $(CH_3)_2Hg$). However, in spite of conceptual differences between TGM and GEM, they have often been used without clear distinction. This was allowable to a degree as the predominant fraction of TGM (usually in excess of 99%) is often represented by GEM under normal conditions. GEM is relatively inert under atmospheric conditions, only slightly soluble and also quite volatile, whereas the oxidized Hg forms found in the atmosphere are both soluble and involatile, thus they are efficiently scavenged and consequently deposited by liquid atmospheric water, such as rain and fog droplets, but also deliquesced aerosol particles. The dispersion of GEM on global scale therefore, depends on the rate of its oxidation in the atmosphere as this determines its long atmospheric lifetime (generally >1 year), limiting local emission controls from protecting all environments. Several international initiatives and programs [i.e., the United Nations Environment Program (UNEP)] have also made a tremendous effort in identifying and quantifying Hg pollution across the globe, especially the ‘‘hot-spots’’, aimed at reducing risk of exposure to this neurotoxin pollutant. Policy makers are working toward a worldwide effort for supporting the constructing an accurate global Hg budget and to model the benefits or consequences of changes in Hg emissions, for example, as proscribed by the Minamata Convention. Anticipating a global policy, in 2010 the European Commission began a five-year project called the Global Mercury Observation System (GMOS, www.gmos.eu) to create a coordinated global network to gaps in emissions monitoring and in the spatial coverage of environmental observations, mostly in the tropical regions and Southern Hemisphere, thus adequate for improving models and making policy recommendations (Sprovieri et al., 2016a,b). To date the GMOS network consists of more 43 monitoring stations worldwide distributed including high altitude and sea level monitoring sites, and located in climatically diverse regions, including polar areas (Sprovieri et al., 2016a,b). One of the major outcomes of GMOS has been an interoperable e-infrastructure developed following the Group on Earth Observations (GEO) data sharing and interoperability principles which allows us to provide support to UNEP for the implementation of the Minamata Convention (i.e., Article 22). GMOS activities are currently part of the GEO strategic plan (2016–2025) within the flagship on “tracking persistent pollutants”. The overall goal of this flagship is to support the development of

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

GEOSS by fostering research and technological development on new advanced sensors for in situ and satellite platforms, in order to lower the management costs of long-term monitoring programs and improve spatial coverage of observations. Since automated measurement methods of Hg require power, argon gas, and significant operator training, they are difficult to apply for understanding Hg air concentrations and deposition across broad regional and global scales. Therefore, the lack of an inexpensive, stand-alone, low power, low-maintenance sensor is a primary technical issue to be solve for the sustainability of a global network such as GMOS. Previous research highlighted that Hg-concentration levels in air vary greatly across different environmental locations, remote as the Polar Regions, background or rural, and urban locations with an average range between 1.5 ngm^{-3} (GEM) and 1 pgm^{-3} (GOM and PBM), depending on the speciation. Hence, for the determination of atmospheric Hg also at such low levels, sampling and analytical methods should be sensitive enough to quantify the concentration profiles of diverse Hg species in each respective environmental setting to better understand their environmental behavior and patterns. Fortunately, many advances made in analytical methodologies have made it possible to study atmospheric Hg in different environmental locations. However, several limitations and difficulties have still experienced in Hg analysis, as most methods cannot yet directly or accurately determine minor Hg species (Gustin et al., 2013). Hence, efforts should be continued to secure further the reliability, the traceability, and the accuracy of Hg levels measured in air. Current air monitors are amply sensitive to detect the global background but are costly, complicated configuration, electricity requirements and high maintenance. This limits the scientific research community's ability to long-term measure atmospheric Hg concentrations worldwide. Sampling and analysis of atmospheric Hg is made most commonly as GEM/TGM because of their greater abundance, even if both manual and automatic methods have been currently developed for different Hg forms to suit the measurement and monitoring application. The most common sampling method employed relies on adsorption on gold amalgam and then, either directly or indirectly, through a stepwise process of thermal desorption and final detection [usually by cold-fiber atomic absorption spectroscopy (CVAAS) or cold-fiber atomic fluorescence spectroscopy (CVAFS)]. However, our knowledge presents currently several gaps to be solved. Firstly, The atmospheric chemistry of Hg remains poorly understood, especially the oxidation pathways by which GEM is converted to GOM, the reduction pathway which converts GOM back to GEM, and the gas-particle partitioning. This is partially due to the need for identification of the chemical forms of oxidized Hg in the atmosphere and methods to measure these compounds individually. In addition, the limitations and potential interferences with our current measurement methods have not been adequately investigated, thus alternate methods to measure atmospheric Hg are needed. Given the uncertainty and restrictions associated with

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

automated and/or semi-automated Hg measurements (Gustin et al., 2013; Pirrone et al., 2013), and above all, responding to the technical needs of an expanding Hg global observation network, we developed a reliable, sensitive, and inexpensive surface for atmospheric Hg detection. In particular, we investigated and demonstrated the utility of composite nanofibrous electrospun layers of titania decorated with gold nanoparticles (AuNPs) to obtain nanostructured materials capable of adsorbing GEM as a useful alternative system for making regional and global estimates of air Hg concentrations. Methods and new sampling systems previously developed, such as passive samplers, have been used to understand long-term global distribution of persistent organic pollutants (POPs) (Harner et al., 2003; Pozo et al., 2004). Other passive samplers for both TGM and GOM collection on the basis of diffusion have been constructed using a variety of synthetic materials (i.e., gold and silver surfaces, and sulfate-impregnated carbon) and housings (Lyman et al., 2010; Gustin et a., 2011; Zhang et al., 2012; Huang et al., 2014). However, because of the differences in design of passive samplers, ambient air Hg concentrations quantified by various samplers may not be comparable. In addition, sampling rates (SRs) using the same passive samplers may depend on environmental conditions and atmospheric chemistry at each site. Moreover, it has been also highlighted that the performance of passive samplers may be influenced by meteorological factors (e.g., T °C, RH, wind speed) therefore inducing bias for the result of passive sampling (Plaisance et al.,. 2004; Sderstrm et al., 2004). On the other hand, incentive for developing simple and cost-effective samplers that are capable of monitoring over an extended period and require no technical expertise for deployment of these systems also at remote locations is now obvious. In this work we describe an alternative approach adopted in the place of conventional ones demonstrating that the combination of gold affinity for Hg with a nanoscale sized framework of titania provided the chance to create promising sensors for environmental monitoring in real time, characterised by high sensitivity to the analyte. The novel sensor is a relatively simple and low cost method for measurement of the most abundant Hg form in ambient air (TGM/GEM) due to reusable parts and simple deployment steps. Further, we have evaluated the applicability of this measurement technique with respect to real environmental conditions highlighting future directions of research on airborne Hg determination. The TGM/GEM sensor surface described here could be deployed in a global network such as GMOS; a permanent network of ground based monitoring sites and observations of Hg and/or related species on a global scale and with remote sensors would in fact be highly desirable. These data are needed to test and validate model processes and predictions, understand the source-receptor relationships, understand long-term changes in the global Hg cycle, and at least, would help policy makers to set regulations for different areas. The sensor features are related to the nanofibrous scaffold of titania capable of growing up gold

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

Atmospheric Chemistry and Physics Discussions

nanostructures by photocatalysis, tunable in size and shape. Such a nanostructured layer, fabricated by electrospinning technology, firstly improves sensor features with respect to those of compact films, by enhancing the global number of binding sites of analyte-sensor and reducing some bulk drawbacks. Secondly, the combination of metal oxides and metal nanostructures, improves the sensitivity, allows sensor to work at room temperature, tunes selectivity towards different gas species by adjusting the surface to volume ratio of nanosized structures and affect sensor lifetime. Morphological, optical, electrical aspects and sensing measurements of fibers of GEM in air have been reported and discussed. When designed, the resulting Hg ad-absorbent material was expected to be suitable for novel Hg sensors fabrication, since a similar nanofibrous scaffold doped with AuNPs was described in literature as filtering systems capable of adsorbing and removal Hg fiber from the environment with an efficiency of about 100% (Y Yuan et al., 2012). In fact, in previous works **(**Macagnano et al., submitted**,** Macagnano et al., 2015a**)** the authors reported the ultra-high sensitivity of the sensor, capable to detect up to dozens ppt, despite of a long time necessary to reveal the analyte at these concentrations, in air. In this work the chance to apply the sensor in polluted sites and in real time has been presented and described.

2 Materials and methods

2.1. Chemicals

All chemicals were purchased from Sigma-Aldrich and used without further purification: polyvinylpyrrolidone (PVP, Mn 1,300,000), titanium isopropoxide (TiiP, 99.999%), gold(III) chloride hydrate ($HAuCl_4$, 99.999%), anhydrous ethanol ($EtOH_a$) and glacial acetic acid ($AcAc_g$). Ultrapure water ($5{\cdot}5\ 10^{-8}$ S cm^{-1}) was produced by MilliQ-EMD Millipore.

2.2. Electrospinning technology

Electrospinning is a widely used technique for the electrostatic production of nanofibers, during which an electric field is used to make polymer fibers with diameters ranging from 2 nm to some micrometres from polymer solutions (or melts). It is currently the most economic, versatile, and efficient technology to fabricate highly porous membranes made of nano and/or micro fibers also for sensors (Macagnano et al., 2015b). It is based on the application of a high voltage difference between a spinneret ejecting a polymeric solution and a grounded collector. The jet of solution is accelerated and stretched by the external electric field while travelling towards the collector, leading to the creation of continuous solid fibers as the solvent efiberates. The electrospinning apparatus used in the present study (designed and assembled in CNR laboratories) comprised a home-made clean box equipped with temperature and humidity sensors, a syringe pump (KDS 200, KD Scientific) and a grounded rotating cylindrical collector (45 mm diameter), a high voltage oscillator

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

Atmospheric Chemistry and Physics Discussions

(100 V) driving a high voltage (ranging from 1 to 50 kV) and a high power AC-DC (alternative current-direct current) converter. Electrospinning solution (7.877×10^{-5} M), was prepared by dissolving PVP in $EtOH_a$ and stirring (2 hours). A 2 ml aliquot of 1:4 (w/v) solution of TiiP solved in 1:1 (v/v) mixture of $AcAc_g$ and $EtOH_a$ was freshly prepared and added to 2.5 ml PVP solution under stirring in order to obtain a 1.95 (w/w) TiiP/PVP final ratio. Both mixtures were prepared in a glove box under low humidity rate (<7% RH). The syringe filled with the TiiP/PVP solution and housed in the syringe pump, was connected to a positive DC-voltage (6 kV), and set to a 15 cm far grounded rotating collector. The substrates were fixed through suitable holders onto the collector (600 rpm, 21 °C and 35% RH) and processed (feed rate 150 ml h^{-1}) for 20 min to obtain scaffolds for sensors. After deposition, PVP/TiO_2 composite nanofibers were left for some hours at room temperature to undergo fully self-hydrolysis of TiiP [Li et al., 2003]. And then annealed under oxygen atmosphere (muffle furnace) using a thermal ramp from room temperature up to 550 °C (1 °C min^{-1}, 4 h dwell time) in order to remove PVP and crystallize the metal oxide (*anatase*).

2.3 Transducers: interdigitated electrodes

The transducer adopted in the present work to convert the physico-chemical interactions of analytes with the different polymer fibers in an electrical signal was an interdigitated electrode (IDE) [Bakir et al,1973; James et al., 2013]. Specifically, the transducer consisted of 40 pairs of electrodes (150 nm in electrode thickness, 20 μm in gap and electrode width and 5620 μm in length) was manufactured in CNR laboratories through a standard photolithographic process (lift-off procedure), then followed by Ti sputtering and Pt efiberation, suitable to generate the electrodes of the size reported above, on a 4 in. oxidised silicon wafer. After electrospinning deposition all the electrical signals of the resulting chemoresistors were recorded by an electrometer (Keithley 6517 Electrometer).

2.4 Titania nanofibers

Upon calcination, the diameters of fibers extraordinarily shrunk: mean diameters of fibers were estimated through image analyses to be approximately within the range of 60–80 nm. Specifically, the resulting fibers appeared fine and rough at surface, with a fairly homogeneous fabric. The absence of beads and the good quality of the long and continuous fibers was confirmed through SEM micrographs. A highly porous and dense network of nanofibers covering the electrodes was observed, showing interconnected void volumes (porosity) and high surface-to-volume ratios (specific surface area). Zampetti et al., (2013) reported that such a fibrous layer showed a 99% of pores having an area less than 10 μm^2, with more than 80% pores being <1 mm^2.

2.5 AuNPs/TiO_2NFs photocatalytic decoration

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

Atmospheric Chemistry and Physics Discussions

Exploiting the photocatalytic properties of TiO_2, gold nanoparticles were selectively grown, under UV-light irradiation, on the electrospun titania nanofibers through the photoreduction of $HAuCl_4$ in the presence of an organic capping reagent (PVP). Thus the resulting fibrous scaffolds were dipped into an aqueous solution containing $HAuCl_4$ and PVP ($1.5 \cdot 10^{-3}$M and 0.1M respectively) and exposed to UV light irradiation for specified intervals (UV lamp (365 nm) (Helios, Italquartz). Depending on the gold nanoparticles size that were forming in photocatalysis, the dip-solution changed from light yellow to purple. Samples were rinsed extensively with water and then air-dried. Before morphological, electrical and sensing measurements, samples were heated at 450 °C per 1 h to eliminate the PVP traces. Morphological characterization were provided by scanning electron microscopy (SEM) (Jeol, JSM 5200, 20 keV) with pictures captured at 5 kV accelerating voltage. AFM (atomic force microscopy) micrographs were taken in tapping mode using 190Al-G tips, 190 kHz, 48N/m (Nanosurf FlexAFM). TEM (C-TEM, control transmission electron microscopy) micrographs were performed at 200 keV with an analytical double tilt probe. TEM specimen were prepared by gently scraping at first the TiO2 nanofibrous layer electrospun onto the silicon support and then collecting the nanofibers, through adhesion upon contact with holy carbon thin film. UV-Vis spectra were provided by Spectrophotometer UV-2600 (Shimadzu), analysing quartz slices coated with nanofibers. These substrates were able to collect fibers by electrospinning (20 min), and then were subjected to calcination according to the described above procedure, and then UV irradiation in the aqueous solution. The fibrous layer stayed stuck to the substrate if the thickness was thin enough. Longer depositions caused curling of fibers during the calcination process.

2.5 Measurement set-up

The sensor was placed in a suitable PTFE-made measurement chamber (0.7 ml volume) connected to an electrometer (Keithley 6517 Electrometer) capable of measuring the current flowing through the IDE, when a fixed potential was applied to it, and to send data to a PC. Dynamic measurements were carried out at room temperature both using: (i) 4 channel MKS 247 managing four MKS mass flow controllers (MFC), set in the range 0–200 sccm and (ii) Environics S4000 (Environics, Inc.) flow controller, containing three MFCs supplying different flow rates (up to 500, 250 and 25 sccm, respectively), managed by its own software. Pure air (5.0) (Praxair–Rivoira, Italy) was used as gas carrier. A homemade PTFE (polytetrafluoroethylene) permeation tube filled with a suitable amount of Hg^0 was included within such a delivery system to get set dilutions of Hg-saturated vapours. The tube was immersed in a thermostatically controlled bath, thus the desired Hg^0 concentration delivered to the sensor was achieved by both tuning the temperature of the permeation tube and the dilution flow. The Hg^0 concentration was checked by Tekran®2537A analyser. Responses were calculated as $\Delta I/I_0$, where ΔI was the current variation and I_0 was the current when synthetic pure

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

Atmospheric Chemistry and Physics Discussions

dry air was flowed. Sensor was restored after a quick thermal shot at 450°C under flow of pure air (450°C).

3 Results and Discussion

Nonwoven mats made of PVP and amorphous TiO2 were obtained by the combination of electrospinning and sol-gel techniques (Fig. 1). The deposition occurred for 20 min on oxidised silicon wafers and IDEs, properly fixed on the surface of a conducting rotating collector to form nanofibrous layers characterized by high surface areas and relatively small pore sizes (Zampetti et al.,2013). By changing the deposition time, both thickness and consistence of the mats changed: one hour deposition provided the formation of a thicker white and soft fabric easily peeled off (Fig.1), hygroscopic and soluble in both water and polar solvents; a 20 min deposition generated a fibrous film adhering to substrates, too thin to be weeded. The calcination process caused a complete degradation of PVP with formation of crystalline TiO_2 (*anatase*) and a significant shrinkage of fibers dimension (60-80 nm diameter, 5-40 nm grain sixe). Exploiting the photocatalytic properties of titania (*anatase*), a tunable decoration of fibers with gold nanoparticle could be achieved by dipping the fibrous mats in a proper aqueous solution ($HAuCl_4$, PVP) under UV light irradiation (Li et al, 2004; Macagnano et al., 2015). The photocatalytic reaction was proved by the color of the solution (red purple from light yellow) (Fig.1). Changing both UV irradiation exposure time and PVP concentration, as capping reagent, morphology, size and density of gold nanoparticles could be tuned [Macagnano et al., submitted**]**.

Figure 1. Sketch of an electrospinning set-up comprising a syringe and a grounded rotating cylinder collector where the samples take place for their coverage (*left*); a piece of a nanofibrous fabric of TiiP/PVP peeled from the substrate after 1 hour of electrospinning deposition (*centre*) a red-purple aqueous solution of $HAuCl_4$/PVP after UV-light irradiation treatment holding a piece of the nanofibrous fabric of TiO_2 (*anatase*) obtained after TiiP/PVP annealing (*right*)

In the present work, among a series of differently coated fibrous layers, only the fibrous nanocomposites that were conductive at room temperature were selected and then their electrical and sensing features investigated. The controlled gold deposition was due to the photo-excited electrons on the surface of TiO_2 nanofibers that were able to reduce the gold ions thus inducing gold deposition (Fig. 2, *the sketch*). The capping reagent war responsible of the shape of the particles.

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

Atmospheric Chemistry and Physics Discussions

262 The surfaces of nanofibers observed in SEM micrographs (Fig.2, *right*) appeared densely decorated
263 with globular nanoparticles. In C-TEM image (Fig.2, *inset*) the gold nanoparticles appeared darker
264 and spherical or quasi-spherical. The single particles size were ranging between 2 and 20 nm, with a
265 7.8±3 nm average diameter [Macagnano et al., submitted]. Gold nanoparticles grew directly onto
266 the nanofibers and their adhesion appeared relatively strong (despite due to van der Waals forces),
267 since they both resisted to water rinsing and fibers scratching for TEM analyses.
268

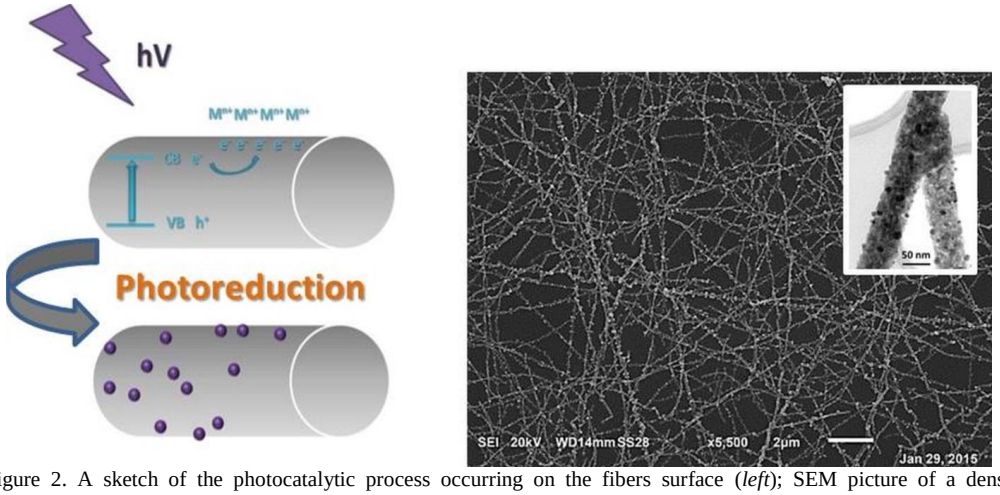

Figure 2. A sketch of the photocatalytic process occurring on the fibers surface (*left*); SEM picture of a dense nanofibrous network of AuNPs/TiO_2 coating a silicon wafer (*right*); a C-TEM micrograph of fibers finely decorated with gold nanoparticles (*the darkest ones*) fixed without using any additional linker (*inset*).
269

270 After photocatalytic process, the white porous mat became purple-violet. As seen in the spectrum of
271 the AuNP/TiO_2 system, a characteristic absorbance band appeared at around 543 nm, which
272 corresponded to the surface plasmon resonance (SPR) of the AuNPs (Sun et al, 2003). A red
273 shifting and broadening of the absorbance band was observed with the increasing in AuNP size and
274 fiber loading, respectively (data not shown). The colour is strictly depending on the size of the
275 nanoparticles, and then their agglomeration at the solid state. According to Bui et al. (2007), such a
276 band broadening phenomenon is due to the electric dipole–dipole interactions and coupling
277 occurring between the plasmons of neighbouring particles, whereas nanoparticle agglomeration
278 phenomena occurred.
279

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

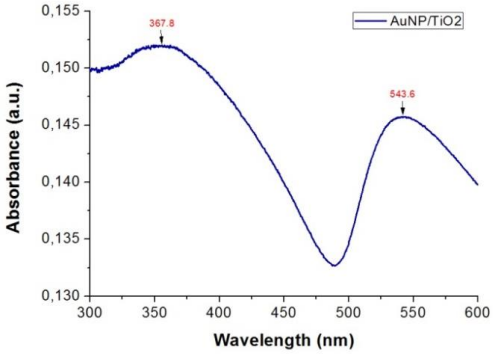

Figure 3. UV-Vis spectrum of a titania nanofibrous network after gold decoration (TiO_2: 367.8 nm; Au NPs: 543.6 nm)

Due to these features, UV-Vis absorption spectroscopy has been used in literature as a technique to reveal the changes in size, shape and aggregation of metal nanoparticles in liquid suspension after exposure to heavy metals, as Hg^0 (Morris et al., 2002). Both blue-shifted wavelength and its extent were proportional to the amount of Hg^0 that entered the liquid suspension. Similarly, when the gold decorated nanofibers of titania, collected on a quartz slice, were exposed to Hg^0 vapours (2 ppm) in air for 15 min, a significant blue shifting was reported (~ 3 nm) (Fig. 4) due to the atomic adsorption of GEM on the surface. The nanoparticles could be regenerated by heating the sample at 550°C for 3 minutes to remove Hg^0. The recovery of the AuNPs was stated by the achieving of the original values of wavelength. The regeneration could be done for dozen times without any deterioration. Similarly, in chemoresistors, the TiO_2 nanofibrous layers attached to the substrates (Fig. 5), changed colour after photocatalytic treatment.

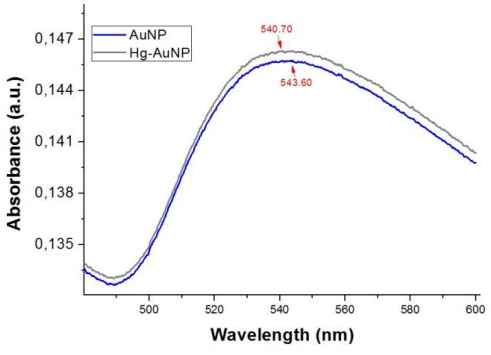

Figure 4. UV-Vis spectra of AuNPs/TiO_2nanofibers before (*blue*) and after a 15 min exposure to 2 ppm of Hg^0 (*gray*)

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

Atmospheric Chemistry and Physics Discussions

293 The IDE layout (Fig.5) show a set of interdigitated electrodes which occupies an area
294 approximately 3x5 mm, completely coated with the sensitive fibers, and two bonding pads (2x2
295 mm) that will be connected to the electrometer (DC voltage). Such a planar interdigitated electrode
296 configuration is the most commonly used for conductometric sensing applications.
297

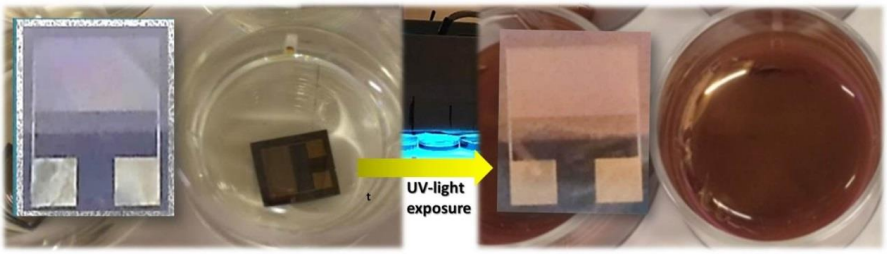

Figure 5. Chemosensor fabrication and final structure: IDE dipped (*left*) and exposed to UV-light (*right*) for gold decoration

298

299 Figure 6 depicts a Current–Voltage (I-V) curve of a chemosensor, under synthetic dry air. However
300 curve shape was unaltered when air or nitrogen were flushed over the fibers (Macagnano et al.,
301 2015), suggesting that oxygen concentrations poorly affected the electrical properties of such a
302 chemoresistor. The resistance value of IDE coated with TiO_2 nanofibers before photocatalysis,
303 resulted to be too high at room temperature to contribute straight to the final current. The resulting
304 linear shape (Ohmic behaviour) within the selected voltage range (−3V to +3V) showed a constant
305 resistance value for the sensor. The very low value of resistance (~1.2 kΩ) provided the chance to
306 work at low voltage, with consequent effects on the energy consumption as well as lifetime of the
307 material. Moreover, the linearity of I–V curve let us suppose that the sensing scaffold had a good
308 adhesion to the metal electrodes. The electron conductivity has been described by the percolation
309 model **(**Macagnano et al, submitted; Muller et al., 2003**)** since the titania at room temperature was
310 supposed like an insulating organic matrix. When it is metal doped, the electron conductivity is
311 ruled by thermally activated electron tunneling from one metal island (gold nanoparticles) to the
312 other. However, the conductivity of the nanocomposite is lower than that of pure metal (gold)
313 because the electron mean free path is greatly reduced due to the presence of the dielectric (the
314 titania crystals). The electrical features, such as the reproducibility of the fabrication process, of this
315 conductive device have been previously investigated by the authors (Macagnano et al, submitted;
316 Macagnano et al., 2015), showing encouraging results for the development of a low cost sensor for
317 mercury detection. However, in spite of the high sensitivity (LOD: 2ppt) of the sensor, too long
318 response time was necessary to detect traces of Hg^0, when compared to the current monitoring
319 instrumentations (Ghaedi et al., 2006; Sanchez-Rodas et al., 2010; Ferrua et al., 2007). Extremely

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

Atmospheric Chemistry and Physics Discussions

encouraging resulted if compared to other sensors currently involved in detecting mercury in air (Drelich et al., 2008; Kabir et al., 2015; Sabri et al., 2009; Mohibul Kabir et al., 2015; Raffa et al., 2006; James et al., 2012-2013; Chemnasiri et al., 2012; Sabri et al., 2011; Keebaugh et al., 2007; Crosby, 2013; McNicholas et al., 2011). The long time in response was supposed to be in part due to the layout of the measuring system, since the sensor was housed in a quartz bottle of 100 mL volume. In fact, an additional time was expected to be caused by the adsorption of the Hg^0 traces from the surrounding environment (measuring chamber) up to achieve a sufficient number of mercury atoms adsorbed on the surface sensor to be electrically revealed. In the present study the measuring chamber was designed in order to reduce the volume (0.7 mL) and to expose the fibers to the gas entry (Fig. 7). Such a measuring layout was designed to allow the fibrous network to be exposed to the mercury atoms as delivered into the sensor chamber.

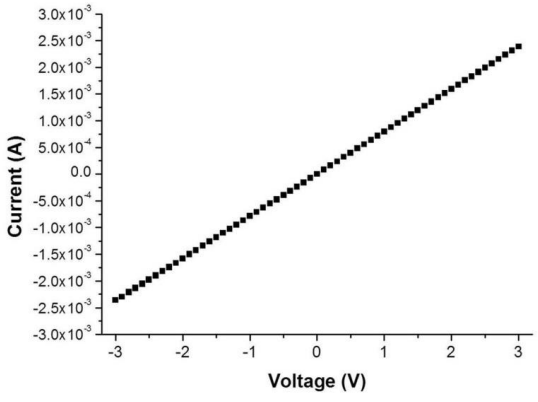

Figure 6. Chemosensor current-voltage curve

Sensing measurements, i.e. current (or resistance) changes, were provided in continuous. The sensor measurements, that were the electrical signals reported when interaction between the sensing layers and the analytes were happening, resulted in a change of the whole current (or resistance, i.e. I =V/R) according to Ohm's law. The sensor was exposed to a flow of Hg^0 in air with a concentration of 800 ppb for 1 min (Fig. 7, *right*), and then air was used to clean the sensor surface. A rapid decrease in current was recorded ($1.056 \cdot 10^{-7}$ $A \cdot s^{-1}$) when Hg^0 entered the measuring chamber. The current curve trend slightly changed when clean air was flowed, stabilizing at about the current values reached for Hg^0 adsorption. Due to the strong affinity between Au and Hg^0, a 3 min-thermal treatment was necessary to remove mercury from layer and get the same starting current value.

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

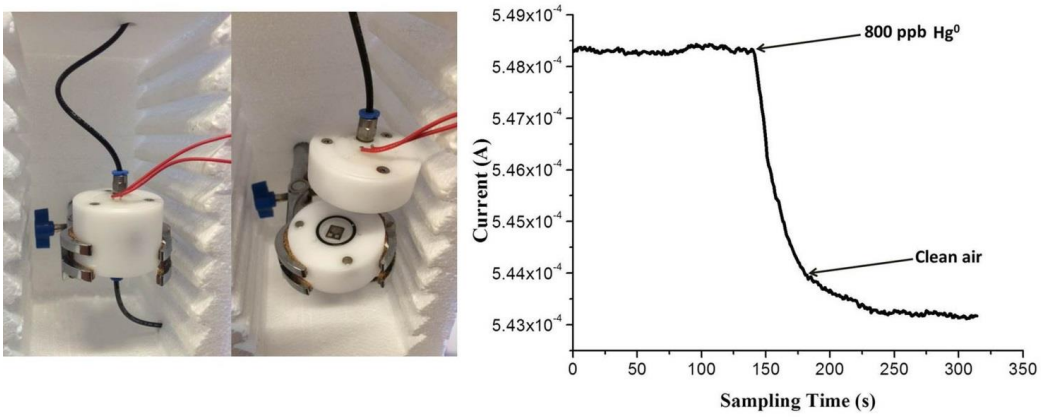

Figure 7. Homemade measurement chamber to house the chemosensor for laboratory experiments (left); plot depicting the transient response curve to 800 ppb Hg^0 (V=0.3 V)

344

345

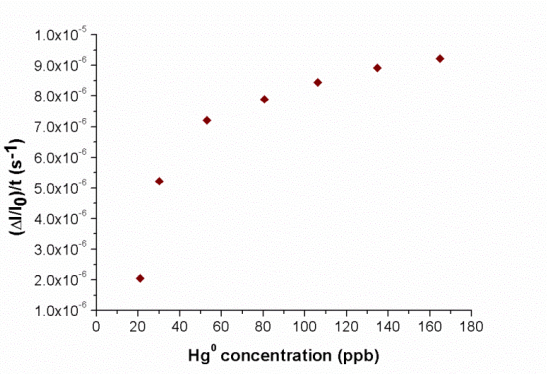

Figure 8. The normalized sensor response rate to the increasing concentration of vapour elemental mercury

346

347 Figure 8 depicts the normalized sensor response rate, i.e. the normalized current change per second,
348 toward the increasing concentration of GEM (ranging between 20 and 160 ppb). The resulting
349 logarithmic curve describes how the Hg^0 concentration affects the response time: small variations of
350 Hg^0 concentration up to 80 ppb are able to deeply change the response rate, on the contrary higher
351 concentration seem to affect only slightly this sensing feature. Since a strong relationship is
352 recorded between the concentration and the response time when the content of mercury in
353 environment is low, is possible to correlate the slope of the transient responses within the first
354 minutes of the sensor response to definite concentrations of Hg^0 in air. Figure 9 depicts the linear
355 fitting of 10 min-sensor responses when increasing concentrations of mercury were flowed onto the
356 sensor. Related data were reported in Table 1.

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

Atmospheric Chemistry and Physics Discussions

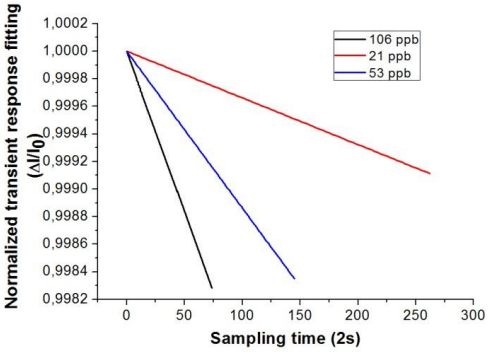

Figure 9. Linear fitting of the normalized sensor response within the first ten minutes

357

358

359

Table 1. Linear fitting parameters of 10 min-sensor responses to 21 ppb≤ [Hg^0] ≤106ppb

ppb	$(\Delta I/I_0)s^{-1}$	SE(±)	R^2
21	-7.12602E-10	1.75521E-11	0.86
33	-1.50647E-9	1.05521E-10	0.91
39	-1.78067E-9	1.02615E-10	0.91
40	-1.85901E-9	1.01833E-10	0,92
53	-2.44657E-9	4.24993E-11	0.91
70	-3.19082E-9	2.55882E-11	0.93
106	-4.83599E-9	2.67462E-10	0.88

360

361 A linear relationships has been reported between the response rate and the concentration of Hg,
362 according to the following equation (1):

363 (1) $y = (-4.56226E^{-11}) \cdot [Hg^0],\ [Hg^0] < 100\ ppb;\ SE: \pm 1.504E^{-12};\ R^2 = 0.99675$

364

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

Atmospheric Chemistry and Physics Discussions

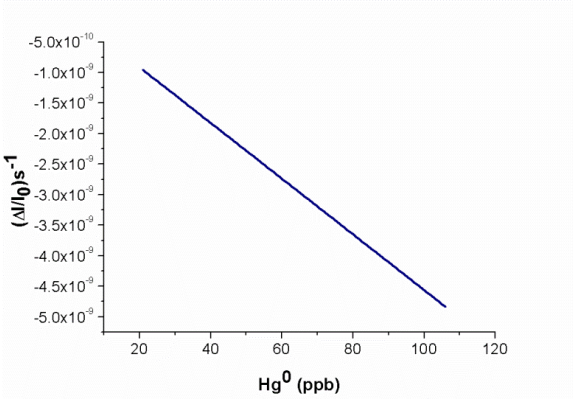

Figure 10. Linear relationships between the normalized response time and the Hg^0 concentration, within the range of 20 and 100 ppb.

Therefore when the concentration of Hg increased, the response curve slope changed too linearly, allowing a limit of detection of about 1 ppb, when the sensor is exposed to air polluted with Hg^0 per 10 min. For what concerns main interfering compounds, since at room temperature and in dark condition the measured current is supposed to be due to AuNPs decorating titania fibers, only chemical compounds interacting with gold are expected to be mostly responsible of the current changes (i.e. halides and sulphides). Thus in a blend of other chemicals, this sensor has been designed as a pretty selective sensor, being able to greatly decrease the environmental disturbances allowing the investigator/manufacturer to design and then fabricate easier strategies to prevent contaminations from environment (selective filtering systems or coatings). Among common potential contaminants authors investigated previously water vapour influence (%RH) reporting no-effects on the electrical signals **(**Macagnano et al., 2015**)**.

4 Conclusions

The adopted sensing strategy has focused on the strong affinity of mercury to gold combined to the nanostructures properties. Exploiting the photocatalytic properties of electrospun titania nanofibers, a novel conductometric sensor has been designed and fabricated to detect GEM in air. Electrospinning technology has been used successfully to create a 3D-framework of titania covering the electrode sensing area of the properly designed chemoresistors (IDEs). AuNPs have been grown on TiO_2 nanofibers exploiting the photocatalytic properties. Such a sensor was able to work at room temperature and was highly sensitive to Hg^0. Since it is composed of titania and gold, it sounds to be robust and resistant to common solvents and VOCs commonly in the air. The

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

short thermal treatments (450°C, 3min), necessary to desorb mercury from AuNPs, didn't seem to affect the lifetime of the device. Depending on the strategy of sampling, a sensing device based on such a chemosensor, could be designed for real applications, specifically for real time monitoring of polluted sites. Few minutes of sampling of air are sufficient to quantify the concentration of mercury in the air, in the range between 20-100 ppb (LOD: 1 ppb), without using traps or gas carriers. However further investigations are necessary also to assess the effects of physical parameters of the environment, such as temperature fluctuations and UV-light, as well as chemical ones, such as volatile organic compounds and gas (like halides and sulphides) which are well known interfering of the adsorption process of the Hg^0 on gold.

Acknowledgments

The activity is part of the International UNEP-Mercury Programme (UNEP-Mercury Air Transport and Fate Research (UNEP-MFTP) within the framework Global Mercury Observation System, funded by EC as part of EC FP7. Furthermore, authors gratefully thank Mr. Giulio Esposito and Mr. A. Capocecera for their support in the use of laboratory instrumentations.

References

Bui, M.P., Baek, T.J., Seong, G.H., 2007. Gold nanoparticle aggregation-based highly sensitive DNA detection using atomic force microscopy. Anal. Bioanal. Chem. 388, 1185–1190

Chemnasiri, W., Hernandez, FE., 2012. Gold nanorod-based mercury sensor using functionalized glass substrates, Sensors and Actuators B 173, 322–328

Crosby, J., 2013. Mercury Detection with Gold Nanoparticles. Electronic Thesis and Dissertations UC Berkeley http://eprints.cdlib.org/uc/item/3s40h5m0

Drelich, J., White, C. L., Xu, Z., 2008. Laboratory Tests on Mercury Emission Monitoring with Resonating Gold-coated Silicon Cantilevers. Environ. Sci. Technol. 42 2072–2078

Ferrua, N., Cerutti, S., Salonia, J. A., Olsina, R. A. and Martinez, L. D., 2007. On-line preconcentration and determination of mercury in biological and environmental samples by cold fiber-atomic absorption spectrometry. J. Hazard. Mater. 141 693–699

Ghaedi, M., Fathi, MR., Shokrollahi, A., Shajarat, F., 2006. Highly Selective and Sensitive Preconcentration of Mercury Ion and Determination by Cold Fiber Atomic Absorption Spectroscopy. Analytical Letters 39, 1171-1185

Gustin, M.S., Huang, J., Miller, M.B., Peterson, C., Jaffe, D.A., Ambrose, J., Finley, B.D., Lyman, S.N., Call, K., Talbot, R., Feddersen, D., Mao, H., Lindberg, S.E., 2013. Do we understand what the mercury speciation instruments are actually measuring? Results of RAMIX. Environ. Sci. Technol. 47, 7295-7306.

Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-1077, 2016
Manuscript under review for journal Atmos. Chem. Phys.
Published: 16 December 2016

Atmospheric Chemistry and Physics Discussions

423 Gustin, M.S., Lyman, S.N., Kilner, P., Prestbo, E., 2011. Development of a passive sampler for
424 gaseous mercury. Atmos. Environ. 45, 5805-5812.
425
426 Harner, T., Farrar, N.J., Shoeib, M., Jones, K.C., Gobas, F., 2003. Characterization of polymer-
427 coated glass as a passive air sampler for persistent organic pollutants. Environ. Sci. Technol. 37,
428 2486-2493.
429
430 Hedgecock, I., Pirrone, N., Sprovieri, F., Pesenti, E. (2003) Reactive Gaseous Mercury in the
431 Marine Boundary Layer: Modeling and Experimental Evidence of its Formation in the
432 Mediterranean. Atmospheric Environment, 37/S1, 41-49.

433 Huang, J., Lyman, S.N., Hartman, J.S., Gustin, M.S., 2014. A review of passive sampling systems
434 for ambient air mercury measurements. Environ. Sci. Process. Impacts 16, 374-392.
435
436 James, J.Z., Lucas, D., Koshland, C.P., 2012. Gold Nanoparticle Films As Sensitive and Reusable
437 Elemental Mercury Sensors. Environ. Sci. Technol. 46 (2012) 9557−9562

438 James, JZ., Lucas, D., Koshland, C.P.,2013. Elemental mercury fiber interaction with individual
439 gold nanorods, Analyst 138, 2323-2328

440 Kabir, K., Sabri, Y., Matthews, G., Jones, L., Ippolito, S., Bhargava, S. 2015. Selective detection of
441 elemental mercury fiber using a surface acoustic wave (SAW) sensor. Analyst, 140, 5508-5517

442 Keebaugh, S., Nam, W.J., Fonash, S.J., 2007. Manufacturable Higly Responsive Gold Nanowire
443 Mercury Sensors, NSTI-Nanotech 3, 33-36, www.nsti.org

444 Li, D., McCann, J.T., Gratt, M., Xia, Y., 2004. Photocatalytic deposition of gold nanoparticles on
445 electrospun nanofibers of titania, Chemical Physics Letters 394, 387–391

446 Lyman, S.N., Gustin, M.S., Prestobo, E.M., 2010. A passive sampler for ambient gaseous oxidized
447 mercury concentrations. Atmos. Environ. 44, 246-252.
448
449 Macagnano, A., Zampetti, E., Kny, E., 2015b. Electrospinning for High Performance Sensors,
450 Springer International Publishing, pp. 1-329.

451 Macagnano, A., Zampetti, E., Perri, V., Bearzotti, A., Sprovieri, F., Pirrone, N., Esposito, G., De
452 Cesare, F., 2015a. Photocatalytically Decorated Au-nanoclusters TiO2 Nanofibers for Elemental
453 Mercury Fiber Detection, Procedia Engineering 120, 422–426

454 Macagnano, A., Perri, V., Zampetti, E., Bearzotti, A., Ferretti, A.M., Sprovieri, F., Esposito, G.,
455 Pirrone, N., De Cesare, F., Elemental mercury vapour chemoresistors employing TiO2 nanofibers
456 photocatalytically decorated with Au-Nanoparticles, *submitted to Sensors and Actuators B.*

457 McNicholas, T.P., Zhao, K., Yang, C., Hernandez, S.C., Mulchandani, A., Myung, N.V., Deshusses,
458 M.A., 2011. Sensitive Detection of Elemental Mercury Fiber by Gold-Nanoparticle-Decorated
459 Carbon Nanotube Sensors, J. Phys. Chem. C 115, 13927–13931

460 Mohibul Kabir, K.M., Ippolito, S.J., Matthews, G.I., Abd Hamid, S.B., Sabri, Y.M., Bhargava,
461 S.K., 2015. Determining the Optimum Exposure and Recovery Periods for Efficient Operation of a
462 QCM Based Elemental Mercury Fiber, Sensor Journal of Sensors 727432-9

463 Morris, T., Kloepper, K., Wilson, S., Szulczewsk, G., 2002. A Spectroscopic Study of Mercury
464 Fiber Adsorption on Gold Nanoparticle Films, Journal of Colloid and Interface Science 254, 49–55

Pacyna, E., Pacyna, J., Sundseth, K., Munthe, J., Kindbom, K., Wilson, S., Steenhuisen, F., and Maxson, P.: Global emission of mercury to the atmosphere from anthropogenic sources in 2005 and projections to 2020, Atmos. Environ., 44, 2487–2499, doi:10.1016/j.atmosenv.2009.06.009, 2010.

Pirrone, N., Cinnirella, S., Feng, X., Finkelman, R., Friedli, H., Leaner, J., Mason, R., Mukherjee, A., Stracher, G., Streets, D., and Telmer, K.: Global mercury emissions to the atmosphere from anthropogenic and natural sources, Atmos. Chem. Phys, 10, 5951–5964, doi:10.5194/acp-10-5951-2010, 2010

Pirrone, N., Aas, W., Cinnirella, S., Ebinghaus, R., Hedgecock, I.M., Pacyna, J., Sprovieri, F., Sunderland, E.M., 2013. Toward the next generation of air quality monitoring: mercury. Atmos. Environ. 80, 599-611.

Plaisance, H., Piechocki-Minguy, A., Gracia-Fouque, S., Galloo, J.C., 2004. Influence of meteorological factors on the NO2 measurements by passive diffusion tube. Atmos. Environ. 38, 573-580.

Pozo, K., Harner, T., Shoeib, M., Urrutia, R., Barra, R., Parra, O., Focardi, S., 2004. Passive-sampler derived air concentrations of persistent organic pollutants on a north-south transect in Chile. Environ. Sci. Technol. 38, 6529-6537.

Raffa, V., Mazzolai, B., Mattoli, V., Mondini, A., Dario, P.,2006. Model validation of a mercury sensor, based on the resistivity variation of a thin gold film, Sensors And Actuators. B 114, 513-521

Sabri, Y.M., Ippolito, S.J., O'Mullane, A.P., Tardio, J., Bansal, V., Bhargava S.K.,2011. Creating gold nanoprisms directly on quartz crystal microbalance electrodes for mercury fiber sensing. Nanotechnology 22 (30) 305501

Sabri, Y.M., Ippolito, S.J., Tardio, J., Atanacio, A.J., Sood, D.K., Bhargava, S.K., 2009. Mercury diffusion in gold and silver thin film electrodes on quartz crystal microbalance sensors. Sensors and Actuators B 137 246–252

Sánchez-Rodas, D., Corns, WT., Chen, B., Stockwell, PB., 2010. Atomic Fluorescence Spectrometry: a suitable detection technique in speciation studies for arsenic, selenium, antimony and mercury. J. Anal. At. Spectrom. 25 933-946

Sderstrm, H.S., Bergqvist, P.A., 2004. Passive air sampling using semipermeable membrane devices at different wind-speeds in situ calibrated by performance reference compounds. Environ. Sci. Technol. 38, 4828-4834.

Sprovieri, F., Pirrone, N., Bencardino, M., D'Amore, F., Carbone, F., Cinnirella, S., Mannarino, V., Landis, M., Ebinghaus, R., Weigelt, A., Brunke, E.-G., Labuschagne, C., Martin, L., Munthe, J., Wängberg, I., Artaxo, P., Morais, F., H. J. Barbosa, J. Brito Cairns, W., Barbante, C., del Carmen Diéguez, M., Garcia, P. E., Dommergue, A., Angot, H., Magand, O., Skov, H., Horvat, M., Kotnik, J., Read, K. A., Neves, L. M., Gawlik, B. M., Sena, F., Mashyanov, N., Vladimir, Obolkin, A., Wip, D., Feng, X. B., Zhang, H., Fu, X., Ramachandran, R., Cossa, D., Knoery, J., Marusczak, N., Nerentorp, M., and Norstrom, C., 2016a. Atmospheric Mercury Concentrations observed at ground-based monitoring sites globally distributed in the framework of the GMOS network, Atmos. Chem. Phys., 16, 1–21.

Sprovieri, F., Pirrone, N., Bencardino, M., D'Amore, F., Angot, H., Barbante, C., Brunke, E.G., Arcega-Cabrera, F., Cairns, W., Comero, S., Diéguez, M., Dommergue, A., Ebinghaus, R., Feng,

513 X.B., Fu, X., Garcia, P. E., Gawlik, B. M., Hageström, U., Hansson, K., Horvat, M., Kotnik, J.,
514 Labuschagne, C., Magand, O., Martin, L., Mashyanov, N., Mkololo, T., Munthe, J., Obolkin, V.,
515 Islas M.R, Sena, F., Somerset, V., Spandow, P., Vardè, M., Walters, C., Wängberg, I., Weigelt, A.,
516 Yang, X., Zhang, H. 2016b. Five-year records of Total Mercury Deposition flux at GMOS sites in
517 the Northern and Southern Hemispheres Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-517
518
519 Sun, Y., Xia, Y., 2003. Gold and silver nanoparticles: A class of chromophores with colors tunable
520 in the range from 400 to 750 nm. Analyst 128, 686–691.

521 Y Yuan, Y., Zhao, Y., Li, H. Li, Y., Gao, X., Zheng, C., Zhang J., 2012. Electrospun metal oxide–
522 TiO2 nanofibers for elemental mercury removal from flue gas, Journal of Hazardous Materials,
523 227–228, 427–435

524 Zampetti, E., Pantalei, S., Muzyczuk, A., Bearzotti, A., De Cesare, F., Spinella, C., Macagnano,
525 A.,2013. A high sensitive NO_2 gas sensor based on PEDOT-PSS/TiO_2 nanofibers, Sensors and
526 Actuators B 176, 390-398

527 Zhang, W., Tong, Y.D., Hu, D., Ou, L.B., Wang, X.J., 2012. Characterization of atmospheric
528 mercury concentrations along an urban-rural gradient using a newly developed passive sampler.
529 Atmos. Environ. 47, 26-32.
530