A smart nanofibrous material for adsorbing and detecting elemental mercury in air

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Abstract. The combination of the affinity of gold for mercury and nanosized frameworks has allowed for the design and fabrication of 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 gaseous elemental mercury (GEM) traces from the environment. The electrical properties of the resulting chemosensors were measured. A few minutes of air sampling were sufficient to detect the concentration of mercury in the air, ranging between 20 and 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 and very stable (due to the peculiar structure), requiring low power, low maintenance, and simple equipment.

1 Introduction

Mercury (Hg) is released into the atmosphere through human activities, predominantly fossil fuel combustion, and naturally, for example, from soil outgassing, volcanoes, and evaporation from the sea (Pirrone et al., 2010; Pacyna et al., 2010). One of the more troublesome issues in recent years has been to quantify not only the strength of emission sources but also the effects of the 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 (gaseous oxidized mercury, GOM; particle-bound mercury, PBM) and their chemical and physical characteristics (Lyman et al., 2010; Sprovieri et al., 2016, 2017). To be precise, total gaseous mercury (TGM) is mainly comprised of GEM with minor fractions of other volatile species (e.g., HgO, HgCl2, HgBr2, CH3HgCl, or (CH3)2Hg). However, in spite of the 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 quite volatile, whereas several 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 a 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 in protecting all environments. Several international initiatives and programs (i.e., the United Nations Environment Programme; UNEP)
have also made a tremendous effort to identify and quantify Hg pollution across the globe, especially the “hot-spots”, to reduce the risk of exposure to this neurotoxic pollutant. Policy makers are working toward a worldwide effort to support the construction of 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 5-year project called the Global Mercury Observation System (GMOS; www.gmos.eu) to create a coordinated global network to fill the gaps in emission monitoring and in the spatial coverage of environmental observations, mostly in the tropical regions and the Southern Hemisphere, thus adequately improving models and making policy recommendations (Sprovieri et al., 2016, 2017). To date, the GMOS network consists of more than 43 monitoring stations worldwide, including high-altitude and sea-level monitoring sites located in climatically diverse regions, including the polar areas (Sprovieri et al., 2016, 2017). 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 project to track persistent pollutants. The overall goal of this flagship project is to support the development of GEOSS (Global Earth Observation System of Systems) 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 the spatial coverage of observations. Since automated measurement methods for Hg often require power, carrier gases like argon, 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, and low-maintenance sensor is a primary technical issue to be solved for the sustainability of a global network such as GMOS. Previous research has highlighted the fact that Hg concentration levels in air vary greatly across different environmental locations; these include remote polar regions, background or rural, and urban locations with an average range between 1.5 ng m\(^{-3}\) (GEM) and 1 pg m\(^{-3}\) (GOM and PBM) depending on the speciation. Hence, for the determination of atmospheric Hg 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 in analytical methodologies have made it possible to study atmospheric Hg in different environmental locations. However, several limitations and difficulties are 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 further secure the reliability, traceability, and accuracy of Hg levels measured in air. Current air-monitoring devices are amply sensitive to detect the global background but are costly and high maintenance with complicated configurations and electricity requirements. A further limitation is the ultralow levels of ambient mercury in the atmosphere. The typical background gaseous elemental mercury (GEM) level of 1.5 ng m\(^{-3}\) is equivalent to 168 parts per quadrillion by volume (ppq\(_v\)). There is no other atmospheric compound being measured routinely, continuously, and automatically at this ultralow concentration. These features limit the scientific research community’s long-term ability to measure atmospheric Hg concentrations worldwide. Sampling and analysis of atmospheric Hg is conducted most commonly as GEM and/or TGM because of their greater abundance, even though both manual and automatic methods have been 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, there are several gaps in our current knowledge 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 that converts GOM back to GEM, and the gas–particle partitioning. This is partially due to the need for the identification of the chemical forms of oxidized Hg in the atmosphere and the 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 automated and/or semi-automated Hg measurements (Gustin et al., 2013; Pirrone et al., 2013), particularly in 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. Previously developed methods and new sampling systems, such as passive samplers, have been used to understand the 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 al., 2011; Zhang et al., 2012; Huang et al., 2014). However, because of
the differences in the design of passive samplers, ambient air Hg concentrations quantified by various samplers may not be comparable. In addition, the sampling rates (SRs) using the same passive samplers may depend on the environmental conditions and atmospheric chemistry at each site. It has also been highlighted that the performance of passive samplers may be influenced by meteorological factors (e.g., $T$ °C, RH, and wind speed), therefore inducing bias to the results of passive sampling (Plaisance et al., 2004; Sderstrm and Bergqvist, 2004). On the other hand, an incentive for developing simple and cost-effective samplers that are capable of monitoring over an extended period and require no technical expertise even at remote locations is now clear. In this work, we describe an alternative approach adopted in the place of conventional ones, demonstrating that the combination of the affinity of gold for Hg and a nanoscale-sized framework of titania provided the chance to create promising sensors for environmental monitoring in real time characterized by high sensitivity to the analyte. The novel sensor is a relatively simple and low-cost method for the measurement of the most abundant Hg form in ambient air (TGM or 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 for 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 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 would help policy makers to set regulations for different areas. The sensor features are related to the nanofibrous scaffold of titania capable of growing gold nanoaggregates through photocatalysis that are tunable in size and shape. Such a nanostructured layer, fabricated with electrospinning technology, firstly improves sensor features with respect to those of compact films, by enhancing the global number of analyte–sensor binding sites and reducing some bulk drawbacks. Secondly, the combination of metal oxides and metal nanostructures improves the sensitivity, allows the sensor to work at room temperature, tunes the selectivity towards different gas species by adjusting the surface-to-volume ratio of nanosized structures, and affects the sensor lifetime. Morphological, optical, and electrical aspects, as well as the sensing measurements of GEM fibers in air have been reported and discussed. When designed, the resulting Hg adsorbent or absorbent material was expected to be suitable for novel Hg sensor fabrication, since a similar nanofibrous scaffold doped with AuNPs was described in the literature as a filtering system capable of adsorbing and removing Hg vapor from the environment with an efficiency of nearly 100 % (Yuan et al., 2012). In previous work (Macagnano et al., 2017, 2015a), the authors reported a high sensitivity of the sensor capable of detecting up to dozens of ppt, despite the long time needed 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 (St. Louis, MO, USA) 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$_4$), and glacial acetic acid (AcAc$_4$). Ultrapure water ($5.5 \times 10^{-8}$ Scm$^{-1}$) was produced by Milli-Q EMD (Millipore Corporation, Darmstadt, Germany).

2.2 Electrospinning technology

Electrospinning (ES) 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 several micrometers from polymer solutions (or melts). It is currently the most economic, versatile, and efficient technology to fabricate highly porous membranes made of nanofibers and/or microfibers 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 traveling towards the collector, leading to the creation of continuous solid fibers as the solvent evaporates. The electrospinning apparatus used in the present study (designed and assembled in CNR laboratories) is comprised of a homemade clean box equipped with temperature and humidity sensors, a syringe pump (KDS 200; KD Scientific, Holliston, MA, USA), a grounded rotating cylindrical collector (45 mm diameter), and a high-voltage oscillator (100 V) driving a high-voltage (ranging from 1 to 50 kV) high-power AC–DC (alternating current to direct current) converter. The electrospinning solution (7.877 $\times$ 10$^{-5}$ M) was prepared by dissolving PVP in EtOH$_4$ and stirring (2 h). A 2 mL aliquot of a 1 : 4 ($w/v$) solution of TiiP solved in a 1 : 1 ($v/v$) mixture of AcAc$_4$ and EtOH$_4$ was freshly prepared and added to the 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 a 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 perpendicular to a grounded rotating collector at 15 cm of distance. 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 sca-
folds for the sensors. After deposition, PVP/TiO$_2$ composite nanofibers were left for some hours at room temperature to fully undergo the self-hydrolysis of TiIP (Li et al., 2004) and then annealed under an 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 physiochemical 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); it was manufactured in CNR laboratories through a standard photolithographic process (lift-off procedure) followed by Ti sputtering and Pt evaporation suitable to generate electrodes of the size reported above on a 4 in. oxidized silicon wafer. After electrospinning deposition, all the electrical signals of the resulting chemoresistors were recorded by an electrometer (Keithley Instruments, Inc., Cleveland, OH, USA; model 6517).

2.4 Titania nanofibers

Upon calcination, the diameters of the fibers extraordinarily shrank: the mean diameters of the 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 the 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 99 % of pores with an area of less than 10 µm$^2$ with more than 80 % pores being < 1 mm$^2$.

2.5 AuNPs/TiO$_2$NFs photocatalytic decoration

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). The resulting fibrous scaffolds were dipped into an aqueous solution containing HAuCl$_4$ and PVP (1.5 × 10$^{-3}$ M and 0.1 M, respectively) and exposed to UV light irradiation for specified intervals (UV lamp 365 nm; Helios; Italquartz, Calenzano, Italy). Depending on the gold nanoparticle sizes that were formed in photocatalysis, the dip solution changed from light yellow to purple. Samples were rinsed extensively with water and then air dried. Before the morphological, electrical, and sensing measurements, the samples were heated at 450 °C per 1 h to eliminate the PVP traces. Morphological characterization was provided through scanning electron microscopy (SEM; Jeol, Tokyo, Japan; JSM 5200; 20 keV) with pictures captured at 5 kV of accelerating voltage. AFM (atomic force microscopy) micrographs were taken in tapping mode using 190Al-G tips, 190 kHz, and 48 N m$^{-1}$ (Nanosurf AG, Liestal, Switzerland; FlexAFM). TEM (C-TEM; control transmission electron microscopy) micrographs were performed at 200 keV with an analytical double-tilt probe. TEM specimens were prepared by gently scraping the TiO$_2$ nanofibrous layer electrospun onto the silicon support and then collecting the nanofibers through adhesion upon contact with holey carbon thin film. UV–vis spectra were provided by a spectrophotometer (UV-2600; Shimadzu Corporation, Kyoto, Japan) by analyzing quartz slices coated with nanofibers. These substrates were able to collect fibers by electrospinning (20 min) and were then subjected to calcination according to the procedure described above; the final step was 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 the fibers during the calcination process.

2.6 Measurement setup

The sensor was placed in a suitable PTFE-made measurement chamber (0.7 mL volume) connected to an electrometer (Keithley 6517) capable of measuring the current flowing through the IDE when a fixed potential was applied to it and sending data to a PC. Dynamic measurements were carried out at room temperature using (i) a four-channel MKS 247 managing four MKS mass flow controllers (MFC) set in the range of 0–200 sccm and (ii) an Environics S4000 (Environics, Inc., Tolland, CT, USA) flow controller comprised of three MFCs supplying different flow rates (up to 500, 250, and 25 sccm) managed by its own software. Pure air (5.0: Praxair–Rivoira, Milan, Italy) was used as a gas carrier. A homemade PTFE (polytetrafluoroethylene) permeation tube filled with a suitable amount of Hg$^0$ was included within the delivery system to obtain set dilutions of Hg-saturated vapors. The tube was immersed in a thermostatically controlled bath; thus the desired Hg$^0$ concentration delivered to the sensor was achieved by tuning both the temperature of the permeation tube and the dilution flow. The Hg$^0$ concentration was checked by a Tekran® 2537A analyzer (Toronto, ON, Canada). Responses were calculated as ΔI/I₀, where ΔI was the current variation and I₀ was the current when synthetic pure dry air flowed. The sensor was restored after a quick thermal shot at 450 °C under a flow of pure air.

3 Results and discussion

Nonwoven mats made of PVP and amorphous TiO$_2$ were obtained through the combination of electrospinning and sol-
Figure 1. A sketch of an electrospinning setup comprised of a syringe and a grounded rotating cylinder collector where the samples are placed for coverage (a); a piece of TiP/PVP nanofibrous fabric removed from the substrate after 1 h of electrospinning deposition (b); a reddish purple aqueous solution of HAuCl₄/PVP after UV light irradiation treatment and (c) a piece of the TiO₂ (anatase) nanofibrous fabric obtained after TiP/PVP annealing.

Figure 2. A sketch of the photocatalytic process occurring on the fiber surface (a); an SEM picture of a dense nanofibrous network of AuNPs/TiO₂ coating a silicon wafer (b); and a C-TEM micrograph of fibers finely decorated with gold nanoparticles (the darkest ones) bound without using any additional linker (inset).

gel techniques (Fig. 1). The ES deposition proceeded for 20 min on oxidized silicon wafers and IDEs properly fixed on the surface of a conducting and 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 the thickness and the consistency of the mats changed. More specifically, a 1 h deposition provided the formation of a thicker, white, soft fabric that is hygroscopic, soluble in both water and polar solvents, and easily peeled off (Fig. 1); instead, a 20 min deposition generated a fibrous film adhering to substrates too thin to be weeded and thus preferred for sensor fabrication. The calcination process caused a complete degradation of PVP with the formation of crystalline TiO₂ (anatase) and a significant shrinkage in the fiber dimensions (60–80 nm diameter; 5–40 nm grain size). By exploiting the photocatalytic properties of titania (anatase), a tunable decoration of fibers with gold nanoparticles could be achieved by dipping the fibrous mats in a proper aqueous solution (HAuCl₄; PVP) under UV light irradiation (Li et al., 2004; Macagnano et al., 2015a). The photocatalytic reaction was confirmed by the color change of the solution (reddish purple from light yellow; Fig. 1). By changing both the UV irradiation exposure time and the PVP concentration as a capping reagent, the morphology, size, and density of the gold nanoparticles could be tuned (Macagnano et al., 2017).

In the present work, among a series of differently coated fibrous layers, only the fibrous nanocomposites that were conductive at room temperature were selected; their electrical and sensing features were then investigated. The controlled gold deposition was due to the photo-excited electrons on the surface of TiO₂ nanofibers that were able to reduce the gold ions, thus inducing gold metal deposition (Fig. 2 sketch). The capping reagent was responsible for the shape...
of the particles. The surfaces of the nanofibers, as observed in SEM micrographs (Fig. 2a), appeared densely decorated with globular nanoparticles. In the C-TEM image (Fig. 2, inset) the gold nanoparticles appeared darker with spherical or quasi-spherical shapes. The single particle sizes ranged between 2 and 20 nm with a $7.8 \pm 3$ nm average diameter (Macagnano et al., 2017). Gold nanoparticles grew directly onto the nanofibers, and their adhesion appeared relatively strong (despite van der Waals forces), since they resisted both water rinsing and fiber scratching for TEM analyses.

As a result of the photocatalytic process, the white porous mat became purplish violet. As reported in the spectrum of the AuNP/TiO$_2$ hybrid system, a characteristic absorbance band appeared at about 543 nm, which corresponded to the surface plasmon resonance (SPR) of the AuNPs (Sun and Xia, 2003). A red shifting and broadening of the absorbance band was observed with the increase in AuNP size and fiber loading, respectively (data not shown). The color strictly depends on the size of the nanoparticles and then their agglomeration at the solid state. According to Bui et al. (2007), such a band-broadening phenomenon is due to the electric dipole–dipole interactions and coupling occurring between the plasmons of neighboring particles, whereas nanoparticle agglomeration phenomena occurred.

Due to these features, UV–vis absorption spectroscopy has been used in the literature as a technique to reveal the changes in the size, shape, and aggregation of metal nanoparticles in liquid suspension after exposure to heavy metals as Hg$^0$ (Morris et al., 2002). Both the blue-shifted wavelength and its extent were proportional to the amount of Hg$^0$ that entered the liquid suspension. Similarly, the TiO$_2$ nanofibrous layers coating the metal electrodes of the transducers (Fig. 5) changed color after photocatalytic treatment (from white to pink).

Figure 4. UV–vis spectra of AuNPs/TiO$_2$ nanofibers before (blue) and after a 15 min exposure to 2 ppm of Hg$^0$ (gray).

The IDE layout (Fig. 5) is comprised of a set of interdigitated electrodes that occupies an area approximately $3 \times 5$ mm, is completely coated with the sensitive fibers, and has two bonding pads ($2 \times 2$ mm) that will be connected to the electrometer (DC voltage). Such a planar interdigitated electrode configuration is most commonly used for conductometric sensing applications.

Figure 5. The chemosensor fabrication and final structure: IDE dipped (a) and exposed to UV light (b) for gold decoration.

Figure 6 depicts the current–voltage (I–V) curve of a chemosensor under a synthetic dry air flow. However, the curve shape was unaltered when air or nitrogen was flushed over the fibers (Macagnano et al., 2015a), suggesting that oxygen concentrations poorly affected the electrical properties of such a chemoresistor. The resistance value of the IDE coated with TiO$_2$ nanofibers before photocatalysis was too high at room temperature to contribute directly to the final current value. The resulting linear shape (ohmic behavior) within the selected voltage range (from $-3$ to $+3$ V) showed a constant resistance value for the sensor. The very low value of resistance ($\sim 1.2$ k) provided the chance to work at low voltage, with consequent effects on the energy consumption as well as the lifetime of the material. Moreover, the linearity of the I–V curve let us suppose that the sensing scaffold
had a good adhesion to the metal electrodes. The electron conductivity has been assumed to occur according to the percolation model (Macagnano et al., 2017; Müller et al., 2003), since the titania at room temperature was expected to be an insulating matrix. When it is metal doped, the electron conductivity is ruled by thermally activated electron tunneling from one metal island (gold nanoparticles) to the other. However, the conductivity of the nanocomposite is lower than that of pure metal (gold) because the electron mean free path is greatly reduced due to the presence of the dielectric (the titania crystals). The electrical features, such as the reproducibility of the fabrication process, of this conductive device have been previously investigated by the authors (Macagnano et al., 2017, 2015a), showing encouraging results for the development of a low-cost sensor for mercury detection. However, in spite of the high sensitivity (LOD: 2 ppt) of the sensor, a too-long response time was necessary to detect traces of Hg$^0$, especially if compared to the monitoring instrumentation (Ghaedi et al., 2006; Sanchez-Rodas et al., 2010; Ferrua et al., 2007) commonly involved in OES detection. The long response time was in part due to the layout of the measuring system, since the sensor was previously housed in a quartz bottle of 100 mL in volume. Additional time is caused by the adsorption of Hg$^0$ traces from the surrounding environment (measuring chamber) up to the achievement of a sufficient number of Hg$^0$ atoms adsorbed on the surface sensor to be electrically revealed. However, this sensor looks extremely encouraging if compared to other sensors currently involved in detecting mercury in air (Drellich 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 and Hernandez, 2012; Sabri et al., 2011; Keebaugh et al., 2007; Crosby, 2013; McNicholas et al., 2011).

Many sensors have been designed and investigated to detect the several forms of mercury. Most of them have exploited the strong affinity between mercury and gold (Ford and Pritchard, 1971; Joyner and Roberts, 1973). Several studies have documented changes in the electrical properties, work function, and resistance of thin gold films upon exposure to various concentrations of mercury vapor. For instance, an array of microcantilevers with different sizes developed by Drellich et al. (2008) could measure different ranges of mercury concentration (between 37 and 700 µg m$^{-3}$) and were capable of revealing up to 10 pg Hg$^0$ adsorbed. However, their sensing system required both a dust-free gas carrier and a heating procedure (350 °C for 20 min) to regenerate the sensors. Gold-based conductometric sensors, too, have been designed and used to reveal mercury vapor through their electrical resistance changes (Raffa et al., 2006), but their sensitivity often seemed poor (about 1 µg m$^{-3}$). Quartz crystal microbalance (QCM) devices, too, have been used as Hg$^0$ vapor sensors (Sabri et al., 2009) due to their high portability, selectivity, and unnecessary sample pretreatments. Their absorptive capacity was improved (700 ng cm$^{-2}$) when the gold electrodes were made rough (more binding sites). However, additionally to the natural affinity of gold for mercury, the increased facility for producing and depositing nanoparticles with noble metals facilitated their use as possible sensors, especially in aqueous environments (Nolan and Lippard, 2008; Chemnasiri and Hernandez, 2012; Ratner and Mandler, 2015; Dong et al., 2015). James et al. (2012) developed a highly sensitive chip working in an SPR mode based on gold nanoparticles (5 nm diameter) to monitor Hg$^0$ vapors. Such a system was able to linearly detect mercury concentrations from 1 to 825 µg m$^{-3}$, but it was strictly related to the flow rate: increasing the flow velocity (and mass transfer rate) increased the peak shift rate. The time resolution was limited by the rate of adsorption, which increased with the Reynolds number; at the greatest flow rate tested (57 LPM), an ambient mercury measurement (1 ng m$^{-3}$) needed 410 h to shift 1 nm, but accelerating the flow rate could reduce the time resolution.

Gold thin film technology has been recently adopted in a commercial portable device to detect mercury (Jerome® J405 mercury vapor analyzer) with a 0.01 µg m$^{-3}$ resolution and a 750 ± 50 cc min$^{-1}$ flow rate (http://www.azic.com/jerome/j405/). A series of scrubbers and filtering systems are able
to reduce the effects of interferents on the gold–mercury interaction. The flow rate and the measuring system layout were key parameters in the proper working of the device. These features also seem to be significant for the ES-based chemosensor.

The measuring chamber was designed to reduce the volume (0.7 mL) and 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.

Sensing measurements, i.e., the current (or resistance) changes, were carried out in a continuous mode. The sensor measurements resulted in a change of the whole current (or resistance, i.e., $I = V/R$) according to Ohm’s law. Firstly, the sensor was exposed to a flow of $\text{Hg}^0$ in air with a concentration of 800 ppbv, for 1 min (Fig. 7, right), and then air was flowed through the measuring chamber to clean the sensor surface. A rapid decrease in current was recorded ($1.056 \times 10^{-7} \text{ A s}^{-1}$) when $\text{Hg}^0$ entered the measuring chamber. The current curve trend slightly changed when clean air entered, quickly stabilizing to about the same current values reached for $\text{Hg}^0$ adsorption. Such an effect was probably due to the $\text{Hg}^0$ still housed inside the delivering tubes. The polluted line contribution was confirmed by further measurements (data not shown), where the slow current decrease completely disappeared when clean air never passed through the part of the tubes that had carried Hg vapors. Due to the strong affinity between Au and $\text{Hg}^0$, a 3 min thermal treatment was necessary to remove mercury from the layer and obtain the same starting current value.

Figure 8 depicts the normalized sensor response rate, i.e., the normalized current change per second, toward the increasing concentration of GEM (ranging between 20 and 160 ppbv). Within this study, the selected flow rate was kept at 50 sccm in order to avoid turbulence effects. The resulting logarithmic curve describes the relationship between $\text{Hg}^0$ concentration and the response time; small variations in $\text{Hg}^0$ concentration up to 80 ppbv, were able to deeply change the response rate. In contrast, a higher concentration seemed to only weakly affect this sensing feature. Thus, since a strong relationship is recorded between the concentration and the

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### Table 1. Linear fitting parameters of 10 min sensor responses to $21 \text{ ppb} \leq [\text{Hg}^0] \leq 106 \text{ ppb}$.

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<th>ppbv</th>
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<td>70</td>
<td>$-3.19082 \times 10^{-9}$</td>
<td>2.55882E-11</td>
<td>0.93</td>
</tr>
<tr>
<td>106</td>
<td>$-4.83599 \times 10^{-9}$</td>
<td>2.67462E-10</td>
<td>0.88</td>
</tr>
</tbody>
</table>
response time of mercury under 80 ppb, it is possible to find a correlation between the slope of the transient responses within the early minutes of the sensor response and definite concentrations of Hg\(^0\) in air. Figure 9 depicts the linear fitting of 10 min sensor responses when increasing concentrations of mercury were flowed over the sensor. The related data are reported in Table 1.

A linear relationship has been reported between the response rate and the concentration of Hg, according to Eq. (1):

\[
y = (\pm 4.56226 \times 10^{-12}) \cdot [\text{Hg}^0], \quad [\text{Hg}^0] < 100 \text{ ppb},
\]

\[
\text{SE} = \pm 1.504 \times 10^{-12}, \quad R^2 = 0.99675.
\]  

Therefore, when the concentration of Hg increased, the response curve slope changed too linearly, allowing a limit of detection of about 1 ppb, for a 10 min exposure (50 sccm). Regarding the main interfering compounds, at room temperature and in dark conditions the measured current should be due to AuNPs decorating titania fibers; only chemical compounds interacting with gold are therefore expected to be mostly responsible for the current changes (i.e., halides and sulfides). Thus in a blend of other chemicals, this sensor has been designed as a relatively selective sensor able to greatly decrease the environmental disturbances, allowing the investigator or manufacturer to design and then implement easier strategies to prevent contamination from the environment (selective filtering systems or coatings). Among the common potential contaminants, the authors previously investigated water vapor influence (% RH) and reported no effects on the electrical signals (Macagnano et al., 2015a).

4 Conclusions

The adopted sensing strategy focused on the strong affinity of mercury for gold combined with the nanostructure properties. By exploiting the photocalytic 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 3-D framework of titania covering the electrode-sensing area of the properly designed chemoresistors (IDEs). By exploiting the photocalytic properties of titania, AuNPs have been grown on nanofibers. Such a sensor was able to work at room temperature and was sensitive to Hg\(^0\), even after a few minutes of exposure to polluted air. Since it is composed of titania and gold, it seems to be robust and resistant to common solvents and VOCs in the air (as reported in the literature). Preliminary results suggest that the short thermal treatments necessary to desorb mercury from AuNPs did not seem to affect the electrical properties of the device. Depending on the sampling strategy, a sensing device based on such a chemosensor could be designed for real applications, specifically for the real-time monitoring of polluted sites. A few minutes of air sampling are sufficient to quantify the concentration of mercury in the air in the range between 20 and 100 ppb, (LOD: 1 ppb), without using traps or gas carriers. Changing parameters such as the flow rate, density, and size of the gold particles is expected to significantly improve the LOD and the response time. Finally, a modified transducer layout could be designed to better exploit the adsorptive capacity of the 3-D nanofibrous framework. However, further investigations are necessary to also assess the effects of the physical parameters of the environment, such as temperature fluctuations and UV light (that should activate the titania surface in adsorbing oxidized Hg), as well as chemical ones, such as volatile organic compounds and gas (like halides and sulfides), which are known to interfere with the adsorption process of Hg\(^0\) on gold.

Data availability. All the data reported within the paper resulted from electrical and optical measurements carried out in the laboratory using innovative materials and devices. The data have been uploaded onto the GMOS central database and are available upon request at http://sdi.iia.cnr.it/geoint/publicpage/GMOS/gmos_historical.zul (GMOS Database, 2014). The GMOS database coordinated by the CNR-IIA is part of the GMOS CyberInfrastructure (CI), which supports advanced data acquisition, storage, management, integration, mining, and visualization. All GMOS stations provide near-real-time raw data that are archived and managed for the QA/QC process by the GMOS-Data Quality Management System (G-DQM), a web-based application which is part of the GMOS-CI, developed by the CNR-IIA in 2011, and is currently ongoing.

Competing interests. The authors declare that they have no conflict of interest.

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