

Cadmium Contaminated Water Detection with Interdigitated Electrodes and Microfluidic System

Suroosh Ali, Vishal Rathee, Jayu Kalambe, Suresh Balpande

Abstract: In this paper, a sensing mechanism based on Microfluidics, for detection of dangerous heavy metals in liquids has been prepared. A pair of interdigitated electrodes (IDE's) was fabricated on a glass substrate using a silver ink constituting nano-particle size molecules. The impedance between the pair of IDEs was measured for different samples with varied cadmium concentrations. The capability of these IDE's for detecting very low to low concentrations (Picomolar to Millimolar) of Cadmium was explored. The response curve based on Impedance Spectroscopy Analysis of the IDE's with the help of a Digital impedance testing equipment was plotted for detection of Picomolar, Nanomolar, Micromolar and Millimolar concentration levels of Cadmium sulfide compound in water. The averaged out results obtained showed coherently decreasing impedance values with increasing Cadmium sulfide concentrations in water.

Index Terms: Impedance Spectroscopy, Interdigitated electrodes, Micro-fluidic sensing mechanism, Water dissolved Cadmium

I. INTRODUCTION

Contamination of Water because of hazardous heavy metals is a widespread concern that needs meticulous attention and awareness to retain the safe and unadulterated water with unharmed quality. Millions of children, below the age of five pass away each year as a result of infections and disorders caused by such destructive water intake [1, 2]. Heavy metals in environment are a result of liberation of toxic atomic and nuclear wastes from industries into water-bodies, unintentional spills, farming and mining activities, etc. [3]. While these metals are omnipresent in the surroundings and some are vital nutrients when present in micro level, but all are toxic to biota above some threshold concentrations [4].

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The existing Spectrometric detection methods such as Flame Atomic Absorption Spectrometry (FAAS), Atomic Absorption Spectrometry (AAS) and Graphite Atomic Absorption Spectrometry (GFAAS) are too complex, time-consuming, expensive processes and require hefty setups. This fact accentuates the

need to build a system which will have the ability to detect and indicate the concentrations of heavy metals in a portable, savvy and streamlined way.

Group IIB element of the periodic table, 'Cadmium' is a heavy metal and its excessive omnipresence causes callous dangers to the mankind. It is found that universally about 7000 t/year of cadmium is emitted from varied sources [5, 6]. Along with copper, zinc and lead, cadmium coexists in the form of minerals. It finds widespread utilization in manufacturing methods as an operator which is anticorrosive, as an added substance in PVC items, as a color dye, in nuclear power plants as an absorber of neutrons and in manufacturing of NiCd rechargeable batteries [7]. Manures consisting of phosphate too demonstrate enormous cadmium content. . Despite the fact that few cadmium inherent items can be reused, a major share of the basic cadmium contamination is brought about by dumping and burning cadmium-embedded waste [8-10]. Cadmium exists as compounds of Sulfides, Sulphates, Chlorides, etc. in water. The metal itself is insoluble in water, but its compound illustrates varied solubility. Cadmium Sulfide (CdS), has low solubility, with a reported limit of 1.3 mg/L whereas the Sulfate (CdSO₄) and Chloride (CdCl₂) salts of Cd demonstrate quite high dispersion property in water with reported limit of 608,000 mg/L and 1,680,000 mg/L respectively [11, 12]. Excessive cadmium pesters the neural sensors, human digestive and urinary systems with undesirable physical and mental human wellness. Furthermore, over cadmium consumption causes tumors and cancers to significantly develop because of the solid bonds that the sulfur atoms in the human body make with these metal particles [13, 14]. An Impedance spectroscopy technique is utilized here to detect cadmium which gives the advantage of operations without any labels on the sample, miniaturization friendly and relatively inexpensive for validation and quantitative interpretation of experimental impedances [15]. The sensing mechanism based on IDE's as measuring elements offer vital advantage of requirement of low sample volumes and avoidance to

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monotonous polishing of electrodes. The interdigitated form of electrodes in general raises the sensing power and detection limits towards the contaminant and is appropriate for developing analytical devices and for other chemical science studies [16].

agencies, Food manufacturing industries, Hospitals etc. in urban as well as rural areas.

In this work, Ag based ink was employed to fabricate two paired (4 teeth) Interdigitated electrodes on a substrate i.e. glass which was then sintered for 10 minutes on a heating mantle at about 280°C to acquire the IDE structure. Polydimethylsiloxane (PDMS) was deposited on the glass substrate around the IDE to create a boundary for the obstruction of the liquid flow. The potential of the IDE mechanism was tested by observing the response based on Impedance Spectroscopy for Cd compound based water samples with the assistance of a handheld LCR meter. Sample concentrations of cadmium sulfide (CDS) in DI water prepared to a tune of Picomolar, Nanomolar, Micromolar and Millimolar were analyzed for impedance variations at a frequency of 100 Hz utilizing digital LCR meter.

II. EXPERIMENTATION

A. Processing of Materials for sample preparation

A translucent glass petri-dish having a thickness of 3 mm was used as the base for development of sensing mechanism. A nano molecular silver conductive ink (TEC-IJ-010) from InkTec Inc. was used for preparing metallic electrodes. Heat curable silicon-based Polydimethylsiloxane (PDMS), an organic polymer was bought as two-part silicone elastomer kit (Sylgard® 184) from Dow Corning and was used to create the boundary of IDEs to obstruct the liquid flow. Difference in the properties of the above two materials (Given in Table 1) ensured efficient isolation and non reactivity amongst them.

Table 1: Properties of Silver metal and PDMS

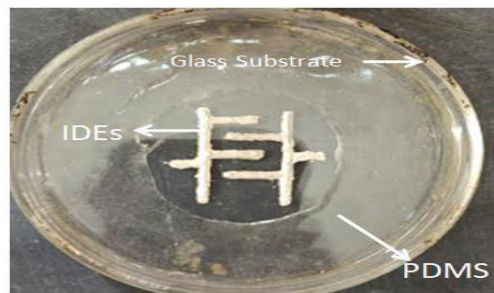
	Silver	PDMS
Density	10.50 g/cm ³	965 kg/m ³
Melting Point	962°C	-49.9 – 40°C
Boiling Point	2212°C	200°C
Hardness	25	46
Thermal Conductivity	419 W/mK	0.15 – 0.2 W/mK

Crystallised Cadmium Sulfide (CdS) was bought from the Chemical Company Sigma–Aldrich® to prepare sample and Agilent’s U1732C Digital Handheld LCR Meter having maximum frequency range upto 10 kHz was used for its impedance measurement purpose. A J-SIL High Precision Micropipette was used for the preparation of millimolar, micromolar, nanomolar and picomolar CdS solutions. The suspension of CdS was done in DI water to form solutions

of 1 to 13 mM, μM, nM and pM concentrations. All the prepared solutions were stored in 100 ml aliquots at 4.44°C before use.

B. Fabrication of Microfluidic Sensing Platform

The Silver ink used to formulate the interdigitated electrodes on the substrate made up of glass was preferred over other metallic pastes like copper/aluminium for its good reliable electrochemical properties over a wide variety of applications and processes. For the fabrication of the electrodes, Gray resin moulds (shown in Fig. 2) were created using 3D printing process. The sensing area consisted of two pairs of IDEs that were 10 mm (10000 μm) long with spacing and width of 4 mm (4000 μm) and 2 mm (2000 μm) respectively, selected keeping futuristic mobility and space constraints for handheld sensor in mind. The glass substrate was first cleaned with acetone to remove the dirt and fingerprints. A PDMS based solution was prepared by taking the PDMS and curing agent in a ratio of 10:1 and pouring it around the fabricated IDE to create a boundary for the obstruction of the liquid flow. The sensor structure was then kept on a heating mantle at around 150°C for about 15 minutes to cure the PDMS



solution [17].

Fig.1: Fabricated IDE on glass substrate



Fig.2: Fabricated Gray resin 3D printed moulds

C. Experimental Setup

The calibration of the wires and probes in the experimental setup (shown in Fig.3) was done before taking measurements with offset null adjusted properly. All the measurements were performed at normal ambient temperature. A simple syringe was used to pour 0.2 ml of the prepared sample drop-wise on the fabricated IDEs. The crocodile clip leads were connected to the handheld digital LCR meter and these leads were connected to the two electrodes of the IDE to measure the impedance values. The magnitude of impedance of the device was measured at a frequency of 100 Hz using LCR meter. The lowest frequency of 100 Hz was selected as it is less likely to show up any stray reactances and higher frequencies may modify the effective value of the impedance. The response of the sensing mechanism was observed and plotted.

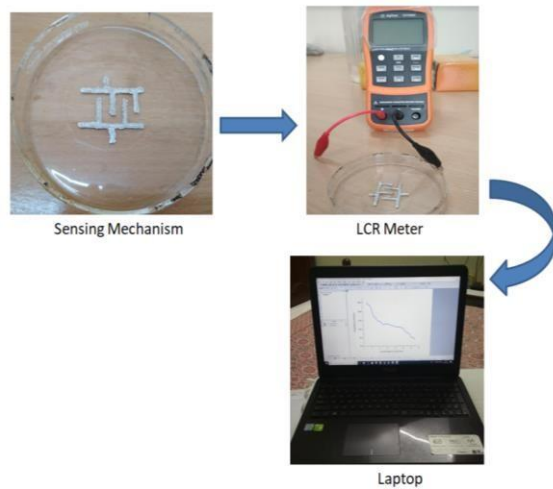


Fig.3. Experimental Setup

III. RESULTS

The fabricated sensing mechanism was tested for different concentrations of CdS in DI water. Figure 4, 5, 6 &

7 show the dynamic impedance response curves of the mechanism for the four different concentrations of CdS (pM to mM) having a step-size of 10^3 determining the cadmium concentration present from 0.187798×10^{-9} gm to 0.014446 gm with an impedance variation ranging from 343-163 k Ω , 485-275 k Ω , 1600-750 k Ω and 420-70 k Ω respectively. The decrement in the value of impedance with linearly increasing sample concentration (CdS dissolved in DI water) arises due to the increase in capacitance and decrease in resistance values of the interdigitated electrodes, since impedance is inversely

proportional to capacitance and directly proportional to the resistance between the electrodes.

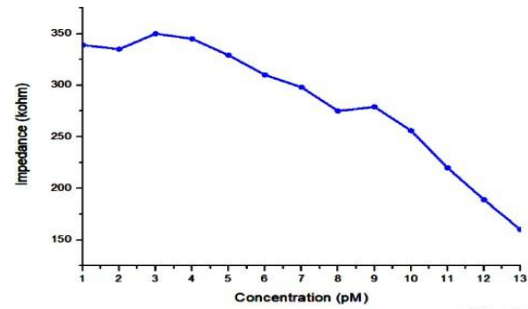


Fig.4: Picomolar CdS concentration

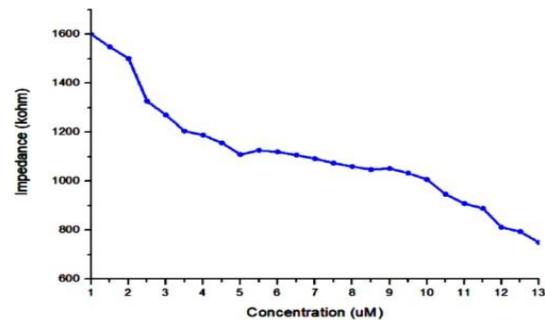


Fig.5: Nanomolar CdS concentration

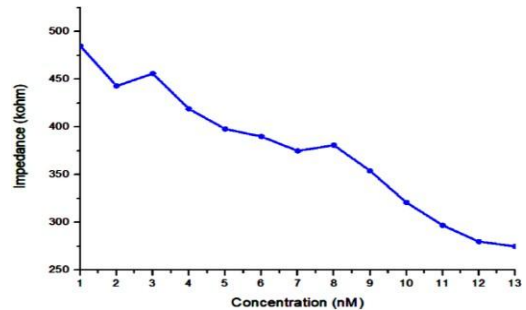


Fig.6: Micromolar CdS concentration

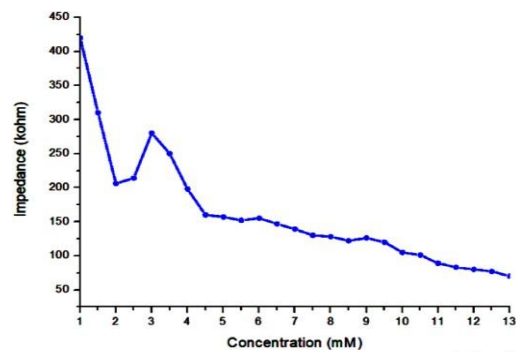


Fig.7: Millimolar CdS concentration

IV. CONCLUSION

In this work, a Microfluidic Sensing mechanism for detection of a standalone suspended heavy metal Cadmium was successfully investigated with the help of Silver ink-based interdigitated electrodes. The mechanism's potential to differentiate between different concentrations of prepared CdS was determined. The mechanism's impedance response curves showed an easily measurable impedance range between 1600 kΩ to 70 kΩ with standard deviations of 61.90, 56.89, 196.41 & 64.18 respectively for increasing concentrations of Cd. The results indicate coherence with study of gradual decrease in impedance with increasing sample concentration while some of the hikes were observed in between measurements due to environmental effects, presence of static electric field and lack of automation in the measurement setup

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