



Tungsten oxide electrode for measurement of ultralow liquid flow velocity



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ABSTRACT

This study demonstrated the application of a tungsten oxide electrode to the measurement of ultralow liquid flow velocity based on streaming potential, using a pH glass electrode and temperature sensor to compensate for changes in open circuit potential induced by pH and temperature, respectively. A change in flow velocity on the surface of the tungsten oxide electrode caused the expected shift in its open circuit potential. Experimental results showed that the open circuit potential was linear relative to the logarithmic flow velocity, and the flow sensor exhibited excellent reproducibility and a large dynamic range, with the ratio between measurable maximum and minimum flow velocity surpassing 60:1. Moreover, a higher sensitivity was observed in the alkaline solution relative to the acidic solution, indicating that the alkaline solution is more suitable for this type of measurement. In conclusion, the tungsten oxide flow sensor provides a cheap, simple, adaptable, and energy-saving solution for the measurement of ultralow liquid flow velocity.

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1. Introduction

Flow measurement is an important issue in industrial and agricultural production, national defense, trade, and various aspects of people's livelihood. Currently, electromagnetic [1], ultrasonic [2], differential pressure [3], vortex [4,5], turbine [6], and thermal [7] flowmeters are the main sensing instruments adopted for the flow measurement.

The flow velocities of liquids used in the semiconductor industry and chemical analyses are usually very small. However, conventional flowmeters listed above usually cannot measure low flow velocity because of inadequate sensitivity, and they also do not have sufficient long-term stability. Therefore, there is a strong demand for a new sensor that can measure low flow velocities of liquids.

Many sensors have been developed in order to measure very low flow velocity because of demand in the semiconductor, medical, and chemical industries. Takamoto et al. improved the ultrasonic flowmeter so that it could reliably measure a flow rate of less than 1 mL/min with a standard deviation of approximately 0.01 mL/min in a glass tube with diameter of 0.5 mm [8]. Some thermal flow velocimeters, which comprise heater elements and temperature sensors, have demonstrated a capability for measuring low liquid flow velocities. For instance, Dinh et al. used a hotwire to

measure a low flow rate of 0.05 mL/min; in this flow channel, the flow velocity was lower than 3 mm/s [9]. A flow velocity measurement method using the photothermal effect has been developed to realize a small, noninvasive microflow velocimeter, with a dynamic range of 25–300 $\mu\text{L}/\text{min}$ in a microchannel; the minimum flow velocity that could be measured was ~ 3 cm/s [10]. There are other types of microflow measurement methods, including microparticle imaging velocimetry, in which flow velocities are obtained from the motion of tracers dispersed in the fluid; a bulk velocity of approximately 50 $\mu\text{m}/\text{s}$ could be measured [11]. However, the tracer particles must be introduced into the solution for the flow measurement and they can be a disturbance to chemical reactions. Another technology uses an integrated micro-opto-fluidic flow-sensor, in which a fiber cantilever transduces flow rate into optical transmission that could measure a minimum flow rate of ~ 7 $\mu\text{L}/\text{min}$ [12]. An ultrathin curved-up microcantilever flow sensor has been developed and calibrated for measurement of small flow velocities based on the changes in electrical parameters induced by pressure; experimental results showed that the microcantilever could measure flow velocities in the range of 0–23 cm/s [13]. Laser systems using fiber Bragg gratings and doped fibers have been developed for use as sensors capable of measuring flow rates in a linear range from 0 to 200 mL/s, with a resolution of approximately 8 mL/s in a pipe with diameter of 15 mm [14].

Although these methods meet the requirement of measuring low flow velocities, ultralow flow velocities could not be measured reliably, and most of these methods require complex systems

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and data processing. Meanwhile, efficient, stable measurement of ultralow flow velocity in open channels remains a challenge.

Park and Choa [15] have reported measurement of liquid flow rate by self-generated electrokinetic potential on the microchannel surface of a solid. In this paper, we propose the use of a tungsten oxide electrode as an indicator electrode and an Ag/AgCl electrode as a reference electrode to sensor ultralow liquid flow velocities based on streaming potential. As the core of flow sensor, the tungsten oxide electrode has a bilayer structure consisting of a compact inner layer of WO_3 and a loosely bound outer hydrated layer of $WO_3 \cdot xH_2O$. Lillard et al. has reported a corresponding decrease in the thickness of the surface layer under hydrodynamic effect [16]. In addition, our previous work demonstrated a dramatic change in the open circuit potential of the tungsten oxide electrode under the hydrodynamic effect. The new open circuit potential of the tungsten oxide electrode could remain stable under the fixed rotation speed of a magnetic rotor, gradually recovering when the magnetic stirrer was turned off. Repeated cycles revealed that tungsten oxide could maintain its stability under hydrodynamic conditions where oxide dissolution had altered the potential signals [17]. Thus, the tungsten oxide electrode could be used to measure liquid flow velocity. The responses of open circuit potentials to flow velocities were studied in solutions of different pH. Therefore, to obtain accurate flow velocity, a pH glass electrode and a temperature sensor were used to compensate for changes in open circuit potential induced by pH and temperature.

2. Materials and methods

2.1. Chemicals and reagents

In this study, 0.05 M potassium acid phthalate (pH 4), 0.025 M mixed phosphate (pH 6.86), 0.01 M borax (pH 9.18) and dilute hydrochloric acid were used as measuring solutions. All solutions were prepared by using deionized water (electric resistance of 18.2 MX) supplied by a MilliQ water purification system (Millipore, Billerica, MA).

2.2. Electrode preparation

A Teflon-coated tungsten electrode with a diameter of 2 mm (purity >99.99%) was purchased from Tianjin Aidahengsheng Technology Corporation (China). Then, the tungsten electrode was abraded with grinding paper, finely polished with 0.5- μ m aluminum oxide particles, and finally rinsed with distilled water. For producing the tungsten oxide electrode, we electrochemically oxidized the tungsten electrode in a 0.1 M H_2SO_4 solution by cycling the potential between 1 and 2 V vs. a saturated calomel electrode for 20 cycles at a scan rate of 20 mV/s. Subsequently, the electrode was immersed in the same solution for 12 h, washed with distilled water, and dipped in a buffer solution of pH 6.86 [17,18].

2.3. Experimental setup and signal acquisition

The sensor section of the system consisted of a tungsten oxide electrode, a pH glass electrode, an Ag/AgCl electrode, and a temperature sensor (PT1000). As seen in Fig. 1, the temperature sensor, pH glass electrode, and Ag/AgCl electrode were placed in the storage tank. The tungsten oxide electrode was immersed and fixed in a flow cell with an inner diameter of 10 mm. The solution in the storage tank was pumped to the flow cell by a peristaltic pump (BT50S, Baoding LeadFluid Technology, China), and then flowed back to the storage tank. The adjustable range of average flow velocity in the flow cell was 0.08 to 40 mm/s with a resolution of 0.08 mm/s. The PT1000 sensor was the temperature sensor used to continuously monitor the temperature of solution. The tungsten

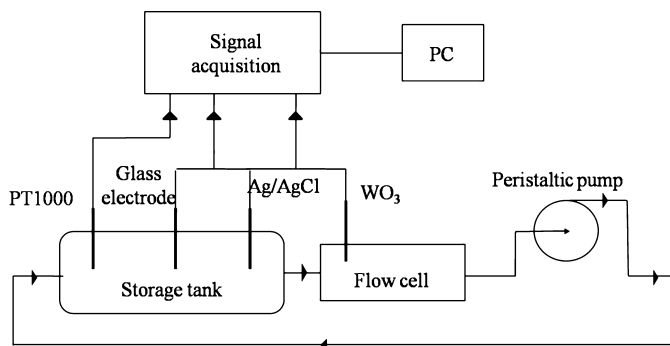


Fig. 1. Experimental setup.

oxide electrode and pH glass electrode were used as an indicator electrode separately, and the Ag/AgCl electrode was used as a reference electrode, the open circuit potentials between the indicator and reference electrodes were measured. The potentiometric measurements were performed on an electrochemical station (CHI760, CH Instruments, USA). All the data presented below have been compensated according to the temperature characteristics of the tungsten oxide electrode.

3. Results and discussion

3.1. Mechanism and characteristics of flow sensor based on the tungsten oxide electrode

The tungsten oxide electrode has a bilayer structure consisting of a compact inner layer of WO_3 and a loosely bound outer hydrated layer of $WO_3 \cdot xH_2O$. Lillard et al. [16] has reported a corresponding decrease in the thickness of the surface layer under the hydrodynamic effect, our previous work also demonstrated a dramatic and positive change in the open circuit potential of the tungsten oxide electrode under the hydrodynamic effect [17].

In order to investigate the mechanism of flow sensor, the change of potentials were measured for the flow velocity of 1.6 mm/s in solutions of different pH which contain the point of zero charge (pzc). The experimental result is seen in Fig. 2. When the tungsten oxide electrode was immersed in the solution of pH above the point

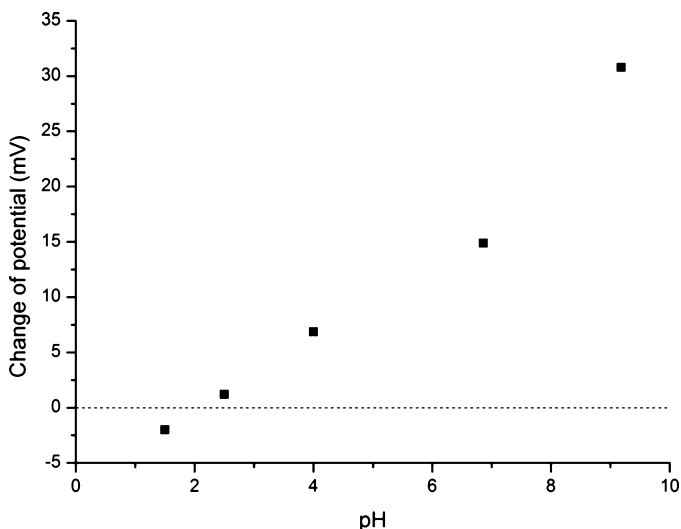


Fig. 2. The change of potentials for the flow velocity of 1.6 mm/s in solutions of different pHs.

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