NON-INVASIVE DRY ELECTRODES FOR EEG

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Abstract - This paper presents a non-invasive dry electrode for electroencephalogram (EEG) acquisition. These electrodes can be integrated in a wearable braincap for wireless EEG that can be used for a long time by patients avoiding the contact gel. This wearable braincap will allow mobility to the patients, while simultaneously having their electrical brain activity monitored. The fabricated dry electrodes are based on iridium oxide (IrO_x) deposited by reactively pulsed sputtering. The electrode employs an array of microtips that penetrate the outer skin layer (i.e. stratum corneum) that is 10 μ m thick. The microtips (100-200 μ m deep) were micromachined through a bulk-micromachining process based on a KOH solution.

Keywords: dry electrodes, IrO_x electrode.

I - Introduction

The electrodes are key elements for acquiring EEG signals. The high demand on non-invasive EEG electrodes for monitoring, diagnostics and treatment of patients with neural diseases, such as epilepsy, is driving the research of electrodes fabricated in different materials and with very low impedance.

The Ag/AgCl standard electrodes present a low impedance in the interface skin-electrode and good stability through its life span. However, these electrodes require a previous preparation in the outer skin (i.e. stratum comeum) and a gel to provide a good contact between electrode and skin to minimize the insulating effect of the stratum comeum layer. The standard Ag/AgCl gel electrodes have limited use because they suffer from dehydration, which leads to a modified electrode impedance and consequently, generates noise and other artifacts in the measured signals. Furthermore, the gel can cause skin irritation and support bacterial growth [1]. A dry Ag/AgCl electrode was proposed for acquiring biopotentials with promising results [2]. Nevertheless, despite the good electrochemical characteristics, the AgCl is toxic and presents an infection risk because it dissolves in contact with the skin [3].

This paper presents a non-invasive dry electrode for EEG. The brain electrical activity occurs between neurons as well as in muscles. The neuroscience field has been demanding invasive electrodes that are implanted in single and multiple recording sites [4]. Electrodes with microtips are presented in Figure 1(b). The microtips length is about 100-200 μ m for providing the penetration of the outer skin layer (i.e. *stratum corneum*) that is 10 μ m thick.



Figure 1: Application of biopotential electrodes: (a) standard EEG electrode; (b) EEG dry electrode with microtips.

II – Design

The design of the dry electrode consists in a pyramidal structure. The penetration of the microtips in the skin requires a specific pressure [5]. The pressure in the microtip, when an axial load is applied, is given by:

$$P = \frac{F}{A} \tag{1}$$

where *P* is the pressure resulted in the solid structure (i.e. microtip), *F*, the perpendicular force applied, and *A*, the section where the force is applied. The force necessary to insert and remove the electrode is about 10 N [5]. The Figure 2 shows the pairs of action-reaction forces, established in the electrode when inserted in skin (the distribution of the force is uniform on the electrode through the 4x4 microtips array - 0.625 N per microtip). The base of the microtip structure is subjected to a pressure of 15.6 MPa. Therefore, every microtip achieves a maximum pressure endured by silicon at 165 μ m from the base (red section in Figure 2). The average height of the pyramidal structures is about 200 μ m.

III – Fabrication

A. Etching

Each microtip in the dry electrode has a three dimensional structure, which is fabricated by a wetetching process in silicon through a liquid solution of potassium hydroxide (KOH). The microtips desired shape is obtained by the undercut effect in the etching, where the planes of fast etching are revealed. It was used a [100]-type silicon wafer with a thickness of $500 \,\mu\text{m}$. Also, two layers of silicon nitride (SiN) on the top and on bottom were used as masks for the etching. The Figure 3 shows the microtips fabrication steps.



Figure 2: Pairs of action-reaction forces on a electrode with an array of 4x4 microtips.



Figure 3: Fabrication of the microtips.

In the Figure 3(a), a mask, with 200 μ m square openings, was used to define the microtips by lithography process. The photoresist was removed in the second step (b) from the uncovered areas. Then, the opened window in the SiN layer allowed the exposition to its elimination in the third step (c) whereas the photoresist is removed afterwards (d). The Si wafer is etched during the step (e). The remaining areas with SiN worked out as a protection layer for the etching process of the Si wafer. The etching solution of KOH was by 30% at a temperature of 87 °C, ensuring an etching rate of 1.6 μ m/min. Finally, the remaining layer of SiN was removed (f).

B. IrO_x Sputtering

The iridium oxide (IrO_x) electrodes were deposited by means of pulsed magnetron sputter using Ar/O₂ plasma prior to which, a Ti adhesion layer (with a thickness of 50 nm) was deposited on the substrate [6]. The IrO_x resistance in a 270 nm thick film was $349 \times 10^{-6} \Omega$ cm. The film thicknesses were determined via liftoff process with a *Tencor Pa-10* profilometer. The thin film resistance was measured in a classic fourpoint probe system.

IV – Experimental Results

A. Electrode Microtips Fabrication

The Figure 4 shows the emerged microtips by optical microscopy every 30 minutes through the undercut effect on the etch process.



Figure 4: Top view of etching progression stages with 30 minutes intervals.

The etching process employed is anisotropic, with an angle of 54.7° between the [111] and the [100] planes. The microtip pyramidal structure due to the anisotropic process is presented in the Figure 5. One can observe the etching of square corners to an hexagonal shape as its final structure. The final microtip was 150-200 µm wider with a height around 100-200 µm.



Figure 5: Microtip structure with a pyramidal shape.

B. Electrode Characterization

The deposition sessions were done in a *Nordiko NS* 2550 sputtering system and pulse generator *ENI RPG-100* at 180 W. The plasma uses argon (Ar) at 100 sccm of flow with a pressure of 10^{-4} Pa. The target of pure iridium (Ir) has a resistivity of 5.3×10^{-8} Ωm.

The Table 1 lists the characteristics of the thin-films made of IrO_x . The resistivity of the thin-films is one of the main characteristic for determining the quality of the electrode, and it was measured via the *van der Pauw* method [7].

Table 1: *The session parameters in the sputtering depositions at frequency of 50 kHz.*

O ₂ flow (sccm)	2	3.5	6.5	10
Deposition rate (nm/min)	13.5	14	33	28
Thickness (nm)	265	349	526	199
Resistivity (Ωm×10 ⁻⁷)	1.926	1.664	2.487	4.914

It would be expected that the increase of O_2 would result in an increase of the films resistivity. Such result would be expected as the increase of O_2 decreases the film purity. However, around 3.5 sccm of O_2 , a lower film resistivity was obtained.

The SEMs in the Figure 6 may induce a justification for the lower resistivity at 3.5 sccm of O_2 flow.



Figure 6: SEM images in cut-view of IrO_x thin-film deposited at different O_2 flows: (a) 2 sccm; (b) 3.5 sccm; (c) 6.5 sccm; (d) 10 sccm.

Apparently, the transition point between stable and unstable thin-film structures happens with an O_2 flow around 3.5 sccm. It appears that in the region of this O_2 flow, the film permittivity increases ensuring lower resistivity than the other samples. Under this flow, the structure is very consolidated which leads to an increased resistivity. Above this flow, the irregular surface induces the highest resistivity obtained. Consequently, 3.5 sccm was the selected flow for coating the electrodes under evaluation.

The O_2 in IrO_x allows the utilization of these electrodes for other application besides recording. As IrO_x has four states of oxidation in reversible chemical reactions, it enables functional stimulation [8].

The Figure 7 shows the variation of current density versus the applied voltage in the IrO_x thin-films when submerged in a NaCl solution with 0.9 % concentration.

These results allow the evaluation of the electrical activity for stimulation of the IrO_x dry electrodes. The results showed good electrochemical properties for 2 and 3.5 sccm O_2 flows.

The charge capacity versus the O_2 flow was plotted in Figure 8. The maximum charge capacity occurs for 3-4 sccm of O_2 flow, for a sputtering frequency of 50 kHz.



Figure 7: Current density vs. applied voltage for four different O_2 flows at 50 kHz sputtering frequency.



Figure 8: Comparison between charge capacity vs. O_2 flow at 50 kHz.

V – Conclusions

This paper proposes a non-invasive IrO_x dry electrode for EEG acquisition fabricated in bulkmicromachining technology. This electrode can avoid the skin preparation with gel due to its microtips structure, which penetrates through the *stratum corneum* layer.

After the fabrication and the characterization of the electrodes, the optimum deposition conditions were obtained for O_2 flows between 2-3.5 sccm and at 50 kHz sputtering frequency. Also, the developed microtips showed a good mechanical behaviour and strength to penetrate the skin.

The iridium oxide had an excellent overall performance not only for biopotential recordings but also functional stimulation application due to charge delivery capacity, low-constant transition impedance and lowresistivity. Thin-films made of IrO_x demonstrated longterm mechanical stability and good corrosion resistance.

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