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**Polymer microchip impedance spectroscopy through two parallel planar embedded microelectrodes: Understanding the impedance contribution of the surrounding polymer on the measurement accuracy**

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## **Abstract**

The present work describes a new methodology for contact free impedance of a solution in a polymer microchip taking into account the role played by the surrounding polymer on the impedance accuracy. Measurements were carried out using a photoablated polyethylene terephthalate (PET) microchannel above two embedded microband electrodes. The impedance diagrams exhibit a loop from high frequencies to medium frequencies (1 MHz to 100 Hz) and a capacitive behaviour at low frequencies (100 Hz to 1 Hz). The impedance diagrams were corrected by eliminating from the global microchip response the contribution of the impedance of the PET layer between the two microband electrodes. This operation enables a clear observation of the impedance in the microchannel solution, including the bulk solution contribution and the interfacial capacitance related to the surface roughness of the photoablated microchannel. Models for the impedance of solutions of varying conductivity showed that the capacitance of the polymer–solution interface can be modeled by a constant phase element (CPE) with an exponent of 0.5. The loop diameter was found to be proportional to the microchannel resistivity, allowing a cell constant around  $4.93 \cdot 10^5 \text{ (m}^{-1}\text{)}$  in contactless microelectrodes configuration.

*Keywords:* Microchip; Contact free impedance; Constant-phase element; Polyethylene terephthalate; Photoablation.

## 1. INTRODUCTION

The development of miniaturization technology is increasing, due to the well-known advantages and the crucial need for micrometer scale measurements with improved detection sensitivity [1-3]. Hence, micro-total analysis systems ( $\mu$ TAS) are usually designed for a well-defined analytical task [4]. A wide range of detection techniques has been employed in microfluidic devices, such as optical detection, laser induced fluorescence, luminescence, electrochemistry, surface plasmon resonance, and mass-spectroscopy [5-7]. However, electrochemical detection (ECD) offers specific advantages for microfluidic systems due to its small size, portability, low cost, high sensitivity, and high selectivity using a proper choice of detection potential and/or electrode material [8]. An additional advantage of ECD is the simplicity of the instrumentation, resulting in low electrical power requirements for in-field use [9-12]. Electrochemical impedance spectroscopy (EIS) is a sensitive analysis method for biological samples [13-16]. Indeed, the possibility to integrate impedance analysis into existing lab-on-a-chip designs offers great potential to obtain new insights and to achieve a profound understanding of biological processes, which are not accessible through conventional analysis. With this method, the bias of the sensing electrode potential can be controlled, even though some unwanted problems such as bubble generation, microelectrode passivation, and/or contamination can occur. To overcome these limitations, use of electrodes that do not contact electrolyte has been investigated, mainly in capillary electrophoresis with electrical detection. It is interesting to stress that non-contact conductivity measurements using external tubular electrodes all around the capillary was introduced in 1998 by Zemmann *et al* [17] due to a problem encountered with electrical coupling between the high voltage applied in the capillary and the conductivity cell detection located at the end-column. However, in the literature, the signal excitation is high (5 - 10 V) due to high

impedance of silica capillary with pasted electrodes. In this case the linearity of the system is not maintained during measurement.

The growth of microfluidic technologies with widely varying substrate requirements has led to an increased interest in polymeric materials due to their good bio-chemical performance and low cost. Recently, Gamby *et al.* [10,13] described microchannels and contact-free microelectrodes using a microfabrication process based on photoablation of polyethylene terephthalate (PET). This process is an attractive alternative to other expensive, laborious, and time-consuming methods for microchannel fabrication such as photolithography approaches. However, when choosing a polymer-based substrate, the properties of the material are critical for both the fabrication process and the successful application of the device. Therefore, we propose in the presented work, a thorough study based on a sensitive non-contact technique in order to understand the physical phenomena taking place within the PET microsystem. Measurements were undertaken of dielectric response to a perturbation of voltage between two microelectrodes embedded in the dielectric polymer. Simulation and experimental results were used throughout the paper to illustrate the effectiveness of such an approach. This approach can be employed to detect any kind of molecule. Further development to identify the type of adsorption that governs the protein-PET system and to a better understanding of the factors that govern this adsorption can be also foreseen.

## **2. MATERIALS AND METHODS**

The microchip device used in this work was described elsewhere [10,13]. Some modifications have been made and will be outlined hereafter. The set-up is displayed in Fig. 1. Impedance measurements were carried out through a photoablated polyethylene terephthalate (PET) microchannel having a trapezoidal cross-section shape with a depth of 45  $\mu\text{m}$ , a top width of 100  $\mu\text{m}$  and a length of 1.4 cm. The separation distance between both microchannels was 200  $\mu\text{m}$

center to center and 120  $\mu\text{m}$  edge to edge. The electrode fabrication was achieved using a carbon ink loaded with gold nanoparticles, thermally laminated at 135°C and a pressure of 2 bar by a polyethylene (10  $\mu\text{m}$ )/polyethylene terephthalate (25  $\mu\text{m}$ ) (PE/PET) layer. The total thickness of the laminated layer is equal to 35  $\mu\text{m}$ . The separation distance in the PET band between the two planar microelectrodes and the main microchannel was equal to 5  $\mu\text{m}$  and the detection surface area per microelectrode was 80  $\mu\text{m}$  x 100  $\mu\text{m}$ .

Electrochemical impedance spectroscopy measurements were performed by applying an ac voltage with constant amplitude (0.1 V) through the microelectrodes with a frequency ranging from 1 MHz to 1 Hz. The current measured is thus related to the total impedance of the sensor. These measurements were performed using a frequency response analyzer (FRA 1255B, Solartron U.K.) together with a dielectric interface 1296 (Solartron, U.K) which extends the frequency range from 1 MHz to 10 Hz. Data analysis was performed using the commercial software SMaRT. The effect of NaCl solutions on the resistance in the microchannel was studied at various concentrations (0 -  $10^{-2}$  M). To analyze these results, a physical model was used as is described in Fig. 2. The data obtained were treated using a home-made software SIMAD which allowed fitting both frequency-dependent analytical expressions and equivalent electrical circuits.

### **3. RESULTS AND DISCUSSION**

The results of impedance measurements between the two band microelectrodes depend on the surface state of the flow channel and on adsorption taking place on the walls of the flow channel. However, in the previous study [10], the impedance between the microelectrodes and the microchannel through the bulk of the PET substrate was not considered since it has been reported that the resistance is large enough that it could be disregarded. Thus, impedance measurements performed here are undertaken on the device when it is empty or filled with a sodium chloride

solution in order to reach a better understanding of the PET impedance contribution in the global response.

### 3.1 EIS measurements in an empty microchannel

Polyethylene terephthalate is a dielectric material [18]. Its capacitive behavior between the two microelectrodes can be evaluated using

$$C_{2,PET} = \frac{\varepsilon_0 \varepsilon_r S_2}{d_2} \quad (1)$$

where  $C_{2,PET}$  is the geometric capacity in the PET thickness,  $\varepsilon_0$  is the vacuum permittivity ( $8.854 \cdot 10^{-12}$  F m<sup>-1</sup>),  $\varepsilon_r$  is the relative permittivity of the PET film (3.4),  $d_2$  is the average distance separation between the two electrode edge to edge (120  $\mu$ m) and  $S_2$  is the vertical surface of the microelectrodes ( $2 \cdot 10^{-7}$  m<sup>2</sup>). This derivation gives a value of  $4.5 \cdot 10^{-14}$  (F). Equation 1 does not take into account the horizontal part of the two microelectrodes shown in figure 1.

Impedance measurements in the case of empty microchannel in the frequency range from 1 MHz to 1 Hz and with 0.1 V are presented in figure 3. The Nyquist diagram shows a capacitive behavior over the entire frequency range, but the representative line is not exactly vertical. It is interesting to stress that the conventional RC circuit is not convenient for modelling impedance response through the dielectric PET layer. The impedance  $Z_2(\omega)$  between the two microelectrodes for the empty microchannel has a capacitance behavior represented by constant phase element (CPE), defined as

$$Z_{2,PET}(\omega) = \frac{1}{(j\omega)^{\alpha_{2,PET}} Q_{2,PET}} \quad (2)$$

where  $Q_{2,PET}$  is the CPE constant ( $\Omega^{-1} s^\alpha$ ) and  $\alpha_{2,PET}$  is the CPE exponent.

As described by Orazem *et al.* [19], the log-log representation of the imaginary part of impedance vs frequency is suitable for a CPE analysis. The  $\alpha_{2,PET}$  value corresponding to the slope of the curve plotted in fig 3d is 0.99. The extracted value of the CPE element ( $Q_{2,PET}$ ) in the PET thickness of 120  $\mu\text{m}$  is estimated to  $8.5 \cdot 10^{-13} (\Omega^{-1} \text{s}^\alpha)$ .

The slight difference in value between the CPE element  $Q_{2,PET}$  and  $C_{2,PET}$  is due to the capacitive effect of the horizontal surface of the microelectrodes as mentioned above. The average value of 0.85 pF value could be considered to be the overall background (or stray) capacitance.

In the following, the impedance  $Z_{2,PET}$  will be determined by using equation 2 with  $\alpha_{2,PET} = 0.99$  and  $Q_{2,PET} = 8.5 \cdot 10^{-13} \Omega^{-1} \text{s}^\alpha$ .

From the knowledge for an empty microchannel  $Z_2(\omega)$ , the impedance in the microchannel ( $Z_1(\omega)$  in Fig. 2) can be deduced from the global impedance measurement  $Z_G(\omega)$ . Then in the following work, all results will be represented by the impedance  $Z_1(\omega)$ .

### 3.2 EIS measurements in microchannel filled with NaCl solution:

The global impedance  $Z_G(\omega)$  between the two microelectrodes, in the case of the microchannel being filled with a solution, depends on the different parts which constitute the microchip. In this configuration,  $Z_G(\omega)$  can be expressed as the combination between  $Z_1(\omega)$  and  $Z_2(\omega)$  (Eq. 3):

$$\frac{1}{Z_G(\omega)} = \frac{1}{Z_1(\omega)} + \frac{1}{Z_2(\omega)} \quad (3)$$

where  $Z_1(\omega)$  corresponds to a combination of impedances in the microchannel with the microchannel interface/PET and the impedance of the 5- $\mu\text{m}$  thick. The impedance  $Z_1(\omega)$  is equal to

$$Z_1(\omega) = 2Z_{1,PET}(\omega) + \frac{R_s}{1 + j\omega C_s R_s} + 2Z_{\text{int}}(\omega) \quad (4)$$



In the microchannel:  $R_S$  with units of  $\Omega$  denotes the solution resistance in the microchannel which depends on the solution conductivity and on the geometry of microchannel, and  $C_S$  in F represents the solution capacitance, which depends on its relative permittivity,  $\epsilon_{r,solution}$ , usually equal to water permittivity in the case of a dilute solutions ( $\epsilon_{r,water} = 78$ ). In a typical measuring cell ( $d = 1$  cm), the distance between the two electrodes is large enough to neglect the capacitance of the solution ( $C_S$ ). However, this is not the case in the present microchip configuration, where the distance between the microelectrodes is only 120  $\mu\text{m}$ .

In the PET layer of 5  $\mu\text{m}$ : the geometric capacitance calculated by equation 5 for the PET layer of 5  $\mu\text{m}$  ( $C_{1,PET}$ ) is equal to  $1.4 \cdot 10^{-14}$  F. This capacitance is depends on the PET permittivity, PET thickness ( $d_1 = 5 \cdot 10^{-6}$  m), and surface contact area ( $S_1 = 8 \cdot 10^{-9}$  m<sup>2</sup>), (see in Fig. 1):

$$C_{1,PET} = \frac{\epsilon_0 \epsilon_r S_1}{d_1} \quad (5)$$

Taking into account the results obtained in the case of the PET thickness of 120  $\mu\text{m}$ , the impedance for the PET thickness of 5  $\mu\text{m}$  is given by

$$Z_{1,PET}(\omega) = \frac{1}{(j\omega)^{\alpha_{1,PET}} Q_{1,PET}} \quad (6)$$

where  $Q_{1,PET}$  in  $\Omega^{-1} \text{s}^\alpha$  is the modeled capacitive behavior of the PET thickness of 5  $\mu\text{m}$  and  $\alpha_{1,PET}$  is the CPE exponent.

At the PET/ microchannel interface:

The impedance in the PET/ microchannel interface has a CPE behavior given by

$$Z_{\text{int}}(\omega) = \frac{1}{(j\omega)^{\alpha_{\text{int}}} Q_{\text{int}}} \quad (7)$$

where  $Q_{\text{int}}$  is the CPE constant at the interface and  $\alpha_{\text{int}}$  is the CPE exponent.

Experimentally,  $Z_1(\omega)$  is obtained from equation 3 where  $Z_2(\omega)$  is the experimental values of the impedance for an empty microchannel and  $Z_G(\omega)$  is the experimental values of the impedance for a filled microchannel with sodium chloride solution.

The real and imaginary parts of  $Z_1(\omega)$  are calculated from

$$Z'_1 = \frac{[(Z'_2 Z'_G - Z''_G Z''_2)(Z'_2 - Z'_G) + (Z''_2 - Z''_G)(Z''_2 Z'_G + Z''_G Z'_2)]}{(Z'_2 - Z'_G)^2 + (Z''_2 - Z''_G)^2} \quad (8)$$

and

$$Z''_1 = \frac{[(Z''_2 Z'_G + Z''_G Z'_2)(Z'_2 - Z'_G) - (Z''_2 - Z''_G)(Z'_2 Z'_G - Z''_G Z''_2)]}{(Z'_2 - Z'_G)^2 + (Z''_2 - Z''_G)^2} \quad (9)$$

where  $Z'_G$ ,  $Z''_G$ ,  $Z'_1$ ,  $Z''_1$ ,  $Z'_2$  and  $Z''_2$  denote the real and imaginary parts of the impedances  $Z_G(\omega)$ ,  $Z_1(\omega)$  and  $Z_2(\omega)$ , respectively.

Electrochemical impedance spectroscopy measurements with a voltage of 0.1 V and frequency ranging from 1 MHz to 1 Hz were performed for microchannels filled with water and various concentrations of NaCl ( $5 \cdot 10^{-4}$  M to  $10^{-2}$  M). The goal was to extract the values of the solution resistance in the microchannel included in the  $Z_1(\omega)$  expression by varying the solution conductivity in the microchannel.

The corrected impedance diagrams plotted by eliminating the PET layer impedance (of 120  $\mu\text{m}$ ) contribution from the global microdevice response are shown in Fig. 4. Nyquist plots exhibit a loop from very high frequencies to medium frequencies (1 MHz to 100 Hz) and a capacitive behavior at low frequencies (see Figs. 4a and b). One can observe that the diameter of the loop is inversely related to the microchannel conductivity.

The highest resistance corresponding to the largest diameter loop obtained was recorded in the case of water (water conductivity measured:  $3 \mu\text{S cm}^{-1}$ ); whereas, the lowest value was obtained

for the  $10^{-2}$  M NaCl solution (conductivity measured:  $1194 \mu\text{S cm}^{-1}$ ). This effect is also clearly shown in Fig. 4c. For a frequency range from 10 kHz to 1 Hz, the real part of the measured impedance decreased linearly with the microchannel conductivity.

The evolution of the log-log plot representation of the imaginary part (Fig. 4c) shows that, in the high frequency range from 1 MHz to 50 kHz (filled with  $10^{-2}$  M NaCl solution) and from 1 MHz to 1 kHz (filled with water), all slopes are estimated to 0.99, which is close to a pure capacitive behavior. Moreover, the cut-off frequency varies and depends on solution conductivity.

It is important to stress that, at the intermediate frequency (5 kHz to 300 Hz for NaCl  $10^{-2}$  M and 1 kHz to 300 Hz for water), the slope of imaginary part was found equal to 0.5. This behavior is characteristic of diffusion impedance or of the impedance of a rough interface.

The diffusion impedance analysis could be complicated in case of anomalous behavior [20-23]. But in the present case, the hypothesis of diffusion impedance could be disregarded due to the fact that the microelectrodes were not in contact with electrolyte and the current was not faradaic. It is important to notice that diffusion impedance is a part of faradaic impedance. Only resistivity, permittivity and species adsorption may be considered for the present system, and, as a consequence, only the interfacial impedance of rough surface was used in the impedance analysis. As a proof of that, scanning electron microscopy (SEM) micrographs of the photoablated PET surface presented in Fig. 5 highlight a microstructured surface constituted by the well-known microhills and microvalleys region on the microchannel surface due to photoablation of crystalline and amorphous regions, respectively [24-26]. This observation supports the conclusion that the interfacial impedance including the photoablated PET microchannel should not be represented as a pure capacitance. To take into account this characteristic, the De Levie model [27] for rough surface was used and adapted by different

authors [28-30] in order to determine impedance in rough or porous surfaces. In our case, this model could be also adapted for a microstructured surface.

In order to study the influence of the CPE exponent ( $\alpha$  parameter) and to validate the hypothesis that the De Levie model applies in the present case, the  $Z_1(\omega)$  impedance model was calculated for different values of the  $\alpha_{CPE}$  parameter. A comparison between the experimental data and  $Z_1(\omega)$  impedance model is presented in Figure 6 with the assumed value of  $\alpha_{CPE}$  as a parameter.

The value of  $\alpha_{CPE}$  has a strong influence on the impedance in a frequency range that extends from 1 MHz to 1 Hz for the imaginary and real parts. The best correspondence between model and experiment was found with  $\alpha_{CPE} = 0.5$ .

As demonstrated by Keiser *et al* [31], for any pore shape, the impedance of a rough surface at high frequencies approaches  $45^\circ$  in the Nyquist plot (corresponding to a slope of 0.5 in the log-log representation of the imaginary part of the impedance as a function of frequency). The geometry of the microstructured surface is close to the shape numbered 5 in reference [31] and can be seen to be a good approximation for a photoablated microchannel. If the impedance corresponds to the high frequency model in the entire frequency range, its representation is a CPE as in equation 7 with  $\alpha_{CPE} = 0.5$ . This indicates that the CPE element for a photoablated surface microchannel is proportional to the square root of the capacitance of a non photoablated one. Then, we shall have to determine the value of  $Q_{int}$  for NaCl at various concentrations.

In the Nyquist diagram, we observed at low frequencies a capacitive branch due to the increase of the imaginary part of the impedance for frequencies lower than 100 Hz. This behavior is influenced by the electrical conductivity of the microchannel solution as displayed in Fig. 4. Referring to the proposed electrical circuit, this behavior defines the interface PET/microchannel. In the knowledge that the PET surface has functional groups that can form electrostatic interactions with ionic species, it is obvious that high concentrations of NaCl

increase the capacitive effect recorded at low frequencies. The use of this impedance model in Eq. 4 provides an excellent fit to the EIS spectra shown in Fig. 4.

A Set of fitted parameters is reported in table 1. The model provided a very good fit to the experimental data, as was confirmed by different graphical representations of the model and data values. The values of the CPE element ( $Q_{1,PET}$ ) obtained for water and for a NaCl solution displayed in table 1 show a slight difference that can be related to the microchannel composition. Moreover, the fitted value of  $Q_{1,PET}$  is much higher than the geometric capacitance  $C_{1,PET}$ . Conversely to the slight difference observed between  $Q_{2,PET}$  and  $C_{2,PET}$ , in the case of CPE element  $Q_{1,PET}$ , the difference arose from the value of the surface contact area  $S_I$  which was underestimated, particularly on the side of the microchannel. The estimated value of  $C_{1,PET}$  could be seen as a smaller value due to the fact that only the vertical streamlines from the  $S_I$  to microchannel section displayed in fig.1 were considered.

The  $R_S$  values obtained with the fitting procedure show that there is a relationship between resistivity and solution resistance, given as

$$K = \frac{R_S}{\rho} \quad (10)$$

where  $K$  in  $m^{-1}$  is the cell constant microchannel.

A plot of the solution resistivity,  $\rho$ , against the length of loop diameter,  $R_S$ , gives rise to straight lines with a slope that is comparable to a cell constant,  $\kappa$ . The cell constant was found to be equal to  $4.93 \cdot 10^5 m^{-1}$  (see Fig. 7). This latter observation supports the model used in this paper.

#### 4. Conclusion

In this study, we have demonstrated that contact-free impedance measurements can be successfully implemented in an insulated polymer for flexible microchips. The impedance technique has the advantage that it can carry out *in situ* characterization to obtain several physical parameters. Moreover, a new approach involving a new data processing for the analysis of the contact-free impedance through two parallel planar microelectrodes is demonstrated. It is pointed out that the resistivity through the PET layer cannot be neglected and must therefore be taken into account in the global microchip response. Indeed, the photoablated PET surface should be considered to be a microstructured surface, characterized by a constant phase element with a typical exponent of 0.5. The impedance model proposed here is inspired by the classical equation established by De Levie which leads to accurate and efficient fitting of our experimental data. This study allowed a superior understanding of the global response of the PET microchip impedance. These steps are crucial in order to choose and optimize the experimental conditions for biosensor development in any kind of flexible polymer microchips [32].

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**Table 1:**

[NaCl] (mol dm <sup>-3</sup> )	$\rho$ ( $\Omega$ cm)	$Q_{1,PET}$ ( $\Omega^{-1} s^a$ )	$\alpha_{1,PET}$	$R_s$ ( $\Omega$ )	$C_s$ (F)	$Q_{int}$ ( $\Omega^{-1} s^a$ )	$\alpha_{int}$	$\chi$ (%)
0 (water)	$3.2 \cdot 10^5$	$25 \cdot 10^{-11}$	0.99	$4.5 \cdot 10^8$	$3 \cdot 10^{-13}$	$0.6 \cdot 10^{-9}$	0.5	4.8
$5 \cdot 10^{-4}$	$10^4$	$14 \cdot 10^{-11}$		$5 \cdot 10^7$	$4 \cdot 10^{-13}$	$0.9 \cdot 10^{-9}$		3.4
$10^{-3}$	$5 \cdot 10^3$			$2.5 \cdot 10^7$		$1 \cdot 10^{-9}$		3.1
$5 \cdot 10^{-3}$	$1.8 \cdot 10^3$			$9 \cdot 10^6$		$1.8 \cdot 10^{-9}$		2.8
$10^{-2}$	840			$5 \cdot 10^6$		$2.4 \cdot 10^{-9}$		2.6

**Table 1:** Fitted parameters values according to the equivalent circuit described in Fig. 2 and results of Fig. 4. The values of the resulting  $\chi^2$  statistic were obtained with errors of fitted values,  $\sigma = 0.01$ , for frequency ranging from 1 MHz to 1 Hz.

Figure

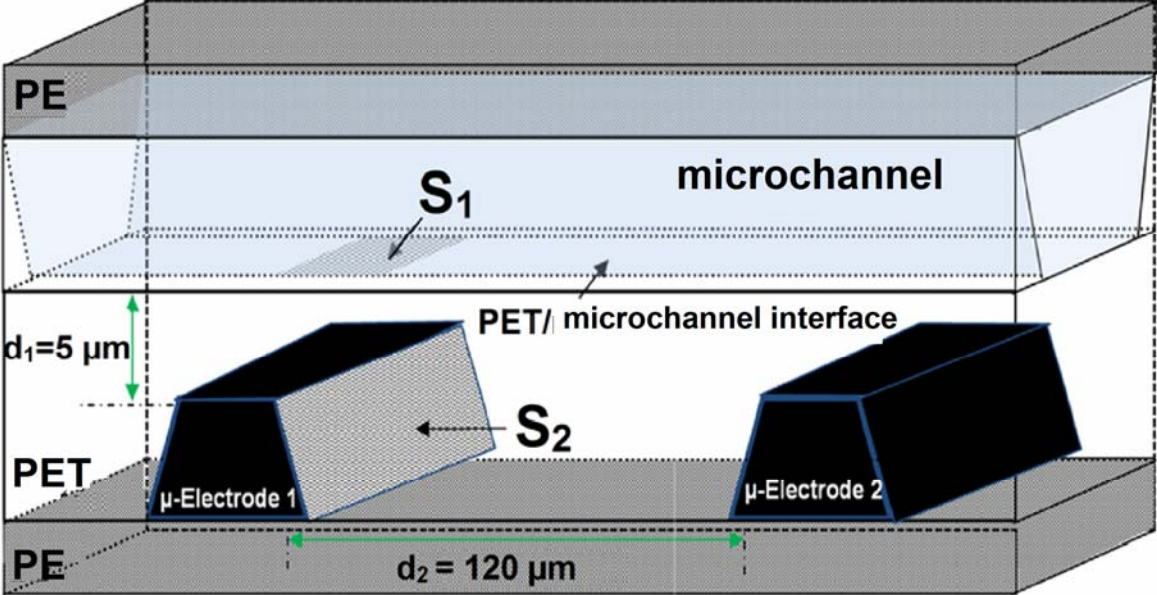
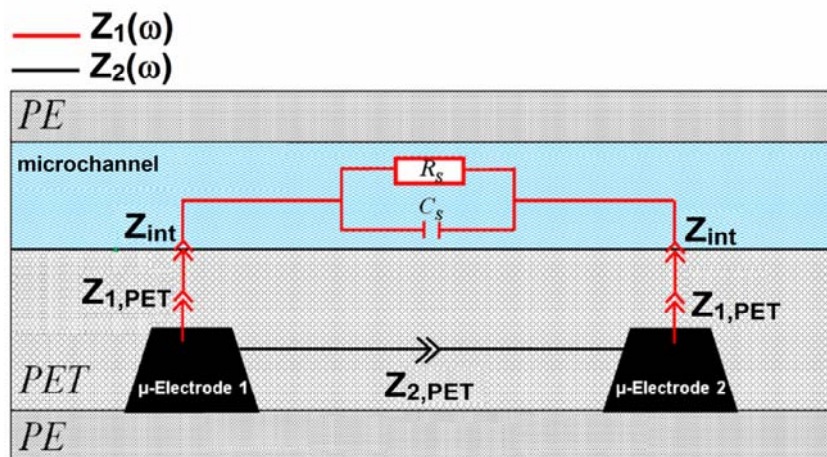
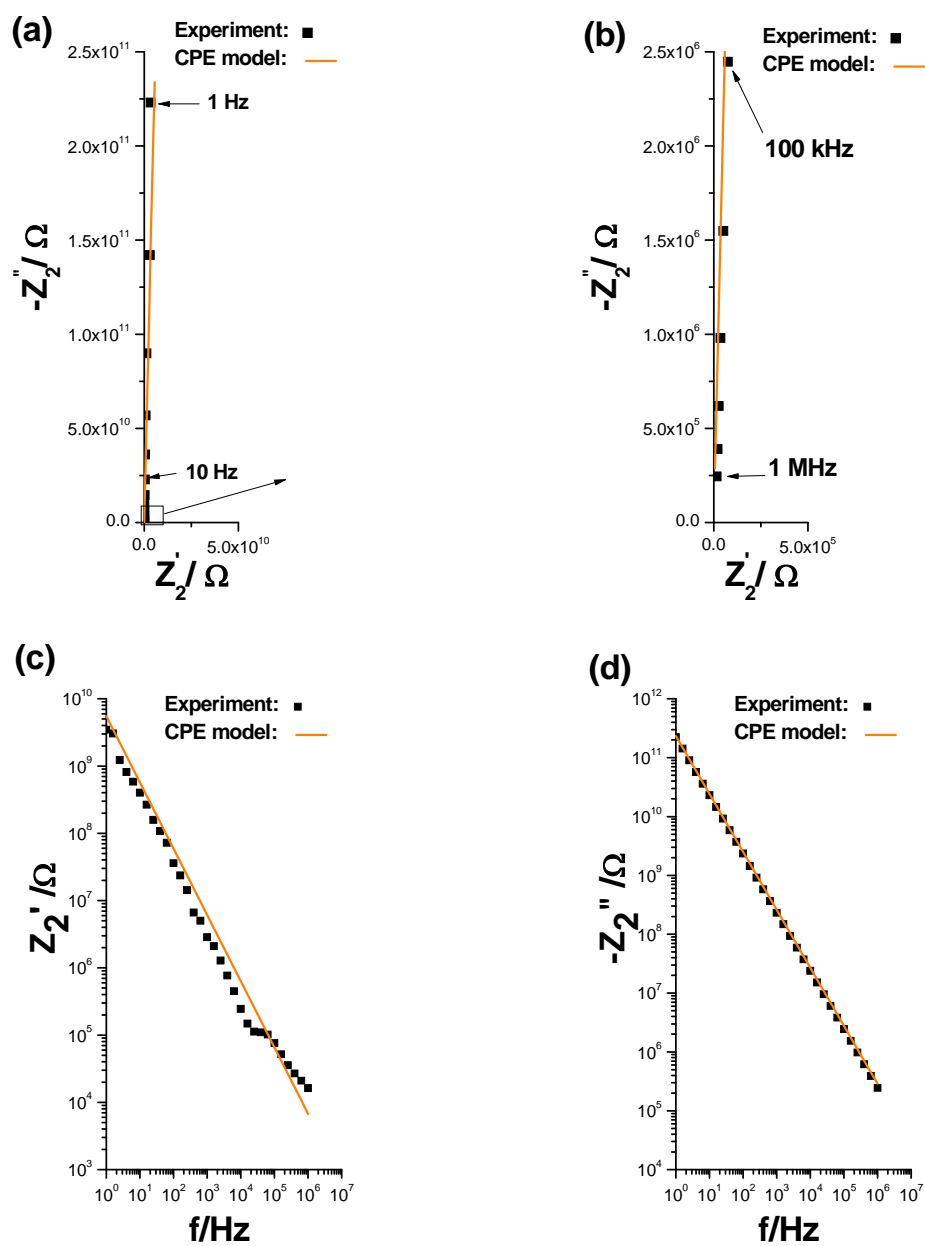


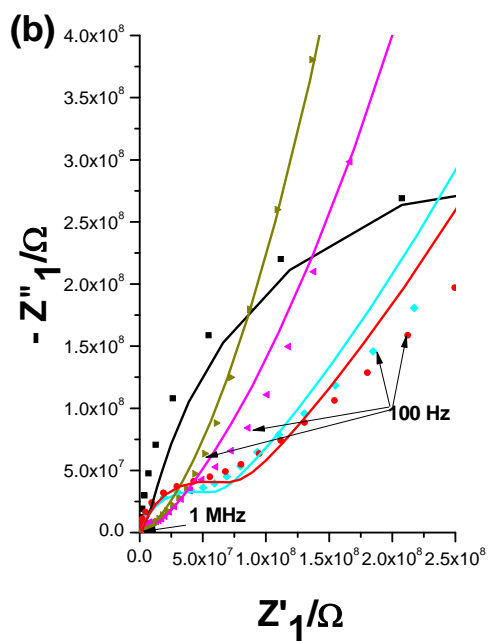
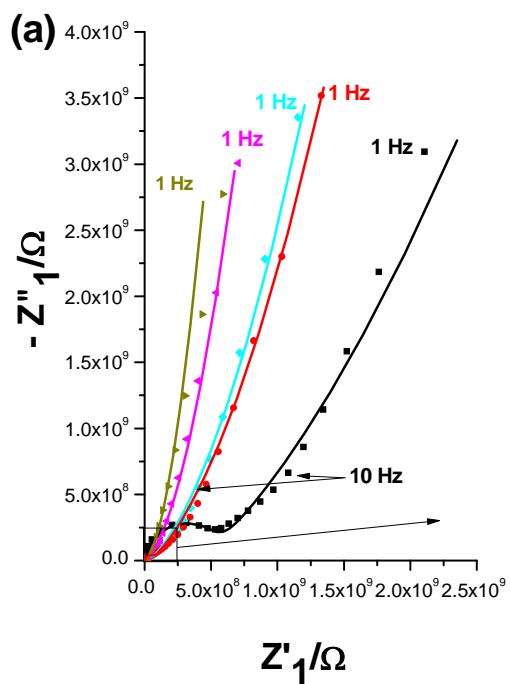
Fig. 1: Schematic representation of PET-microchannel-microelectrodes interface.

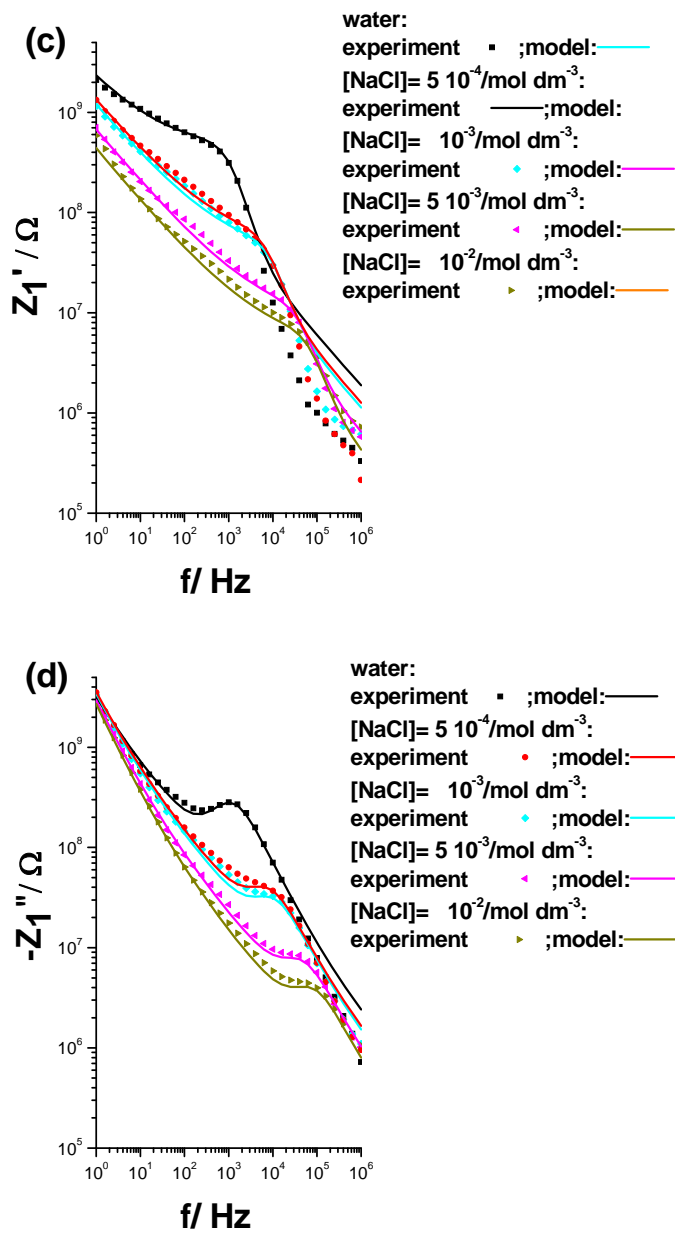


**Fig. 2:** Representation of the microchip by an equivalent electrical circuit.



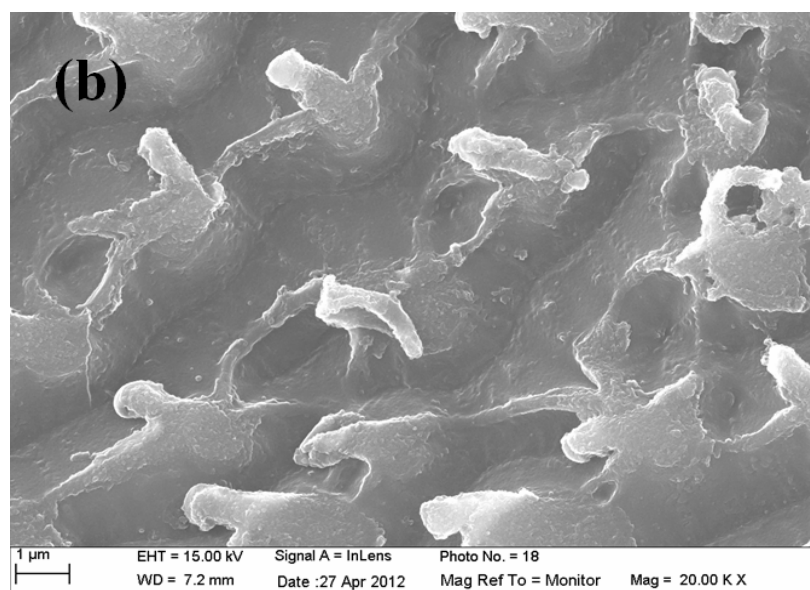
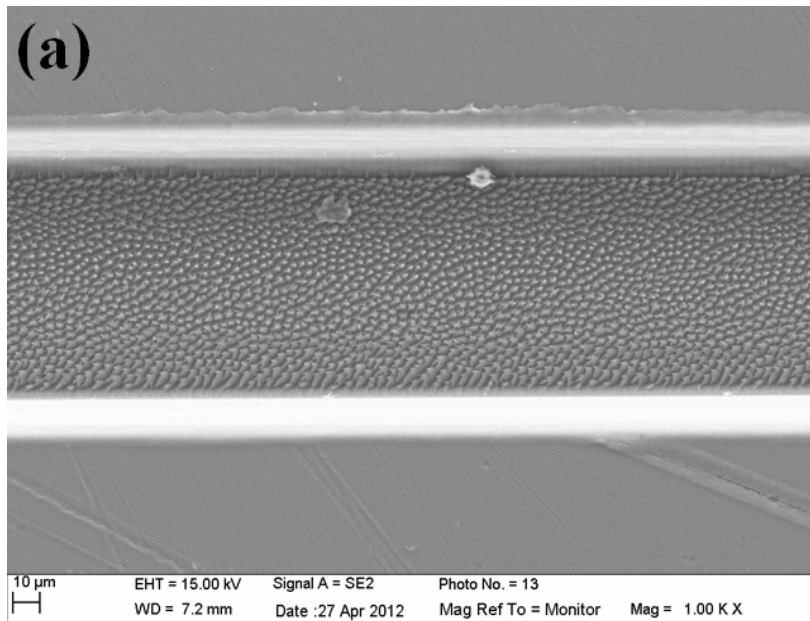
**Fig. 3:** Impedance diagrams; (a) Nyquist plots, (b) Nyquist plots magnification, (c) Real part vs frequency, (d) Imaginary part vs frequency; measured between 1 MHz to 1 Hz and 0.1 V in the case of an empty microchannel.



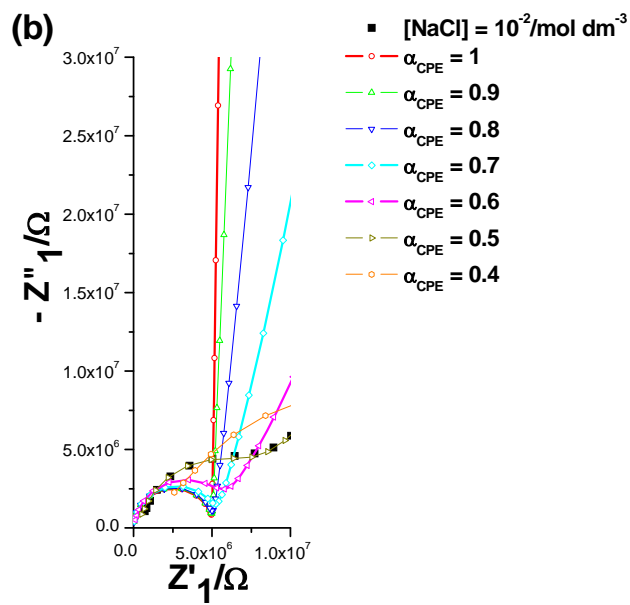
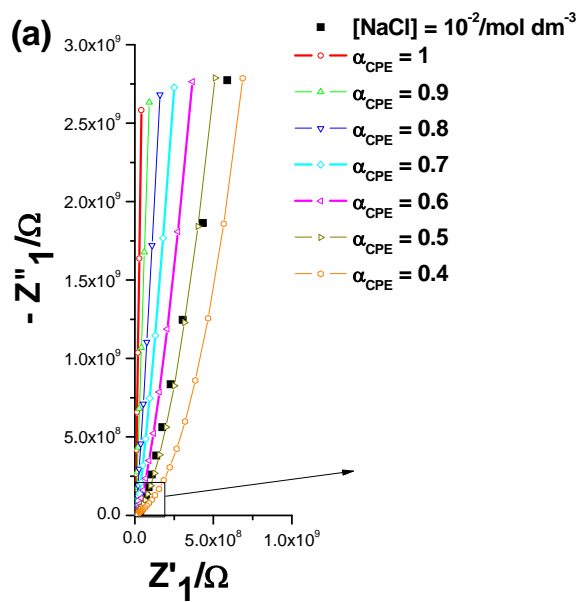


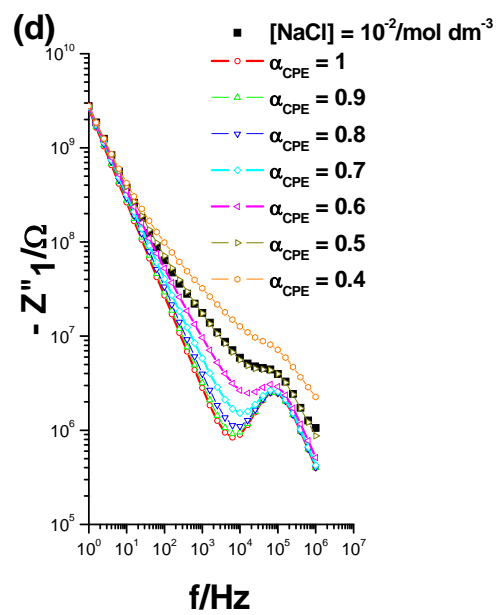
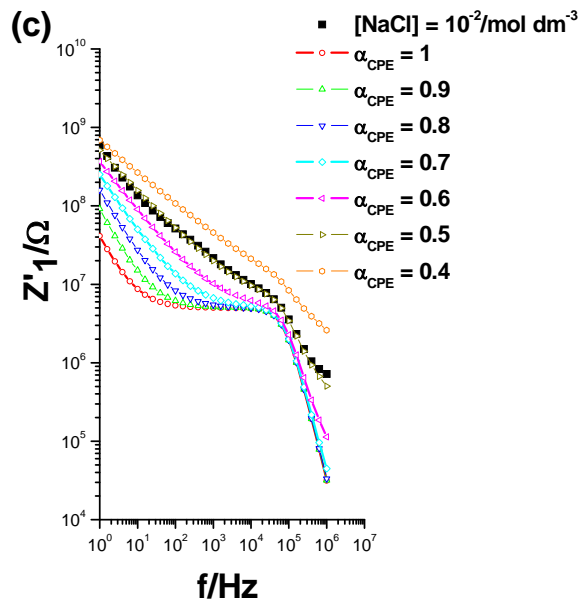
**Fig. 4:** Experimental impedance diagrams and modeling diagrams  $Z_1(\omega)$  in the microchip measured for an empty microchannel, in the presence of water in the microchannel and in the presence of NaCl solution at various concentrations, between 1 MHz to 1 Hz at 0.1V. (a) Nyquist plots. (b) Nyquist plots magnification. (c) Real part vs. frequency. (d) Imaginary part vs frequency.



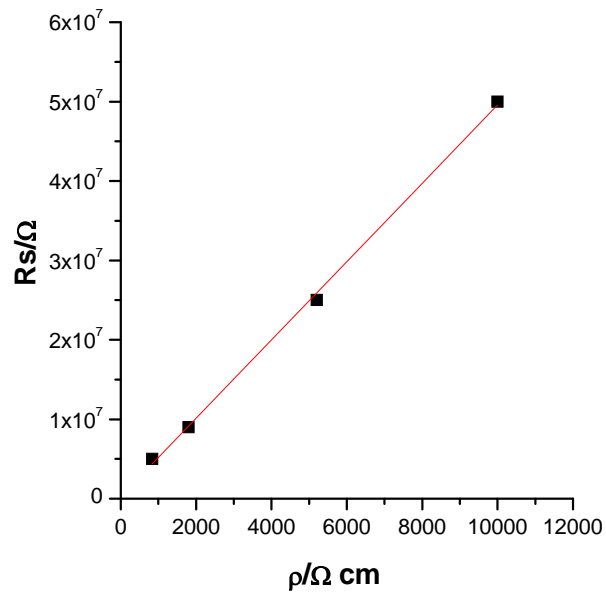


**Fig. 5:** (a) SEM micrograph of the photoablated PET microchannel. (b) Microstructured aspect of the bottom microchannel.





**Fig. 6:** Experimental impedance diagrams and the CPE model at the microchip interface, measured in the case of microchannel filled with NaCl at  $10^{-2}$  M. Between 1 MHz to 1 Hz and 0.1 V: (a) and (b) Nyquist plots; (c) real part; (d) imaginary part.



**Fig. 7:** Plot of loop diameter in Nyquist plot versus solution resistivity