

A microfluidic device with removable packaging for the real time visualisation of intracellular effects of nanosecond electrical pulses on adherent cells

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The biological mechanisms induced by the application of nanosecond pulsed electric fields (nsPEFs: high electrical field amplitude during very short duration) on cells remain partly misunderstood. In this context, there is an increasing need for tools that allow the delivering of such pulses with the possibility to monitor their effects in real-time. Thanks to miniaturization and technology capabilities, microtechnologies offer great potential to address this issue. We report here the design and fabrication of a microfluidic device optimized for the delivery of ultra short (10 ns) and intense (up to 280 kV cm⁻¹) electrical pulses on adherent cells, and the real time monitoring of their intracellular effects. Ultra short electric field pulses (nsPEFs or nanopulses) affect both the cell membrane and the intracellular organelles of the cells. In particular, intracellular release of calcium from the endoplasmic reticulum was detected in real time using the device, after exposure of adherent cells to these nsPEFs. The high intensity and spatial homogeneity of the electric field could be achieved in the device thanks to the miniaturization and the use of thick (25 μm) electroplated electrodes, disposed on a quartz substrate whose transparency allowed real time monitoring of the nsPEFs effects. The proposed biochip is compatible with cell culture glass slides that can be placed on the chip after separate culture of several days prior to exposure. This device allows the easy exposure of almost any kind of attached cells and the monitoring in real time while exposed to nsPEFs, opening large possibilities for potential use of the developed biochips.

I. Introduction

Cell electroporation is a well-known technique that requires a short duration and an intense electric field to destabilize the plasma membrane. This technique is now mature and used in the field of cancer therapies in order to enhance the efficiency of targeted delivery of materials like drugs, antibodies or genes.^{1–3}

Electroporation results from an interaction between the applied electric field and the cell plasma membrane, which increases its permeability. For a circulating cell (assumed to be spherical) the transmembrane potential Ψ_m is time dependent, following the charging of the plasma membrane, eqn (1).^{4,5}

$$\Psi_m = f \cdot E \cdot R \cdot \cos(\theta) \cdot [1 - e^{-\frac{t}{\tau_m}}] \quad (1)$$

where R represents the cell radius, f is function of the cell geometry, and θ stands for the angle between the electric field E

and the cell radius vector. The time constant τ_m , characterizes the dynamics of the charge of the membrane.⁵ The case of adherent cells is more complex, and some simulation study on cell tissues can be found in ref. 6.

In the case where conventional electroporation pulses are applied (such pulses having a few 100 μs duration), the transmembrane potential reaches $\Psi_m = f \cdot E \cdot R \cos(\theta)$, once $t \gg \tau_m \approx$ order of μs. In this case, the cell membrane shields the electrical field that does not penetrate inside the cell, as long as the membrane is not permeabilised.^{7,8}

With the recent technical progresses in the field of pulsed power equipment, the use of ultra short duration pulses (typically 3 to 10 ns) and high intensity (up to 100 kV cm⁻¹), also called nanosecond pulsed electric fields (nsPEF), has been investigated. The nsPEFs, implying a high frequency electric field spectrum (from DC and above 100 MHz), are able to penetrate inside the cells as the plasma membrane is short-circuited at these working frequencies ($t \ll \tau_m$). Hence, nsPEFs are able to disturb the membrane of intracellular organelles as internal vesicles, mitochondria⁹ or endoplasmic reticulum.^{10,11} Nevertheless, we are at the beginning of this very exciting area of manipulating cell interior whose mechanisms remain misunderstood.

In this context, researchers need tools allowing the real time visualization of cell perturbations induced by the application of

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nsPEFs. In particular, the microtechnologies offer high miniaturization capabilities allowing the generation of highly intense nanosecond electric fields, suitable to penetrate the intracytoplasmic compartment. Moreover, the use of transparent material to microfabricate such miniaturized devices allows real time imaging of the effects of such nanopulses.

In particular, the nanoporation of adherent cells requires a particular attention, as these cells need to be cultured and attached on a substrate to be in proper conditions mimicking usual cell environment, prior to nPEFs exposure. While many studies exist in the literature regarding the electroporation of circulating cells on a chip,^{12–16} including our work,¹⁷ very few developments of micromachined devices adapted to the electrical treatment of adherent cells can be found. Hence, we propose in this paper an original approach where adherent cells cultures, prepared *ex situ* on conventional glass slides, can be reported on our microfluidic device, thanks to a specific design. With this approach, one given microdevice can be re-used and easily cleaned for many experiments, where cells are cultured on glass slides prepared separately in large numbers. This concept of removable packaging is here advantageously exploited for adherent cells exposure to nsPEFs and for the real time monitoring of the effects of the nsPEFs, focusing on the generation of internal calcium concentration peaks resulting of membranes nanoporation.

II. Microfluidic biochip design

In this section we describe the design of the new microfluidic device (Fig. 1(a)) that was specifically developed for the observation and study of the effects of nsPEFs on adherent cells.

A. Impedance matching aspects

Due to the high voltages (up to 10 kV) applied to obtain the high field amplitude of the nsPEFs, the impedance matching with the

nanopulser is crucial in order to prevent any damages of the electronic components. Indeed, the level of reflections of the transmitted energy back to the nanopulser depends on this matching, which in addition directly affects the transfer of energy to the cells.^{18–20}

As the electric field is applied to the biological medium with thick electroplated gold electrodes, the impedance is mainly governed by the physical properties of this medium. The propagation along the electrodes can be neglected for the considered frequency spectrum of the nsPEFs which energy is mainly located in the range (0–100 MHz), when 10 ns duration pulses are considered.^{18–20} The medium is mainly conductive at low frequencies (for $f < \frac{\sigma}{2\pi\epsilon}$) and capacitive for higher frequencies, where σ and ϵ are the conductivity and permittivity of the medium, respectively.

The impedance matching cannot be achieved for all frequencies but can be optimized in a large frequency bandwidth, where the medium is considered as a pure conductance, and which includes the main energy frequency range of the nanopulses. To do so, the geometrical parameters of the device (thickness t of the electroplated electrodes, the distance d between these electrodes, the length L and height H of the micro channel) are adjusted to get the required 50 Ω impedance for the 0 to 200 MHz frequency bandwidth, suitable for the delivery of 10 ns nanopulses with the available nanopulser generator described in the experimental section.^{17–21} While d and t are set to $d = 150\ \mu\text{m}$ and $t = 25\ \mu\text{m}$, respectively, L and H depend on the chosen architecture (see next section).

B. Removable packaging architecture

For adherent cells to be exposed to the nsPEFs solicitations, it is mandatory to let the cells incubate and adhere on the substrate prior to field exposure. At least 4 h incubation in sterile conditions, at 37 °C–5% CO₂, is required. A cell culture directly

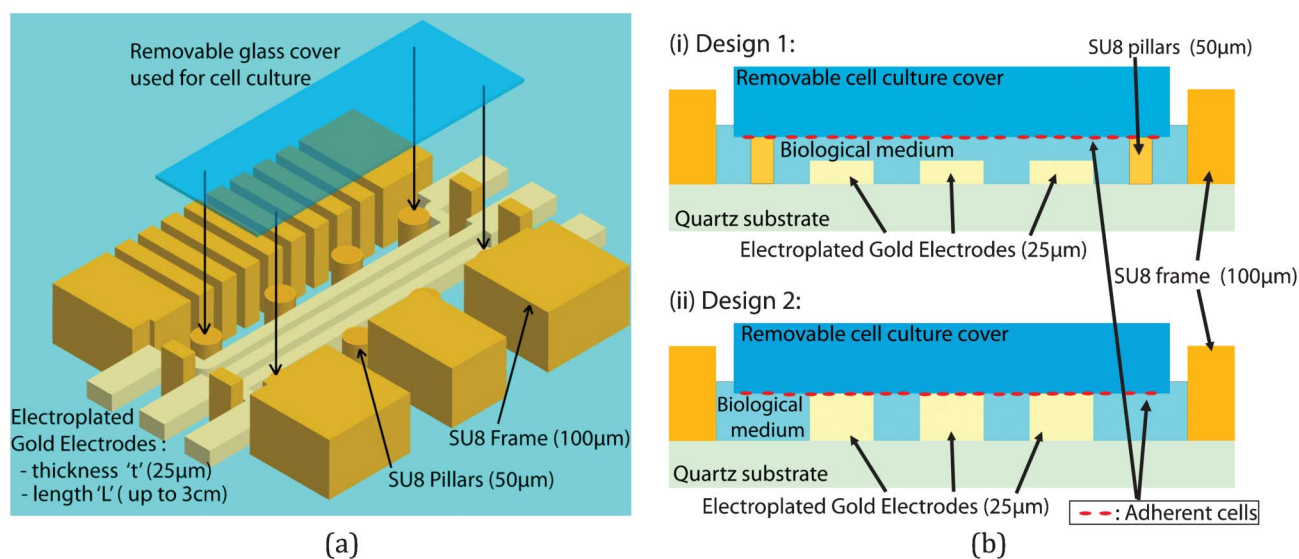


Fig. 1 (a) Schematic representation of the device, involving electroplated gold electrodes in a transmission line configuration, and a thick SU8 resist frame for the centering of the removable glass cover. (b) The two designs of the microfluidic device with removable packaging for cells exposure to nsPEFs: (i) Design with 25 μm thick electrodes, and 50 μm thick resist micro-pillars to support the culture slide (ii) Design with 25 μm thick electrodes and no pillars (arbitrary scales).

performed within the microfluidic device is quite achievable but leads to several constraints: i) only one or two exposures per day are possible, as 4 h are required for the adhesion and spreading of the cells; ii) a specific cleaning of the device is necessary after each experiment. The lifetime of the device is thus decreased, due to this cleaning procedure required.

In order to avoid the drawbacks mentioned above, we developed a specific packaging, well adapted to our case where adherent cells are exposed to nsPEFs within the microfluidic device. The solution that we propose involves the possibility of using a removable coverslip. Indeed, this strategy presents the advantage of allowing the culture on several cover slides in parallel while the biochip is available for short term (or even very short term) experiments. The cleaning of the biochip is highly facilitated as the cell adhesion is not promoted within the channel surfaces. Thus, the device lifetime is increased.

The structure that we developed is presented in Fig. 1(b). Two different configurations have been investigated, each of them including an external frame in thick resist (SU8, Microchem) that allows the positioning of the 12 mm \times 32 mm culture cover slides above the metallic electrodes. Notches have been included in the SU8 layer to facilitate the slide removal after the nsPEFs application. Considering the first design (Fig. 1(b)i), cells are separated from the electrodes, and disposed at a distance controlled by micropillars height (H) which are microfabricated in thick resist. Hence, this structure includes 25 μ m-high gold electrodes, and 50 μ m-high SU8 micro-pillars.

The second design (Fig. 1(b)ii), which was finally chosen for the experiments, possesses the same 25 μ m thick electrodes on which the removable slide covered with adherent cells is directly apposed. In this case, the length of the electrodes was adjusted ($L = 3$ cm) to achieve the impedance matching, taking into account the thickness ($t_c \approx 7$ μ m) of the cell layer cultured on the cover slide, and the conductivity ($\sigma = 1.55$ S m $^{-1}$) of the medium used. The thickness of the cell layer was estimated from the geometrical properties of DC-3F cells (the cell type used in our experiments), and was confirmed by the actual measured impedance (see next section). Once the culture slide is positioned in the SU8 frame, two spring strips are used to maintain it on the biochip.

C. Spatial homogeneity of the applied electric field pulses

The culture slide being positioned above the metallic electrodes, as shown in Fig. 1(b), the amplitude of the electric field depends on the location within the channel section. Hence, it is necessary to quantify the electric field attenuation to validate the feasibility of the cell nanoporation in this configuration. In Fig. 2(a), we compared the field distribution in the two design cases (with or without the 50 μ m resist pillars), using finite element analysis simulations performed with Comsol. We considered the thickness t_c of the cell layers.

This analysis shows that the variation of the electric field in a section of the microchannel remains quite acceptable in both structures, but that the best homogeneity (Table 1) is achieved with design 2 (without micro-pillars). Design 2 (25 μ m thick electrodes, no thick resist pillars) seems to be the most appropriate design, if the field homogeneity criteria in the microchannel is considered as the most important criterion (Fig. 2(a)ii, 2(b), and Table 1).

Nevertheless, the design 1 (involving the 50 μ m resist pillars) has the advantage of preventing any contact between the cells and the electrodes. Indeed, such contact might alter the cell viability (due to electric field edge effects visible on Fig. 2(b)ii).

III. Materials and methods

A. Fabrication process

Both proposed designs were fabricated on a commercial quartz substrate (ACM, France), as it is transparent, has good dielectric properties and presents good adhesion properties. The materials used for the fabrication are biocompatible. The metallic electrodes are made of electroplated gold as it is biocompatible and has a good conductivity ($\sigma = 4.1 \times 10^7$ S m $^{-1}$). The microfluidic chamber is patterned with a thick photosensitive resist (SU8, MicroChem), broadly used in bio-microfluidic devices.

The process started with a quartz substrate (from ACM), which was pre-metallized with a Cr/Au layer whose thickness was 30 nm/150 nm, respectively. After a cleaning of the wafer with successively acetone, isopropanol, water baths and desorption at 150 $^{\circ}$ C, the gold thickness was increased up to 25 μ m. The gold was electroplated using a KAu(CN) $_2$ bath and a current density of 1.5 mA cm $^{-2}$. The gold electrodes were then patterned using UV photolithography of S1805 resist, followed by gold and chromium chemical etching.

The microchamber was then fabricated on the electrodes by the structuring of an SU8 layer using photolithography. For the SU8 pillars design (prototype 1), a double SU8 layer was used to define the 50 μ m thick pillars and the 100 μ m thick SU8 frame. The first SU8 layer (SU8 2025) was spin coated to reach a thickness of 50 μ m (1700 rpm for 30 s), baked (3 min at 65 $^{\circ}$ C and 15 min at 95 $^{\circ}$ C), exposed to UV (390 mJ cm $^{-2}$) and post exposure baked (3 min at 65 $^{\circ}$ C and 15 min at 95 $^{\circ}$ C). The second SU8 layer (SU8 2075) was then spin coated to reach a final thickness of 100 μ m (2100 rpm for 30 s). The resist was baked (3 min at 65 $^{\circ}$ C and 15 min at 95 $^{\circ}$ C), exposed again (600 mJ cm $^{-2}$), post exposure baked (3 min at 65 $^{\circ}$ C and 15 min at 95 $^{\circ}$ C) and the two layers were simultaneously developed. Considering prototype 2 (without the pillars), only one layer of SU8 (SU8 2075) was spin coated to directly reach the targeted thickness of 100 μ m (2100 rpm for 30 s). The resist was then baked (5 min at 65 $^{\circ}$ C and 20 min at 95 $^{\circ}$ C), exposed to UV (600 mJ cm $^{-2}$) and post exposure baked (3 min at 65 $^{\circ}$ C and 15 min at 95 $^{\circ}$ C). The SU8 layer was finally developed to define the microfluidic chamber. Fig. 3 shows photographs of the fabricated biochips.

B. Cell preparation

DC-3F cells (Chinese hamster lung fibroblast cells) were grown in MEM (Minimum Essential Medium) supplemented with 10% fetal bovine serum, 100 U/mL penicillin and 100 μ g mL $^{-1}$ streptomycin. Cells were propagated at 37 $^{\circ}$ C in a humidified 5% CO $_2$ atmosphere. Cells were routinely passed every two days. All the chemicals were purchased from Invitrogen (Cergy Pontoise, France).

For the biological experiments, cells were cultured on glass slides for at least 4 h. In order to visualize the effect of nsPEFs on living cells, the slides were incubated for 30 min with the

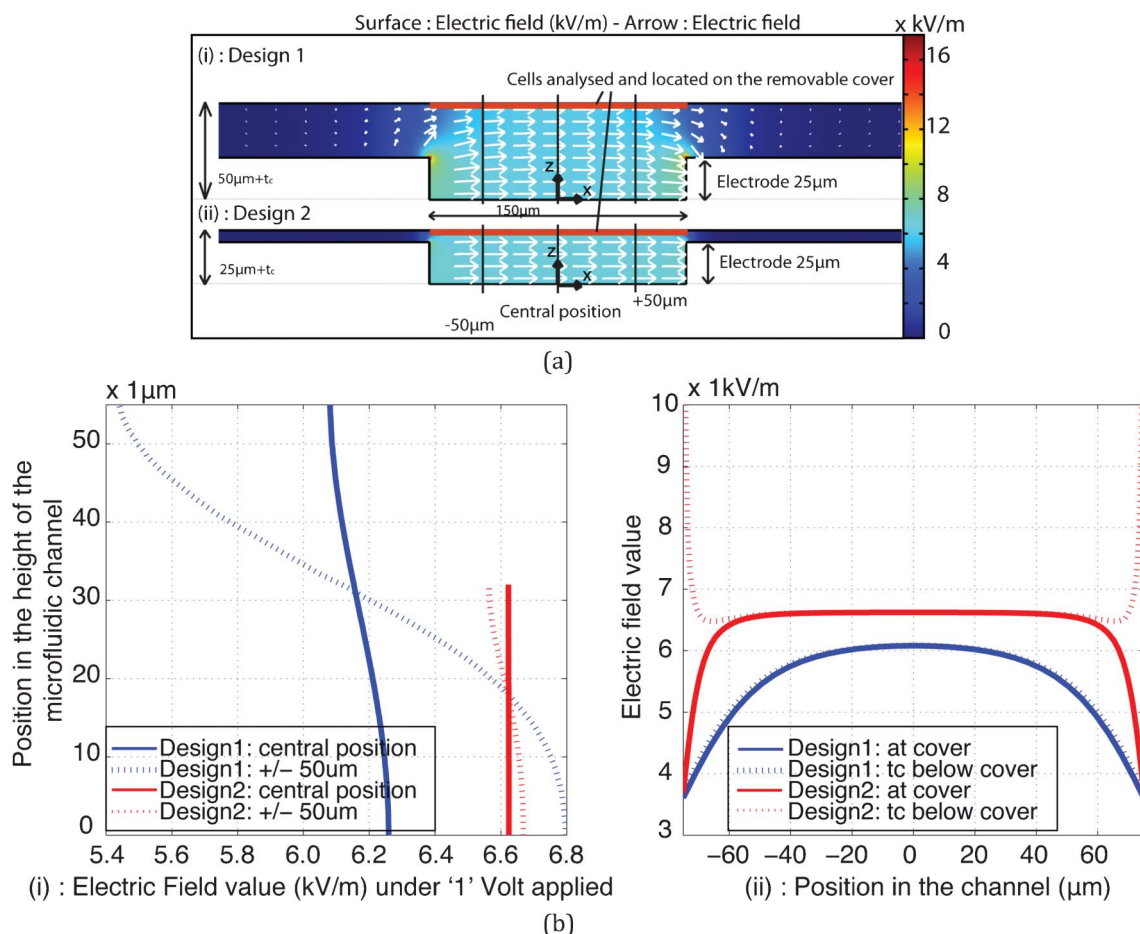


Fig. 2 (a) Electric field calculation by finite element analysis for the two designs developed for the nanoporation of adherent cells: (i) Design with the 50 μm resist pillars and (ii) Design without the micro-pillars. The cell layer is taken into account (estimated at $t_c = 7\text{ }\mu\text{m}$). (b) Homogeneity of the electric field within the nanoporation channel (for 1 V): (i) depending on the z position in the microfluidic chamber and (ii) depending on the x position of cells on the glass slide (from one electrode side to the other). Blue curves correspond to design 1 (with micro-pillars), while red curves correspond to design 2 (without pillars). (b) displays the electric field dependence on the z position (Fig. 2(b)i) and the x position (Fig. 2(b)ii) in the microfluidic chamber. The cell layer cultured on the cover slide is confined between altitudes $z = 50\text{--}57\text{ }\mu\text{m}$ in the case of prototype 1 and $z = 25\text{--}32\text{ }\mu\text{m}$ for prototype 2 (providing the cell thickness does not exceed $t_c = 7\text{ }\mu\text{m}$).

fluorescent calcium marker Fluo-4 AM ($\lambda_{\text{ex}} = 496\text{ nm}$, $\lambda_{\text{em}} = 515\text{ nm}$) in a humidified atmosphere with 5% CO_2 at 37 $^\circ\text{C}$. The slides were washed three times with PBS and placed on top of the chip (the cells being thus located inside the channel). Fluo-4 AM dye possesses acetoxymethyl groups (AM) allowing the molecule to enter the cells and preventing calcium binding outside of the cell. Once the marker is inside the cytoplasm, the AM groups are cleaved by endogenous esterases, resulting in both the trapping of the fluorophore and its activation. If the cytosolic calcium

concentration increases, Fluo-4 fluorescence increases. Hence, monitoring the fluorescence of exposed cells allows us to know whether organelle membranes or plasma membranes are affected by the nsPEF and become permeable to the calcium ions. Cells were exposed to a single 10 ns pulse of 280 kV cm^{-1} .

DC-3F cell images were taken with a Zeiss AxioCam Hrc camera on a Zeiss Axiovert S100 epifluorescence inverted microscope controlled through the Axio Vision 4.6 software (Carl Zeiss, Germany).

Table 1 Spatial homogeneity of the electric field

	Homogeneity of the electric field as a function of the z position in the microchamber		Homogeneity of the electric field as a function of the x position in the microchamber, over 90% of the distance between the electrodes	
	Central position (0 μm)	Side position ($\pm 50\text{ }\mu\text{m}$)	In contact with the cover	t_c (7 μm) below the cover
Design 1	2.85%	21.9%	28.9%	27.1%
Design 2	0.02%	1.55%	10.6%	2.01%

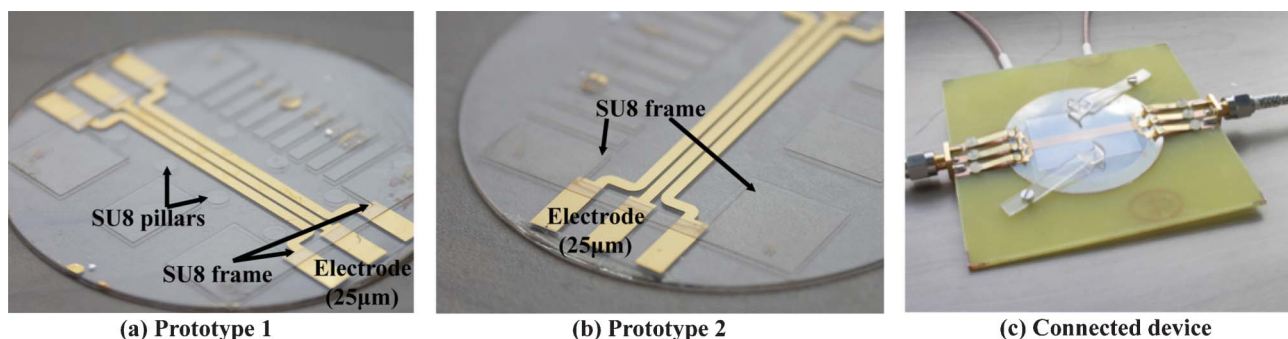


Fig. 3 Photographs of the fabricated microfluidic devices. (a) prototype 1, with SU8 microstructured pillars and (b) prototype 2 without pillars; (c) general view of the device connected to the PCB (printed circuit board), with two springs to maintain the culture slide.

C. Electrical set-up: nanosecond pulse generator and measurement

A commercial generator purchased from FID (FID GmbH, Model FPG 10-ISM10, Burbach, Germany) with an output impedance of $50\ \Omega$ was used to treat the cells. It generates trapezoidal monopolar pulses with a full-width at half maximum of 10 ns.^{18–20} The output voltage magnitude goes from 3 kV to 10 kV. The rising edge lasts between 0.4 and 0.6 ns. The falling edge lasts between 1 and 1.5 ns.

In order to characterize the incident and reflected signals in the exposure set-up, a tap-off (245 NMFFP-100, Barth Electronics Technology, USA) was connected between the generator and the biochip using two transmission lines.²² The tap-off is a three-port device. The impedance of the mainline (*i.e.* the nsPEF-biochip line) is $50\ \Omega$ whereas the impedance of the measurement port (*i.e.* the connection to the oscilloscope) is $4950\ \Omega$, allowing a voltage measurement with a 1 : 100 ratio. A Wavemaster 808 Zi 8 GHz oscilloscope (LeCroy, Courtaboeuf, France) was used for the measurements.

IV. Results and discussion

Optimum architectures of the device have been investigated in order to ensure a good homogeneity of the cells exposure conditions within the microfluidic channel. Indeed, for this study concerning nsPEFs effects on adherent cells, that are attached on glass cover slides reported on the nanoporation device, nsPEFs should be very well controlled and spatially homogeneous.

As shown by finite element analysis, the homogeneity of the electric field is enhanced when the culture slide is directly reported on the $25\ \mu\text{m}$ thick electrodes (prototype 2), as far as we examine cells that are not directly in contact with the electrodes. The $25\ \mu\text{m}$ thick electrodes structure combined to $50\ \mu\text{m}$ resist pillars (Prototype 1) gives satisfying but lower homogeneity for the electric field, as explained in section II.C. Therefore the prototype 2 ($25\ \mu\text{m}$ thick electrodes, no pillars) has been employed to achieve the results presented thereafter.

To validate the possibility of exposing cells reported on the top of the electrodes, biological experiments have been carried out and in particular, we study the dynamics of the $[\text{Ca}^{2+}]_i$ in response to the application of nsPEF. DC-3F cells were loaded with the calcium marker Fluo-4 AM. Cells were exposed to a single 10 ns pulse of $280\ \text{kV cm}^{-1}$. The profile of applied nsPEFs is reported in Fig. 4. As predicted, a good impedance matching is achieved; around 5% of the nsPEF is reflected back to the

nanopulser which gives an effective impedance of the chip equal to $45\ \Omega$. This slight difference could be interpreted as an underestimation of the cell layer height or some inhomogeneity in this height.

Concerning the cell exposure, recording started 100 s before the pulse to visualize the baseline of the fluorescence, and continued for 10 min after the nanopulse delivery. Fig. 5 reports typical results achieved under the experimental conditions reported here. As seen in Fig. 5(a), an increase in the Fluo-4 fluorescence in the presence of extracellular calcium, indicating a raise in the cytosolic calcium concentration, was observed. After 400 s the fluorescence was back to the baseline showing that the cells can recover this massive calcium release, probably through the activation of pumps and transporters.

The kinetics of calcium response is reported on Fig. 5(b) showing the immediate increase in fluorescence following the nanopulse delivery to the cells with a maximum value reached about 40 s after the nsPEF application. All the cells responded at the same time but the response amplitude was different from cell to cell (from a 0.5 fold increase to a 4 fold increase). Thus the nsPEFs applied to the DC-3F under the microdevice-associated experimental conditions actually generate a cytoplasmic calcium concentration transient increase.

Experiments were also performed in the absence of external calcium using this microdevice (cells treated in S-MEM, data not shown). It was still possible to generate calcium peaks showing that in this case, the internal calcium stored in the endoplasmic reticulum was involved in the generated cytosolic calcium peaks. This result is consistent with the work of Vernier *et al.* who showed in 2003¹⁰ that nanopulses were able to permeabilize the cells endoplasmic reticulum. However, according experiments carried out by Silve *et al.*,²¹ proving that it is possible to permeabilize the plasma membrane of the DC-3F cells with one single 10 ns trapezoidal monopolar pulse (at $40\ \text{kV cm}^{-1}$), the calcium peaks reported in Fig. 5 could also be the consequence of the presence of external calcium in the reported experiments.

V. Conclusion

An original microfluidic device dedicated to the application of nsPEFs to adherent cells and their real time monitoring has been developed and characterized in this work. The proposed device involves a removable packaging. Indeed, with such a packaging, the preparation of cell cultures on standard commercial slides

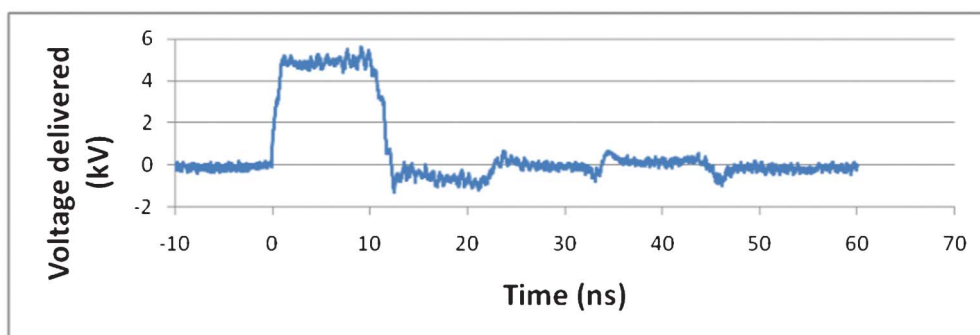


Fig. 4 recording of the applied nsPEFs.

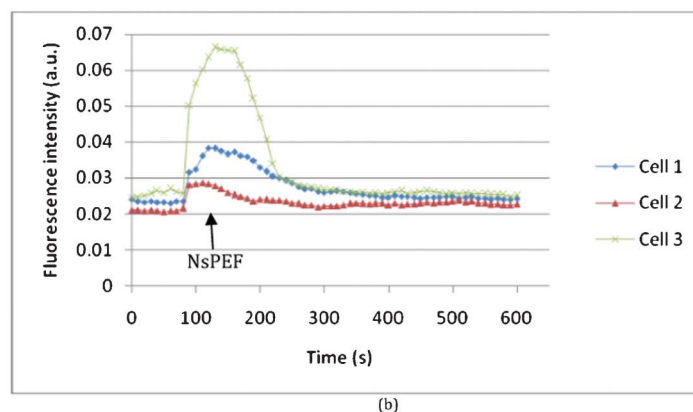
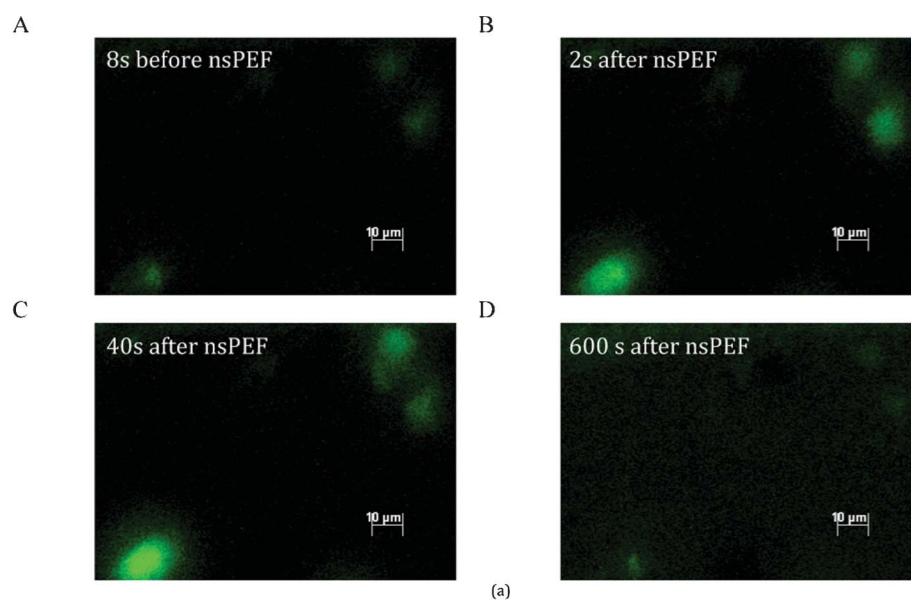


Fig. 5 (a) Real time imaging of a calcium peak from cells imaged every 10 s with a 500 ms exposure. Attached DC-3F cells were loaded with the calcium marker Fluo-4 (5 μ M). A single 10 ns pulse of 280 kV cm^{-1} was delivered. A: 8 s before the nanopulse. B: 2 s after the nanopulse. C: 40 s after the nanopulse. D: 600 s after the nanopulse and (b) kinetics of the calcium response of the three cells observed in the previous figure.

remains possible, these slides being afterwards reported on the device for the nanopulses exposition and cell monitoring experiments. Two approaches have been compared in terms of microfabrication process and electric field homogeneity achieved in the microfluidic chamber. The efficiency of our device has been demonstrated with a successful study of the dynamics of the

$[\text{Ca}^{2+}]_i$ in response to the application of nsPEF. The effect of extremely high amplitude and short duration electric pulses (280 kV cm^{-1} , 10 ns) that could be applied for the first time to adherent cells were recorded in real time in our microdevice. Kinetics of calcium responses show an increase in fluorescence due to the nanopulse application.

These promising results pave the way for the development of micro and nano systems well adapted to a better understanding of nanoporation effects on living cells.

Acknowledgements

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