High-Density Redox Amplified Coulostatic Discharge-Based Biosensor Array

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Abstract—High-density biosensor arrays are essential for many cutting-edge biomedical applications including point-of-care vaccination screening to detect multiple highly contagious diseases. Typical electrochemical biosensing techniques are based on the measurement of sub-pA currents for micrometer-sized sensors requiring highly sensitive readout circuits. Such circuits are often too complex to scale down for high-density arrays. In this paper, a high-density 4,096-pixel electrochemical biosensor array in 180-nm CMOS is presented. It uses a coulostatic discharge sensing technique and interdigitated electrode (IDE) geometry to reduce both the complexity and size of the readout circuitry. Each biopixel contains an interdigitated microelectrode with a 13-aA low-leakage readout circuit directly underneath. Compared to standard planar electrodes, the implemented IDEs achieve a maximum amplification factor of 10.5 × from redox cycling. The array’s sensor density is comparable to state-of-the-art arrays, all without augmenting the sensors with complex post-processing. The detection of anti-Rubella and anti-Mumps antibodies in human serum is demonstrated.

Index Terms—Biosensor, electrochemical biosensor, high-density array, interdigitated electrode (IDE), low-leakage switch.

I. INTRODUCTION

NUMEROUS biomedical applications rely on high-density biosensor arrays, which consist of thousands of individually addressable miniature sensors on a single substrate. One interesting application is the simultaneous detection of a wide range of humoral antibodies either for checking the immune system for the presence of antibodies created in the body post-vaccination, i.e., vaccination screening, or scanning the complete antibody profile for signs of illness as is the case in immunosignaturing (IMS) [1]–[3]. For the former, a single device capable of measuring multiple analytes would make rapid and comprehensive verification of immunization possible. For the latter, rather than directly sensing the disease antigen(s), IMS measures the immune system’s response to the disease, i.e., the patient’s antibody profile, which is amplified rapidly by white blood cells to several orders of magnitude higher concentration than the antigen itself. This shift in focus toward monitoring a set of antibodies not only leads to accurate and early diagnosis, but also allows for the tracking of disease progression [4]. For example, the antibody profile of an individual infected with rhinovirus for the first time is vastly different than the profile during all subsequent infections of the same virus. This occurs because during the initial infection the body has yet to determine the appropriate antibody to target the virus, for which it produces a wide variety of combinations.

To enable these technologies, instead of running several targeted tests for all the possible antibodies, which would require impractical amounts of time, reagents, and biological samples, a single unguided assay can be run. As illustrated in Fig. 1(a), this single assay can be performed by using an array of densely packed sensors, which are functionalized to detect a large set of antibodies in an individual.

Current high-density array technologies use optical detection (i.e., fluorescent dyes attached to the analyte) thus requiring complex microarray imaging equipment that is too bulky and impractical for point-of-care (POC) applications where measurements are made in remote areas away...
from the resources of centralized labs. While electrochemical detection is known to improve the size and scalability of biosensors [5], most implementations still require a potentiostat with an extremely sensitive transimpedance amplifier to measure the minute signals associated with microelectrodes, and such designs typically only scale to a \(~100 \times 100~\mu\text{m}^2\) pixel area [6]–[19]. For higher-density implementations, many potentiostat-based arrays either have specially fabricated sensors to increase sensitivity (i.e., amplify the signal) [10], [20]–[22] or implement parts of the measurement circuitry outside of the array to decrease the pixel size [8], [9]. Nonetheless, neither approach addresses the fundamental difficulty of measuring small currents with decreased sensor size.

In this paper, extended from [23], the design and validation of an integrated high-density biosensor array for vaccination screening, that also enables POC IMS, is presented. It leverages an alternative and less-used electrochemical detection method, coulostatic discharge [23]–[27], to significantly reduce the complexity and size of the readout circuitry. Fig. 1(a) shows an illustration of the array functionalized with different capture proteins to simultaneously detect multiple disease biomarkers. Each biopixel transduces capture events into an electrical signal \(I_{\text{sig}}(t)\), whose magnitude is related to the biomarker concentration. Unlike chronoamperometry, where this current is measured directly, the current is used to discharge the sensor’s innate double-layer capacitance [Fig. 1(b)], translating the measurement to changes in \(v_{\text{out}}(t)\). This technique effectively transforms a minute current measurement to a much simpler voltage-over-time measurement. As the sensor’s intrinsic capacitance is on the order of tens of picofarads and \(I_{\text{sig}}(t)\) is \(~1~\text{pA}/\mu\text{M}\), the discharge rate of \(v_{\text{out}}(t)\) is on the order of \(1~\text{V/s}/\mu\text{M}\), which greatly relaxes and simplifies the readout circuitry requirements. Hence, this technique along with electrochemical amplification from interdigitated electrodes (IDEs) allows for all the sensors and circuitry to be packed densely enough for high-density array applications using only the features available in a standard CMOS process.

The remainder of the paper is organized as follows. Section II describes the coulostatic discharge sensing principle, and Section III discusses the design of the biopixel circuitry. Sections IV and V present characterization and biological measurement results, respectively. Comparisons are made in Section VI, and conclusions are drawn in Section VII.

## II. Sensing Principle

Coulostatic discharge is an electrochemical technique developed independently by both Reinmuth and Wilson [24] and Delahay [27] in 1962, which uses the inherent double-layer capacitance between an electrode and an electrolytic solution to convert the signal current to a voltage that changes slowly over time. Specifically, the measurement is performed by charging up this capacitance to a particular voltage and then letting it discharge through the electrochemical cell. To use this method for biomarker detection, the surface of the electrodes must be coated with capture molecules that give the sensor specificity (i.e., the ability to distinguish a specific molecule from others).

The assay steps used in this paper are as follows: First, each biopixel is functionalized by immobilizing capture proteins on the gold-plated sensor using a standard method of dropcasting an excess amount of proteins to saturate the surface completely. The sensor’s surface is known as the working electrode (WE), which is where the biochemical detection occurs that is subsequently transduced into an electrical signal [Fig. 2(a-i)]. A single sensor contains two WEs separated by an insulating material (e.g., oxide in an IC). Each type of capture protein binds to a specific target antibody due to the binding affinity of the antibody-antigen complex.

Next, the biological sample is added to the sensors and incubated so that any antibody biomarkers present in the sample bind to their specific capture protein [Fig. 2(a-ii)]. After washing to remove any unbound molecules, a secondary antibody that binds to the bound antibodies is added to the array, effectively sandwiching the biomarkers [Fig. 2(a-iii)]. This secondary antibody is conjugated with an enzyme, alkaline phosphatase (ALP), that reacts with a \(p\)-aminophenyl phosphate (pAPP) substrate, thus generating a by-product redox molecule, \(p\)-aminophenyl (pAP), that is detected by the biopixel [Fig. 2(a-iv)]. Detection occurs when pAP approaches a WE biased at a specific voltage, known as the oxidation potential, and reacts to form quinonimine (QI) by transferring electrons to the WE. QI, in turn, reacts at the second WE biased at the reduction potential converting back into pAP by receiving electrons. Thus, this shuttling of electrons creates a current proportional to the concentration of the biomarker in the sample. An important distinction to make here is...
that actual full-scale IMS relies on *in situ* printing a large number of randomly-generated peptides onto the array of sensors [2], [28]–[30]. To use this array for IMS technology, peptides would be used in place of proteins to detect the antibody profile [31]. The underlying sensing mechanism is the same for both cases, so successful operation of the protein-based assay implies that the array can be used for IMS.

The resulting signal would typically be measured directly using standard amperometric techniques and current-based readout circuitry (e.g., as is done in a glucometer). However, in coulostatic discharge, a potential is applied to the electrode only briefly allowing a build-up of charge on the sensor’s intrinsic capacitance $C_{dl}$. This capacitance, known as double-layer capacitance, is formed from the layers of ions and charged molecules that assemble at the interface between the electrode and ionic solution, and exists between the electrode and the bulk solution [Fig. 2(b)]. After the source supplying the potential to the electrode is disconnected, $C_{dl}$ is discharged through the electrochemical cell by the current generated due to the redox reactions, thus slowly decreasing the voltage of the electrode at a rate related to the biomarker concentration.

As implied by Fig. 2(b), near the beginning of the discharge phase the sensor behaves as an $RC$ circuit, where the capacitance is $C_0$ and the resistance is determined by the initial redox current, which is related to the concentration of the redox molecules. Hence, the discharge rate can be written as

$$\frac{dv_{out}(t)}{dt} = \frac{I_{sig}(t)}{C_{dl}}$$  (1)

where $I_{sig}(t)$ is the current generated after opening the switch. In practice, the discharge curve is nonlinear due to the voltage dependence of both $I_{sig}(t)$ and $C_{dl}$. As described by the Nernst–Planck equation [32], $I_{sig}(t)$ depends on the concentration gradient of the redox molecules around the electrode

$$I_{sig}(t) = \frac{nFAD_oC_{ox}(t)}{\delta}$$  (2)

where $n$ is the number of electrons transferred per reaction, $F$ is the Faraday constant, $A$ is the area of the electrode, $D_o$ is the diffusion coefficient of the redox molecule, $C_{ox}(t)$ is the concentration of the redox molecule at the electrode surface, and $\delta$ is the width of the diffusion layer, which, for microelectrodes, is simply the distance between electrode fingers. Also, the Nernst equation links the voltage of an electrode, i.e., WE1 in Fig. 2, to the concentration of redox molecules at its surface as follows

$$V_{WE}(t) = E_0 + \frac{RT}{nF} \ln \left( \frac{C_{ox}(t)}{C_{ox,lim} - C_{ox}(t)} \right)$$  (3)

where $E_0$ is the standard potential of the redox species, $R$ is the universal gas constant, $T$ is the absolute temperature, and $C_{ox,lim}$ is the total concentration of the oxidant and reductant at the electrode surface [33]. Furthermore, the capacitance $C_{dl}$ is also a function of the electrode voltage. Specifically, $C_{dl}$ is given by

$$C_{dl}(t) = \frac{\varepsilon_0\varepsilon_r}{\lambda_D} \cosh \left( \frac{qV_{WE}}{2kT} \right)$$  (4)

where $\varepsilon_0\varepsilon_r$ is the dielectric constant, $\lambda_D$ is the Debye length, which is a measure of how far the electric field extends into the solution, $kT/q$ is the thermal voltage, and $V_{WE}$ is the potential of the electrode [32]. Note that the presence of $C_{dl}$, which is on the order of 10 pF–1 nF, obviates the need for an explicit capacitor in each biopixel. This capacitance is also at least one order of magnitude larger than the capacitance formed by the metal electrodes alone, so it dominates the sensor capacitance.

Fig. 3(a) shows simulated discharge curves obtained using (2)–(4) and known redox coefficients. Due to the nonlinear nature of these curves, the optimum range at which to sample the voltage signal is not obvious. To examine the tradeoffs of different sampling times, a noise model of the biopixel including the buffer and sensor’s equivalent noise...
Contributions, $i_{n,t}(t)$ and $i_{c,t}(t)$, is shown in Fig. 3(b), where the noise only affects the measurement after the switch is open. As the measurements are taken by subtracting two samples of the buffer’s output taken at different times (i.e., right after the switch opens and after a certain amount of time), thus implementing correlated double sampling (CDS) [34], the buffer in this model is connected only to the sensor. Also, charge injection is cancelled by measuring this difference.

As described above, $I_{lep}(t)$ integrates onto $C_{d}$, thus creating the characteristic discharge curves illustrated in Fig. 3(a). The sensor’s impedance is modeled as $C_{d}$ in parallel with the charge transfer resistance $R_{ct}$, which is a measure of how readily the sensor surface reacts with the redox molecules. It follows from Fig. 3(b) and the previous discussion that the voltage noise power can be computed as

$$
\sigma(\tau)^2 = \int_{1/\tau}^{\infty} \left( \frac{R_{ct}}{1 + j 2 \pi f C_{d} R_{ct}} \right)^2 (S_{e,n}(f) + S_{b,n}(f)) df
$$

(5)

where $\tau$ is the time at which the buffer output is measured, $S_{e,n}(f)$ and $S_{b,n}(f)$ are the one-sided power spectral densities (PSDs) of the noise current signals of the sensor and buffer, respectively, and $C_{d}$ is assumed to be constant. The noise contributed by the sensor is highly dependent on several biological and chemical factors. Measurement results show that this noise source, which can be modeled as having a white component and a $1/f$ component, is considerably more dominant than the noise contributed by the buffer, so the buffer design in this system is constrained practically only by the biopixel area.

As shown in Fig. 3(c), for large values of $\tau$ the signal component in $v_{out}(\tau)$ increases, but the noise variance also increases due to both the white and $1/f$ components of $S_{e,n}(f)$. In contrast, for small values of $\tau$, the signal component in $v_{out}(\tau)$ decreases, but the $1/f$ noise barely affects the measurement. Therefore, a range of suitable sampling times as well as an optimal point that maximizes the signal-to-noise ratio (SNR) can be found. Using estimated values for $R_{ct}$ and $C_{d}$, as well as measured noise PSDs for the buffer and sensor, the noise variance for different values of $\tau$ between 10 ms and 100 s was computed. The computed SNR for a 100-nM concentration for different scenarios is shown in Fig. 3(d). As can be seen from Fig. 3(d), the SNR is maximized at $\tau \cong 1.2$ s.

Fig. 3(e) shows that the same trend exists across various concentrations. Depending on the desired resolution, concentration range, and targeted measurement speed, the sampling time $\tau$ can be reduced. For instance, in applications such as IMS, where coarse or even binary detection is sufficient, a smaller sampling time can be chosen to decrease the time required to scan an entire array. Also, by relating the noise level back to current using $C_{d}$ and the sampling time, a current of $\sim$600 fA is expected at $\tau \cong 1$ s, so the switch should have a leakage smaller than this value.

III. SYSTEM DESIGN

A. Architecture

Fig. 4 shows a block diagram of the implemented biosensor array, where $V_{WE1}$ and $V_{WE2}$ are the input voltages applied to the working electrodes WE1 and WE2, respectively, $v_{1}, v_{2}, \ldots, v_{64}$ are the array outputs, and the readout circuit below each biopixel comprises a unity-gain buffer and a low-leakage switch controlled by the digital signals $r$ and $e$. As each $v_{j}$ output is shared among the 64 pixels from its respective column, a 6-bit decoder with outputs $d_{1}, d_{2}, \ldots, d_{64}$ is used to select the row of the array whose buffer outputs are connected to the array outputs. Given that all the buffers in a column are connected to the same output, the higher the biopixel is in the array, the longer the routing to the output pad is, so the signal sees a different delay to the output depending on the position of the biopixel. However, due to the slow-varying nature of the coulstatic discharge technique, these delays negligibly affect the measurements and are accounted for in the settling time. Nonetheless, in larger arrays, where the number of biopixels might force the use of smaller sampling times, this issue might need to be addressed. In such cases, this problem could be circumvented by dividing the array into subsections with separate outputs, so that the biopixel-to-output-pad paths are more uniform across the array.

The second working electrode WE2 is shared by all the pixels within the array, and it is always connected to $V_{WE2}$. In contrast, each biopixel has its own WE1, and $V_{WE1}$ is applied to this electrode only when the low-leakage switch is closed. Not shown in the figure are the counter electrode, implemented as a wide gold-plated strip across the center of the array, and the reference electrode, an external Ag/AgCl wire. Together, these electrodes, controlled by off-chip circuitry for flexibility, set the potential of the solution and provide a common reference voltage for the WEs.

A bias current is generated by a constant-$g_{m}$ reference located at the corner of the array and an off-chip resistor. This current is copied to 16 current mirrors at the top of the array, and each distributes this current down the columns to 16 local bias-voltage generators along the length of the chip. Each one of these bias-voltage generators (256 in total) biases the buffers of a $4 \times 4$ quadrant of biopixels to reduce area overhead.

To assess the performance of the low-leakage switch, test pixels were implemented on the right side of the array with a
dedicated output $v_{test}$. Each test pixel is almost identical to a biopixel, but instead of having an IDE connected to the input of its buffer, it has a 100 fF, 1 pF, or 10 pF MIM capacitor with its other terminal connected to ground. Instead of a low-leakage switch, some of these test pixels have a standard switch consisting of a PMOS transistor with its body tied to the supply voltage.

B. Sensor Design

Since IMS arrays of 330,000 pixels have been shown to be able to diagnose multiple illnesses reliably, the eventual target number of sensors for a future full-sized electrochemical IMS array is on the order of 100,000. Therefore, assuming the array is fabricated using the full reticle size and that 10% is used for control and output circuitry overhead, the maximum size for each pixel would be $\sim 100 \times 100 \, \mu m^2$. Specifically, an area of $42 \times 42 \, \mu m^2$ per sensor would allow for an array of exactly 330,000 pixels. Guided by this analysis, the electrode geometries were designed using similar pixel areas and constrained by the minimum width and spacing rules of the process.

The microelectrode sensors were fabricated using the top metal layer. Each biopixel consists of two IDEs designed to amplify the signal using an electrochemical amplification technique known as redox cycling. Redox cycling is the effect when a reversible redox pair repeatedly diffuses between two electrodes biased at different potentials, one at the reduction potential and the other at the oxidation potential of the pair, transferring electrons through redox reactions at the two electrodes. Hence, a single redox molecule can contribute multiple times to the overall current. The redox current signal $I_{rc}$ obtainable from a planar IDE is given by the following empirical equation

$$I_{rc} = \alpha \left( 0.64 \log \left( 2.6 \left( 1 + \frac{W}{G} \right) \right) - 0.19 \frac{G}{W + G} \right)^2$$

where $W$ and $G$ are the finger width and the gap between fingers, respectively, and

$$\alpha = n N F D o C_{dl} b$$

where $N$ is the number of fingers and $b$ is length of the fingers [35]–[38]. It follows from (6) that for a given area, a smaller gap width and a larger number of fingers provide higher amplification. As shown in Fig. 5, four different electrode designs were fabricated. Electrodes A.1 and A.2 have finger and gap widths of 5 and 5 $\mu m$, whereas electrodes B.1 and B.2 have finger and gap widths of 2 and 3 $\mu m$. In a standard phosphate-buffered saline (PBS) buffer solution, the $C_{dl}$ of these designs range from 24 to 180 pF. According to (6), the amplification factors of A.1, A.2, B.1, and B.2 are 2.25$x$, 5.27$x$, 4.93$x$, and 9.41$x$, respectively. Given that (6) is strictly applicable to 2-D IDE geometries, the estimated amplification factors are expected to underestimate the measured values.

Often, physical channels, trenches, or walls are fabricated directly on the top of or around the sensor to increase the cycling efficiency, which is a measure of a sensor’s ability to keep the same redox molecules shuttling between the two electrodes without them diffusing away from the sensing area [20], [21], [35]. While these structures are all effective in increasing the amplification, they require complex and customized post-processing steps, thus making the fabrication more time consuming and expensive. To avoid these additional complex post-processing steps while still increasing the signal amplification, an alternative approach that is CMOS foundry compatible was taken. Specifically, the passivation layer directly above each sensor was removed to create 3-D structures. By waiving design rule check errors meant to protect the bottom layers of the chip from over etching (the top metal is often used as an etch stop), structures that take advantage of the height of the electrodes were created. Given that the passivation was opened across the entire IDE, the etchant can carve down the gaps between the fingers. As shown in Fig. 6(a), this allows for the formation of 3-D trenches between the two electrodes that increase the collect-
Fig. 7. (a) Biopixel readout circuit. (b) Ultra-low-leakage switch operation. The frequency of the reset signals is 1 Hz with reset early leading by 1 μs.

IV. CHARACTERIZATION MEASUREMENTS

A 5 × 5 mm² chip (Fig. 8) was fabricated in a 1P4M 180-nm CMOS silicon on insulator (SOI) process. The array contains 64 × 64 biopixels split into four quadrants, each with a different IDE design. Each unit biopixel is only 45 × 45 μm².

A. Electrical Characterization

The entire chip consumes a maximum of 95 mW (23 μW per biopixel) from a 2.5-V supply. The offset of the buffer averaged across the entire array was measured to be 1.78 ± 0.16 mV. The input-referred total integrated noise of the buffer is 33 μV rms (100-kHz bandwidth) with a 1/f noise corner of 120 Hz.

To assess the performance of the low-leakage switch, discharge tests were run using the test pixels. There are two different types of test pixels: one with a standard PMOS switch
Fig. 9. (a) Test pixel schematics. (b) Average leakage measurement results from the test structures with 0.5 V applied across each (left) and the average leakage of all the test structures at different sensor voltages (n = 3).

and the other, which is identical to the one used in the array, with a body-driven switch [Fig. 9(a)]. Separate pixels of each type are attached to known MIM capacitors in place of the top metal electrodes. For each of these tests, up to 0.5 V was placed across the two WEs. Just as in a normal discharge test, the switch opens and charge begins to leak from the capacitor into the switch causing the voltage measured to decrease over time. The slope of the measured voltage over time and the known capacitance size was then used to calculate the leakage current the test sensor experiences. The body-driven switch leakage was measured to be 13 aA, which is less than that of the standard PMOS switch with the body tied to $V_{DD}$, which was 195 aA [Fig. 9(b)]. The body-driven switch had better performance across the entire voltage range.

The results suggest that a standard switch could be used instead of a low-leakage switch in this application. This is the case because the array was designed in an SOI process. However, the biopixel topology is meant to be compatible with any standard CMOS process, where standard switches might not perform as well. For instance, in a standard 180 nm process, according to simulations, the leakage current of a typical PMOS switch is $\sim 1.3 \text{ pA}$, whereas that of the low-leakage switch is $\sim 4 \text{ fA}$.

B. Sensor Preparation

For electrochemical compatibility, an electroless nickel immersion gold (ENIG) plating process (MICRO, Stapleton Technologies Inc.) was used to plate the exposed top metal aluminum electrodes [7], [12], [40], [41]. After a thorough cleaning of the surface with acid, alkali, and deoxygenation cleaners, the entire chip was put through a double zincate process to prepare the exposed aluminum electrodes for the subsequent electroless nickel and gold plating steps. The combined thickness of these layers is usually around 3 $\mu$m. Since the electrodes in this work are interdigitated instead of a single pad of metal, the spacings and features of the IDEs are small enough that the plating can create unwanted shorts between the two WEs. Furthermore, even collections of small metal particles left over from the plating process can cause individual IDEs to have much lower resistances. Hence, different plating times of both the nickel and gold steps were experimented with and evaluated by measuring the resistance between the two halves of the IDE [Fig. 10(b)]. A total plating time of $\sim 5$ min was found to be optimal due to its high resistance while still maintaining a robust and even coverage of gold across the array.

C. Electrochemical Characterization

To evaluate the performance of the sensors as well as the gold plating, a chip with bare gold electrodes was used to measure the redox molecule Ferro/Ferricyanide as a proxy for the actual assay. After cleaning the chip by sonicating in isopropyl alcohol, the chip was mounted in a socket designed to create a $\sim 10 \mu$L well over the sensors. External Ag/AgCl and Pt electrodes were dipped into this well to form the reference and counter electrodes. Next, coulostatic discharge was run using the in-pixel circuitry to measure the sensors in various concentrations of Ferri/Ferricyanide in both
single WE, as seen in Fig. 11(a), and dual WE modes to compare the same sensor with and without redox cycling. In single-electrode mode, both sides of the IDE are shorted together, instead of biased independently, effectively making them into a single electrode with combined area where no redox cycling can occur. In dual electrode mode, the IDE operates as intended with one electrode biased at 200 mV and the other at 0 mV relative to \( V_{CM} \), allowing for the shuttling of the redox molecules. Using these measurements, the amplification factor was determined by calculating the ratio of the signals between the dual and single electrode modes. As shown in Fig. 11(b), the average amplification factors for each design are 5.33 ± 1.2, 8.1 ± 1.5, 6.06 ± 2.1, and 10.5 ± 2.1, respectively. These values are slightly larger than the theoretical values due to the 3-D trench and nanowell structures. IDEs with the same gap and finger widths but greater number of fingers have higher amplification. Smaller gap size also increases the redox cycling, as expected. Furthermore, the large variation in amplification can be explained as either a result of uneven plating of gold in the trenches or variability in the formation of the trenches themselves. The latter seems more likely since the plating procedure is widely used while the etching between fingers is unconventional and not guaranteed by the foundry.

V. BIOLOGICAL MEASUREMENT RESULTS

For the biological tests, only a portion of the array that has the same electrode design, A.2, was used to allow for a fair comparison between tests. To demonstrate a bioassay, 2 \( \mu \)g (66 pmol) of Rubella virus capsid protein (ab74574, Abcam) in PBS was dropcast on the surface of the gold sensor array using 20 \( \mu \)g of Traut’s Reagent (26101, ThermoFischer Pierce) and blocked with 1% bovine serum albumin (37525, ThermoFischer Scientific). Mouse anti-Rubella antibodies (ab34749, Abcam) were subsequently added and incubated for one hour. For the secondary antibody, 1 \( \mu \)g of rabbit anti-mouse secondary antibodies linked with ALP (ab6729, Abcam) was used. Lastly, 6-M pAPP substrate (sc-281392, Santa Cruz Biotechnology) in a 0.1-M glycine buffer pH 8.4 was added and allowed to incubate for 10 min. The ALP enzyme reacts with the pAPP substrate producing pAP, an electrochemically active molecule that shuttles electrons between the two fingers of the sensor. Fig. 12(a) shows measurement results for both cyclic voltammetry, measured with a 25-mV/s scan rate from \(-0.2\) to \(0.3\) V for 3 cycles, and coulostatic discharge. Both successfully detect the presence of anti-Rubella antibodies. The experiment was then repeated for anti-Rubella spiked into human serum (HS-20, Omega Scientific, Inc.). Fig. 12(b) shows the average discharge rates for serum with and without the antibody.

Next, a multi-biomarker assay meant to simultaneously detect both anti-Rubella and anti-Mumps (ab9880, Abcam) antibodies was performed using the array. The same portion of the array used for the previous Rubella-only test was split into two, with each part functionalized with either Rubella or Mumps protein (ab74560, Abcam). Four different chips were used for this experiment, all functionalized exactly as described above. Each chip was given a different test sample of serum spiked with 1.3-\( \mu \)M Rubella antibodies, 2.3-\( \mu \)M (ab34749, Abcam) and 1.3-\( \mu \)M Mumps antibodies (ab74560, Abcam).
Mumps antibodies, both, or neither. As seen in Fig. 12(c), the parts of the array that are exposed to their corresponding antibody show a higher slope than those that are not. Although there exists a large chip-to-chip variation, likely due to disparity in functionalization or test conditions between the different chips, the array can still distinguish between each of these different samples to detect the presence of either biomarker, thereby demonstrating its capability to monitor vaccinations.

VI. Comparison

Table I compares this work to other integrated electrochemical biosensor arrays, and Fig. 13 plots their pixel areas and number of devices per pixel with different markers to signify those that have special post-processing and/or have measurement circuitry external to the array. Due to redox cycling, this work achieves a comparatively small pixel area and high sensor density (400 pixels/mm²) without any special post-processing, which others need to increase sensitivity. While augmenting sensors with additional structures and materials is effective, it requires complex fabrication steps that are much more difficult and expensive to produce and scale than an array built purely with a standard CMOS process. To the best of our knowledge, this work is the highest density amperometric biosensor array that does not require additional post-processing steps. Furthermore, coulostatic discharge greatly decreases the number of devices required in the measurement circuitry (~12) allowing for the circuit to reside completely within the area of a pixel. In fact, the two arrays based on coulostatic discharge have the lowest number of devices that fit completely within a pixel. Hence, rather than occupying a considerable amount of area with circuit blocks external to the array, this work makes efficient use of the chip area as illustrated by the total density calculation in Table I.

VII. Conclusion

This paper presents a scalable coulostatic discharge-based high-density biosensor array designed to miniaturize multiple antibody measurement technology. Using coulostatic discharge rather than standard amperometry, the measurement circuitry in each biopixel is simplified to just a low-leakage switch and output buffer, thereby minimizing the pixel area and increasing the density of the overall array. Furthermore, by optimizing on-chip IDE geometries and waiving fabrication design rules to create 3-D structures, a signal amplification of ~10× was achieved without any complex, costly, and time-consuming post-processing of the sensors. Using this array, Rubella antibody was detected in human serum and simultaneous measurement of both Rubella and Mumps antibodies was possible on the same chip. These tests demonstrate this array’s promise for use in full-scale IMS technology for rapid and accurate POC screening and diagnosis.

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