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# A tight squeeze: geometric effects on the performance of three-electrode electrochemical-aptamer based sensors in constrained, *in vivo* placements†

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Electrochemical, aptamer-based (EAB) sensors are the first molecular monitoring technology that is (1) based on receptor binding and not the reactivity of the target, rendering it fairly general, and (2) able to support high-frequency, real-time measurements *in situ* in the living body. To date, EAB-derived *in vivo* measurements have largely been performed using three electrodes (working, reference, counter) bundled together within a catheter for insertion into the rat jugular. Exploring this architecture, here we show that the placement of these electrodes inside or outside of the lumen of the catheter significantly impacts sensor performance. Specifically, we find that retaining the counter electrode within the catheter increases the resistance between it and the working electrode, increasing the capacitive background. In contrast, extending the counter electrode outside the lumen of the catheter reduces this effect, significantly enhancing the signal-to-noise of intravenous molecular measurements. Exploring counter electrode geometries further, we find that they need not be larger than the working electrode. Putting these observations together, we have developed a new intravenous EAB architecture that achieves improved performance while remaining short enough to safely emplace in the rat jugular. These findings, though explored here with EAB sensors may prove important for the design of many electrochemical biosensors.

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## Introduction

The ability to measure the concentrations of clinically relevant molecular targets continuously and in real time in the living body would significantly advance our ability to diagnose, monitor, treat, and understand disease. Towards this goal, we are developing electrochemical aptamer-based (EAB) sensors: a reagentless, reversible molecular sensing platform that performs well *in vivo*. In these, the binding of a specific target induces a conformational change in an electrode-attached, redox-reporter-modified aptamer (Fig. 1a), resulting in an

easily detectable change in the redox reporter's electron transfer kinetics that we typically monitor using square wave voltammetry.<sup>1–6</sup> Because this conformation-linked signaling mechanism does not rely on any specific chemical or enzymatic reactivity of the target itself, the development of EAB sensors against new targets is relatively easy. With this signal transduction mechanisms that mimics those found in naturally occurring chemoperception systems, EAB sensors also support real-time, high-frequency measurements in complex sample matrices, including unprocessed bodily fluids.<sup>2,7–10</sup> Building on these advantages, EAB sensors have already been used to measure the rise and fall of more than a half-dozen therapeutics,<sup>1,2,6,11</sup> drugs of abuse, and metabolites<sup>5</sup> in real time in the blood of live rats. The resulting high-frequency, real-time molecular measurements have even been shown to support closed-loop, feedback-controlled drug delivery,<sup>1,12</sup> an advance that could usher in a new era in which pharmacological therapies can be delivered with unprecedented precision.<sup>13</sup>

In our prior work, we have deployed *in vivo* EAB sensors as a 3-electrode system consisting of a gold wire working electrode modified with the target-recognizing aptamer, a chloride-anodized silver wire reference electrode (RE), and a platinum wire counter electrode (CE). These are individually insu-

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**Fig. 1** (a) Electrochemical aptamer-based (EAB) sensors are comprised of redox-reporter-modified, conformation-shifting aptamers attached to a gold electrode *via* a self-assembled monolayer. The phrase “conformation-shifting” refers to an aptamer that has been reengineered such that target binding reversibly alters its shape, changing in turn the rate of electron transfer of the attached redox reporter. (b) EAB sensors support molecular measurements *in vivo*. To do so in the rat jugular, we have used bundled, catheter-enclosed sensors. In one architecture, for example, a gold wire working electrode, a platinum wire counter electrode, and a chloride-anodized silver wire reference electrode are staggered within a 20-gauge catheter such that the counter electrode remains within the catheter, and half of the reference electrode and all of the working electrode extend into the lumen of the vein. (c) This is then surgically inserted into the external jugular vein of an anesthetized rat. (d) Using this and related architectures we have monitored the plasma concentrations of multiple drugs and metabolites in live rats.<sup>1,2,5,6,11</sup> Shown here, for example, are 14 s-resolved plasma vancomycin measurements performed using the described architecture employed in the rat jugular after an intravenous injection of the drug.

lated using heat-shrink polytetrafluoroethylene (PTFE) and then bundled together in a staggered fashion and placed in a catheter. To date, we have published work with *in vivo* EAB sensors using two specific arrangements following this general scheme. In one, 75  $\mu\text{m}$  diameter wires were used for all three electrodes and only the gold working electrode extends out of

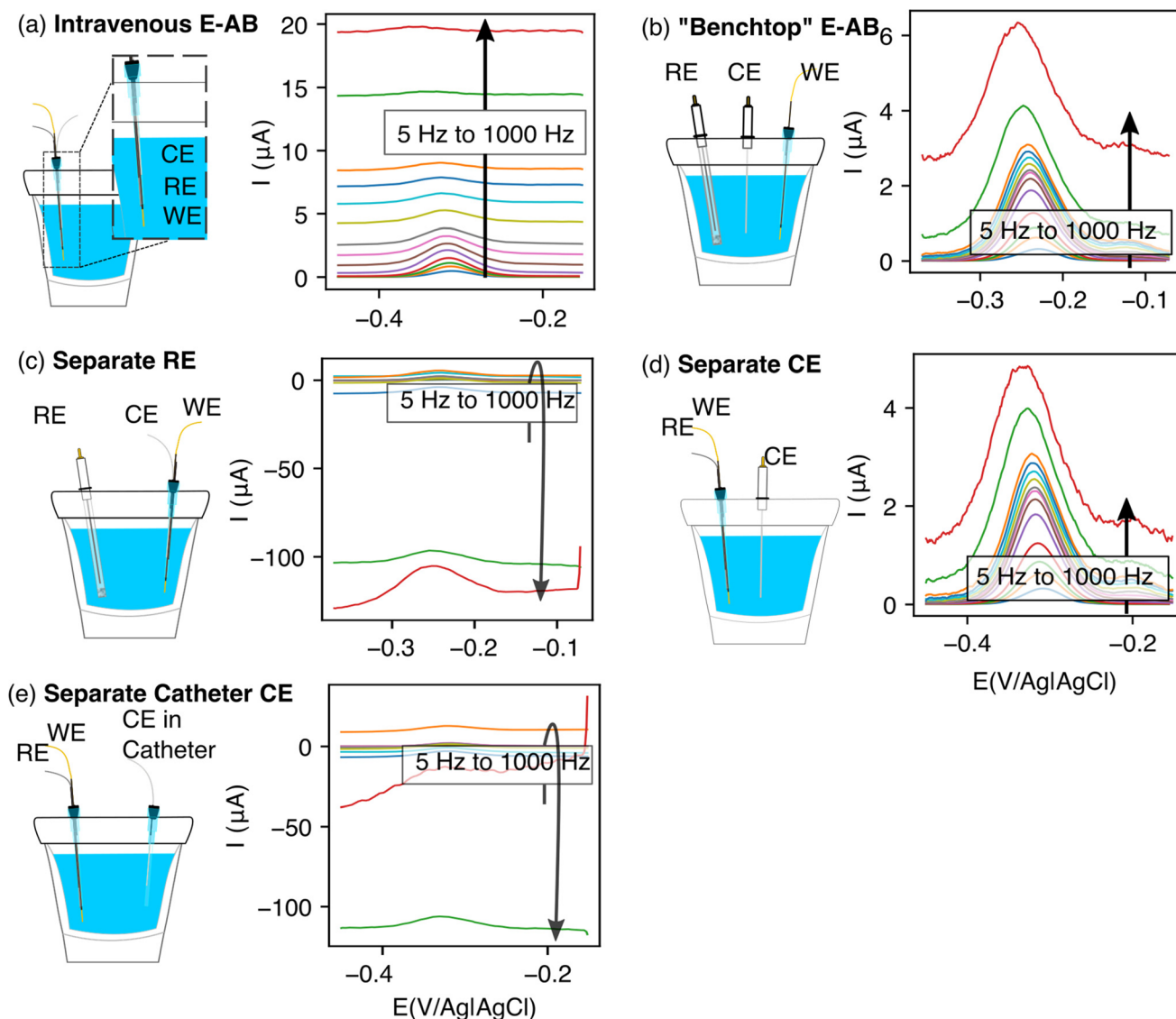
the catheter and into the vein.<sup>1,2,5,6,11,14</sup> In a second, we employed 200  $\mu\text{m}$  diameter wires placed in the catheter (Fig. 1b) such that half of the reference electrode and all of the working electrode extend out of the catheter and into the vein (Fig. 1c).<sup>15</sup> In both cases, we retained the counter electrode within the catheter, so as to reduce the length of the sensor and thus avoid interfering with the heart.

As extensive prior literature demonstrates and as shown here, the *in vivo* EAB architectures we have employed support unprecedentedly highly-time resolved measurements of plasma drug and metabolite levels (*e.g.*, Fig. 1d). This said, however, several aspects of the fabrication of intravenous EAB sensors remain unexamined, suggesting that their performance could be further improved. Specifically, here we explore the extent to which the relative placement of the three electrodes in an intra-catheter EAB sensor alters the device’s performance, with the expectation that better understanding this will guide us towards improved signal-to-noise ratios and smaller, less invasive devices.

## Results and discussion

The performance of EAB sensors depends strongly on whether their electrodes are placed inside or outside of a catheter. To see this, we first used *in vitro* experiments to compare a “bundled-electrodes” intravenous architecture, in which the counter and reference electrodes are retained within a catheter and the working electrode extends out of the lumen, with a sensor comprised of three physically separated, catheter-free electrodes. Doing so, we find that the baseline currents associated with the bundled architecture are (1) high and (2) a relatively strong function of square wave frequency (Fig. 2a). This behavior contrasts with that seen in the “benchtop” EAB sensor where the reference and counter electrodes are separated from the bundle (Fig. 2b). The high baselines associated with the bundled sensor reduce the peak currents, suggesting the details of the placement of EAB electrodes within a catheter may harm sensor performance.

We next set out to determine which of the three electrodes in an EAB sensor are responsible for the high baseline currents associated with our initial bundled architecture. As our first experiment, we characterized a sensor in which the counter and working electrodes were placed in the catheter, but the reference electrode (here a commercial, glass-frit Ag/AgCl reference) was physically separated. Under these conditions, the baseline currents we observe are as high and as strongly frequency dependent as those seen with the initial bundled architecture (Fig. 2c). Following this, we separated the counter electrode, leaving the reference and the working electrode in the catheter. With this case, we recover the low baselines and reduce the frequency dependence observed for the separated electrodes architecture (Fig. 2d). Building on these observations, we next separated the counter electrode but placed it in a second catheter. The baseline currents we observed for this architecture were again as high and as strongly frequency



**Fig. 2** The performance of EAB sensors is strongly dependent on which of their three electrodes are placed inside or outside of the catheter. (a) For example, our initial "bundled" architecture, in which all of the working electrode and half of the reference electrode extend out of a 20-gauge catheter but the counter electrode remains within, produces high and strongly frequency dependent baseline currents. (b) In contrast, when we employ three, physically separated electrodes (including, here, a commercial Ag/AgCl reference), the baseline is lower and less frequency dependent. (c) When only the reference electrode is separated, the resulting baselines are again strongly frequency dependent. (d) When we separate only the counter electrode from the bundle, however, we recover the behavior seen for the fully separated electrodes geometry. (e) Finally, when we place the separated counter electrode in a 20-gauge catheter, the baselines once again become strongly frequency dependent.

dependent as those of the initial bundled architecture (Fig. 2e).

The poorer performance seen when the counter electrode is placed inside a catheter arises due to the high solution resistance this architecture produces. To see why this may be true, consider the working principles of a 3-electrode electrochemical cell. In the absence of redox reactions, this cell can be modeled as a series RC circuit comprised of a resistor of resistance  $R_s$ , which captures the resistance of the solution between the working and counter electrodes, and a capacitor of capacitance  $C_{dl}$ , which captures the capacitance of the electric double layer (Fig. 3a and b).<sup>16</sup> How rapidly the capacitor

charges (*i.e.*, the time it takes for the formation of the double layer) depends on a time constant given by the product  $R_s C_{dl}$ . This, in turn, determines the current observed at a specific time after the potential is applied, which defines the baseline current at a given square wave frequency (in square wave voltammetry, the current is sampled at the end of each square wave). Specifically, smaller time constants (*i.e.*, faster charging) decrease the baseline current seen at higher square wave frequencies (*i.e.*, shorter time delays before current sampling). In contrast, longer time constants increase the baseline currents. For architectures including a catheter-enclosed counter electrode, the small cross section of the solution within the cath-



**Fig. 3** We can explain the differences in electrochemical behavior between the bundled and the separate sensor architectures by equating the electrochemical cell to an equivalent electrical circuit. (a) When no redox reactions are present, the RC circuit model can be used to describe a 3-electrode electrochemical system. (b) It consists of a capacitor representing the double layer at the working electrode interface ( $C_{dl}$ ) and solution resistance ( $R_s$ ) between the counter and working electrode. (c) For the bundled architecture, in which the counter electrode is retained within the catheter, the  $R_s$  between it and the working electrode is high due to limited thickness of the fluid layer within the catheter.

eter ensures that  $R_s$  is much larger, increasing the product  $R_s C_{dl}$  and, in turn, the baseline currents increase substantially as a function of square wave frequencies (Fig. 4a).

Characterization of our various sensor architectures using chronoamperometry and impedance spectroscopy supports

the above conclusions. Specifically, using chronoamperometry to measure the sensor's temporal response to a potential jump we find that, as expected given the above arguments, the time constants observed when the counter electrode is free in solution are shorter than those seen when the counter electrode is



**Fig. 4** As evident with multiple electrochemical techniques, the electrochemical behavior of EAB sensors depends strongly on whether the counter electrode is positioned inside or outside of the catheter. (a) For example, employing square wave voltammetry (here at 100 Hz and 25 mV amplitude), the baseline current is significantly higher, and the peak height is smaller when the counter electrode is retained within the catheter (blue) versus when it is placed outside of the catheter (red). (b) In chronoamperometry, we apply a  $-0.45$  V potential step and observe the resulting current transiently. When the counter electrode is placed inside the catheter (blue curve), the response time constant of the electrode is notably reduced. (c) Differences in the Nyquist plot seen when we characterized our architectures using electrochemical impedance spectroscopy further supports our arguments. Specifically, when the counter electrode is placed inside the catheter (blue curve), the x intercept, which indicates the solution resistance ( $R_s$ ), increases 60-fold (Table 1). In contrast, the increase in the imaginary impedance (on the y axis) is so similar for both cases resulting in  $C_{dl}$  values that are within error of one another (Table 1).

contained within a catheter (Fig. 4b). Using electrochemical impedance to distinguish between the resistive components (arising due to  $R_s$ ) and capacitive components (arising due to  $C_{dl}$ ) of our sensors produces a similar picture.<sup>17</sup> Specifically,  $R_s$  increases and  $C_{dl}$  is largely unchanged when we move the counter electrode from free in solution to constrained within a catheter. In a Nyquist plot, in which the imaginary impedance is plotted as a function of the real impedance (Fig. 4c), we find that the  $x$  intercept, which is indicative of  $R_s$ , is significantly higher when the counter electrode is inside the catheter than when it is outside. In contrast, the capacitive contributions remain unchanged (Table 1).

With our realization that sensor performance improves when the counter electrode is placed outside the catheter, we next designed a 2<sup>nd</sup> bundled sensor architecture in which the reference electrode is placed inside of the catheter and the counter and working electrodes both extend

beyond the catheter tip (Fig. 5). As noted above, however, a potential problem associated with such architectures is that, using the 1.5 cm counter electrode length we usually employ, the bare electrodes extend at least 2 cm beyond the catheter tip, risking interactions between the counter electrode and the heart (Fig. 1c). Thus motivated, we next explored the extent to which we can reduce the length of the exposed counter electrode without unduly degrading sensor performance. To our surprise, given the generally held requirement for three electrode cells that the counter electrode must be larger than the area of the working electrode,<sup>18–20</sup> we found no significant differences in sensor performance when we shortened the counter electrode from 1.5 cm to 3 mm (Fig. 5). We believe that this (to us unexpected) result arises due to the thiol passivating layer on our working electrode, which reduces the double layer capacitance of the working electrode. Irrespective of its origins, however, the practical upshot of this observation is that we can employ a short enough counter electrode to ensure that the sensor does not reach the heart.

Building on the above observations we next designed an intravenous sensor architecture employing a fully exposed, 6 mm-long counter electrode (Fig. 6a). Unfortunately, however, we observed transient shorting when the potential was first applied to the sensor. We suspect that this occurred because the tight fit of these wires inside the catheter leaves the PTFE shrink-tubing we used to insulate the electrodes susceptible to microtears, leading to short-circuiting between the reference and working electrodes. Fortunately, replacing the 200  $\mu\text{m}$  diameter silver and 250  $\mu\text{m}$  diameter platinum wires with 125  $\mu\text{m}$  diameter wires (Fig. 6a) eliminated this effect.

Extending the counter electrode outside of the catheter improves the signal-to-noise of intravenous EAB sensors. To illustrate this, we monitored plasma vancomycin levels upon a 30 mg kg<sup>-1</sup> intravenous dose using both our initial architecture and the above-described, new architecture (Fig. 6a). The 1.0  $\mu\text{M}$  noise (defined as the standard deviations of the fluctuations around the baseline prior to dosing) associated with the new architecture under these experimental conditions is approximately 3 times smaller than that we see for our first bundled architecture (Fig. 6b). One source of this discrepancy is the fact that voltammograms collected in the rat jugular using this new architecture exhibit less noise associated with the heartbeat (Fig. 6c). Given the results reported here, we suspect that this noise occurs as the animal's pulse dilates and constricts the vein and, thus, the catheter, causing the  $R_s$  to vary periodically for our first bundled architecture.

The effects we observe may be relevant to 3-electrode EAB sensor deployments beyond the bundled intravenous sensors explored here. For example, there has been growing interest in less-invasive EAB sensors deployed as subdermal or intradermal microneedles.<sup>21,22</sup> In these scenarios, there is limited liquid or limited electrolyte present, which could similarly lead to high resistances between the counter electrode and working electrode. Consistent with

**Table 1** Parameters obtained from fitting Electrochemical Impedance Spectra when the counter electrode is located either inside or outside of the catheter

Parameter	Counter electrode inside the catheter	Counter electrode outside the catheter
$R_s$	$6540 \pm 350 \Omega$	$104 \pm 4 \Omega$
$C_{dl}$	$0.49 \pm 0.02 \mu\text{F}$	$0.50 \pm 0.02 \mu\text{F}$
$\chi^2$	1.0313	0.0091524



**Fig. 5** Reducing the counter electrode size by 15-fold does not significantly affect the performance of EAB sensors. Specifically, we obtain near identical voltammograms whether using a (a) 1.5 cm counter electrode or (b) one that is just 1 mm in length. For results at intermediate counter electrode lengths, see Fig. S1.† Insets illustrate the length of counter electrode exposed in each geometry.



**Fig. 6** Building on the results reported here we have designed a new intravenous EAB architecture. (a) Altering the relative placement of the three electrodes to ensure that the counter electrode is placed outside the catheter significantly reduces the noise we see when we place these sensors in the rat jugular. (b) To illustrate this, here we have monitored plasma vancomycin concentrations after a  $30 \text{ mg kg}^{-1}$  intravenous injection. The baseline noise decreases from 2.3% when the counter electrode is enclosed in the catheter to 0.8% in the new architecture, which converts to standard deviations in concentration of  $3.5 \text{ }\mu\text{M}$  and  $1.0 \text{ }\mu\text{M}$  respectively. (c) Looking closely at the square wave voltammograms, we see that there is a reduction in noise associated with the animal's pulse.

our findings, as a result of the limited electrolyte present, others have measured high signal to noise when such EAB sensors are placed in the dermal tissue of a rat,<sup>21</sup>

suggesting that reducing the resistance between the working and counter electrodes could improve the performance of such sensors.

## Conclusions

Here we have shown that changing the relative placements of their three electrodes can significantly improve the performance of catheter-bound *in vivo* sensors. Specifically, the voltammogram baselines were lower and peak heights higher for architectures in which the counter electrode emerges from the lumen of the catheter. We attribute this difference to a reduction in the resistance between the working electrode and counter electrodes. We have also shown that the EAB counter electrode can be significantly smaller than the working electrode without harming sensor performance. Finally, we combined these two observations to create a new, intravenous EAB sensor architecture that achieves significantly improved performance when placed in the jugular veins of live rats.

## Conflicts of interest

The authors declare the following competing financial interest(s): One author (K. W. P.) has a financial interest in and serves on the scientific advisory boards of two companies attempting to commercialize EAB sensors. Following the completion of this work, two authors (K. L. and J. G.) became employees of a company attempting to commercialize EAB sensors.

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