



## PassStat, a simple but fast, precise and versatile open source potentiostat



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### ABSTRACT

This work presents 4 open source potentiostat solutions for performing accurate measurements in cyclic voltammetry and square wave voltammetry at a low price. A very simple and easy to reproduce analogic board (c.a. 10 €) was driven either by a Teensy card from the company PJRC under an Arduino/Python software solution (39 €) or by an Analog Discovery 2 device from Digilent (less than 300 €). A smartphone Bluetooth Android interface was also created to circumvent the use of a computer. We demonstrated that our scheme is suitable for measurements in classical electrochemical conditions but also to carry out experiments with ultramicroelectrodes. We could thus reach a noise resolution of less than 1 pA. Scan rates of 8000 Vs<sup>-1</sup> with ohmic drop compensation were also achieved. The device is suitable for teaching purposes but also for experiments in a participative science context on the ground, or countries with lower financial possibilities.

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## Specifications table

Hardware name	<i>PassStat</i>
Subject area	<ul style="list-style-type: none"> <li>Chemistry and biochemistry</li> <li>Educational tools and open source alternatives to existing infrastructure</li> <li>General</li> </ul>
Hardware type	<ul style="list-style-type: none"> <li>Measuring physical properties and in-lab sensors</li> <li>Other: Performing electrochemical measurements (cyclic voltammetry and square wave voltammetry)</li> </ul>
Closest commercial analog	There are many commercial potentiostats available (Autolab, PalmSens, Biologic, Origalys, CH Instruments)

(continued on next page)

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(continued)

Hardware name	<i>PassStat</i>
Open source license	CERN-OHL-S v2 for the hardware GNU GPL v3 for the software Creative Commons v4 for design files and documentation
Cost of hardware	40–303 € depending on the configuration
Source file repository	<a href="https://zenodo.org/record/5719382#.YZwj17rjJhE">https://zenodo.org/record/5719382#.YZwj17rjJhE</a> <a href="https://ohwr.org/project/passstat">https://ohwr.org/project/passstat</a>
OSHW certification UID (OPTIONAL)	FR000017

## Hardware in context

Analytical electrochemistry is a domain in continuous expansion because electrochemical measurements are simple, cheap to implement, and can be realized in a point of care approach [1]. Many analytes are electroactive or at least involved in reactions with electroactive species, so that electrochemistry is relevant for environmental, health or food areas among others. Specific electrode preparation is sometimes necessary to reach ultralow concentrations but in many cases non modified electrodes are already sufficient. Some commercial potentiostats present excellent detection limits and are compact enough to be easily transported. Nevertheless, they are still black boxes that are impossible to be repaired by the end user in case of damage. Their relatively high cost prevent their use in resource-limited or within participative science contexts. Therefore, many interesting cheap home-made and open source systems have been proposed for educational purposes or developing countries [2–11]. In some cases, some devices such as the one proposed by Matsubara reach characteristics close to commercial systems [11]. The heart of a potentiostat is an analytical device that should handle a three electrode configuration, meaning that no current should flow through the reference electrode (RE) while precisely controlling the potential of the working electrode (WE). For that, several operational amplifiers (OAs) are usually used [12]. Those are however very cheap and compact components. Additionally, a generator to impose electrode potential and an acquisition card or processor to register the data are also mandatory. Some open-source devices specifically develop all these elements, which optimizes compactness. For example Whitesides et al., Leech et al. and Rajendran et al. independently developed potentiostats that can be remotely controlled through wifi with a smartphone [3,7,9]. Nevertheless, the benefit in size is at the expense of board simplicity and reparability which may discourage new and not trained users to build their own apparatus. Many DIY devices rely on the well-known Arduino Uno microcontroller. Nevertheless, no digital-analytical converter (DAC) is available in this device so that this function either have to be implemented electronically or thanks to another specific component to accurately control the electrode potential and generate the ramp in cyclic voltammetry for example [4]. To face these issues, the Rodeostat [13] proposed by the company Irodeo (250\$) and the SweepStat proposed by Glasscott et al. took benefit from the Teensy 3.2 card provided by PJRC that offer DAC outputs and is compatible with the Arduino environment [14]. Nevertheless, the Rodeostat scheme is based on surface mounted component and thus can only be repaired in a well-equipped electronic laboratory. The Sweepstat proposes a rather simple scheme that takes benefit from use of a quad amplifier but it only works in a two electrode configuration. All open source potentiostats are compared in the recent paper by Matsubara (see section J in SI) [11].

Both to propose a low cost but performant device and for educational and intellectual purposes, we studied how to produce the simplest potentiostat scheme working in three electrode configuration. This led us to propose an easy to understand and cheap (<10€) electronic card from which several configurations can be implemented by simply displacing some switches. The first alternative to run the potentiostat is to use positive and negative power supplies, to apply the potential perturbation with a function generator and to acquire the signal with an oscilloscope. In this case, the potentiostat scheme can be resumed to three OAs, three resistors and two capacitors. In this paper, we took benefit from the device Analog Discovery 2 proposed by Digilent (academic price 294€) that proposes all these features in a single USB driven device. The second and cheapest option stands again on the Teensy 3.2 or 3.6 cards (price 24 or 39 €). The potentiostat power supply is directly provided by the USB voltage available from the Teensy. Since USB port power pins are 0 and 5 V, and that no negative tension is available from the computer, the electronic scheme was conceived to handle this problem. An additional Bluetooth module and an external battery can be added easily for remote control using an Android smartphone. Moreover, we wished to examine and push the performances of our system either towards low current (pA) detection or fast scan (several thousands  $Vs^{-1}$ ) voltammetry that we commonly use in our laboratory. This aspect was not treated in previous papers excepted in the work of Matsubara [11]. Below, we first depict the different possible implementations of the potentiostat. We then present representative results obtained in cyclic voltammetry or square wave voltammetry. We took ferrocene as common reference electroactive entity, and paracetamol as typical example of drug analysis application [15–25]. We named our potentiostat the PassStat, after the French electrochemical YouTube video series “Le Courant Passe” produced by the Societe Chimique de France to catch student interest [26]. All the electronic schemes and driving softwares are provided in Zenodo repository.

## Hardware description

- Cheap and repairable potentiostat card
- Low noise level down to 1 pA
- Fast scan rates up to  $8000 \text{ Vs}^{-1}$
- Can be driven by Teensy or Analog Discovery 2 devices

For designing our potentiostats, we took advantage of the TI LMC 6484 quad OA as first explored by Glasscott et al. [14]. This component displays very low input currents at the expense of its bandwidth that is limited to 1 MHz which is less than the one required to reach scan rates above  $5 \times 10^4 \text{ s}^{-1}$  (this was not the purpose of the present work). Having all amplifiers in a same component greatly simplifies the electronic scheme, particularly regarding the power supplies. In the following, we provide one figure per possible implementation. The printed circuit board measured only  $5 \times 5 \text{ cm}^2$ . We also propose standard values for the electronic components to probe the system in a “standard configuration” paragraph. We noted, as often observed that measurements are improved when realized in a Faraday cage (mandatory for currents lower than  $1 \mu\text{A}$ ). Even better results were obtained when the power supply of the laptop computer was disconnected (in that case the Faraday cage should be connected to the ground of the computer). A Faraday cage is however not mandatory for standard measurements with millimetric electrodes. The different schemes are provided by increasing complexity. Each labeled component has the same role in various configurations. Jumpers on the printed circuit board card allow to switch between different configurations. Table 1 summarizes the main characteristics of the different configurations.

**Table 1**  
Main characteristics of different PassStat configurations.

Name	Potential Range (V)	Microcontroller	Interface	Maximum scan rate	Remote control	Price (€)
PassStat 1	5 V with AD2, up to 15 V	Analog Discovery 2	Waveforms from Digilent	$8000 \text{ Vs}^{-1}$	no	310
PassStat 2.0	1.6 V	Teensy 3.6	Arduino/Python	$66 \text{ Vs}^{-1}$	no	50
PassStat 2.1	2.4 V	Teensy 3.6	Arduino/Python	$100 \text{ Vs}^{-1}$	no	50
PassStat 2.2	2.4 V	Teensy 3.6	Android Studio	$100 \text{ Vs}^{-1}$	Bluetooth	70

### *PassStat 1.0: Simplest and fastest design.*

This first setup, which represents the classical scheme [12,27,28,28–30], is presented in Fig. 1. Here, only three resistors and two capacitors (to stabilize the amplifiers) are used. Amplifier OA3 is used as follower in order to ensure that no current flows through the reference electrode while OA2 adjusts the potential of the counter electrode. The gain of the current-tension converter OA4 can be tuned thanks to resistor  $R_6$ , and capacitor  $CF_3$  in parallel can be adjusted to filter the noise while keeping  $R_6C_6$  low compared to the characteristic timescale of the measurements to keep the signal integrity.

In this work, this setup was implemented with an Analog Discovery 2. This device includes one positive and one negative power supplies ( $V^+ = +5\text{V}$  maximum,  $V^- = -5\text{V}$  minimum), two signal generators and two scope channels. For a larger compliance supplying up to  $\pm 15 \text{ V}$  is possible (but necessitates two additional power supplies). The generator output was sent to the potentiostat to provide the electrode potential (*i.e.* the ramp in cyclic voltammetry). It was also sent to the first scope channel to measure the potential although this is not completely necessary.

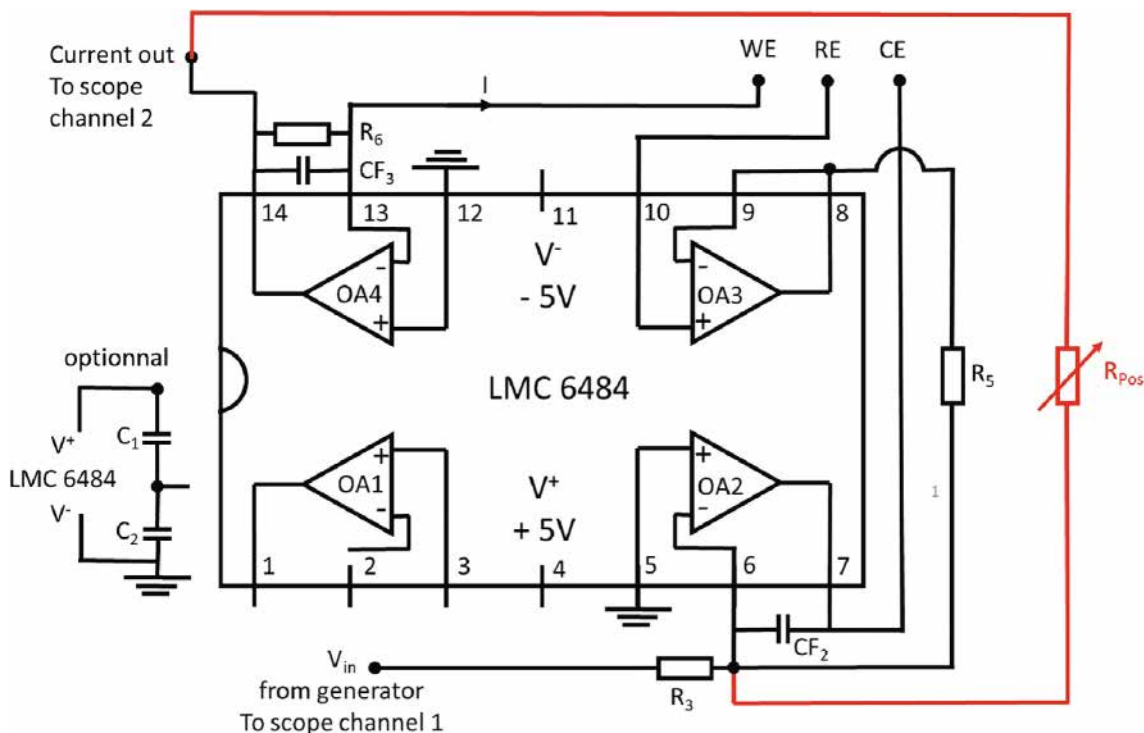
This simple setup allows to benefit from the whole bandwidth of the potentiostat. On a test with a resistor, we noted that with  $CF_2 = 15 \text{ pF}$ ,  $CF_3 = 3.3 \text{ pF}$  and  $R_6 = 10 \text{ k}\Omega$  there is no apparent amplitude diminution at  $40,000 \text{ Vs}^{-1}$ . In these conditions the phase shift translates into a temporal delay between the current and potential of  $0.5 \mu\text{s}$ . Peak potentials would then be altered by  $5 \text{ mV}$  at  $10,000 \text{ Vs}^{-1}$ . These estimations were however carried out without ohmic drop compensation that is usually necessary at large scan rates and may alter the bandwidth (see below for experimental results on ferrocene).

*Standard configuration:*  $R_3 = R_5 = 10 \text{ k}\Omega$ .  $CF_2 = 15 \text{ pF}$ .  $CF_3 = 1 \text{ nF}$ .  $R_6 = 1 \text{ M}\Omega$ . This configuration is suitable for probing a  $1 \text{ mM}$  ferrocene solution in acetonitrile with a  $0.5 \text{ mm}$  diameter platinum electrode at scan rates between  $0.05$  and  $5 \text{ Vs}^{-1}$ . For larger scan rates or electrode diameters, the current will saturate. Then diminish  $R_6$  to  $100$  or  $10 \text{ k}\Omega$  and  $CF_3$  progressively down to  $3 \text{ pF}$  for increasing the bandwidth.

Although this configuration is the most powerful and versatile, the final price of the device may be limiting. In addition, programming the generator for cyclic voltammetry and chronoamperometry is not very difficult. For more complicated potential ramps such as those used in square wave voltammetry or differential pulse voltammetry for example, a specific program should be elaborated (see section E in the work of Matsubara) [11]. The subsequent configurations solve these issues by using a low price interface card together with a dedicated software.

### *Ohmic drop compensation*

Ohmic drop occurs when large electrodes or relatively fast scan rates are used. It is due to the non-negligible electrolyte resistor between the working and reference electrodes. This effect distorts the signal but can be compensated by using small electrodes and adding electronically in real time a tension proportional to the current output to the voltage ramp. Further



**Fig. 1.** PassStat 1.0 scheme. The  $R_{\text{Pos}}$  potentiometer is optional and devoted to ohmic drop compensation. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

information about ohmic drop compensation may be found in the literature [12,28,29,31,32]. Only PassStat 1.0 configuration can integrate ohmic drop compensation. For that, a potentiometer should be added between pins 14 and 6 as shown by the red add-ons in Fig. 1. The potentiometer value should take into account the solution resistor (see analytical calculations below). When ohmic drop compensation gets close to 100%, oscillations appear at the initial and inversion potentials, as explained in refs [27–32].

#### PassStat 2: Plug and play low cost configuration with a Teensy card

In the following several configurations of the potentiostat are available for driving with a Teensy 3.2 or 3.6 card. These cards provide the advantage of having one (3.2) or 2 (3.6) analog outputs that can be used for defining accurately the electrode potential. They work with the Arduino IDE after installation of the Teensyduino add-on. A Python software with a graphical interface was also programmed to send the parameters (for example number of cycles, scan rate, potential excursion in CV) and collect the data. However, the potentiostat scheme needs to be adapted because the Teensy card does not provide  $V^+$  and  $V^-$  power supplies unlike the analog discovery. There are however a 5 V output (in fact the tension provided by the USB supply that stands near 5 V) and a 3.3 V one. To face this problem of dissymmetrical tensions, we added on the board voltage dividers to produce intermediate voltages for the positive input  $E^+$  of the amplifiers OA1, OA2 and OA4 that are thus not any more connected to the ground. Capacitors  $C_1 = C_2 = 47 \mu\text{F}$  and  $C_3 = C_4 = 1 \mu\text{F}$  damp the possible fluctuations of the power supply. In our conditions, they could be removed without alteration of the data but we consider that it is safer to include them.  $V^-$  is then connected to ground and  $V^+$  to 3.3 (PassStat 2.0) or 5 V (PassStat 2.1 and 2.2). We show below that the electrode potential, thus  $V_{\text{W}} - V_{\text{RE}}$  can be accurately controlled by this way.

The Teensy analog output is 12 bits over a 3.3 V range so that each potential increment is 0.8 mV (PassStat 2.0) or 1.2 mV (PassStat 2.1 and 2.2). Such values are precise enough for most electrochemical experiments. The input resolution for the current can be chosen up to 16 bits. The Teensy card allows applying and measuring data as fast as 12  $\mu\text{s}$ . This limits the scan rate to 66  $\text{Vs}^{-1}$  for PassStat 2.0 and 100  $\text{Vs}^{-1}$  for PassStat 2.1 and 2.2. Faster scan rate could be achieved using potential steps larger than 0.8 or 1.2 mV but this limit is sufficient for most analytical electrochemistry experiments. Since USB voltage is not precisely 5.00 V on all computers, a calibration procedure, detailed below, is necessary if accurate measurements are desired. A precise voltage reference stage can also be used in case voltage fluctuations are observed but this was found not necessary in our case.

#### PassStat 2.0: Simple but compliance limited to 1.6 V

Here, the 3.3 V output is connected to  $V^+$ .  $V^-$  is connected to the analog ground as displayed in Fig. 2. Resistors  $R_9$  and  $R_{10}$  (1  $\text{k}\Omega$ ) are used to provide a 1.65 V tension for the  $E^+$  entry of OA2 and OA4. This configuration may be used when the best

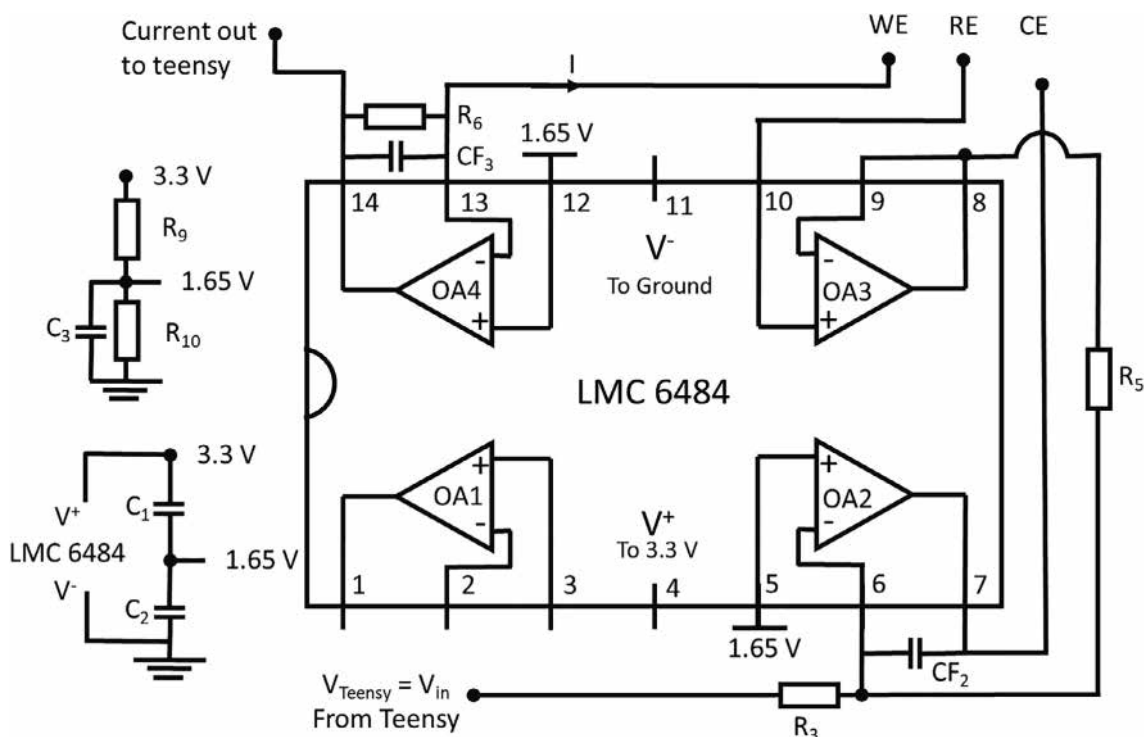


Fig. 2. PassStat 2.0 scheme.

current and potential precisions are desired (0.8 mV on the DAC and ADC instead of 1.2 mV). It is for example suitable with ultramicroelectrodes but may be limited if the potential excursion should be extended beyond  $\pm 1.6$  V or for large electrodes for which the counter electrode needs a larger compliance.

**Standard configuration:**  $R_9 = R_{10} = 1$  k $\Omega$ .  $R_3 = R_5 = 10$  k $\Omega$ .  $CF_2 = CF_3 = 100$  pF.  $R_6 = 100$  M $\Omega$ . This configuration is suitable for probing a 1 mM ferrocene solution in acetonitrile with a 12.5  $\mu$ m radius electrode at 10 mVs $^{-1}$ . A steady state CV should be observed. For smaller electrodes or concentrations, increase  $R_6$  up to 1 G $\Omega$ .

#### PassStat 2.1: Extended compliance to 2.4 V

In order to benefit from the maximum compliance possible with the Teensy card,  $V^+$  is now connected to the  $V^{in}$  pin of the Teensy card that is powered directly by the USB lead (an external battery could also be used). The analog output range is extended thanks to an inverting voltage amplifier stage implemented with OA1, the first amplifier of LMC6484 with resistors  $R_1$  (10 k $\Omega$ ) and  $R_2$  (15 k $\Omega$ ). Calculations detailed below show that  $E^+$  of this OA should be poised to 2 V, thus another voltage divider was implemented thanks to resistors  $R_{11}$  (1.5 k $\Omega$ ) and  $R_{12}$  (1 k $\Omega$ ). The voltage precision on the DAC and ADC is now thus  $0.8 \times 1.5 = 1.2$  mV.

Nevertheless, tension at the analog inputs of the Teensy should not be above 3.3 V to avoid irreversible damage to the card. To face this issue, voltage at the output of the current-tension converter ( $V_{14}$ , OA4) is converted back to a 0–3.3 V range thanks to another voltage divider (resistors  $R_7 = 510$   $\Omega$  and  $R_8 = 1$  k $\Omega$ ). Another input was designed for supplying an additional perturbation through resistor  $R_4$  in view of future impedance measurements but this has not been implemented yet. This scheme is displayed in Fig. 3.

**Standard configuration:**  $R_1 = 10$  k $\Omega$ .  $R_2 = 15$  k $\Omega$ .  $R_4 =$  not connected.  $R_7 = 510$   $\Omega$ ,  $R_8 = 1$  k $\Omega$ ,  $R_{11} = 1.5$  k $\Omega$ .  $R_{12} = 1$  k $\Omega$ .  $R_9 = R_{10} = 1$  k $\Omega$ .  $R_3 = R_5 = 10$  k $\Omega$ .  $CF_2 = CF_1 = 15$  pF.  $CF_3 = 1$  nF.  $R_6 = 1$  M $\Omega$ . This configuration is suitable for probing a 1 mM ferrocene solution in acetonitrile with a 0.5 mm diameter platinum electrode at scan rates between 0.05 and 5 Vs $^{-1}$ .

We underline here that Teensy 3.5 analog/digital inputs are 5 V tolerant, giving the possibility to remove resistors  $R_7$  and  $R_8$  but this was not tested in the present study.

#### PassStat 2.2: Remote control by Bluetooth for Android smartphone

The teensy card may integrate a remote control via Bluetooth with the RX and TX pins. Here, we chose the HC05 module to establish a serial communication with a smartphone. In this case, an Arduino program should be first uploaded in the Teensy. Then, the Teensy can be disconnected from the computer and powered by a 5 V battery. An application was developed with Android Studio to establish communication. Like for a computer, all parameters can be entered from the phone and the voltammogram is displayed after the acquisition.

**Standard configuration:** identical to PassStat 2.1.

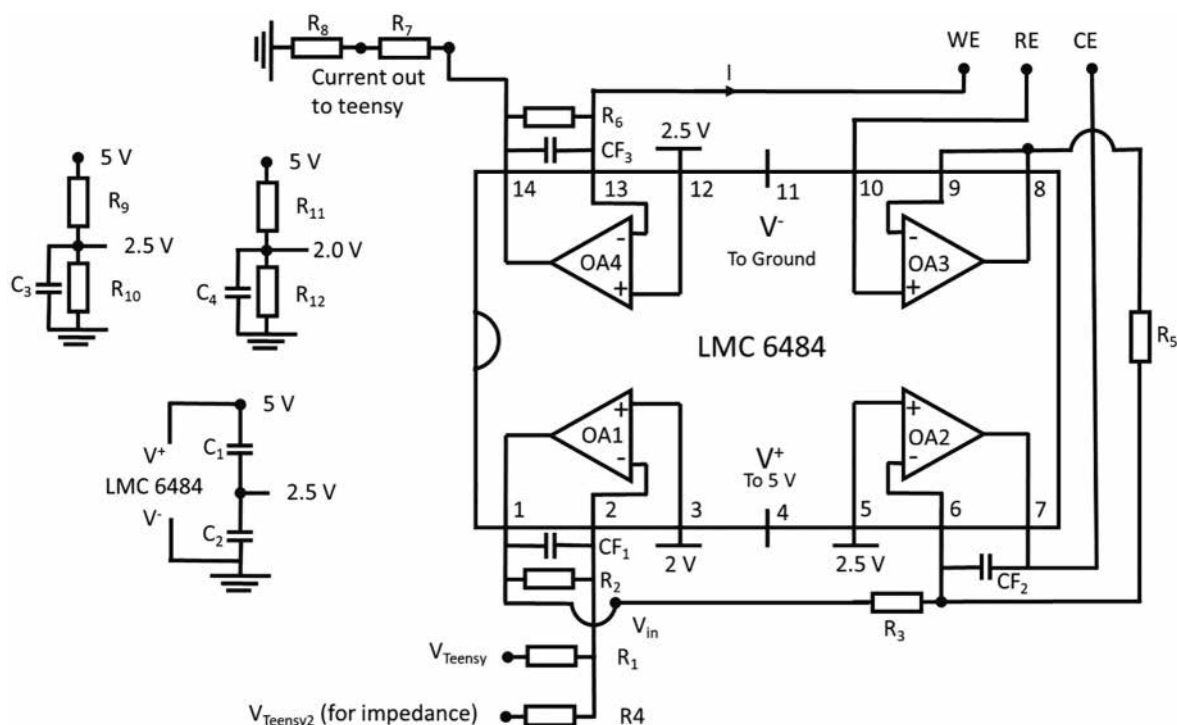


Fig. 3. PassStat 2.1 scheme.

Analytical formulation for the different configurations

We below develop the analytical formulations for the different potentiostat configurations, considering ideal operational amplifiers. We tried to write this section to be understandable by readers not familiar with electronics. The role of capacitors CF<sub>1</sub> to CF<sub>3</sub> that are present to minimize noise and stabilize the system is then neglected. We consider that a tension V<sub>in</sub> is entered at the potentiostat input. This tension is applied either directly to R<sub>3</sub> from the Teensy A22 (or A14 for Teensy 3.2) or Analog Discovery outputs (PassStat 1.0 and 2.1) or is obtained by conversion from the Teensy A22 pin tension through OA1.

With ideal amplifiers in linear regime we have:

- E<sup>+</sup> = E<sup>-</sup> the voltages are equal at both inputs
- i<sup>+</sup> = i<sup>-</sup> = 0 no current flows through the input

In our scheme, the working electrode potential is kept constant at the ground (or departed ground).

Control of the working electrode potential

PassStat 1.0

The tension and currents are labeled after the pins of the LMC 6484 component.

In this classical formulation, we have by application of node law at point 6:

$$(V_{in} - V_6) / R_6 = - (V_8 - V_6) / R_5 \text{ since } I_6 = 0.$$

Here V<sub>6</sub> = V<sub>5</sub> = V<sub>12</sub> = V<sub>13</sub> = 0 (virtual ground).

With R<sub>6</sub> = R<sub>5</sub> we have V<sub>8</sub> = - V<sub>in</sub> = V<sub>9</sub> = V<sub>10</sub> the tension applied to the reference electrode.

Thus V<sub>13</sub> - V<sub>10</sub> = E = V<sub>in</sub> is the working electrode potential.

Ohmic drop compensation

Because of ohmic losses in solution, the voltage applied to the faradaic impedance is in fact not E but E - R<sub>s</sub>I, where R<sub>s</sub> is the solution resistor between the working and reference electrodes. Ohmic drop is particularly problematic with solutions of poor conductivity and/or at relatively large scan rates. Electronic compensation allows to circumvent this problem, with



some limitations explained in references [25–29]. Here positive feedback is operated thanks to variable resistor  $R_{pos}$ . It was implemented only onto PassStat 1.0 that provides access to large scan rates.

Application of node law on pin 6 leads to:

$$V_{in}/R_3 + V_{14}/R_{pos} + V_{10}/R_5 = 0.$$

Hence with  $R_3 = R_5$ .

$$E = V_{13} - V_{10} = 0 - V_{10} = V_{in} + R_5 V_{14}/R_{pos} = V_{in} + (R_6 R_5/R_{pos}) \cdot I.$$

By diminishing  $R_{pos}$  the feedback is increased. It reaches 100% compensation for  $R_6 R_5/R_{pos} = R_s$  where  $R_s$  is the solution resistor. In practice, near 100% compensation, an oscillatory behavior appears. Capacitor  $CF_3$  damps these oscillations, at the expense of a bandwidth reduction. The damped oscillations appear at the potential inversion on Fig. 13 (see below).

#### PassStat 2.0

$$(V_{in} - V_6)/R_6 = -(V_8 - V_6)/R_5 \text{ since } I_6 = 0.$$

Here  $V_6 = V_5 = V_{12} = V_{13} = 1.65 \text{ V}$  (deported virtual ground created by voltage divider made with  $R_9$  and  $R_{10}$ ).

$$V_8 = V_9 = V_{10} = -V_{in} R_5/R_6 + V_6(1 + R_5/R_6) = -V_{in} + 2 V_6 \text{ with } R_5 = R_6.$$

$$\text{Thus } V_{13} - V_{10} = E = V_{in} - V_6.$$

Since the Teensy card analog output ranges from 0 to 3.3 V, setting  $V_6$  to 1.65 V allows to reach electrode potentials between  $-1.65$  and  $+1.65 \text{ V}$ . The compliance is thus limited.

#### PassStat 2.1 and 2.2

Here,  $V_2 = V_3 = 2 \text{ V}$  thanks to the bridge divider made with  $R_{11}$  and  $R_{12}$ .

$$(V_1 - V_2)/R_2 = -(V_{Teensy} - V_2)/R_1 \text{ since } I_6 = 0.$$

$$\text{Hence } V_1 = V_{in} = V_2(1 + R_2/R_1) - V_{Teensy} R_2/R_1.$$

The analogic output of the Teensy ranges from 0 to +3.3 V. With  $R_2/R_1 = 1.5$   $V_1 = V_{in}$  ranges from 5 to 0.05 V.

Next.

$$V_6 = V_5 = V_{12} = V_{13} = 2.5 \text{ V}.$$

Here  $V_6 = V_5 = V_{12} = V_{13} = 1.65 \text{ V}$  (deported virtual ground created by voltage divider made with  $R_9$  and  $R_{10}$ ).

$$V_8 = V_9 = V_{10} = -V_{in} R_5/R_6 + V_6(1 + R_5/R_6) = -V_{in} + 2 V_6 \text{ with } R_5 = R_6.$$

$$\text{Thus } V_{13} - V_{10} = E = V_{in} - V_6.$$

Therefore the accessible electrode potential  $E = V_{13} - V_{10}$  ranges from  $-2.45$  to  $+2.5 \text{ V}$ . The compliance is increased compared to PassStat 2.0.

For the Teensy card, the correct command to send in order to apply the desired potential to the working electrode is calculated in the Python software. Similarly the corresponding currents are directly provided by the program.

#### Current reading

##### PassStat 1.0

Here  $V_{14} = R_6 \cdot I$  is the tension directly read by the oscilloscope.

##### PassStat 2.0

Here  $V_{14} = R_6 \cdot I + V_{13}$  is the tension directly read by the A0 analog input of the Teensy.  $V_{14}$  is limited by the supply voltages thus ranges between 0 and 3.3 V. If  $R_6 \cdot I$  would be too large or too low, the signal would then simply saturate.

##### PassStat 2.1 and 2.2

$V_{14} = R_6 \cdot I + V_{13}$  as above, but here may range between 0 and 5 V. A0 input should be inferior to 3.3 V otherwise the card would be damaged. Hence, we used an additional voltage divider (resistors  $R_7$  and  $R_8$  to limit the maximum tension to 3.3 V.

#### Design files

Software and firmware: Python and android studio are open source softwares.

#### Design files summary

Design file name	File type	Open source license	Location of the file
PassStatDesign.zip	CAD files of the circuit	Creative commons v4	<a href="https://zenodo.org/record/5719382#.YZwJ17rjJhE">https://zenodo.org/record/5719382#.YZwJ17rjJhE</a>
Arduino files	Arduino files to be loaded in the Teensy 3.2 or 3.6 card	GNU GPL v3	<a href="https://zenodo.org/record/5719382#.YZwJ17rjJhE">https://zenodo.org/record/5719382#.YZwJ17rjJhE</a>
Software Python		GNU GPL v3	<a href="https://zenodo.org/record/5719382#.YZwJ17rjJhE">https://zenodo.org/record/5719382#.YZwJ17rjJhE</a>
Software Android		GNU GPL v3	<a href="https://zenodo.org/record/5719382#.YZwJ17rjJhE">https://zenodo.org/record/5719382#.YZwJ17rjJhE</a>
Android app		GNU GPL v3	<a href="https://zenodo.org/record/5719382#.YZwJ17rjJhE">https://zenodo.org/record/5719382#.YZwJ17rjJhE</a>

This design file contains all the necessary files to implement or modify the potentiostat design with the open source software KiCad.

*Bill of materials*

**Bill of materials summary**

Designator	Component	Number	Cost per unit - currency	Total cost - currency	Source of materials	Material type
Capacitor 1 $\mu$ F	C1, C2	2	0.20 €	0.40 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Electrolytic
Capacitor 47 $\mu$ F	C3, C4	2	0.20 €	0.40 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Electrolytic
Capacitor 15 pF	CF1, CF2	2	0.10 €	0.20 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Ceramic
Capacitor 1 nF	CF3	1	0.10 €	0.10 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Ceramic
Connector 01x03	J1, J2	2	0.05 €	0.10 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Metal
Connector 01x03	J3, J4	2	0.05 €	0.10 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Metal
Jumper	JP1, JP2	2	0.05 €	0.10 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Metal
Jumper	JP3, JP4	2	0.05 €	0.10 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Metal
Resistor 10 k $\Omega$	R1, R3, R4, R5	4	0.10 €	0.40 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Composite
Resistor 15 k $\Omega$	R2	1	0.10 €	0.10 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Composite
Resistor 1 k $\Omega$	R8, R9, R10, R12	4	0.10 €	0.40 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Composite
Resistor 1.5 k $\Omega$	R11	1	0.10 €	0.10 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Composite
Resistor 1 M $\Omega$	R6	1	0.10 €	0.10 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Composite
Resistor 510 $\Omega$	R7	1	0.10 €	0.10 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Composite
Quad CMOS Rail-to-Rail Input and Output Operational Amplifier, DIP-14/SOIC-14	U1	1	3.21 €	3.21 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Semi-conductor
Dual In Line Support IC Sockets	U1	1	0.273 €	0.273 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Composite
		1 row of 20	0.742 €	0.742 €	<a href="https://fr.rs-online.com/web/">https://fr.rs-online.com/web/</a>	Composite
Printed Circuit Board		1	4.30 € (price for 20)	4.30 €	<a href="https://www.eurocircuits.com/">https://www.eurocircuits.com/</a>	Polymer
Teensy 3.2		1	26.39 €	26.39 €	<a href="https://www.lextronic.fr/">https://www.lextronic.fr/</a>	Hardware
Teensy 3.6		1	39.60 €	39.60 €	<a href="https://www.lextronic.fr/">https://www.lextronic.fr/</a>	Hardware
Analog Discovery 2		1	294 €	294 €	<a href="https://www.lextronic.fr/">https://www.lextronic.fr/</a>	Hardware
Bluetooth Module HC05		1	16.90 €	16.90 €	<a href="https://www.lextronic.fr/">https://www.lextronic.fr/</a>	Hardware
$\mu$ USB cable		1	3.00 €	3.00 €	<a href="https://www.lextronic.fr/">https://www.lextronic.fr/</a>	Cable
External Battery NX 5000 mAh (cheapest models will also work)		1	18.70 €	18.70 €	<a href="https://www.1001piles.com/lithium-ion-batterie-externe-universelle-5v-5000mah-103724.html">https://www.1001piles.com/lithium-ion-batterie-externe-universelle-5v-5000mah-103724.html</a>	Hardware

We advise to use DIL support for the quad amplifier but this is not mandatory. IC sockets for R6 and CF<sub>3</sub> should conversely be used since these components need to be adjusted depending on the electrode size and concentration of electroactive compound to be analyzed.

The cheapest prize with a Teensy 3.2 is 40 €. A system working with Bluetooth module and Teensy 3.6 costs 65€ + price of external battery. For fast scan voltammetry with Analog discovery 2 the price is 305 €, which is still much lower than commercial systems.

**Build instructions**

The electronic card is presented in Fig. 4. Jumpers JP1 and JP2 are present to switch between different configurations, and bypass OA1 for PassStat 1.0 and 2.0. They can be replaced by soldered shortcuts if a single configuration is used. Figs. 5–7 present pictures of the different potentiostat implementations (See Fig. 8.).



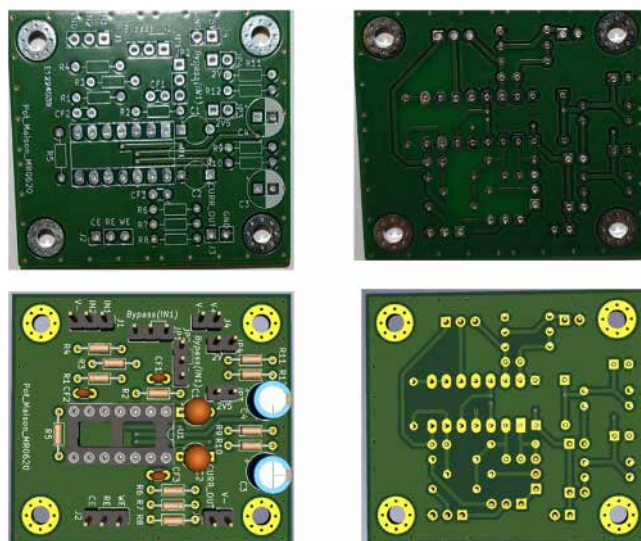


Fig. 4. Electronic card used to realize the different configurations and view extracted from KiCAD.

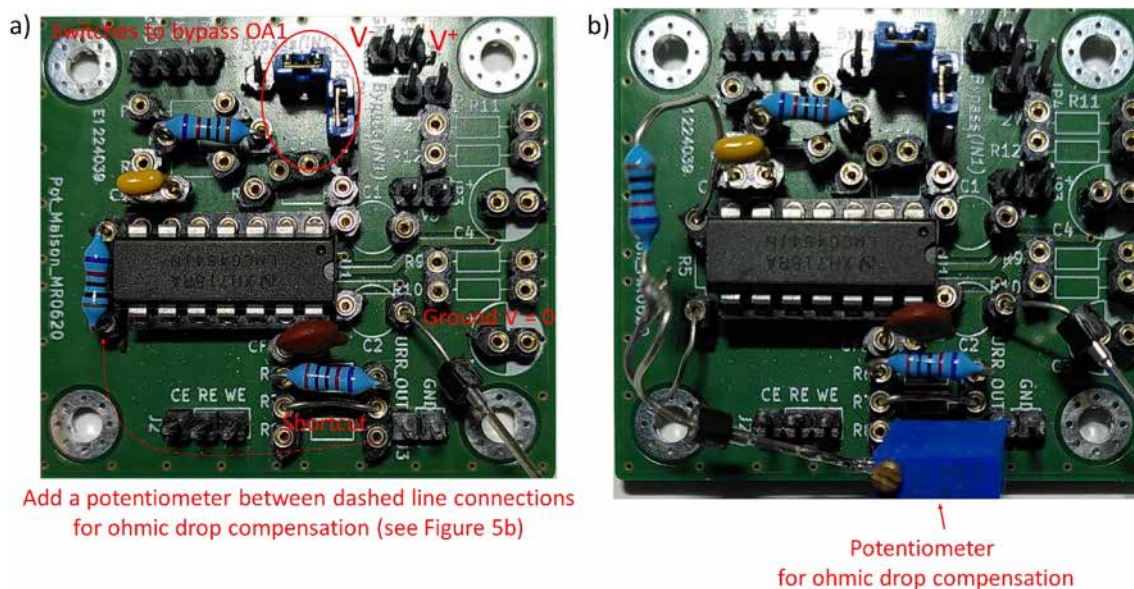


Fig. 5. PassStat 1.0 without (a) and with (b) ohmic drop compensation. For (b) a lead was soldered to  $R_5$  and positive feedback was applied thanks to a 200 k $\Omega$  potentiometer (this value should be adapted for different conditions, see calculations above).

**Operation instructions**

A first program should first be implanted in the Teensy 3.6 card. A computer is necessary for this step. This program contains both CV or SWV methods. When the Teensy is powered, the program waits for parameters arriving on the serial port (methods, potentials, scan rate etc). The orders may be sent either by a computer or by an Android smartphone. For each connection a specific program is provided: CV\_SWV.ino for computer and swv\_cv\_bluetooth.ino for Android Connection with a smartphone is lower (9600 bauds) than with a computer (115200 bauds). The interface was programmed in Python for computer control and with Android Studio for a smartphone.

Should Teensy 3.2 be used instead of Teensy 3.6, A22 should be replaced by A14.



Fig. 6. PassStat 2.0. Ground is deported to 1.65 V thanks to resistors  $R_9$  and  $R_{10}$ .

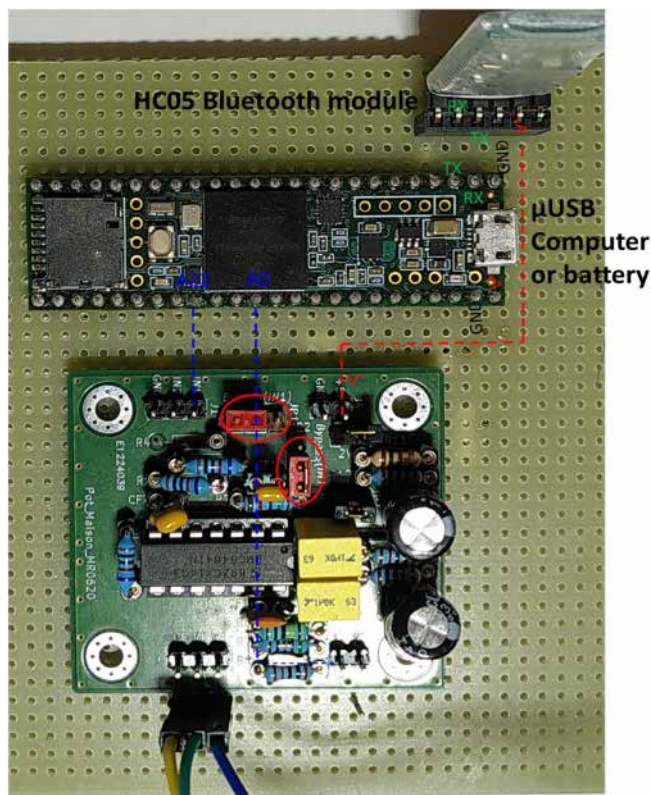
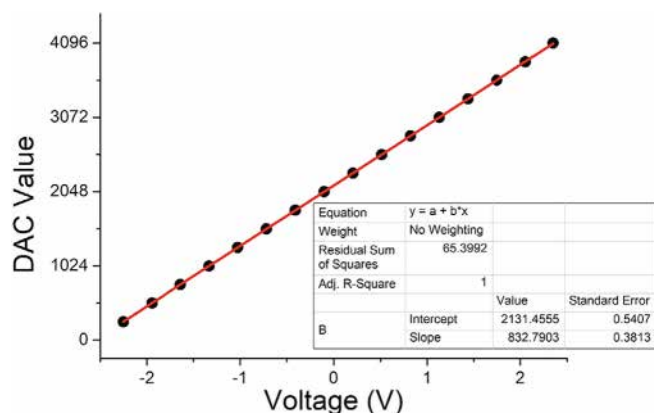


Fig. 7. PassStat 2.2 mounted on a PCB. Power is supplied at the  $\mu$ USB port by a battery to supply the Teensy 3.6 card and the HC05 module. For PassStat 2.1, HC05 is not connected and connection is made directly from a computer.

### Computer control

The Python program is organised with three subprograms. `swv_cyclique.py` should be started by the operator. The port number (that can be read in Arduino program) should be entered prior choosing the method. Then if CV is chosen subprogram `volta.py` is activated whereas `swv.py` is activated for SWV. The potentiostat can be controlled from PC under Windows 7



**Fig. 8.** Calibration curve obtained by measuring the tension applied to a 100 k $\Omega$  resistor with  $R_6 = 100$  k $\Omega$ . DAC = 2131 correspond to an electrode potential of 0 V. It can be entered directly in the program together with the slope 832.8. 12 bits resolution was used here.

or Windows 10, Macintosh or Linux computers. The port board definition however should be defined in the three python programs and this definition should be adapted for each operating system. We deliver the PC version here. To adapt to Linux or Macintosh, activate the correct line by removing the comment symbol (#) and comment the other ones. We provide below an example for the main `swv_cyclique` program and for PC:

```
Line 32: #PORT_BOARD = "/dev/cu.usbmodem" # for MAC CPU.
Line 33: self.PORT_BOARD = "COM" # for PC CPU.
Line 34: #self.PORT_BOARD="/dev/ttyACM" # for linux.
```

See lines 23–25 for [volta.py](#) and lines 31–33 for [swv.py](#).

We installed Anaconda 3 with Spyder 4 but any Python environment should be operative. Note that the libraries `time`, `math` and `pyserial` should be installed in Python (command `pip install pyserial`).

#### Android smartphone control

The program `Potentiostat.apk` can be installed on any Android smartphone that operates with Android 4 or superior (ignore security warnings). It was developed with Android Studio that allows to create an application. Datas are saved in `android/datas/fr.Achit.lecourantpasse`. The code itself is contained in the 4 `MainActivity` files that interact together. Presentation is handled in the `.xml` files. The application should be operative on most screens. Should the reader modify the codes, a new app can be generated in the build tab. The first activity handles the choice of method (as above). The second one concerns CV and the third one SWV. The fourth one displays the voltamogram.

#### Calibration

All the analytical formulations for the different configurations supposed an USB voltage of 5.00 V, but we observed that the effective USB voltage changes from one computer to another. Such variations induce shift on the positive input of the OAs and then on the applied potential and measured current. This does not prevent to perform experiments, but as for any apparatus a calibration procedure is necessary if accurate measurements are desired. For that, a resistor equal to  $R_6$  should be used. The specific Arduino `Teensy_DAC_ADC` program is provided as SI for that purpose.

#### Potential calibration

Here, a correspondence should be established between the digital/analogic scale (from 0 to 4095 for a 12 bits operation) and the applied tension between working and reference electrode (ideally from  $-2.5$  to  $+2.5$  V but in practice a smaller range). For that, DAC values should be sent and the applied tension should be measured with a high impedance voltmeter. A straight line should be observed excepted at the extremities of the potential inversion.

We provide the program `Teensy_DAC_ADC` that allows to send a DAC value to pin A22 and to enter the tension read by the voltmeter. The line provided in Figure S5 was traced by recording the tension by steps of 250 on the DAC from 250 to 4250. The slope and intercept should be entered in the Python and Android programs as explained below.

#### Current calibration

The DAC value corresponding to the intercept should be entered in the program. Check that the tension applied to the calibration resistor is  $0 \pm 0.002$  V. The current is thus 0. Note the ADC value and enter this value in the line that contains `OFFSET_ADC_TEENSY` (see below).



To check the calibration procedure, run a CV between  $-1$  and  $+1$  V on a resistor equal to  $R_6$ . A symmetric straight line should be observed.

This procedure is applicable for 12 bits or 16 bits operation (in this case 4095 should be replaced by 65535).

The following values should be changed in the `volta.py` and `swv.py` programs.

```
self.QUANT_DAC_TEENSY = 1./832.79 # extracted from calibration process.
self.OFFSET_DAC_TEENSY = 2131.45 # extracted from calibration process.
self.QUANT_ADC_TEENSY = 3.3/4095.
self.OFFSET_ADC_TEENSY = 2070.0.
self.COEFF_CONV_TEENSY = 1.51.
```

The following values should be changed in the Android studio first three activities:

```
final double QUANT_DAC_TEENSY=(1/832.79);
final double gain = -1;
final double OFFSET_DAC_TEENSY = 2131.45;
final double OFFSET_ADC_TEENSY = 2070.0;
```

## Validation and characterization

Since PassStat 1.0 works with an arbitrary function generator and an oscilloscope, all potentiostatic techniques may be implemented. For PassStat 2, the present software version includes cyclic voltammetry and square wave voltammetry. Additional techniques will be implemented in the future.

### Materials and methods

The counter electrodes were platinum wires. The reference electrode was either a home-made AgCl/Ag electrode, or a platinum wire (for low currents or high speed). The home-made working electrodes were either a 0.25 mm radius Pt disk (standard conditions), a 9B pencil lead from Cretacolor (paracetamol), a 2  $\mu\text{m}$  radius Pt disk (low currents) or a 15,200  $\mu\text{m}^2$  gold ball.

Ferrocene (Aldrich), tetrabutylammonium hexafluorophosphate (Alfa Aesar), Citric acid (Aldrich) and acetonitrile (VWR) were used without further purification. Doliprane 1000 mg tablets from Sanofi were used to prepare a 1 mM paracetamol solution with 0.100 M citric acid as supporting electrolyte.

Analogue Discovery 2 from Digilent was used for PassStat 1.0.

The Teensy 3.6 acquisition card and HC05 Bluetooth module were purchased from Lextronic.

LMC 6484 operational amplifier and other electronic components were all purchased from radiospares.

The circuit was designed with KiCad 5.1.0, an open source software for electronic design. The circuit may also be implemented onto test breadboards.

### Test on a dummy cell

To probe the electronic system prior to perform real electrochemical experiments or to identify a problem in case of failure, it may be useful to use a test circuit called a dummy cell. Here we are using  $R = 10$  k $\Omega$  in series with  $C = 1$   $\mu\text{F}$  and a scan rate of 2  $\text{Vs}^{-1}$ . Counter and reference electrodes are connected at the same place (shortcircuit). At the potential start or inversion, exponential variations are observed up to plateaus for which the capacitive current is:  $i_c = C v$  as represented in Fig. 9.

### Classical electrochemical conditions.

Fig. 10 presents cyclic voltammograms acquired in the range 0.05–100  $\text{Vs}^{-1}$  using a 0.25 mm radius platinum electrode with PassStat 2.1 for a 1 mM ferrocene solution in acetonitrile, in presence of 0.10 M tetrabutylammonium hexafluorophosphate (TBAPF<sub>6</sub>) as supporting electrolyte.  $C_F$  was 1 nF in all cases.  $R_6$  was 1 M $\Omega$  from 0.05 to 5  $\text{Vs}^{-1}$  and 100 k $\Omega$  above 5  $\text{Vs}^{-1}$ . Plotting peak current against  $v^{1/2}$  displays a good linearity as expected for purely diffusive behavior (see Fig. 10). A deviation is however observed at 100  $\text{Vs}^{-1}$  because at this scan rate the signal is altered both by ohmic drop effects and by the too high value of the  $R_6 C_F$  low pass filter value (0.1 ms with  $R_6 = 100$  k $\Omega$  and  $C_F = 1$  nF, see section 3.3 for optimization at important  $v$ ).

In the same conditions, a square wave voltammogram was acquired at an equivalent scan rate of 0.1  $\text{Vs}^{-1}$ , potential steps of 20 mV and potential increment of 3 mV (see Fig. 11). A very smooth and well-resolved curve is observed, validating the potential of the device towards analytical purposes.

In addition, to demonstrate the efficiency of the device for low cost experiments towards identification of potential fake drug analysis, we analyzed an aqueous solution prepared from a paracetamol tablet with a 9B pencil lead as working electrode. Various pencil lead electrodes have indeed been proposed as cheap electrode material in view of electroanalysis [33–35]. Here, paracetamol concentration was 1 mM and we choose 0.10 M citric acid as supporting electrolyte. Many different conditions are described in the literature [15–24], and we found that an acid medium provided a better reproducibility. Moreover, citric acid can be purchased easily so that preparing this electrolyte may be easier than other

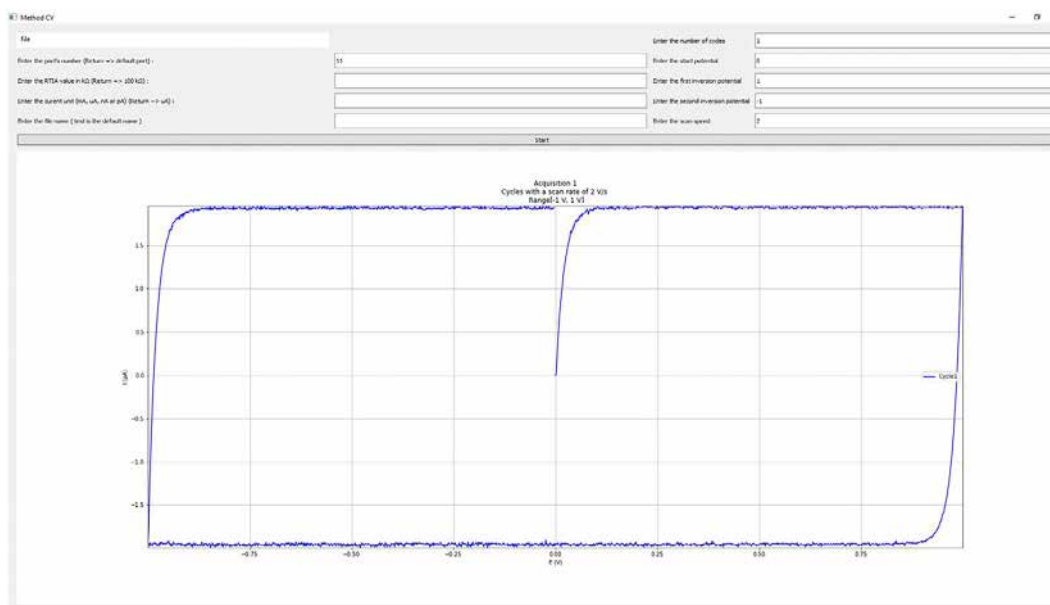


Fig. 9. Voltammogram obtained at  $2 \text{ Vs}^{-1}$  for a dummy cell with  $R = 10 \text{ k}\Omega$  and  $C = 1 \text{ }\mu\text{F}$ . Screenshot of the software window.

classical buffers in non-ideal experimental conditions. This experiment was performed thanks to remote control with a smartphone. Fig. 12 confirms that no additional noise is added in these conditions.

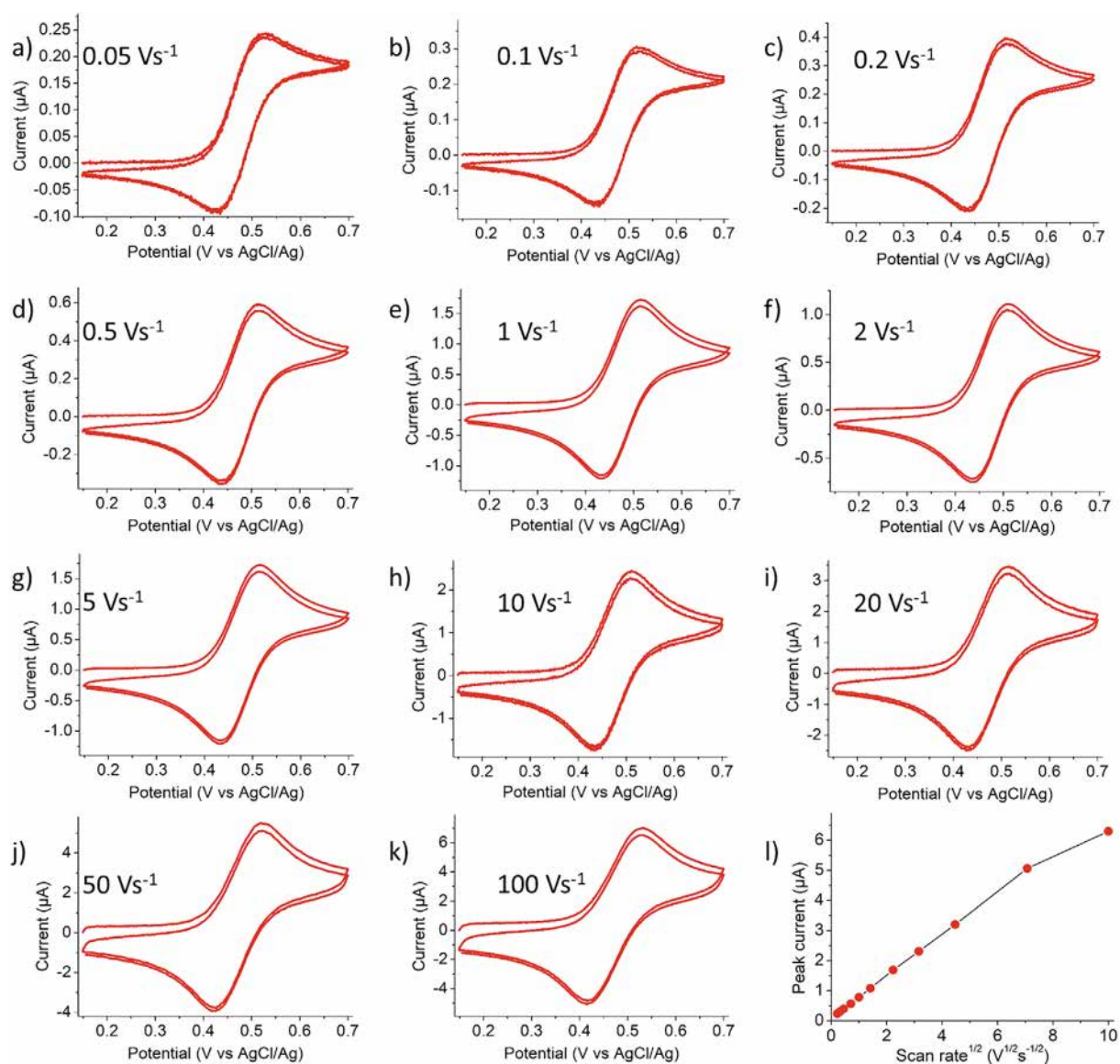
### Detecting low currents

When using ultramicroelectrodes (UMEs), *i.e.* electrodes of micrometric or nanometric dimensions, the CV shifts from transient to steady state at low scan rates. Ultramicroelectrodes are useful to increase signal/noise ratio in analytical chemistry. Additionally, UMEs allow to work with low amounts of supporting electrolyte [28]. Hence, we resorted here onto a solution containing only  $25 \text{ }\mu\text{M}$  of ferrocene and  $2.5 \text{ mM}$  of TBAPF<sub>6</sub> in CH<sub>3</sub>CN. Since very low currents are then concerned, resistor  $R_6$  should be increased to  $1 \text{ G}\Omega$  here. Fig. 13 presents the cyclic voltammogram obtained at  $0.01 \text{ Vs}^{-1}$  for a  $2 \text{ }\mu\text{m}$  radius platinum electrode with the PassStat 2.1 configuration for which  $R_6$  was set to  $1 \text{ G}\Omega$  and  $CF_3$  to  $100 \text{ pF}$ . A current plateau of only  $72 \text{ pA}$  is observed. In these conditions, the RMS noise level was found to be less than  $0.5 \text{ pA}$  as shown in the inset of Fig. 13. The current quantification step with these parameters is only  $0.075 \text{ pA}$  with a 16 bits resolution, a value only slightly slower than the noise level. This second experiment demonstrates that even if in most of the case a 12 bits resolution is sufficient, this is not the case in such extreme conditions.

Finally, we underline that here  $R_6CF_3 = 100 \text{ ms}$  so that the temporal resolution is still excellent.  $CF_3$  may be further diminished if dynamic events such as impacts of biological exocytotic release should be caught. There is moreover still room for optimization, notably by using PassStat 2.0 configuration and/or oversampling the data.

### Fast scan voltammetry

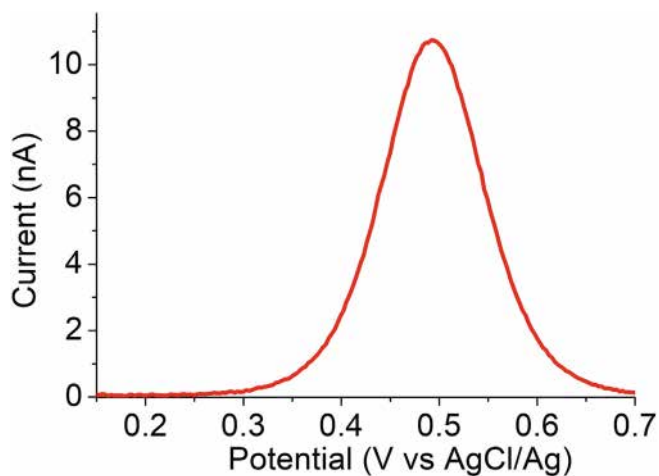
Another interesting property of UMEs is to give access to low time scales (nanoseconds in the best conditions) [32]. We evaluated the PassStat 1.0 configuration in the fast scan range with a gold ball ultramicroelectrode produced by melting a  $12.5 \text{ }\mu\text{m}$  radius gold wire. The electrode area was estimated to be  $15200 \text{ }\mu\text{m}^2$  in a calibration procedure. In the black CV of Fig. 14, realized for a  $1 \text{ mM}$  solution of ferrocene in acetonitrile containing  $0.10 \text{ M}$  TBAPF<sub>6</sub>, the capacitive plateau is not attained immediately because of ohmic drop within the electrochemical cell at  $8000 \text{ Vs}^{-1}$ . Even if UMEs drastically reduced ohmic losses, the large current densities pertaining to large  $v$  finally alter the signal. To face this problem, ohmic drop compensation was applied by adding a positive feedback with the potentiometer  $R_{\text{Pos}}$  displayed in red in Fig. 5b. At 100% compensation, *i.e.* when the equivalent feedback resistor equals the solution one, the red curve shows oscillations at the limits of the potential ramp. The current peaks are then no more distorted by the ohmic drop, as long as positive feedback occurs quickly enough. Our previous reports explain in detail interactions between electrode size, electrolyte composition and electronic set-up so as to carry out ohmic drop compensation with a maximum accuracy.



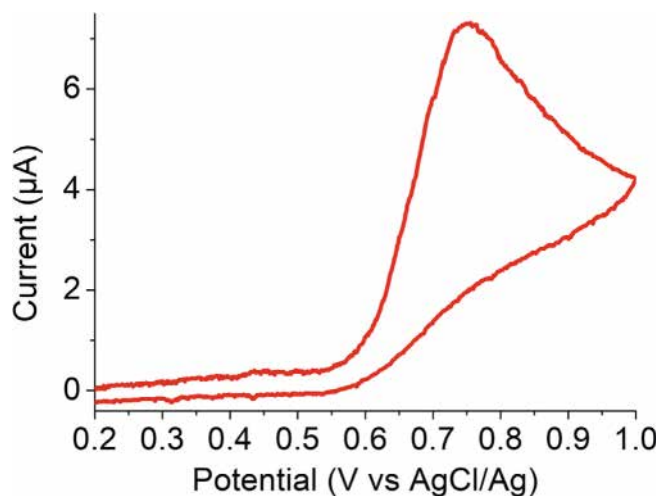
**Fig. 10.** A–k) Cyclic voltammogram obtained at different scan rates for a 0.25 mm radius Pt electrode in acetonitrile containing 1 mM ferrocene and 0.10 M TBAPF<sub>6</sub> with PassStat 2.1. l) Peak potential versus scan rate.

### Conclusion and perspectives

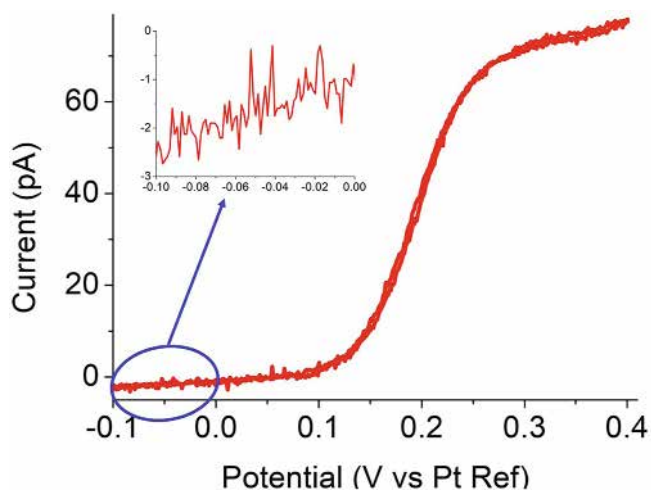
We presented above several possible schemes to build a low cost but powerful potentiostat for analytical electrochemistry. The electronic board is easy to implement and repair, and can be realized at a low cost. The simplicity of the scheme stems from the use of a quad amplifier component. The python and android interfaces are fully open source. Only a minimal training is necessary to use this potentiostat, and electrochemical performances are close to those of commercial potentiostats so that it could be suitable for measurements at a research level [36]. Should one need to adapt the design to specific needs additional elements could be easily implemented. Those could be independent power supplies precisely regulated to apply voltages independent from the USB or battery ones, additional filters and amplification stages or specific faster amplifiers [11,31]. Future implementations could add a bipotentiostat control for measurements with two working electrodes and electrochemical impedance spectroscopy [37,38]. The Analog Discovery 2 or Teensy card are furthermore useful to add further possibilities, for example control of a rotating disk electrode or couple electrochemistry with other techniques such as spectroscopic ones or pH sensing for example [39]. We invite the reader to check software and hardware updates. The present work is thus the first stone to develop other electrochemical applications for example analytical measurements on the ground [40].



**Fig. 11.** Square wave voltammogram obtained at  $0.1 \text{ Vs}^{-1}$  for a 0.25 mm radius Pt electrode in acetonitrile containing 1 mM ferrocene and 0.10 M TBAPF<sub>6</sub> with PassStat 2.1.  $R_6 = 1 \text{ M}\Omega$  and  $CF_3 = 1 \text{ nF}$ .

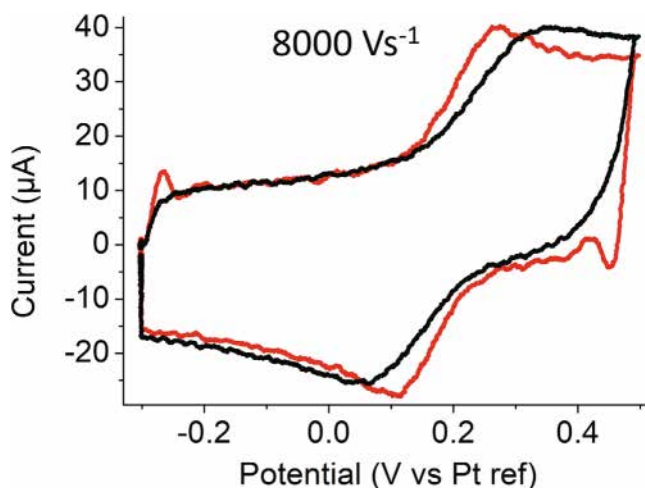


**Fig. 12.** Cyclic voltammogram obtained on a 9B pencil lead at  $0.1 \text{ Vs}^{-1}$  in an aqueous solution containing 0.10 M citric acid as supporting electrolyte and 1 mM paracetamol originating from a pharmaceutical tablet with PassStat 2.2 (smartphone application).  $R_6 = 100 \text{ k}\Omega$ ,  $CF_3 = 1 \text{ nF}$ .



**Fig. 13.** Cyclic voltammogram obtained with a  $2 \mu\text{m}$  radius Pt ultramicroelectrode in an acetonitrile solution containing 25  $\mu\text{M}$  of ferrocene and 2.5 mM tetrabutylammonium hexafluorophosphate at  $10 \text{ mVs}^{-1}$  with PassStat 2.1.  $R_6 = 1 \text{ G}\Omega$ ,  $CF_3 = 100 \text{ pF}$ .





**Fig. 14.** Cyclic voltammogram obtained with a gold ball ultramicroelectrode in an acetonitrile solution containing 1 mM of ferrocene and 0.10 M tetrabutylammonium hexafluorophosphate at  $8000 \text{ Vs}^{-1}$  without (black) and with (red) ohmic drop compensation.  $R_6 = 10 \text{ k}\Omega$ ,  $CF_3 = 3.3 \text{ pF}$ . (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

### CRediT authorship contribution statement

**Mélícia Caux:** Investigation. **Anis Achit:** Software. **Kethsovann Var:** Investigation. **Gabriel Boitel-Aullen:** Investigation. **Daniel Rose:** Conceptualization. **Agnès Aubouy:** review and editing. **Sylvain Argentieri:** Review and editing. **Raymond Campagnolo:** Conceptualization. **Emmanuel Maisonhaute:** Conceptualization, Supervision, writing, review and editing.

### Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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### References

- [1] R.G. Compton, C.E. Banks, *Understanding Voltammetry*, Imperial College Press, 2010, 10.1142/p726.
- [2] A. Nemiroski, D.C. Christodouleas, J.W. Hennek, A.A. Kumar, E.J. Maxwell, M.T. Fernández-Abedul, G.M. Whitesides, Universal mobile electrochemical detector designed for use in resource-limited applications, *PNAS* 111 (33) (2014) 11984–11989, <https://doi.org/10.1073/pnas.1405679111>.
- [3] A. Ainla, M.P.S. Mousavi, M.-N. Tsaloglou, J. Redston, J.G. Bell, M.T. Fernández-Abedul, G.M. Whitesides, Open-source potentiostat for wireless electrochemical detection with smartphones, *Anal. Chem.* 90 (10) (2018) 6240–6246, <https://doi.org/10.1021/acs.analchem.8b00850>.
- [4] Y.C. Li, E.L. Melenbrink, G.J. Cordonier, C. Boggs, A. Khan, M.K. Isaac, L.K. Nkhonjera, D. Bahati, S.J. Billinge, S.M. Haile, R.A. Kreuter, R.M. Crable, T.E. Mallouk, An easily fabricated low-cost potentiostat coupled with user-friendly software for introducing students to electrochemical reactions and electroanalytical techniques, *J. Chem. Educ.* 95 (9) (2018) 1658–1661, <https://doi.org/10.1021/acs.jchemed.8b00340>.
- [5] A.A. Rowe, A.J. Bonham, R.J. White, M.P. Zimmer, R.J. Yadgar, T.M. Hobza, J.W. Honea, I. Ben-Yaacov, K.W. Plaxco, M. Wanunu, CheapStat: An open-source, “do-it-yourself” potentiostat for analytical and educational applications, *PLoS ONE* 6 (9) (2011) e23783, <https://doi.org/10.1371/journal.pone.0023783>.
- [6] P. Irving, R. Cecil, M.Z. Yates, MYSTAT: a compact potentiostat/galvanostat for general electrochemistry measurements, *HardwareX*. 9 (2021) e00163, <https://doi.org/10.1016/j.ohx.2020.e00163>.
- [7] C. Mercer, R. Bennett, P. Conghaile, J. Rusling, D. Leech, Glucose biosensor based on open-source wireless microfluidic potentiostat, *Sensors Actuators B-Chemical*. 290 (2019) 616–624, <https://doi.org/10.1016/j.snb.2019.02.031>.
- [8] G.N. Meloni, Building a microcontroller based potentiostat: a inexpensive and versatile platform for teaching electrochemistry and instrumentation, *J. Chem. Educ.* 93 (7) (2016) 1320–1322, <https://doi.org/10.1021/acs.jchemed.5b00961>.
- [9] S.T. Rajendran, E. Scarano, M.H. Bergkamp, A.M. Capria, C.-H. Cheng, K. Sanger, G. Ferrari, L.H. Nielsen, E.-T. Hwu, K. Zór, A. Boisen, Modular, lightweight, wireless potentiostat-on-a-disc for electrochemical detection in centrifugal microfluidics, *Anal. Chem.* 91 (18) (2019) 11620–11628, <https://doi.org/10.1021/acs.analchem.9b02026>.
- [10] M.D.M. Dryden, A.R. Wheeler, D.T. Eddington, DStat: a versatile, open-source potentiostat for electroanalysis and integration, *PLoS ONE* 10 (10) (2015) e0140349, <https://doi.org/10.1371/journal.pone.0140349>.
- [11] Y. Matsubara, A small yet complete framework for a potentiostat, galvanostat, and electrochemical impedance spectrometer, *J. Chem. Educ.* 98 (10) (2021) 3362–3370, <https://doi.org/10.1021/acs.jchemed.1c00228>.
- [12] A.J. Bard, L.R. Faulkner, *Electrochemical Methods: Fundamentals and Applications*, Wiley India Ltd., New Delhi, 2004.
- [13] <https://iorodeo.com/collections/cheapstat-open-source-potentiostat>, (n.d.). <https://iorodeo.com/collections/cheapstat-open-source-potentiostat>.
- [14] M.W. Glasscott, M.D. Verber, J.R. Hall, A.D. Pendergast, C.J. McKinney, J.E. Dick, SweepStat: a build-it-yourself, two-electrode potentiostat for macroelectrode and ultramicroelectrode studies, *J. Chem. Educ.* 97 (1) (2020) 265–270, <https://doi.org/10.1021/acs.jchemed.9b00893>.

- [15] I. Baranowska, M. Koper, The preliminary studies of electrochemical behavior of paracetamol and its metabolites on glassy carbon electrode by voltammetric methods, *Electroanalysis* 21 (10) (2009) 1194–1199, <https://doi.org/10.1002/elan.200804536>.
- [16] S. Chitravathi, N. Munichandraiah, Voltammetric determination of paracetamol, tramadol and caffeine using poly(Nile blue) modified glassy carbon electrode, *J. Electroanal. Chem.* 764 (2016) 93–103, <https://doi.org/10.1016/j.jelechem.2016.01.021>.
- [17] H. Gorcay, G. Turkoglu, Y. Sahin, H. Berber, Electrochemical determination of paracetamol by a novel derivative of formazan modified pencil graphite electrode, *IEEE Sens. J.* 14 (8) (2014) 2529–2536, <https://doi.org/10.1109/JSEN.2014.2311296>.
- [18] N.T. Hart, W.C. Lane, L. de la Garza, Electrochemical quantification of acetaminophen: an engaging cyclic voltammetry laboratory for the quantitative analysis course, *J. Chem. Educ.* 97 (8) (2020) 2254–2259, <https://doi.org/10.1021/acs.jchemed.9b01127>.
- [19] A. Kassa, M. Amare, R.M. Mahfouz, Electrochemical determination of paracetamol, rutin and sulfonamide in pharmaceutical formulations by using glassy carbon electrode – a review, *Cogent Chem.* 5 (1) (2019) 1681607, <https://doi.org/10.1080/23312009.2019.1681607>.
- [20] B.R. Liyarita, A. Ambrosi, M. Pumera, 3D-printed electrodes for sensing of biologically active molecules, *Electroanalysis* 30 (7) (2018) 1319–1326, <https://doi.org/10.1002/elan.201700828>.
- [21] S.F. Mbokou, M. Pontié, J.-P. Bouchara, F.M.M. Tchieno, E. Njanja, A. Mogni, P.Y. Pontalier, I.K. Tonle, Electroanalytical performance of a carbon paste electrode modified by coffee husks for the quantification of acetaminophen in quality control of commercialized pharmaceutical tablets, *Int. J. Electrochem.* 2016 (2016) 1–10, <https://doi.org/10.1155/2016/1953278>.
- [22] D.J. Miner, J.R. Rice, R.M. Riggan, P.T. Kissinger, Voltammetry of acetaminophen and its metabolites, *Anal. Chem.* 53 (14) (1981) 2258–2263, <https://doi.org/10.1021/ac00237a029>.
- [23] H. Montaseri, P. Forbes, Analytical techniques for the determination of acetaminophen: a review, *TRAC-Trends Anal. Chem.* 108 (2018) 122–134, <https://doi.org/10.1016/j.trac.2018.08.023>.
- [24] B. Suchacz, M. Wesolowski, Voltammetric quantitation of acetaminophen in tablets using solid graphite electrodes, *Anal. Methods* 8 (16) (2016) 3307–3315.
- [25] I. Navarro, D. Gonzalez-Arjona, E. Roldan, M. Rueda, Determination of paracetamol in tablets and blood plasma by differential pulse voltammetry, *J. Pharm. Biomed. Anal.* 6 (6–8) (1988) 969–976, [https://doi.org/10.1016/0731-7085\(88\)80118-3](https://doi.org/10.1016/0731-7085(88)80118-3).
- [26] (n.d.), <https://www.youtube.com/playlist?list=PL4-5Rjd2oUwrvTe3DuxuikS14xa0ijpy>.
- [27] C. Amatore, E. Maisonhaute, G. Simonneau, Ohmic drop compensation in cyclic voltammetry at scan rates in the megavolt per second range: access to nanometric diffusion layers via transient electrochemistry, *J. Electroanal. Chem.* 486 (2) (2000) 141–155, [https://doi.org/10.1016/S0022-0728\(00\)00131-5](https://doi.org/10.1016/S0022-0728(00)00131-5).
- [28] C. Amatore, S. Arbault, E. Maisonhaute, S. Szunerits, L. Thouin, Electrochemistry at ultramicroelectrodes: Small and fast may be useful, in *Trends In Molecular Electrochemistry*, eds J. L. Pombeyro and C. Amatore, Fontis Media, Lausanne. (2004) 385–411.
- [29] C. Amatore, Y. Bouret, E. Maisonhaute, H.D. Abruña, J.I. Goldsmith, Electrochemistry within molecules using ultrafast cyclic voltammetry, *C. R. Chim.* 6 (1) (2003) 99–115, [https://doi.org/10.1016/S1631-0748\(03\)00019-5](https://doi.org/10.1016/S1631-0748(03)00019-5).
- [30] C. Amatore, S. Arbault, E. Maisonhaute, S. Szunerits, L. Thouin, Electrochemistry at ultramicroelectrodes: Small and fast may be useful, in: A. Pombeyro, C. Amatore (Eds.), 2004: pp. 385–411.
- [31] C. Amatore, C. Lefrou, F. Pflüger, Online compensation of ohmic drop in submicrosecond time resolved cyclic voltammetry at ultramicroelectrodes, *J. Electroanal. Chem.* 270 (1–2) (1989) 43–59, [https://doi.org/10.1016/0022-0728\(89\)85027-2](https://doi.org/10.1016/0022-0728(89)85027-2).
- [32] C. Amatore, E. Maisonhaute, When voltammetry reaches nanoseconds, *Anal. Chem.* 77 (15) (2005) 303 A–311 A, <https://doi.org/10.1021/ac053430m>.
- [33] N. Vishnu, M. Gandhi, S. Badhulika, A.S. Kumar, Tea quality testing using 6B pencil lead as an electrochemical sensor, *Anal. Methods* 10 (20) (2018) 2327–2336.
- [34] A. Izadyar, D. Arachchige, H. Cornwell, J. Hershberger, Ion transfer stripping voltammetry for the detection of nanomolar levels of fluoxetine, citalopram, and sertraline in tap and river water samples, *Sensors Actuators B-Chemical.* 223 (2016) 226–233, <https://doi.org/10.1016/j.snb.2015.09.048>.
- [35] W. Sim, N. Kutrakul, P. Khunkaewla, A. Schulte, Three-electrode 30–60  $\mu$  L mini-cell for ecologically conscious analytical voltammetry with common macro- and microelectrodes, *ACS Sustainable Chem. Eng.* 8 (2020) 5082–5090, <https://doi.org/10.1021/acssuschemeng.9b07034>.
- [36] C. Amatore, D. Genovese, E. Maisonhaute, N. Raouafi, B. Schöllhorn, Electrochemically driven release of picomole amounts of calcium ions with temporal and spatial resolution, *Angew. Chem. Int. Ed.* 47 (28) (2008) 5211–5214, <https://doi.org/10.1002/anie.200705274>.
- [37] T. Touzalin, S. Joiret, I.T. Lucas, E. Maisonhaute, Electrochemical tip-enhanced Raman spectroscopy imaging with 8 nm lateral resolution, *Electrochem. Commun.* 108 (2019), <https://doi.org/10.1016/j.elecom.2019.106557>.
- [38] G. Boitel-Aullen, L. Fillaud, F. Huet, I. Nierengarten, B. Delavaux-Nicot, J.-F. Nierengarten, E. Maisonhaute, Electron transfer inside a decaferrocenylated rotaxane analyzed by fast scan cyclic voltammetry and impedance spectroscopy, *Chemelectrochem* 8 (18) (2021) 3506–3511, <https://doi.org/10.1002/celec.202100738>.
- [39] T. Touzalin, S. Joiret, E. Maisonhaute, I.T. Lucas, Complex electron transfer pathway at a microelectrode captured by in situ nanospectroscopy, *Anal. Chem.* 89 (17) (2017) 8974–8980, <https://doi.org/10.1021/acs.analchem.7b01542>.
- [40] L. Pujol, D. Evrard, K. Groenen-Serrano, M. Freyssinier, A. Ruffien-Cizsak, P. Gros, Electrochemical sensors and devices for heavy metals assay in water: the French groups' contribution, *Front. Chem.* 2 (2014), <https://doi.org/10.3389/fchem.2014.00019>.



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