Electrochemical impedance of electrodiffusion in charged medium under dc bias

Juhyun Song, Edwin Khoo, and Martin Z. Bazant

1 Department of Chemical Engineering, Massachusetts Institute of Technology, 77 Massachusetts Avenue, Cambridge, Massachusetts 02139, USA
2 Department of Mathematics, Massachusetts Institute of Technology, 77 Massachusetts Avenue, Cambridge, Massachusetts 02139, USA

(Received 14 May 2019; published 4 October 2019)

An immobile charged species provides a charged medium for transport of charge carriers that is exploited in many applications, such as permselective membranes, doped semiconductors, biological ion channels, as well as porous media and microchannels with surface charges. In this paper, we theoretically study the electrochemical impedance of electrodiffusion in a charged medium by employing the Nernst-Planck equation and the electroneutrality condition with a background charge density. The impedance response is obtained under different dc bias conditions extending above the diffusion-limiting bias. We find a transition in the impedance behavior around the diffusion-limiting bias and present an analytical approximation for a weakly charged medium under an overlimiting bias.

DOI: 10.1103/PhysRevE.100.042204

I. INTRODUCTION

Transport of charged species relative to the surrounding medium is driven by gradients in their concentrations (i.e., diffusion) and in electric potential (i.e., migration or drift), which is therefore called electrodiffusion. The Nernst-Planck equation [1,2], also known as the drift-diffusion or the diffusion-migration equation, has successfully described electrodiffusion in various fields, including batteries [3], fuel cells [4], semiconductors [5], solar cells [6], ion-exchange membranes [7–10], as well as biological systems [11–13], although its theoretical validity is restricted to dilute systems [14–16]. It is solved either under the electroneutrality condition in the thin-double-layer limit [17–19], or more generally, under the electric potential that is determined self-consistently from the charge density via the Poisson equation governing electrostatics [16,20,21]. In the latter case, the model is called the Poisson-Nernst-Planck (PNP) set of equations, whose steady-state solution leads to the Poisson-Boltzmann distribution and the Gouy-Chapman double-layer model [22]. Recently, its transient response was thoroughly reviewed by Bazant and colleagues [16]. Its impedance response has also been studied extensively by Macdonald and Franceschetti [19,21,23–27], Buck and Brumley [28–30], Jannik and Maier [31–33], Moya and Horno [34–36], and others [37–39] for various boundary conditions. When there is an excess amount of supporting electrolyte or when there is a large disparity in the mobilities of the charge carriers, the electrodiffusion models reduce to the neutral diffusion equation, or Fick’s law, exhibiting Warburg behavior in its impedance response [19,40,41].

In many applications, there can be an immobile charged species that electrostatically interacts with the charge carriers. For example, in doped semiconductors, the dopants become ionized and provide immobile charges, either positive if the dopants are donors or negative if acceptors [5]. Permselective membranes, often used in desalination and chemical separations, typically have charged functional groups attached onto the polymer backbone chain [10,15,34,42,43]. In addition, ion channels in cell membranes are proteins that have a pore structure when open, whose inner surface consists of ionized groups [11–13,44,45]. The immobile charged species effectively provides a charged medium for the transport of charge carriers. Therefore, as suggested by Teorell, Meyer, and Sievers [8–10] for membranes, the immobile charge species can be considered as a background charge density that is added to the charge density of the mobile charge carriers either in the electroneutrality condition or the Poisson equation. Even for microchannels or porous media where the majority of charge carriers are out of the double layer on their local surface, it was recently shown that the surface charge can be treated as a homogenized background charge density in upscaled macroscopic transport models [46].

The electrical response of electrodiffusion in the presence of an immobile charged species has been studied for the aforementioned applications in various setups, including steady state [17,18,47–50], transient [50–55], and impedance [15,34,56–58]. When the immobile species has the opposite charge to the active charge carrier that carries the current at boundaries, the current may exceed the diffusion limit in steady state [17,18]. A depletion region is formed near the sink boundary where the concentration of charge carriers in the bulk electrolyte diminishes. The overlimiting current is sustained by either an extra conductivity provided by the charge carriers screening the immobile charges or by electroosmotic circulation and instability if the medium is fluid, in the depletion region [18,59]. The transient response under an overlimiting current was thoroughly studied by Zangle et al. [52–55], Mani and Bazant [60], and Yaroshchuk [61], where they showed concentration polarization propagating as a shock wave. Also, Yan et al. [62] and Khoo and Bazant [50] have recently studied the linear sweep voltammetry of...
electrodiffusion in a charged medium with different boundary conditions. On the other hand, although there are a few studies on the impedance response in the context of a permselective membrane [56–58], a solar cell [6], and a microchannel [63,64], a general theory of the electrodiffusion impedance in a charged medium is not yet available to the best of our knowledge.

In this paper, we present a theoretical model for the electrochemical impedance of electrodiffusion in the presence of an immobile charged species and study its behavior under different dc biases. We consider a system that has two symmetric, oppositely charged charge carriers, which can be binary electrolytes in a liquid solution, electrons and holes in a semiconductor, or electrons and cations in a mixed conductor. No reaction or generation in the bulk is assumed, as well as negligible convection, as a first attempt to focus on the effect of the immobile charged species and the dc bias. Their contributions could be important in some applications, for which the model in this paper should be modified accordingly [6,18,65,66]. We consider two cell configurations:

(i) a reservoir configuration in which one side of the system is exposed to a reservoir that maintains constant concentrations and potential, and the other side is in contact with a selective boundary that accepts only the active charge carrier, and

(ii) a symmetric configuration in which both sides are in contact with the selective boundaries. Our model isolates the response of the bulk electrodiffusion from the contributions of the interfaces and displacement current. For many applications where the electric double layer is much thinner than the system length scale (i.e., thin-double-layer limit) [16,67], the interfacial impedance appears well separated in frequency from that of the bulk transport [19,27]. The displacement current also contributes at much higher frequencies. For a linearized response such as impedance, their contributions could be obtained separately and added to the present model to examine the total cell impedance for the entire frequency range, as shown in our following study [68].

Beginning in Sec. II, we will first set up governing equations and boundary conditions, nondimensionalize variables, and then introduce a small perturbation to the system. In Sec. III, the zero-order terms are solved analytically to study the steady-state behavior. The first-order terms are then solved in Sec. IV, given the zero-order solutions, which provides the impedance response. We investigate the solutions for different combinations of immobile charge density and dc bias including those in the overlimiting regime. Asymptotic expressions for low- and high-frequency limits are obtained for the small- and large-bias regimes. Lastly, we also propose and validate a two-zone approximation for a large bias above the limiting current in Sec. V.

II. MODEL

Consider a one-dimensional system from \( x = 0 \) to \( x = L \), containing two oppositely charged mobile species and another charged immobile species. It could be an unsupported liquid binary electrolyte in a permselective membrane with charged functional groups, or in a porous medium or a microchannel with surface charges. It could also be a solid-state system, such as a semiconductor or a mixed ion-electron conductor containing ionized dopants. The Nernst-Planck equation is employed to describe electrodiffusion of the charge carriers. Assuming no generation and negligible convection,

\[
\frac{\partial c_{\pm}}{\partial t} = - \frac{\partial F_{\pm}}{\partial x}, \tag{1}
\]

\[
F_{\pm} = -D \left( \frac{\partial c_{\pm}}{\partial x} \pm \frac{z e c_{\pm}}{k_B T} \frac{\partial \phi}{\partial x} \right), \tag{2}
\]

where \( c_{\pm}(t, x) \) and \( F_{\pm}(t, x) \) are the concentrations and the fluxes, respectively, of the positive and the negative charge carriers, and \( \phi(t, x) \) is the electric potential. \( t \) and \( x \) are the time and the position variables. \( e, k_B, \) and \( T \) are the electron charge, the Boltzmann constant, and the absolute temperature, respectively. Here we assumed a symmetric charge number \( z \) and a symmetric chemical diffusivity \( D \). Although \( D \) is generally a function of concentration, we consider it constant for simplicity. For a porous medium or a microchannel, \( D \) is the effective diffusivity corrected by the porosity and the tortuosity [69] and \( \phi \) is the electric potential in the solution phase.

Assuming thin double layers, we isolate the quasineutral bulk and adopt the local electroneutrality condition to replace the Poisson equation. The immobile charged species provides as a background charge density in the neutrality condition

\[
0 = z e c_{+} - z e c_{-} + \rho, \tag{3}
\]

where \( \rho \) is the charge density of the immobile species. For a porous medium or microchannel, Eq. (3) means macroscopic electroneutrality, which is valid as long as the pores are larger than a couple of nanometers [70,71]. Then \( \rho \) becomes the homogenized density of the surface charge; \( \rho = a_{i} \sigma_{i}/\epsilon_{p} \), where \( a_{i} \) is the volumetric area, \( \sigma_{i} \) is the surface charge, and \( \epsilon_{p} \) is the porosity (unity for a straight microchannel) [46,50,72]. This condition constrains \( c_{+} \) and \( c_{-} \) to vary exactly in phase with each other, keeping a constant offset \( \rho/z e \) in bulk. The results in this paper are illustrated by assigning the positive charge carriers to be the active carrier. This choice is arbitrary and the results are symmetric in \( \rho \), if the negative charge carriers become the active carriers.

Let us define the dimensionless variables \( \tilde{x} = x/L, \tilde{t} = D t/L^{2}, \tilde{\rho} = \rho/2ze c_{0}, \tilde{\phi} = \phi/ze d_{0} T, \) and \( \tilde{c} = c_{-}/c_{0} = c_{+}/c_{0} + 2\tilde{\rho} \), where \( c_{0} \) is the initial anion concentration at equilibrium. Then by Eqs. (2) and (3), the dimensionless fluxes (\( \tilde{F}_{\pm} = LF_{\pm}/Dc_{0} \)) become

\[
\tilde{F}_{+} = -\frac{\partial \tilde{c}}{\partial \tilde{x}} - (\tilde{c} - 2\tilde{\rho}) \frac{\partial \tilde{\phi}}{\partial \tilde{x}} , \tag{4}
\]

\[
\tilde{F}_{-} = -\frac{\partial \tilde{c}}{\partial \tilde{x}} + \tilde{c} \frac{\partial \tilde{\phi}}{\partial \tilde{x}} . \tag{5}
\]

Upon plugging the flux equations into the conservation equations given by Eq. (1), we add and subtract the conservation equations of the positive and the negative charge carriers. Thus, the following dimensionless governing equations are obtained for \( \tilde{c}(\tilde{t}, \tilde{x}) \) and \( \tilde{\phi}(\tilde{t}, \tilde{x}) \):

\[
\frac{\partial \tilde{c}}{\partial \tilde{t}} = \frac{\partial^{2} \tilde{c}}{\partial \tilde{x}^{2}} - \tilde{\rho}^{2} \frac{\partial \tilde{\phi}}{\partial \tilde{x}^{2}} , \tag{6}
\]

\[
0 = \frac{\partial}{\partial \tilde{x}} \left( \tilde{c} - \tilde{\rho} \frac{\partial \tilde{\phi}}{\partial \tilde{x}} \right) . \tag{7}
\]
The first equation describes conservation of the total number of charge carriers, whereas the latter equation describes conservation of the net electric charge. By assuming the local neutrality condition in Eq. (3), we are able to reduce the number of concentration variables and drop the time derivative for the conservation of net electric charge.

As for the boundary conditions, we consider two common cell configurations as shown in Fig. 1. In both configurations, a potential ($-V = -zeV/k_BT$) is applied at the selective boundary at $x = 1$, where only the active carrier (the positive charge carrier in this paper) can pass the current and the other carrier is completely blocked ($F_1 = 0$). Since we are isolating the bulk electrodiffusion, the boundary conditions are defined inside the transport medium, not involving charge-transfer kinetics or double-layer charging dynamics at the interfaces. The influence of interfacial impedance is discussed in detail elsewhere by Macdonald using the Chang-Jaffe kinetics [73] and by ourselves using the Butler-Volmer kinetics [68]. Therefore, in this work, the boundary conditions at $\bar{x} = 1$ are

$$\bar{\phi} = -\bar{V},$$

$$\frac{d\bar{c}}{d\bar{x}} - \bar{c} \frac{d\bar{\phi}}{d\bar{x}} = 0.$$  

In the reservoir configuration, Fig. 1(a), an ideal reservoir maintains $\bar{c}$ and $\bar{\phi}$ constant on the other side ($\bar{x} = 0$). We set $\bar{\phi}$ at $\bar{x} = 0$ to be zero as a reference value. Therefore, the boundary conditions for the reservoir configuration at $\bar{x} = 0$ are

$$\bar{c} = 1,$$  

$$\bar{\phi} = 0.$$  

In the symmetric configuration, Fig. 1(b), the boundary at $\bar{x} = 0$ has the same semiblocking condition ($F_1 = 0$) as well, identical to the boundary at $\bar{x} = 1$. The boundary conditions for the symmetric configuration at $\bar{x} = 0$ are

$$\frac{d\bar{c}}{d\bar{x}} - \bar{c} \frac{d\bar{\phi}}{d\bar{x}} = 0,$$  

$$\bar{\phi} = 0.$$  

The two blocking conditions for the negative charge carrier in the symmetric configuration do not specify a unique solution. To obtain a unique solution, the total number of the negative charge carriers should be conserved, which adds the following integral constraint for the symmetric configuration:

$$\int_0^1 \bar{c} d\bar{x} = 1.$$  

Regardless of configuration, the dimensionless system is governed by two dimensionless parameters, $\bar{\rho}$ and $\bar{V}$, which represent the charge density of the immobile species and the magnitude of the applied bias, respectively. It is also possible to specify the applied bias by the current ($\bar{j}$) as well. Disregarding the displacement current, the current comes from the charge carrier fluxes: $j = ze(F_1 - F_0)$. When scaled by the diffusion-limiting current ($j_{\text{lim}} = 2zeDc_0/L$), the dimensionless current ($\bar{j} = j/j_{\text{lim}}$) can be obtained by

$$\bar{j} = - (\bar{c} - \bar{\rho}) \frac{\partial \bar{\phi}}{\partial \bar{x}}.$$  

$j$ is constant throughout the system domain due to the local neutrality condition in Eq. (3). In this paper, we refer to $\bar{j}$ to specify the magnitude of the dc bias rather than $\bar{V}$, because it gives a more relevant scale to electrodiffusion.

The system is perturbed by a small sinusoidal stimulus with a frequency $\omega = \omega L^2/D$ in either $\bar{V}$ or $\bar{j}$ to calculate the impedance. When the amplitude is small enough, variables can then be expanded according to the perturbation theory:

$$\bar{c}(\bar{x}, t) = c^{(0)}(\bar{x}) + \epsilon c^{(1)}(\bar{x}) e^{i\omega t} + O(\epsilon^2),$$

$$\bar{\phi}(\bar{x}, t) = \phi^{(0)}(\bar{x}) + \epsilon \phi^{(1)}(\bar{x}) e^{i\omega t} + O(\epsilon^2),$$

$$\bar{j}(t) = j^{(0)} + \epsilon j^{(1)} e^{i\omega t} + O(\epsilon^2),$$

$$\bar{V}(t) = V^{(0)} + \epsilon V^{(1)} e^{i\omega t} + O(\epsilon^2),$$  

where $\epsilon$ is an arbitrary small number that represents the amplitude of the AC perturbation. Other variables may be expanded in the same manner as well. Upon substitution of the expanded variables into Eqs. (6)–(15), the collection of $O(1)$ terms can be first solved for the reference steady state under a dc bias. Given the steady-state solution, the collection of $O(\epsilon)$ terms can be solved for the perturbation around the steady state, which is then used for calculating the impedance.

III. STEADY STATE

Impedance is measured by applying a perturbation around a reference steady state. By achieving different steady states by dc bias, we can study nonlinear behavior of charge carriers via impedance spectroscopy [74]. To interpret such results, we need to incorporate the steady state in an impedance model. Although the steady-state solution under dc bias has already been studied in Refs. [18,50], we revisit its behavior for completeness of the paper. The steady-state solution is obtained by solving the $O(1)$ terms, and steps involved in obtaining an analytical solution are presented in Appendix A. For the reservoir configuration, the solution is

$$\bar{j}^{(0)} = 1 - (e^{-\bar{V}^{(0)}} + \bar{\rho} V^{(0)}).$$
Concentration and potential distributions for \( \tilde{\rho} = -0.01 \) under different dc biases. For the symmetric configuration: (d) current-voltage curves for different \( \tilde{\rho} \). (e) Concentration and (f) potential distributions for \( \tilde{\rho} = -0.01 \) under different dc biases.

\[
\tilde{c}^{(0)} = \begin{cases} 
-\tilde{\rho}W_0 \left[ -\frac{1}{\tilde{\rho}} \exp \left( \frac{(j^{(0)} \tilde{x} - 1)}{\tilde{\rho}} \right) \right], & \text{if } \tilde{\rho} < 0, \\
1 - j^{(0)} \tilde{x}, & \text{if } \tilde{\rho} = 0, \\
-\tilde{\rho}W_{-1} \left[ -\frac{1}{\tilde{\rho}} \exp \left( \frac{(j^{(0)} \tilde{x} - 1)}{\tilde{\rho}} \right) \right], & \text{if } \tilde{\rho} > 0,
\end{cases}
\]

\[
\tilde{\phi}^{(0)} = \log(\tilde{c}^{(0)}),
\]

where \( W_0 \) and \( W_{-1} \) are the principal (upper) and the lower branches of the Lambert W function, respectively [75].

Figure 2(a) shows \( j^{(0)} \) versus \( V^{(0)} \) for \( \tilde{\rho} \in \{-0.1, -0.01, 0, 0.01, 0.1\} \). Without the immobile charge (\( \tilde{\rho} = 0 \)), \( j^{(0)} \) saturates at the diffusion-limiting current (\( j^{(0)} = 1 \)). If the immobile species and the active carriers are cocharged (\( \tilde{\rho} > 0 \)), \( j^{(0)} \) is limited by \( 1 - \tilde{\rho}[2 - \log(2\tilde{\rho})] \), at which \( \tilde{c}^{(0)} \) reaches \( 2\tilde{\rho} \) at \( \tilde{x} = 1 \). When charge-transfer kinetics is considered, the current limit cannot be reached with a finite potential bias, since the charge-transfer resistance diverges due to the diminishing concentration at the boundary. On the other hand, if the immobile species and the active carrier are countercharged (\( \tilde{\rho} < 0 \)), \( j^{(0)} \) can exceed the limiting current above which it keeps a constant slope of \( -\tilde{\rho} \). In the overlimiting regime, the current is sustained by the active carriers that screen the immobile charge.

In Figs. 2(b) and 2(c), \( \tilde{c}^{(0)} \) and \( \tilde{\phi}^{(0)} \) are plotted under a varying dc bias for \( \tilde{\rho} = -0.01 \). As a bias is applied, a concentration gradient develops and it becomes steeper as \( j^{(0)} \) increases. Below the limiting current, \( c^{(0)} \) drops almost linearly along \( \tilde{x} \), while \( \phi^{(0)} \) shows a steeper drop closer to the boundary at \( \tilde{x} = 1 \). On the other hand, above the limiting current, the depletion region appears near the boundary at \( \tilde{x} = 1 \), where \( \tilde{c}^{(0)} \) diminishes and \( \tilde{\phi}^{(0)} \) shows a steep linear decrease. Therefore, the contribution of electric migration should be larger compared to that of diffusion in the depletion region. Also, notice that the depletion region grows as \( j^{(0)} \) increases. In the Supplemental Material [76], a MATLAB script is attached which plots the steady-state solution for the reservoir configuration with a given combination of \( \tilde{\rho} \) and \( j^{(0)} \). When \( \tilde{\rho} \) becomes more negative, the transition in \( \tilde{c}^{(0)} \) and \( \tilde{\phi}^{(0)} \) becomes smoother, which makes the depletion region less distinctive. An analytical approximation for the overlimiting regime is discussed in Sec. V.

For the symmetric configuration, an analytical solution can also be obtained. Steps involved in solving the \( O(1) \) terms are presented in Appendix A. The solution for the symmetric configuration is

\[
\tilde{c}^{(0)} = \alpha(1 - e^{-\tilde{\rho}V^{(0)}}) - \tilde{\rho}V^{(0)} ,
\]

\[
\tilde{\phi}^{(0)} = \log \left( \frac{\tilde{c}^{(0)}}{\alpha} \right),
\]
where \( \alpha \) is an intermediate parameter such that

\[
\alpha = \frac{(1 + \bar{\rho})(1 - e^{-\rho(0)})}{1 - e^{-2\rho(0)}} + \sqrt{(1 + \bar{\rho})^2(1 - e^{-\rho(0)})^2 - 2\bar{\rho}^2(1 - e^{-2\rho(0)})}{1 - e^{-2\rho(0)}}.
\]  

(23)

It can be inferred by Eq. (22) that \( \alpha = \bar{\rho}(\bar{x} = 0) \).

Figure 2(d) shows \( \bar{I}(0) \) versus \( \bar{V}(0) \) for \( \bar{\rho} \in \{-0.1, -0.01, 0, 0.01, 0.1\} \). While its general behavior is similar to Fig. 2(a), the diffusion-limiting current in the symmetric configuration is twice that of the reservoir configuration. The current is limited by 2 if \( \bar{\rho} = 0 \), or by \( \alpha - \bar{\rho}(2 - \log(2\bar{\rho}/\alpha)) \) if \( \bar{\rho} > 0 \), where \( \alpha \) is implicitly determined by putting \( -\bar{V}(0) = \log(2\bar{\rho}/\alpha) \) in Eq. (23). On the other hand, the overlimiting regime is accessible if \( \tilde{\rho} < 0 \). In Figs 2(e) and 2(f), \( \tilde{\phi}(0) \) and \( \tilde{\xi}(0) \) are plotted under a varying dc bias for \( \bar{\rho} = -0.01 \). When \( \bar{I}(0) > 0 \), \( \tilde{\phi}(0) \) begins to have a gradient and \( \tilde{\xi}(0) \) at \( \bar{x} = 1 \) drops, which results in \( \tilde{\rho}(0) \) building up near the boundary at \( \bar{x} = 0 \) due to the integral constraint in Eq. (14). It results in a steeper gradient, which enhances the transport and makes the diffusion-limiting current twice higher than that of the reservoir configuration. Other than that, \( \tilde{\phi}(0) \) and \( \tilde{\xi}(0) \) show similar behaviors to those of the reservoir configuration [Figs. 2(b) and 2(c)]. In the Supplemental Material [76], a MATLAB script is attached which plots the steady-state solution for the symmetric configuration with a given combination of \( \bar{\rho} \) and \( \bar{I}(0) \). An analytical approximation for the overlimiting regime in the symmetric configuration is discussed in Sec. V as well. The steady-state solutions are used in calculating the \( O(\epsilon) \) perturbation solutions in the next section.

IV. IMPEDANCE

While the \( O(1) \) solution corresponds to the reference steady state under a dc bias, the \( O(\epsilon) \) terms describe the perturbation around it, which leads to the impedance solution. The \( O(\epsilon) \) governing equations are

\[
i\omega\tilde{\epsilon}(1) = \frac{d^2\tilde{\xi}(1)}{dx^2} - \bar{\rho} \frac{d^2\tilde{\phi}(1)}{dx^2},
\]  

(24)

\[
0 = (\tilde{\epsilon}(0) - \bar{\rho}) \frac{d^2\tilde{\phi}(1)}{dx^2} + \frac{d\tilde{\phi}(0)}{dx} \frac{d\tilde{\phi}(1)}{dx} + \frac{d\tilde{\phi}(0)}{dx} \frac{d\tilde{\xi}(1)}{dx} + \frac{d\tilde{\phi}(0)}{dx} \tilde{\epsilon}(1).
\]  

(25)

For the reservoir configuration, the \( O(\epsilon) \) boundary conditions are

\[
\tilde{\xi}(1) = 0 \text{ at } \bar{x} = 0,
\]

(26)

\[
\tilde{\xi}(1) = 0 \text{ at } \bar{x} = 0.
\]

(27)

\[
\tilde{\phi}(1) = -\tilde{\phi}(1) \text{ at } \bar{x} = 1,
\]

(28)

\[
\frac{d\tilde{\xi}(1)}{dx} = \tilde{\epsilon}(1) \frac{d\tilde{\phi}(0)}{dx} + \tilde{\epsilon}(0) \frac{d\tilde{\phi}(1)}{dx} \text{ at } \bar{x} = 1.
\]

(29)

This boundary value problem involves the \( O(1) \) solution derived in the previous section. Since they are not elementary functions, Eqs. (24)–(29) should be solved numerically. We attach our MATLAB script in the Supplemental Material, which solves the boundary value problem and calculates the impedance for any combinations of \( \bar{I}(0) \) and \( \bar{\rho} \) [76]. When the \( O(\epsilon) \) solution is obtained, combined with the \( O(1) \) solution, the impedance \( (\bar{Z} = 2Dc_0/\kappa B TL) \) is calculated by

\[
\bar{Z} = \frac{\tilde{\phi}(1)}{\bar{I}(1)} = \frac{-\tilde{\epsilon}(0) - \bar{\rho} \frac{d\tilde{\phi}(1)}{dx} - \bar{\rho}(1) \frac{d\tilde{\phi}(0)}{dx}}{\tilde{\epsilon}(1) \frac{d\tilde{\phi}(0)}{dx} + \tilde{\epsilon}(0) \frac{d\tilde{\phi}(1)}{dx}}.
\]

(30)

Figure 3 presents \( \bar{Z} \) under different dc biases on the complex plane for the reservoir configuration with \( \bar{\rho} = -0.01 \). Below the diffusion-limiting bias \( \bar{I}(0) < 1 \) as shown in Fig. 3(a), the impedance of bulk electrodiffusion appears as a semicircle that grows in both low-frequency and high-frequency limits with increasing bias. At the high-frequency limit, it is dominated by the \( O(\epsilon) \) conduction with the steady-state conductivity that leads to the pure resistive behavior. The capacitive contribution comes from the fluctuation in \( O(\epsilon) \) conductivity under the steady-state potential gradient, which turns back to the resistive behavior at the lower-frequency limit when the conductivity fluctuates in phase with \( \tilde{V}(1) \) and \( \tilde{I}(1) \). Around the diffusion-limiting bias \( \bar{I}(0) = 1 \), the semicircle becomes suppressed, and then the impedance appears more like a finite-length Warburg element under a bias above the diffusion limit \( \bar{I}(0) > 1 \), as shown in Fig. 3(b). Its overall magnitude also starts converging above the limit. Such response comes from the conduction in the depletion region and the diffusion out of the depletion region, under an overlimiting bias. A detailed discussion of this regime is presented in the next section with an analytical approximation. Without any bias, the impedance shrinks to a pure resistance of \( (1 - \bar{\rho})^{-1} \).

The transition along increasing bias becomes more apparent by plotting the limiting behaviors as functions of the applied bias \( \bar{I}(0) \). Figure 4(a) shows the low-frequency
resistance, $\tilde{R}_L = Z(\omega \to 0)$, and the high-frequency resistance, $\tilde{R}_H = Z(\omega \to \infty)$. Both increase exponentially in the underlimiting regime ($\tilde{j}^{(0)} < 1$) and then converge to $\tilde{\rho}^{-1}$ in the overlimiting regime ($\tilde{j}^{(0)} > 1$). Asymptotic expressions of $\tilde{R}_L$ and $\tilde{R}_H$ are derived in Appendix B for the small-bias limit ($\tilde{j}^{(0)} \ll 1$) and in Appendix C for the large-bias limit ($\tilde{j}^{(0)} \gg 1$):

$$\tilde{R}_L(\tilde{j}^{(0)} \ll 1) \simeq \frac{1 - \tilde{\rho}}{(1 - \tilde{\rho})^2 - \tilde{j}^{(0)}},$$  \hspace{1cm} (31)

$$\tilde{R}_L(\tilde{j}^{(0)} \gg 1) \simeq \frac{C}{1 - \tilde{\rho}},$$  \hspace{1cm} (32)

$$\tilde{R}_H(\tilde{j}^{(0)} \ll 1) \simeq -\frac{(1 - \tilde{\rho})}{\tilde{j}^{(0)}} \log\left(\frac{(1 - \tilde{\rho})^2 - \tilde{j}^{(0)}}{(1 - \tilde{\rho})^2}ight),$$  \hspace{1cm} (33)

$$\tilde{R}_H(\tilde{j}^{(0)} \gg 1) \simeq \left(1 - \frac{1}{\tilde{\rho}}\right) \tilde{j}^{(0)} - 1.$$  \hspace{1cm} (34)

They are compared to the numerical limiting behaviors in Fig. 10 in Appendix A. In addition, Fig. 4(b) shows the local phase angle at the high-frequency limit $\theta_H$. In the underlimiting regime, $\theta_H$ starts by $-90^\circ$, which corresponds to the semicircle shape on the complex plane, Fig. 3(a). It then approaches $-45^\circ$ in the overlimiting regime as the impedance transitions to the finite-length Warburg shape. If $\tilde{\rho} > 0$, $\tilde{R}_L$ and $\tilde{R}_H$ diverge before the diffusion-limiting bias. The maximum current bias in the reservoir configuration is 1 if $\tilde{\rho} = 0$ or $1 - \tilde{\rho}[2 - \log(2\tilde{\rho})]$ if $\tilde{\rho} > 0$, as discussed in Sec. III.

In a similar approach, the impedance for the symmetric configuration can be obtained. Its $O(\epsilon)$ governing equations are the same as the reservoir case, Eqs. (24) and (25). The $O(\epsilon)$ boundary conditions for the symmetric configuration are

$$\phi^{(1)} = 0 \quad \text{at} \quad \tilde{x} = 0,$$  \hspace{1cm} (35)

$$\frac{d\tilde{c}^{(1)}}{d\tilde{x}} = \tilde{c}^{(1)} \frac{d\phi^{(0)}}{d\tilde{x}} + \tilde{c}^{(0)} \frac{d\phi^{(1)}}{d\tilde{x}} \quad \text{at} \quad \tilde{x} = 0,$$  \hspace{1cm} (36)

To solve Eqs. (24), (25), and (35)–(39) with a typical solver for boundary value problems, another field variable is introduced, $\tilde{\psi}^{(1)}(\tilde{x}) = \int_0^1 \tilde{c}^{(1)}(s) ds$, which adds another governing equation, $d\tilde{c}^{(1)}/d\tilde{x} = \tilde{c}^{(1)}$. Then the integral boundary condition, Eq. (39), is replaced by two new boundary conditions for $\tilde{\psi}^{(1)}(\tilde{x})$: $\tilde{\psi}^{(1)}(0) = 0$ and $\tilde{\psi}^{(1)}(1) = 1$. The implementation can be found in our MATLAB script attached in the Supplemental Material [76], which solves the $O(\epsilon)$ problem and calculates the impedance for the symmetric configuration with any combinations of $\tilde{j}^{(0)}$ and $\tilde{\rho}$ [76].

Figure 5 shows $\tilde{Z}$ under different biases for the symmetric configuration with $\tilde{\rho} = -0.01$, and Fig. 6 shows the transition of its limiting behaviors along increasing $\tilde{j}^{(0)}$. Like in the reservoir configuration, $\tilde{Z}$ appears similar to a semicircle on the complex plane below the diffusion-limiting bias ($\tilde{j}^{(0)} < 2$), whose $\tilde{R}_L$ and $\tilde{R}_H$ increase exponentially with increasing $\tilde{j}^{(0)}$, as shown in Fig. 6(a). However, the semicircle is tilted by a small angle at the high-frequency limit due to the integral constraint in the symmetric configuration, Eq. (14). Therefore, $\theta_H$ starts less negative than $-90^\circ$ in Fig. 6(b). Then $\tilde{Z}$ shows a transition around the diffusion-limiting bias ($\tilde{j}^{(0)} = 2$), and it appears like a finite-length Warburg element in the overlimiting regime ($\tilde{j}^{(0)} > 2$), as shown in Fig. 5(b). Similar to the transition observed in the reservoir configuration, $\tilde{R}_L$ and $\tilde{R}_H$ converge to $\tilde{\rho}^{-1}$, and $\theta_H$ converges to $-45^\circ$ in the overlimiting regime. Asymptotic expressions of $\tilde{R}_L$ and $\tilde{R}_H$
for the symmetric configuration are derived in Appendix B for the small-bias limit ($j^{(0)} \ll 2$) and in Appendix C for the large-bias limit ($j^{(0)} \gg 2$):

$$\hat{R}_L(j^{(0)} \ll 2) \approx \frac{1 + \bar{\rho}}{2(1 - \bar{\rho})^2 j^{(0)} - j^{(0)}},$$

$$\hat{R}_L(j^{(0)} \gg 2) \approx \frac{1 - \bar{\rho}}{2(1 - \bar{\rho})^2 j^{(0)} + j^{(0)}},$$

$$\hat{R}_H(j^{(0)} \ll 2) \approx \frac{1 - \bar{\rho}}{2(1 - \bar{\rho})^2 - j^{(0)}},$$

$$\hat{R}_H(j^{(0)} \gg 2) \approx \frac{1 - \bar{\rho}}{2(1 - \bar{\rho})^2 + j^{(0)}}.$$

They are compared to the numerical limiting behaviors in Fig. 10. The current bias $\bar{j}^{(0)}$ is limited by 2 if $\bar{\rho} = 0$, or by $\alpha - \bar{\rho}(2 - \log(2\bar{\rho}/\alpha))$ if $\bar{\rho} > 0$, as discussed in Sec. III, and $\hat{R}_C$ and $\hat{R}_H$ diverge before the limiting current bias. Regardless of $\bar{\rho}$, $\hat{Z}$ becomes a pure resistance of $(1 - \bar{\rho})^{-1}$ without any bias, same as the reservoir configuration.

Notice that our model focuses on the bulk electrodiffusion without considering the contributions of the interfaces and displacement current. Each of these additional contributions would yield another relaxation behavior at higher frequencies, usually well separated from the bulk features [19,27,37,39,41]. Also, if the diffusivities of positive and negative charge carriers are different, another finite-length Warburg appears around the bulk diffusion frequency starting from the unbiased condition. While we have isolated its contribution out in this paper by assuming identical diffusivities, the extra Warburg coming from unequal diffusivities could appear merged with the bulk impedance studied in this paper [19,37,39]. In our following study, we apply the model and solution methods to a more general setup with unequal diffusivities and charge numbers [68].

FIG. 6. Limiting behaviors of $\hat{Z}$ as functions of $\bar{j}^{(0)}$ in the symmetric configuration with $\bar{\rho} = -0.01$: (a) $\hat{R}_L$ and $\hat{R}_H$, and (b) $\theta_H$.

V. TWO-ZONE APPROXIMATION

In this section, we present an analytical approximation for a countercharged system with a low density of the immobile charged species and with a large bias above the diffusion-limiting bias. It is motivated by recognizing that $\tilde{c}^{(0)}(\tilde{x})$ and $\tilde{\varphi}^{(0)}(\tilde{x})$ show two distinct zones under such conditions as shown in Figs. 7(a) and 7(b). The two zones become more apparent by defining the diffusion and the conduction currents, $\tilde{j}_D^{(0)}$ and $\tilde{j}_C^{(0)}$, for the steady state:

$$\tilde{j}_D^{(0)} = -\frac{d\tilde{c}^{(0)}}{d\tilde{x}},$$

$$\tilde{j}_C^{(0)} = -\frac{d\tilde{\varphi}^{(0)}}{d\tilde{x}},$$

which add up to the total current: $\tilde{j}_D^{(0)} + \tilde{j}_C^{(0)} = \tilde{j}^{(0)}$. Figure 7(c) shows how contributions of $\tilde{j}_D^{(0)}$ and $\tilde{j}_C^{(0)}$ change along $\tilde{x}$. From $\tilde{x} = 0$, expanding to a position around the middle in the figure, $\tilde{c}^{(0)}$ drops linearly while $\tilde{\varphi}^{(0)}$ changes little compared to the rest of the space. In this zone, $\tilde{j}_D^{(0)}$ dominates, and we define it as the diffusion zone with length $\tilde{l}_D$ that depends on $\tilde{j}^{(0)}$ and $\tilde{\varphi}$. On the other hand, from $\tilde{x} = \tilde{l}_D$ to 1, $\tilde{c}^{(0)}$ stays near zero while $\tilde{\varphi}^{(0)}$ shows a large linear drop. Since $\tilde{j}_C^{(0)}$ dominates here, we define this zone as the conduction zone with length $\tilde{l}_C$ ($\tilde{l}_D + \tilde{l}_C = 1$), which is referred to as the depletion region in a lot of literature on the overlimiting current. In this approximation, the Nernst-Planck model is reduced to a pure diffusion equation in the diffusion zone and to a pure conduction equation in the conduction zone. We will derive the analytical solutions for steady state as

FIG. 7. Spatial distributions of (a) $\tilde{c}^{(0)}$, (b) $\tilde{\varphi}^{(0)}$, and (c) $\tilde{j}_D^{(0)}$ and $\tilde{j}_C^{(0)}$ at an overlimiting current ($\bar{j}^{(0)} = 2.0$) in the reservoir configuration with $\bar{\rho} = -0.01$. 

well as the impedance in both of the reservoir and symmetric configurations. Its validity is then discussed over a range of \( j^{(0)} \) and \( \rho \) in the last part.

For the reservoir configuration in steady state, the approximated concentration \( \tilde{c}_{TZ}^{(0)} \) is obtained by solving \( d^2 \tilde{c}_{TZ}^{(0)}/d\tilde{x}^2 = 0 \) in the diffusion zone with boundary conditions \( \tilde{c}_{TZ}^{(0)} = 1 \) at \( \tilde{x} = 0 \) and \( \tilde{c}_{TZ}^{(0)} = 0 \) at \( \tilde{x} = \tilde{I}_D \). On the other hand, \( \tilde{c}_{TZ}^{(0)} \) is kept zero throughout the conduction zone. Therefore,

\[
\tilde{c}_{TZ}^{(0)} = \begin{cases} 
1 - \frac{\tilde{x}}{\tilde{I}_D}, & \text{if } 0 \leq \tilde{x} \leq \tilde{I}_D, \\
0, & \text{if } \tilde{I}_D < \tilde{x} \leq 1,
\end{cases}
\] (46)

where \( \tilde{I}_D \) is determined such that the concentration gradient drives \( j^{(0)} \) in the diffusion zone by Eq. (44): \( \tilde{I}_D = 1/j^{(0)} \). The approximated potential \( \tilde{\phi}_{TZ}^{(0)} \) is then obtained by solving the pure conduction equation, \( \tilde{\rho} d\tilde{\phi}_{TZ}^{(0)}/d\tilde{x} = \tilde{j}^{(0)} \), in the conduction zone. Since the potential drop in the diffusion zone is significantly smaller than that in the conduction zone, we set \( \tilde{\phi}_{TZ}^{(0)} = 0 \) in the diffusion zone. Therefore,

\[
\tilde{\phi}_{TZ}^{(0)} = \begin{cases} 
0, & \text{if } 0 \leq \tilde{x} \leq \tilde{I}_D, \\
\frac{\tilde{j}^{(0)}(\tilde{x} - \tilde{I}_D)}{\tilde{\rho}}, & \text{if } \tilde{I}_D < \tilde{x} \leq 1
\end{cases}
\] (47)

\( \tilde{c}_{TZ}^{(0)} \) and \( \tilde{\phi}_{TZ}^{(0)} \) are compared to the exact solutions in Figs. 8(a) and 8(b), respectively. When \( |\tilde{\rho}| \) is small and \( \tilde{j}^{(0)} \) is large above the diffusion limit, the two-zone approximation shows good agreement with the exact steady-state solutions.

The impedance approximation can then be obtained by separately calculating the impedance of the two zones. For impedance of the diffusion zone, we first calculate \( \tilde{Z}_{TZ}^{(1)} \) at \( \tilde{x} = \tilde{I}_D \) given \( \tilde{j}^{(1)} \). \( \tilde{Z}_{TZ}^{(1)} \) is obtained by solving \( d^2 \tilde{Z}_{TZ}^{(1)}/d\tilde{x}^2 = i\tilde{\omega} \tilde{Z}_{TZ}^{(1)} \) in the diffusion zone with boundary conditions \( \tilde{Z}_{TZ}^{(1)} = 0 \) at \( \tilde{x} = 0 \) and \( d\tilde{Z}_{TZ}^{(1)}/d\tilde{x} = -j^{(1)} \) at \( \tilde{x} = \tilde{I}_D \):

\[
\tilde{Z}_{TZ}^{(1)} = -j^{(1)} \tilde{I}_D \frac{\sinh(\sqrt{i\tilde{\omega}}\tilde{x})}{\sqrt{i\tilde{\omega}} \cosh(\sqrt{i\tilde{\omega}}\tilde{x})},
\] (48)

for \( 0 \leq \tilde{x} \leq \tilde{I}_D \) (i.e., in the diffusion zone). To calculate \( \tilde{\phi}_{TZ}^{(1)}(\tilde{x} = \tilde{I}_D) \), we consider the \( O(\epsilon) \) terms of Eqs. (15) and (44): \( (\tilde{c}_{TZ}^{(1)} - \tilde{\rho})d\tilde{\phi}_{TZ}^{(1)}/d\tilde{x} = d\tilde{Z}_{TZ}^{(1)}/d\tilde{x} \). Since \( \tilde{c}_{TZ}^{(0)} \) approaches zero at \( \tilde{x} = \tilde{I}_D \), we can integrate \( d\tilde{\phi}_{TZ}^{(1)}/d\tilde{x} = -\tilde{\rho}^{-1} \) from the unperturbed condition (i.e., \( \tilde{\phi}_{TZ}^{(0)} = 0 \) and \( \tilde{c}_{TZ}^{(0)} = 0 \)), and obtain

\[
\frac{\tilde{\phi}_{TZ}^{(1)}(\tilde{x} = \tilde{I}_D)}{\tilde{c}_{TZ}^{(1)}(\tilde{x} = \tilde{I}_D)} = \frac{1}{\tilde{\rho}}.
\] (49)

Then, impedance in the diffusion zone \( \tilde{Z}_{DZ} \) can be obtained by

\[
\tilde{Z}_{DZ} = \frac{-\tilde{\phi}_{TZ}^{(1)}(\tilde{x} = \tilde{I}_D)}{\tilde{j}^{(1)}} = \frac{-\tilde{\phi}_{TZ}^{(1)}(\tilde{x} = \tilde{I}_D) \tilde{c}_{TZ}^{(1)}(\tilde{x} = \tilde{I}_D)}{\tilde{c}_{TZ}^{(1)}(\tilde{x} = \tilde{I}_D)} = \frac{\tilde{I}_D}{\tilde{\rho} \sqrt{i\tilde{\omega}} \tilde{I}_D}
\] (50)

which turns out to be a finite-length Warburg element scaled by \( \tilde{I}_D/(\tilde{\rho}) \). Also, its characteristic frequency corresponds to
diffusion with length $\tilde{l}_D$. On the other hand, impedance in the conduction zone $\tilde{Z}_{CZ}$ is a pure resistor with a conductivity of $-\rho$ and a length of $l_C$. Therefore, the overall impedance in the two-zone approximation is

$$\tilde{Z}_{TZ} = \tilde{Z}_{CZ} + \tilde{Z}_{DZ} = \frac{\tilde{l}_C}{-\rho} + \frac{\tilde{l}_D}{\rho} \tanh(\sqrt{i\omega}\tilde{l}_D),$$  

(51)

where $\tilde{l}_C = (\tilde{j}^{(0)} - 1)/\tilde{j}^{(0)}$ and $\tilde{l}_D = 1/\tilde{j}^{(0)}$. In Fig. 8(c), $\tilde{Z}_{TZ}$ is compared to the exact $\tilde{Z}$ obtained by a numerical solution obtained in Sec. IV for the reservoir configuration. The two-zone approximation provides an accurate approximation in impedance for a weakly charged system ($\rho = -0.01$) under an overlimiting current bias. It also infers that the finite-length Warburg feature corresponds to the diffusion zone and the resistor corresponds to the conduction zone. As $\tilde{j}^{(0)}$ increases above the diffusion limit, the conduction zone pushes out the diffusion zone, and the finite-length Warburg part becomes smaller while the resistor becomes larger, keeping the overall low-frequency limit the same.

For the symmetric configuration, the approximation can be obtained in a similar approach. For the steady state, $\tilde{c}_{TZ}^{(0)}$ is obtained by solving $d^2\tilde{c}_{TZ}^{(0)}/d\tilde{x}^2 = 0$ in the diffusion zone with boundary conditions $\tilde{c}_{TZ}^{(0)} = \alpha$ at $\tilde{x} = 0$ and $\tilde{c}_{TZ}^{(0)} = 0$ at $\tilde{x} = \tilde{l}_D$. On the other hand, $\tilde{c}_{TZ}^{(0)}$ is kept zero throughout the conduction zone:

$$\tilde{c}_{TZ}^{(0)} = \begin{cases} \alpha \left( 1 - \frac{1}{\tilde{l}_D} \right), & \text{if } 0 \leq \tilde{x} \leq \tilde{l}_D, \\ 0, & \text{if } \tilde{l}_D < \tilde{x} \leq 1, \end{cases}$$  

(52)

where $\tilde{l}_D$ is determined such that the concentration gradient drives $\tilde{j}^{(0)}$ in the diffusion zone by Eq. (44), $\tilde{l}_D = \alpha l_j^{(0)}$. $\alpha$ in the approximation can be obtained as a function of $\tilde{j}^{(0)}$ using the integral constraint in Eq. (14). Since the triangular area under $\tilde{c}_{TZ}^{(0)}$ in the diffusion zone should be 1, $\alpha = 2/l_D$, and therefore, $\alpha = \sqrt{2} l_j^{(0)}$ and $\tilde{l}_D = \sqrt{2} l_j^{(0)}$. The approximated potential $\tilde{\phi}_{TZ}^{(0)}$ is then obtained by solving the conduction equation, $\tilde{d} \tilde{\phi}_{TZ}^{(0)} / d\tilde{x} = \tilde{j}^{(0)}$, in the conduction zone. Since the potential drop in the diffusion zone is relatively small,

$$\tilde{\phi}_{TZ}^{(0)} = \begin{cases} 0, & \text{if } 0 \leq \tilde{x} \leq \tilde{l}_D, \\ \rho (\tilde{x} - \tilde{l}_D), & \text{if } \tilde{l}_D < \tilde{x} \leq 1. \end{cases}$$  

(53)

Though the expression is exactly the same as $\tilde{\phi}_{TZ}^{(0)}$ in the reservoir configuration, Eq. (47), $\tilde{l}_D$ is a different function of $\tilde{j}^{(0)}$ between the configurations. $\tilde{c}_{TZ}^{(0)}$ and $\tilde{\phi}_{TZ}^{(0)}$ of the symmetric configuration are compared to the exact solutions in Figs. 8(d) and 8(e), respectively, where they show good agreement when $|\tilde{\rho}|$ is low and $\tilde{j}^{(0)}$ is large enough above the diffusion limit.

The impedance approximation for the symmetric configuration is obtained also by the similar approach used in the reservoir configuration. First, $\tilde{c}_{TZ}^{(1)}$ is obtained by solving $d^2\tilde{c}_{TZ}^{(1)}/d\tilde{x}^2 = i\omega\tilde{c}_{TZ}^{(0)}$ in the diffusion zone with a boundary condition $d\tilde{c}_{TZ}^{(1)}/d\tilde{x} = -\tilde{j}^{(1)}$ at $\tilde{x} = \tilde{l}_D$ and an integral constraint $\tilde{J}_0^{(1)} \tilde{c}_{TZ}^{(1)} d\tilde{x} = 0$:

$$\tilde{c}_{TZ}^{(1)} = -\tilde{j}^{(1)} \tilde{l}_D \left( \frac{\cosh(\sqrt{i\omega}\tilde{l}_D) - 1}{\sqrt{i\omega}\tilde{l}_D \sinh(\sqrt{i\omega}\tilde{l}_D)} \right),$$  

(54)

for $0 \leq \tilde{x} \leq \tilde{l}_D$ (i.e., in the diffusion zone). Then, as we did for the reservoir configuration, $\tilde{\phi}_{TZ}^{(1)}$ at $\tilde{x} = \tilde{l}_D$ is calculated by solving the $O(\epsilon)$ terms of Eqs. (15) and (44), which turns out to be the same as that in the reservoir configuration shown in Eq. (49). Following the steps described in Eqs. (50) and (51), the impedance approximation for the symmetric configuration is obtained:

$$\tilde{Z}_{TZ} = \tilde{Z}_{CZ} + \tilde{Z}_{DZ} = \frac{\tilde{l}_C}{-\rho} + \frac{\tilde{l}_D}{\rho} \cosh(\sqrt{i\omega}\tilde{l}_D) - \frac{1}{\sqrt{i\omega}\tilde{l}_D \sinh(\sqrt{i\omega}\tilde{l}_D)},$$  

(55)

where $\tilde{l}_C = (\tilde{j}^{(0)} - \alpha)/\tilde{j}^{(0)}$, $\tilde{l}_D = \alpha \tilde{j}^{(0)}$, and $\alpha = \sqrt{2} l_j^{(0)}$. Due to the integral constraint in Eq. (14), $\tilde{Z}_{DZ}$ has a different expression in the symmetric configuration, even though it appears almost identical to the traditional finite-length Warburg element on the complex plane. It leads to the extra term in $R_L$ in Eq. (41) compared to that of the reservoir configuration, which makes $R_L$ still increase in the overlimiting regime. In Fig. 8(f), $\tilde{Z}_{TZ}$ is compared to the exact $\tilde{Z}$ obtained numerically for the symmetric configuration. Similar to the reservoir configuration, the two-zone approximation provides an accurate approximation of $\tilde{Z}$ when $|\tilde{\rho}|$ is small and $\tilde{j}^{(0)}$ is large above the diffusion limit.

The validity of the approximation largely depends on the interphase thickness ($\tilde{\delta}$) indicated in Fig. 7(c), because we assumed that the system can be exhaustively separated into the conduction and the diffusion zones by a sharp interphase between them. If $\tilde{\delta}$ is not small enough, the contribution of the interphase should introduce a significant error in the approximation. We define $\tilde{\delta}$ by a distance between two positions where $\tilde{Z}^{(0)} = 0.9 \tilde{j}^{(0)}/(1 - \tilde{\rho})$ and $\tilde{Z}_C^{(0)} = 0.9 \tilde{j}^{(0)}$. Figures 9(a) and 9(b) show how $\tilde{\delta}$ changes by $\tilde{j}^{(0)}$ and $\tilde{\rho}$ in the overlimiting regime for the reservoir and the symmetric configurations, respectively. When $\tilde{j}^{(0)}$ is small and $|\tilde{\rho}|$ is large, there do not exist any distinct zones where either the conduction or the diffusion current dominate and the interphase thickness is not well defined. Otherwise, $\tilde{\delta}$ decreases in a direction of decreasing $|\tilde{\rho}|$ and increasing $\tilde{j}^{(0)}$. Its slope is steeper along $|\tilde{\rho}|$, and a sharp interface ($\tilde{\delta} < 0.1$) is obtained regardless of $\tilde{j}^{(0)}$, if $|\tilde{\rho}| < 0.01$ for the reservoir configuration or if $|\tilde{\rho}| < 0.02$ for the symmetric configuration, as long as an overlimiting bias is applied.

An error in the approximation is quantified by the average norm of relative residuals ($\Delta$) at a range of frequencies between $10^{-2}$ and $10^6$ spaced logarithmically with two points per decade:

$$\Delta = \frac{1}{N} \sum_{n=1}^{N} \left| \frac{\tilde{Z}(\tilde{\omega}_n) - \tilde{Z}_{TZ}(\tilde{\omega}_n)}{|\tilde{Z}(\tilde{\omega}_n)|} \right|,$$  

(56)

where $N = 17$. Figures 9(c) and 9(d) show $\Delta$ in a range of $\tilde{j}^{(0)}$ and $\tilde{\rho}$ for the reservoir and the symmetric configurations, respectively. $\Delta$ is relatively large above 0.1 in a region where $\tilde{j}^{(0)}$ is small and $|\tilde{\rho}|$ is large, when the two zones are not fully developed and $\tilde{\delta}$ is not defined. Then, it decreases quickly in
a direction of decreasing $|\tilde{\rho}|$ and increasing $\tilde{j}^{(0)}$. A small $\Delta$ ($<0.1$) is obtained even when the interface is somewhat thick with $\tilde{\delta}$ up to 0.2 at least. Therefore, it confirms that the two-zone approximation is valid as long as the two zones are fully developed and the interface is not too thick, which is achieved with a small $|\tilde{\rho}|$ and an overlimiting $\tilde{j}^{(0)}$.

VI. CONCLUSION

The impedance of electrodiffusion in the presence of a charged immobile species is studied by a theoretical model that combines the Nernst-Planck equations and the electroneutrality condition. In particular, its behavior is examined for a range of immobile charge density and applied bias, expanding to the overlimiting regime. The impedance of bulk electrodiffusion is commonly considered as a smaller contribution than those of interface kinetics. Under a dc bias, however, its magnitude increases exponentially, and it could be an important contribution to the overall cell impedance. In a biased condition, a bulk relaxation appears as a semicircle on the complex plane, which is attributed to perturbation in the bulk conductivity under the steady-state electric field.

As the applied dc bias approaches the diffusion limit, the diffusion current dominates in the bulk and the impedance exhibits a transition to a series connection of a resistor and a finite-length Warburg element. When there is no charged immobile species or when they are cocharged to the active charge carriers, the impedance diverges before the diffusion-limiting bias, and an overlimiting bias may not be sustained by electrodiffusion unless promoted by other mechanisms. On the other hand, when the immobile species is countercharged to the active charge carriers, an overlimiting bias can be applied. In the overlimiting regime with a further increasing bias, contribution of the resistor replaces that of the finite-length Warburg element, while the overall magnitude is kept finite by the constant conductivity provided by charge carriers screening the immobile species.

Under an overlimiting bias, two distinct regions are found in the steady-state solution with a sharp interphase. Based on this observation, we present an analytical approximation of the impedance where we assume complete dominance of conduction or diffusion in their respective zones. It attributes the resistor to the conduction zone and the finite-length Warburg element to the diffusion zone. The resistor is taking over the finite-length Warburg under increasing bias, because the conduction zone is replacing the diffusion zone. The approximation is valid when the immobile charge density is low and the bias is larger than the diffusion limit. Combined
with an interface model [68], the model and the approximation can be employed to interpret the impedance spectra of electrochemical cells with a charged immobile species under a dc bias even above the diffusion limit.

ACKNOWLEDGMENTS

The authors acknowledge support by a Martin Family Fellowship and a Kwanjeong Fellowship to J.S., the National Science Foundation (Ph.D.) funded by the Agency for Science, Technology and Research, Singapore (A*STAR) to E.K., and a Professor Amar G. Bose Research Grant to M.Z.B.

APPENDIX A: ANALYTICAL SOLUTIONS IN STEADY STATE

The steady-state solution can be obtained by solving the $O(1)$ terms in the perturbation. Collecting the $O(1)$ terms, the governing equations become

$$0 = \frac{d^2 \tilde{c}^{(0)}}{d \tilde{x}^2} - \tilde{\rho} \frac{d^2 \tilde{\phi}^{(0)}}{d \tilde{x}^2}, \quad (A1)$$

$$0 = \frac{d}{d \tilde{x}} \left[ (\tilde{c}^{(0)} - \tilde{\rho}) \frac{d \tilde{\phi}^{(0)}}{d \tilde{x}} \right]. \quad (A2)$$

Equations (A1) and (A2) can be integrated once by introducing the current ($\tilde{j}^{(0)}$), which is unknown at this stage but will be determined later for each configuration. Since the system is locally neutral, $\tilde{j}^{(0)}$ is constant throughout the domain. The current is entirely carried by the positive charge carrier everywhere in steady state, since the negative charge carrier is blocked at the boundary, $\tilde{x} = 1$. Then the total flux of charge carriers as well as the net charge current should be $\tilde{j}^{(0)}$ in $O(1)$ dimensionless form. Thereby

$$\tilde{j}^{(0)} = \frac{d \tilde{c}^{(0)}}{d \tilde{x}} - \tilde{\rho} \frac{d \tilde{\phi}^{(0)}}{d \tilde{x}}, \quad (A3)$$

$$\tilde{j}^{(0)} = (\tilde{c}^{(0)} - \tilde{\rho}) \frac{d \tilde{\phi}^{(0)}}{d \tilde{x}}. \quad (A4)$$

By subtracting Eq. (A4) from Eq. (A3), we recover the flux of the negative charge carriers, which should be zero:

$$0 = \frac{d \tilde{c}^{(0)}}{d \tilde{x}} - \tilde{c}^{(0)} \frac{d \tilde{\phi}^{(0)}}{d \tilde{x}}. \quad (A5)$$

Equations (A3) and (A5) are more straightforward to solve analytically.

For the reservoir configuration, we first integrate Eq. (A5) and apply the boundary condition at $\tilde{x} = 0$: $\tilde{c}^{(0)} = 1$ and $\tilde{\phi}^{(0)} = 0$. Then we obtain

$$\log(\tilde{c}^{(0)}) = \tilde{\phi}^{(0)}. \quad (A6)$$

Equation (A3) can be integrated using the boundary condition at $\tilde{x} = 0$ again:

$$1 - \tilde{j}^{(0)} \tilde{x} = \tilde{c}^{(0)} - \tilde{\rho} \tilde{\phi}^{(0)}. \quad (A7)$$

Then $\tilde{j}^{(0)}$ can be determined by the boundary condition at $\tilde{x} = 1, \tilde{\phi}^{(0)} = -\tilde{V}^{(0)}$:

$$\tilde{j}^{(0)} = 1 - (e^{-\tilde{V}^{(0)}} + \tilde{\rho} \tilde{V}^{(0)}). \quad (A8)$$

Equations (A6), (A7), and (A8) make up an implicit form of the steady-state solution for the reservoir configuration. They become explicit if $\tilde{\rho} = 0$. To obtain an explicit form for any values of $\tilde{\rho}$, we plug Eq. (A6) into Eq. (A7) and rearrange it to

$$\frac{1}{\tilde{\rho}} \exp \left( \frac{1 - \tilde{j}^{(0)} \tilde{x}}{\tilde{\rho}} \right) = \frac{\tilde{c}^{(0)}}{\tilde{\rho}} \exp \left( \frac{\tilde{c}^{(0)}}{\tilde{\rho}} \right). \quad (A9)$$

Then, we recognize that $(-e^{\tilde{V}^{(0)}}/\tilde{\rho})$ should be the Lambert W function, unless $\tilde{\rho} = 0$. Depending on the sign of $\tilde{\rho}$, a different branch gives a physically meaningful solution. When $\tilde{\rho} = 0$, Eq. (A7) gives the explicit solution for $\tilde{c}^{(0)}$. Collectively, we obtain Eq. (18) as for an explicit form of the steady-state solution for the reservoir configuration for any values of $\tilde{\rho}$.

On the other hand, for the symmetric configuration, we again integrate Eq. (A5). We introduce an unknown parameter $\alpha = \tilde{c}^{(0)}(\tilde{x} = 0)$, which will be determined later. By applying the boundary conditions at $\tilde{x} = 0$, $\tilde{c}^{(0)} = \alpha$, and $\tilde{\phi}^{(0)} = 0$, we obtain

$$\log \left( \frac{\tilde{c}^{(0)}}{\alpha} \right) = \tilde{\phi}^{(0)}. \quad (A10)$$

Equation (A3) can be integrated using the boundary conditions at $\tilde{x} = 0$ again:

$$\alpha - \tilde{j}^{(0)} \tilde{x} = \tilde{c}^{(0)} - \tilde{\rho} \tilde{\phi}^{(0)}. \quad (A11)$$

Then $\tilde{j}^{(0)}$ can be determined by the boundary condition at $\tilde{x} = 1, \tilde{\phi}^{(0)} = -\tilde{V}^{(0)}$:

$$\tilde{j}^{(0)} = \alpha (1 - e^{-\tilde{V}^{(0)}}) - \tilde{\rho} \tilde{V}^{(0)}. \quad (A12)$$

Upon determining $\alpha$, Eqs. (A10), (A11), and (A12) provide an implicit form of the steady-state solution for the symmetric configuration. They become explicit if $\tilde{\rho} = 0$. $\alpha$ can be determined by employing the integral constraint,

$$\int_0^1 \tilde{c}^{(0)} d \tilde{x} = \alpha \int_0^1 e^{-\tilde{V}^{(0)}} \left( \frac{d \tilde{\phi}^{(0)}}{d \tilde{x}} \right)^{-1} d \tilde{\phi}^{(0)} = 1, \quad (A13)$$

where $d \tilde{\phi}^{(0)}/d \tilde{x}$ can be obtained by combining Eqs. (A12) and (A3),

$$-\alpha e^{\tilde{V}^{(0)}} \frac{d \tilde{\phi}^{(0)}}{d \tilde{x}} + \tilde{\rho} \frac{d \tilde{\phi}^{(0)}}{d \tilde{x}} = \alpha (1 - e^{-\tilde{V}^{(0)}}) - \tilde{\rho} \tilde{V}^{(0)}, \quad (A14)$$

$$\frac{d \tilde{\phi}^{(0)}}{d \tilde{x}} = \frac{\alpha (1 - e^{-\tilde{V}^{(0)}}) - \tilde{\rho} \tilde{V}^{(0)}}{\tilde{\rho} - \alpha e^{\tilde{V}^{(0)}}}. \quad (A15)$$

By plugging Eq. (A15) into Eq. (A13) and rearranging the terms,

$$\alpha \int_0^1 (\tilde{\rho} e^{\tilde{V}^{(0)}} - \alpha e^{\til{\tilde{V}^{(0)}}}) d \tilde{\phi}^{(0)} = \alpha (1 - e^{-\til{\tilde{V}^{(0)}}}) - \til{\til{\rho}} \til{\til{V}^{(0)}}. \quad (A16)$$

Upon performing the integrations, Eq. (A16) becomes a quadratic equation for $\alpha$:

$$1 - e^{-2\til{\til{V}^{(0)}}} \alpha^2 + 2(1 + \til{\til{\rho}})(e^{-\til{\til{V}^{(0)}}} - 1) \alpha + 2\til{\til{\rho}} \til{\til{V}^{(0)}} = 0. \quad (A17)$$
Only one of the roots provides a physically meaningful solution. Therefore,
\[
\alpha = \frac{(1 + \tilde{\rho})(1 - e^{-\tilde{V}(0)})}{1 - e^{-2\tilde{V}(0)}} + \sqrt{(1 + \tilde{\rho})^2 (1 - e^{-\tilde{V}(0)})^2 - 2\tilde{\rho} \tilde{V}(0)(1 - e^{-2\tilde{V}(0)})}. 
\]
(A18)

To obtain an explicit form for any values of \( \tilde{\rho} \), we plug Eq. (A10) into Eq. (A11) and rearrange to
\[
\alpha = \frac{\alpha - \tilde{\jmath}(0)}{-\tilde{\rho}} \exp \left( \frac{\tilde{\alpha}(0)}{-\tilde{\rho}} \right). \tag{A19}
\]

The same as the reservoir configuration, we recognize that \((-\tilde{\alpha}(0)/\tilde{\rho})\) should be the Lambert W function, unless \( \tilde{\rho} = 0 \). Depending on the sign of \( \tilde{\rho} \), a different branch gives a physically meaningful solution. When \( \tilde{\rho} = 0 \), Eq. (A11) gives the explicit solution for \( \tilde{\alpha}(0) \). Collectively, we obtain Eq. (21) as for an explicit form of the steady-state solution for the symmetric configuration for any values of \( \tilde{\rho} \).

**APPENDIX B: SMALL-BIAS LIMITS OF \( \tilde{R}_L \) AND \( \tilde{R}_H \)**

Under a small bias, \( \tilde{\phi}(\tilde{x}) \) can be approximated by a linear function. To satisfy the boundary conditions for the reservoir configuration,
\[
\tilde{\alpha}(0)(\tilde{\jmath}(0) \ll 1) \simeq 1 - \frac{\tilde{\jmath}(0)}{1 - \tilde{\rho}} \tilde{x}. \tag{B1}
\]

In the low-frequency limit, the impedance should converge to the local resistance in steady state, i.e., \( \tilde{R}_L = d\tilde{V}(0)/d\tilde{\jmath}(0) \). Under a small bias, \( \tilde{\phi}(0)(\tilde{x}) \) is obtained by integrating the \( O(1) \) terms of Eq. (15): \(-\tilde{\phi}(0)(\tilde{x}) = (\tilde{\omega}(0) - \tilde{\rho})d\tilde{\phi}(0)/d\tilde{x} \), using \( \tilde{\phi}(0) \) in Eq. (B1) and a boundary condition, \( \tilde{\phi}(0) = 0 \) at \( \tilde{x} = 0 \):
\[
\tilde{\phi}(0)(\tilde{\jmath}(0) \ll 1) \simeq (1 - \tilde{\rho}) \log \left( \frac{(1 - \tilde{\rho})^2 - \tilde{\jmath}(0)\tilde{x}}{(1 - \tilde{\rho})^2} \right). \tag{B2}
\]

Then \( \tilde{V}(0) \) is obtained by \(-\tilde{\phi}(0)\) at \( \tilde{x} = 1 \):
\[
\tilde{V}(0)(\tilde{\jmath}(0) \ll 1) \simeq -(1 - \tilde{\rho}) \log \left( \frac{(1 - \tilde{\rho})^2 - \tilde{\jmath}(0)}{(1 - \tilde{\rho})^2} \right). \tag{B3}
\]

Therefore,
\[
\tilde{R}_L(\tilde{\jmath}(0) \ll 1) \simeq \frac{(1 - \tilde{\rho})}{(1 - \tilde{\rho})^2 - \tilde{\jmath}(0)}. \tag{B4}
\]

In the high-frequency limit, \( \tilde{\alpha}(1) \) does not respond to the perturbation, and \( \tilde{\phi}(1) \) is obtained by solving the \( O(\tilde{\rho}) \) terms of Eq. (15) with \( \tilde{\phi}(1) = 0 \): \(-\tilde{\phi}(1)(\tilde{x}) = (\tilde{\omega}(0) - \tilde{\rho})d\tilde{\phi}(1)/d\tilde{x} \). Then, evaluating \(-\tilde{\phi}(1)\) at \( \tilde{x} = 1 \), \( \tilde{V}(1) \) becomes
\[
\tilde{V}(1)(\tilde{\jmath}(0) \ll 1) \simeq -\frac{\tilde{\jmath}(1)(1 - \tilde{\rho})}{\tilde{\jmath}(0)} \log \left( \frac{(1 - \tilde{\rho})^2 - \tilde{\jmath}(0)}{(1 - \tilde{\rho})^2} \right). \tag{B5}
\]

Then, \( \tilde{R}_H(\tilde{\jmath}(0) \ll 1) \) is obtained by
\[
\tilde{R}_H(\tilde{\jmath}(0) \ll 1) = \frac{\tilde{V}(1)(\tilde{\jmath}(0) \ll 1)}{\tilde{\jmath}(1)} \simeq -\frac{(1 - \tilde{\rho})}{\tilde{\jmath}(0)} \log \left( \frac{(1 - \tilde{\rho})^2 - \tilde{\jmath}(0)}{(1 - \tilde{\rho})^2} \right). \tag{B6}
\]

The same approach can be applied to the symmetric configuration under a small bias \( (\tilde{\jmath}(0) \ll 2) \). First, \( \tilde{\phi}(0)(\tilde{x}) \) is approximated by a linear function,
\[
\tilde{\phi}(0)(\tilde{\jmath}(0) \ll 2) \simeq \alpha - \frac{\tilde{\jmath}(0)}{1 - \tilde{\rho}} \tilde{x}. \tag{B7}
\]

where \( \alpha \) can be obtained as a function of \( \tilde{\jmath}(0) \) employing the integral constraint in Eq. (14): \( \alpha(\tilde{\jmath}(0) \ll 2) \simeq 1 + \tilde{\omega}(2 - 2\tilde{\rho}) \). Then \( \tilde{\phi}(0)(\tilde{x}) \) and \( \tilde{V}(0)(\tilde{\jmath}(0) \ll 2) \) are obtained:
\[
\tilde{V}(0)(\tilde{\jmath}(0) \ll 2) \simeq -(1 - \tilde{\rho}) \log \left( \frac{-(\tilde{\jmath}(0) + 2(1 - \tilde{\rho})^2}{\tilde{\jmath}(0) + 2(1 - \tilde{\rho})^2} \right). \tag{B8}
\]

\( \tilde{R}_L \) is obtained by \( d\tilde{V}(0)/d\tilde{\jmath}(0) \). For the symmetric condition under a small bias,
\[
\tilde{R}_L(\tilde{\jmath}(0) \ll 2) \simeq \frac{(1 - \tilde{\rho})}{2(1 - \tilde{\rho})^2 - \tilde{\jmath}(0)} + \frac{(1 - \tilde{\rho})}{2(1 - \tilde{\rho})^2 + \tilde{\jmath}(0)}. \tag{B10}
\]

Then to obtain \( \tilde{R}_H(\tilde{\jmath}(0) \ll 2) \) is obtained by solving \(-\tilde{\phi}(1)(\tilde{x}) = (\tilde{\omega}(0) - \tilde{\rho})d\tilde{\phi}(1)/d\tilde{x} \) for \( \tilde{\phi}(1) \) and evaluating it at \( \tilde{x} = 1 \):
\[
\tilde{V}(1)(\tilde{\jmath}(0) \ll 2) \simeq -\frac{\tilde{\jmath}(1)(1 - \tilde{\rho})}{\tilde{\jmath}(0)} \log \left( \frac{-\tilde{\jmath}(0) + 2(1 - \tilde{\rho})^2}{\tilde{\jmath}(0) + 2(1 - \tilde{\rho})^2} \right). \tag{B11}
\]

\( \tilde{R}_H(\tilde{\jmath}(0) \ll 2) \) is obtained by \( \tilde{V}(1)/\tilde{\jmath}(1) \):
\[
\tilde{R}_H(\tilde{\jmath}(0) \ll 2) \simeq - \frac{(1 - \tilde{\rho})}{\tilde{\jmath}(0)} \log \left( \frac{-(\tilde{\jmath}(0) + 2(1 - \tilde{\rho})^2}{\tilde{\jmath}(0) + 2(1 - \tilde{\rho})^2} \right). \tag{B12}
\]

The small-bias asymptotic expressions of \( \tilde{R}_L \) and \( \tilde{R}_H \) are compared to the numerical limits of the full model in Fig. 10. For both configurations, the asymptotic expressions show good agreement in the small-bias regime, even close to the limiting current.

**APPENDIX C: LARGE-BIAS LIMITS OF \( \tilde{R}_L \) AND \( \tilde{R}_H \)**

Asymptotic expressions under a large bias above the diffusion limit can be obtained from the two-zone approximation in Sec. V. \( \tilde{R}_L \) is again obtained by the local resistance in the steady-state approximation. For the reservoir configuration, \( \phi_{Tz}(0) \) in Eq. (47) is evaluated at \( \tilde{x} = 1 \):
\[
\tilde{V}(0)(\tilde{\jmath}(0) \gg 1) \simeq -\frac{\tilde{\jmath}(0)}{\tilde{\rho}}(1 - \tilde{\jmath}(0)). \tag{C1}
\]
In the high-frequency limit, the diffusion zone contributes little to the impedance. Therefore, $\tilde{R}_H$ comes only from the impedance of the conduction zone, $\tilde{Z}_CZ$. Therefore, 

$$\tilde{R}_H \approx \frac{L_0}{\tilde{\rho}} = \frac{1}{\tilde{\rho}} \left( \frac{j^{(0)}}{\tilde{\rho}} - 1 \right).$$  \hspace{1cm} (C3)

The same approach can be applied to the symmetric configuration under a large bias ($\tilde{V}^{(0)} \gg 2$). Evaluating $\tilde{\Phi}_{TZ}^{(0)}$ in Eq. (53) at $\tilde{x} = 1$,

$$\tilde{V}^{(0)}(\tilde{V}^{(0)} \gg 2) \approx -\frac{j^{(0)}}{\tilde{\rho}} (1 - \tilde{I}_D),$$  \hspace{1cm} (C4)

where $\tilde{I}_D = \sqrt{2/\tilde{V}^{(0)}}$ in the approximation. Then, $d\tilde{V}^{(0)}/d\tilde{j}^{(0)}$ leads to

$$\tilde{R}_L(\tilde{j}^{(0)} \gg 2) \approx \frac{1}{\tilde{\rho}} \left( 1 - \frac{1}{2 \tilde{j}^{(0)}} \right).$$  \hspace{1cm} (C5)

$\tilde{R}_H$ comes only from $\tilde{Z}_CZ$. For the symmetric configuration,

$$\tilde{R}_H(\tilde{j}^{(0)} \gg 2) \approx \frac{1}{\tilde{\rho}} \left( \tilde{j}^{(0)} - \frac{2\tilde{j}^{(0)}}{\tilde{j}^{(0)}} \right).$$  \hspace{1cm} (C6)

The large-bias asymptotic expressions of $\tilde{R}_L$ and $\tilde{R}_H$ are compared to the numerical limits of the full model in Fig. 10. Like the small-bias expressions, the large-bias expressions show good agreement in the overlimiting regime, even close to the limiting current when $\tilde{\rho} = -0.01$. Since they are based on the two-zone approximation (Sec. V), they are valid when $|\tilde{\rho}|$ is low and the bias is above the diffusion limit.

---

[1] M. Planck, Annalen der Physik 275, 161 (1890).
[2] W. Nernst, Zeitschrift für Physikalische Chemie 2U, 613 (1888).
[3] W. Lai and F. Ciucci, Electrochimica Acta 56, 4369 (2011).
[4] A. Z. Weber, R. L. Borup, R. M. Darling, P. K. Das, T. J. Dursch, W. Gu, D. Harvey, A. Kasoglu, S. Litster, and M. M. Mench, J. Electrochem. Soc. 161, F1254 (2014).
[5] W. V. Roosbroeck, Bell Syst. Tech. J. 29, 560 (1950).
[6] J. Bisquert, L. Bertoluzzi, I. Mora-Sero, and G. Garcia-Belmonte, J. Phys. Chem. C 118, 18983 (2014).
[7] I. Rubinstein, Electro-Diffusion of Ions (Society for Industrial and Applied Mathematics, Philadelphia, 1990), Vol. 11.
[8] K. H. Meyer and J. F. Sievers, Helv. Chim. Acta 19, 649 (1936).
[9] T. Teorell, Proc. Soc. Exp. Biol. Med. 33, 282 (1935).
[10] T. Teorell, Prog. Biophys. Biophys. Chem. 3, 305 (1953).
[11] R. S. Eisenberg, Acc. Chem. Res. 31, 117 (1998).
[12] R. S. Eisenberg, J. Membr. Biol. 150, 1 (1996).
[13] R. S. Eisenberg, in Recent Developments in Theoretical Studies of Proteins, edited by R. Elber, Advanced Series in Physical Chemistry Vol.7 (World Scientific, Singapore, 1996), p. 269.
[14] G. J. Newman, D. Bennon, and C. W. Tobias, Berichte der Bunsengesellschaft für Physikalische Chemie 69, 608 (1965).
[15] R. P. Buck, J. Membr. Sci. 17, 1 (1984).
[16] M. Z. Bazant, K. Thornton, and A. Ajdari, Phys. Rev. E 70, 021506 (2004).
[17] E. V. Dydek and M. Z. Bazant, AIChE J. 59, 3539 (2013).
[18] E. V. Dydek, B. Zaltzman, I. Rubinstein, D. S. Deng, A. Mani, and M. Z. Bazant, Phys. Rev. Lett. 107, 118301 (2011).
[19] D. R. Franceschetti, J. R. Macdonald, and R. P. Buck, J. Electrochem. Soc. 138, 1368 (1991).
[20] M. Z. Bazant, K. T. Chu, and B. Bayly, SIAM J. Appl. Math. 65, 1463 (2005).
[21] J. R. Macdonald, J. Chem. Phys. 58, 4982 (1973).
[22] M. Z. Bazant, Electrochemical Energy Systems, Massachusetts Institute of Technology: MIT OpenCourseWare, https://ocw.mit.edu/, Spring 2014.
[23] J. R. Macdonald, J. Chem. Phys. 54, 2026 (1971).
[24] J. R. Macdonald, J. Electroanal. Chem. Interfacial Electrochem. 32, 317 (1971).
[25] J. R. Macdonald, L. R. Evangelista, E. K. Lenzi, and G. Barbero, J. Phys. Chem. C 115, 7648 (2011).
[26] J. R. Macdonald, Phys. Rev. 92, 4 (1953).
[27] J. R. Macdonald, J. Chem. Phys. 61, 3977 (1974).
[28] T. R. Brumleve and R. P. Buck, J. Electroanal. Chem. Interfacial Electrochem. 126, 73 (1981).
[29] R. Buck, J. Electroanal. Chem. Interfacial Electrochem. 23, 219 (1969).
[30] R. P. Buck and C. M. Munday, Electrochimica Acta 44, 1999 (1999).
[31] J. Jannik and J. Maier, J. Electrochem. Soc. 146, 4183 (1999).
[32] J. Jamnik and J. Maier, Phys. Chem. Chem. Phys. 3, 1668 (2001).
[33] J. Jamnik, J. Maier, and S. Pejovnik, Electrochim. Acta 44, 4139 (1999).
[34] A. Moya, Electrochim. Acta 62, 296 (2012).
[35] J. Horno, A. Moya, and C. González-Fernández, J. Electroanal. Chem. 402, 73 (1996).
[36] A. Moya and J. Horno, J. Phys. Chem. B 103, 10791 (1999).
[37] I. Rubinstein and B. Zaltzman, Phys. Rev. E 80, 021505 (2009).
[38] G. Barbero, J. J. da Silva, and A. M. Figueiredo Neto, J. Appl. Phys. 101, 113705 (2007).
[39] W. Lai and S. M. Haile, J. Am. Ceram. Soc. 88, 2979 (2005).
[40] R. P. Buck, J. Electroanal. Chem. Interfacial Electrochem. 210, 1 (1986).
[41] G. Barbero, Phys. Chem. Chem. Phys. 19, 32575 (2017).
[42] A. Mauro, Biophys. J. 2, 179 (1962).
[43] F. G. Helfferich, Ion Exchange (McGraw-Hill, New York, 1962).
[44] B. Eisenberg and W. Liu, SIAM J. Math. Anal. 38, 1932 (2007).
[45] W. N. Green and O. S. Andersen, Annu. Rev. Physiol. 53, 341 (1991).
[46] M. Schmuck and M. Z. Bazant, SIAM J. Appl. Math. 75, 1369 (2015).
[47] W. Liu, SIAM J. Appl. Math. 65, 754 (2005).
[48] P. A. Markowich, SIAM J. Appl. Math. 44, 896 (1984).
[49] A. Singer, D. Gillespie, J. Norbury, and R. S. Eisenberg, Eur. J. Appl. Math. 19, 541 (2008).
[50] E. Khoo and M. Z. Bazant, J. Electroanal. Chem. 811, 105 (2018).
[51] P. M. Biesheuvel, Y. Fu, and M. Z. Bazant, Phys. Rev. E 83, 061507 (2011).
[52] A. Mani, T. A. Zangle, and J. G. Santiago, Langmuir 25, 3898 (2009).
[53] T. A. Zangle, A. Mani, and J. G. Santiago, Langmuir 25, 3909 (2009).
[54] T. A. Zangle, A. Mani, and J. G. Santiago, Chem. Soc. Rev. 39, 1014 (2010).
[55] T. A. Zangle, A. Mani, and J. G. Santiago, Anal. Chem. 82, 3114 (2010).
[56] A. Moya, J. Phys. Chem. C 118, 2539 (2014).
[57] A. Moya, A. Hayas, and J. Horno, Solid State Ionics 130, 9 (2000).
[58] A. Moya and P. Sistat, J. Electrochem. Soc. 166, H25 (2019).
[59] S. Nam, I. Cho, J. Heo, G. Lim, M. Z. Bazant, D. J. Moon, G. Y. Sung, and S. J. Kim, Phys. Rev. Lett. 114, 114501 (2015).
[60] A. Mani and M. Z. Bazant, Phys. Rev. E 84, 061504 (2011).
[61] A. Yaroschuk, Adv. Colloid Interface Sci. 183, 68 (2012).
[62] D. Yan, M. Z. Bazant, P. M. Biesheuvel, M. C. Pugh, and F. P. Dawson, Phys. Rev. E 95, 033303 (2017).
[63] J. Schiffbauer, U. Liel, and G. Yossifon, Phys. Rev. Lett. 110, 204504 (2013).
[64] J. Schiffbauer, U. Liel, and G. Yossifon, Phys. Rev. E 89, 033017 (2014).
[65] P. B. Peters, R. van Roij, M. Z. Bazant, and P. M. Biesheuvel, Phys. Rev. E 93, 053108 (2016).
[66] C. P. Nielsen and H. Bruus, Phys. Rev. E 90, 043020 (2014).
[67] A. MacGillivray, J. Chem. Phys. 48, 2903 (1968).
[68] E. Khoo, J. Song, and M. Z. Bazant (unpublished).
[69] E. Petersen, AIChE J. 4, 343 (1958).
[70] A. Levy and M. Z. Bazant, arXiv:1905.05789.
[71] Z.-X. Luo, Y.-Z. Xing, Y.-C. Ling, A. Kleinhammes, and Y. Wu, Nat. Commun. 6, 6358 (2015).
[72] J. S. Newman and C. W. Tobias, J. Electrochem. Soc. 109, 1183 (1962).
[73] J. R. Macdonald, J. Phys. Chem. A 115, 13370 (2011).
[74] G. Barbero, A. Gliozzi, M. Scalerandi, and A. Scarfone, J. Mol. Liq. 272, 565 (2018).
[75] R. M. Corless, G. H. Gonnet, D. E. G. Hare, D. J. Jeffrey, and D. E. Knuth, Adv. Comput. Math. 5, 329 (1996).
[76] See Supplemental Material at http://link.aps.org/supplemental/10.1103/PhysRevE.100.042204 for attached MATLAB scripts that solve the steady-state and impedance problems.