Poisson–Fermi model of single ion activities in aqueous solutions

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A B S T R A C T

A Poisson–Fermi model is proposed for calculating activity coefficients of single ions in strong electrolyte solutions based on the experimental Born radii and hydration shells of ions in aqueous solutions. The steric effect of water molecules and interstitial voids in the first and second hydration shells play an important role in our model. The screening and polarization effects of water are also included in the model that can thus describe spatial variations of dielectric permittivity, water density, void volume, and ionic concentration. The activity coefficients obtained by the Poisson–Fermi model with only one adjustable parameter are shown to agree with experimental data, which vary nonmonotonically with salt concentrations.

1. Introduction

Comprehensive discussions of theoretical and experimental studies on the activity coefficient of single ions in electrolyte solutions have been recently given by Fraenkel [1], Valkó and Boda [2], and Rowland et al. [3], where more references can also be found. The Poisson–Fermi (PF) model proposed in this letter belongs to the continuum approach that traces back to the simple, elegant, but very coarse theory—the Debye–Hückel (DH) theory. As mentioned by Fraenkel, the continuum theory has evolved in the past century into a series of modified Poisson–Boltzmann (PB) equations that can involve an overwhelmingly large number of parameters in order to fit Monte Carlo (MC), molecular dynamics (MD), or experimental data. Many expressions of those parameters are rather long and tedious and do not have clear physical meaning [1].

The Debye–Hückel model is derived from a linearized PB equation [4]. Extended from the DH model, the Pitzer model [5] is most widely used to describe the thermodynamic properties of multicomponent electrolyte solutions due to its unmatched precision over wide ranges of temperature and pressure [3]. However, the combinatorial explosion of adjustable parameters in the extended DH modeling functions (including Pitzer) can cause profound difficulties in fitting experimental data and independent verification because the parameters are very sensitive to numerous related thermodynamic properties in multicomponent systems [3]. The Poisson–Fermi model proposed here involves only one adjustable parameter.

The ineffectiveness of previous Poisson–Boltzmann models is mainly due to inaccurate treatments of the steric and correlation effects of ions and water molecules whose nonuniform charges and sizes can have significant impact on the activities of all particles in an electrolyte system. Unfortunately, the point charge particles of PB theories have electric fields that are most approximate where they are largest, near the point. PB theories are not an appealing choice for the leading terms in a series of approximations, for that reason. The PF theory developed in our papers [6–10] demonstrates how these two effects can be described by a simple steric potential and a correlation length of ions. The parameters of the PF theory describe distinct physical properties of the system in a clear way [9]. The Gibbs–Fermi free energy of the PF model reduces to the classical Gibbs free energy of the PB model when the steric potential and correlation length are omitted [9]. The PF model has been verified with either MC, MD or double layer data at (more or less) equilibrium [6–8], and nonequilibrium data from calcium and gramicidin channels [9,10].

Here, we apply the PF theory to study the activity properties of individual ions in strong electrolytes. The steric effect of all particles and the interstitial voids that accompany them are described by a Fermi-like distribution that defines the water densities in the hydration shell of a solvated ion and the particle concentrations in the solvent region outside the hydration shell. The resulting correlations produce a dielectric function that shows variations in permittivity around the solvated ion. The experimental concentration-dependent dielectric constant model proposed in [2] is used to define the concentration-dependent Born radii of...
the solvated ion in the present work. The experimental data of the activity coefficients of NaCl and CaCl₂ electrolytes reported in [11] are used to test the PF model.

2. Theory

The activity coefficient \( \gamma_i \) of an ion of species \( i \) in electrolyte solutions describes the deviation of the chemical potential of the ion from ideality \( (\gamma_i = 1) \). The excess chemical potential is \( \mu_i^{ex} = k_B T \ln \gamma_i \), where \( k_B \) is the Boltzmann constant and \( T \) is an absolute temperature. In Poisson–Boltzmann theory, the excess chemical potential can be calculated by [12]

\[
\mu_i^{ex} = \Delta G_i^\text{PB} - \Delta G_i^0, \quad \Delta G_i^\text{PB} = \frac{1}{2} q_i^2 \phi_i^\text{PB}(0), \quad \Delta G_i^0 = \frac{1}{2} q_i^2 \phi_i^0(0),
\]

where the center of the hydrated ion (also denoted by \( i \)) is set to the origin \( 0 \) for convenience in the following discussion and \( q_i \) is the ionic charge. The potential function \( \phi_i^\text{PB}(r) \) of spatial variable \( r \) is found by solving the Poisson–Boltzmann equation

\[
-e_\varepsilon \nabla \cdot \nabla \phi_i^\text{PB}(r) = -\sum_{j=1}^{K} q_j C_j(r) = \rho(r),
\]

(2)

\[
C_j(r) = C_j^B \exp(-\beta_j \phi_i^\text{PB}(r)),
\]

(3)

where the concentration function \( C_j(r) \) is described by a Boltzmann distribution in (3) with a constant bulk concentration \( C_j^B \), \( \beta_j = e_{w} / e_{w0} \), \( e_{w} \) is the dielectric constant of bulk solvent, and \( e_{w0} \) is the vacuum permittivity. The potential \( \phi_i^\text{PB}(r) \) of the ideal system is obtained by setting \( \rho(r) = 0 \) in (2), i.e., all ions of \( K \) species in the system do not electrostatically interact with each other since \( q_j = 0 \) for all \( j \). We consider a large domain \( \Omega \) of the system in which \( \phi_i^\text{PB}(r) = 0 \) on the boundary of the domain \( \partial \Omega \). The ideal potential \( \phi_i^\text{PB}(r) \) is then a constant, i.e., \( \Delta G_i^0 \) is a constant reference chemical potential independent of \( C_j^B \).

For an equivalent binary system, the Debye–Hückel theory simplifies the calculation by analytically solving a linearized equation of (2) so that the potential function \( \phi_i^\text{DH}(r) \) becomes a constant [4]

\[
\phi_i^{\text{DH}} = -\frac{q_i K_i}{4 \pi e_\varepsilon} \frac{1}{k_B T} \left( \sum_{j=1}^{K} q_j C_j(r) \right)^{1/2},
\]

(4)

de pending of the bulk concentration \( C_j^B \), with \( l \) is the Avogadro constant.

The Poisson–Fermi equation proposed in [9] is

\[
e_\varepsilon \left( \frac{\varepsilon}{\varepsilon} \nabla^2 \right) - \nabla \cdot \nabla \phi_i^{\text{PF}}(r) = -\sum_{j=1}^{K} q_j C_j(r) = \rho(r), \quad \forall r \in \Omega,
\]

(5)

\[
C_j(r) = C_j^B \exp(-\beta_j \phi_i^{\text{PF}}(r) + S^{\text{intr}}(r)), \quad S^{\text{intr}}(r) = \ln \frac{\Gamma(r)}{\Gamma_i^B},
\]

(6)

where \( S^{\text{intr}}(r) \) is the ionic potential, \( \Gamma(r) = 1 - \sum_{j=1}^{K+1} \nu_j C_j(r) \) is a void fraction function, \( \Gamma_i^B = 1 - \sum_{j=1}^{K+1} \nu_j C_j^B \) is a constant void fraction, and \( \nu_j \) is the volume of a species \( j \) particle (hard sphere). Note that the PF equation includes water as the last species of particles without the zero charge \( q_{K+1} = 0 \). The polarization of the water and solution is an output of the theory. The water can be described more realistically, for example, as a quadrupole in later versions of the theory. The distribution (6) is of Fermi type since all concentration functions are bounded above, i.e., \( C_j(r) < 1/\nu_j \) for all particle species with any arbitrary (or even infinite) potential \( \phi(r) \) at any location \( r \) in the domain \( \Omega \) [9]. The Boltzmann distribution (3) would however diverge if \( \phi(r) \) tends to infinity. This is a major deficiency of PB theory for modeling a system with strong local electric fields or interactions. The PF Eq. (5) and the Fermi distribution reduce to the PB Eq. (2) and the Boltzmann distribution (3), respectively, when \( l = S^{\text{intr}} = 0 \), i.e., when the correlation and steric effects are not considered.

If the correlation length \( l \gg a_0 \neq 0 \), the dielectric operator \( \varepsilon = \varepsilon_0 (1 - \varepsilon^2 \nabla^2) \) approximates the permittivity of the bulk solvent and the linear response of correlated ions [6,7,13,14], where \( a_i \) is the radius of the ion. The dielectric function \( \varepsilon(r) = \varepsilon_0 (1 + \eta / \rho) \) is a further approximation of \( \varepsilon \). It is found by transforming (5) into two second-order PDEs [6]

\[
e_\varepsilon \left( \frac{\varepsilon}{\varepsilon^2} - 1 \right) \nabla^2 \phi_i^{\text{PF}}(r) = \rho(r)
\]

(7)

\[
\nabla^2 \phi_i^{\text{PF}}(r) = \Psi(r)
\]

(8)

by introducing a density like variable \( \Psi \) that yields a polarization charge density \( \eta = -e_\varepsilon \Psi - \rho \) of water using Maxwell’s first equation [7]. Boundary conditions of the new variable \( \Psi \) on the boundary \( \partial \Omega \) were derived from the global charge neutrality condition [6].

To obtain more accurate potentials at the origin \( 0 \), i.e., \( \phi_i^{\text{PF}}(0) \), we need to consider the size and hydration shell of the hydrated ion \( l \). The domain \( \Omega \) is partitioned into three parts such that \( \Omega = \Omega_{\text{ion}} \cup \Omega_{\text{Sh}} \cup \Omega_{\text{Solv}} \), where \( \Omega_{\text{ion}} \) is the spherical domain occupied by the ion \( l \), \( \Omega_{\text{Sh}} \) is the hydration shell of the ion, and \( \Omega_{\text{Solv}} \) is the rest of the solvent domain as shown in Figure 1. The radii of \( \Omega_{\text{ion}} \) and the outer boundary of \( \Omega_{\text{Sh}} \) are denoted by \( R_{\text{Born}} \) and \( R_S \), respectively, whose values will be determined by experimental data. It is natural to choose the Born radius \( R_{\text{Born}} \) as the radius of \( \Omega_{\text{ion}} \) [12]. We consider both first and second shells of the ion [15,16], the dielectric constants in \( \Omega_{\text{ion}} \) and \( \Omega_{\text{Sh}} \) are denoted by \( \varepsilon_{\text{ion}} \) and \( \varepsilon_{\text{w}} \), respectively.

The PF Eq. (5) then becomes

\[
\varepsilon_0 \left( \frac{\varepsilon_0^2}{\varepsilon_0^2} - 1 \right) \nabla^2 \phi_i^{\text{PF}}(r) = \rho(r)
\]

(9)

\[
\varepsilon_0 \left( \frac{\varepsilon_0^2}{\varepsilon_0^2} - 1 \right) \nabla^2 \phi_i^{\text{PF}}(r) = \rho(r)
\]

\[
\phi_i^{\text{PF}}(r) = \sum_{j=1}^{K+1} q_j C_j(r) \text{ in } \Omega_{\text{ion}}
\]

where \( \delta(r-0) \) is the delta function at the origin, \( l = 0 \) in \( \Omega_{\text{ion}} \), \( l \neq 0 \) in \( \Omega_{\text{Sh}} \), \( \varepsilon_0 = \varepsilon_{\text{ion}} \varepsilon_0 \) in \( \Omega_{\text{ion}} \), and \( \varepsilon = \varepsilon_{\text{ion}} + \varepsilon_{\text{w}} \varepsilon_0 \) in \( \Omega_{\text{Sh}} \). The shell radius \( R_S \) is determined by Eq. (6) as

\[
S^{\text{Sh}}_{\text{Sh}} = \ln \left( \frac{V_{\text{Sh}} - v_{\text{w}} O^w}{V_{\text{Sh}} \Gamma_0^B} \right) = \ln \left( \frac{O^w}{V_{\text{Sh}} \Gamma_0^B} \right) \Rightarrow V_{\text{Sh}} = \frac{\Gamma_0^B}{\varepsilon_{\text{w}}} O^w + v_{\text{w}} O^w
\]

(10)

where \( v_{\text{w}} \) is the volume of a water molecule and \( V_{\text{Sh}} \) is the volume of the hydration shell that depends on the bulk void fraction \( \Gamma_0^B \).
the bulk water density $c_{\text{w}}$, and the total number $q_{\text{i}}^\Omega$ (coordination number) of water molecules occupying the shell of the hydrated ion $i$. Note that the shell volume $V_{\text{sh}}$ varies with bulk ionic concentrations $c_{\text{i}}$. The occupancy number $q_{\text{i}}^\Omega$ is given by experimental data [15,16] and so is the shell volume that of course determines the shell radius $R_{\text{i}}^\text{sh}$.

To deal with the singular problem of the delta function $\delta(\mathbf{r} - \mathbf{0})$ in Eq. (9), we use the numerical methods proposed in [6] to calculate $\phi^\Omega(\mathbf{r})$ as follows:

(i) Solve the Laplace equation $\nabla^2 \phi^\Omega(\mathbf{r}) = 0$ in $\Omega_{\text{ion}}$ with the boundary condition $\phi^\Omega(\mathbf{r}) = \phi^i(\mathbf{r}) = q_i/(4\pi \varepsilon_{\text{ion}} \varepsilon_0 | \mathbf{r} - \mathbf{0} |)$ on $\partial \Omega_{\text{ion}}$.

(ii) Solve the Poisson–Fermi Eq. (9) in $\Omega \setminus \Omega_{\text{ion}}$ with the jump condition $\left[ \mathbf{e} \cdot \nabla \phi^\Omega(\mathbf{r}) \cdot \mathbf{n} \right] = - \phi^\Omega(\mathbf{r}) + \phi^i(\mathbf{r})$ on $\partial \Omega_{\text{ion}}$ and the zero boundary condition $\phi^\Omega(\mathbf{r}) = 0$ on $\partial \Omega$, where $\mathbf{n}$ denotes the jump function across $\partial \Omega_{\text{ion}}$ [6].

The evaluation of the Green function $\phi^\Omega(\mathbf{r})$ on $\partial \Omega_{\text{ion}}$ always yields finite numbers and thus avoids the singularity. Note that our model can be applied to electrolyte solutions at any temperature $T$ having any arbitrary number $(K)$ of ionic species with different size spheres and valences.

3. Results

Numerical values of model notations are given in Table 1, where the occupancy number $q_{\text{i}}^\Omega = 18$ is taken to be the experimental coordination number of the calcium ion $\text{Ca}^{2+}$ given in [15] for all ions $i = \text{Na}^+$, $\text{Ca}^{2+}$, and $\text{Cl}^-$ since the electric potential produced by the solvated ion diminishes exponentially in the outer shell region in which a small variation of $q_{\text{i}}^\Omega$ for $i = \text{Na}^+$ and $\text{Cl}^-$ does not affect numerical approximations too much. Obviously the coordination number may be different for different types of ions and at different concentrations and so on. We were surprised that we can fit experimental data so well using a single experimentally determined occupancy number for all ions and conditions.

As discussed in [2], the solvation free energy of an ion $i$ should vary with salt concentrations and can be expressed by a dielectric constant $\epsilon(c_{\text{i}})$ that depends on the bulk concentration of the ion $c_{\text{i}}$. Following [2], we assume that

$$\epsilon(c_{\text{i}}) = \epsilon_w - \delta_i c_{\text{i}} + (c_{\text{i}})^{3/2} \tag{11}$$

with only one parameter $\delta_i$, whose value is given in Table 1, instead of two in [2]. Note that $\epsilon(c_{\text{i}})$ is a constant when the dimensionless $c_{\text{i}}$ is given. It is not a function of a spatial variable $\mathbf{r}$ like $\epsilon(\mathbf{r})$. The parameter $\delta_i$ represents the ratio of the factor of $c_{\text{i}}$ to that of $(c_{\text{i}})^{3/2}$ in the original formula, where the factors of various electrolytes are taken from various sources of either theoretical or experimental data [2]. Our ratios $\delta_i$ in Table 1 are comparable with those given in [2].

The Born formula of the solvation energy can thus be modified as

$$\Delta \epsilon_i^{\text{Born}}(c_{\text{i}}) = \frac{q_i^2}{8\pi \varepsilon_0 \delta_i c_{\text{i}}^{3/2}} \left( \frac{1}{\epsilon_w} - 1 \right),$$

$$\theta(c_{\text{i}}) = \frac{\epsilon(c_{\text{i}})^{3/2}}{\epsilon_w (\epsilon(c_{\text{i}}) - 1)}, \tag{12}$$

where $R_{\text{i}}^0$ is the Born radius when $c_{\text{i}} = 0$ ($\theta(0) = 1$) and $R_{\text{i}}^{\text{Born}} = \theta(c_{\text{i}}) R_{\text{i}}^0$ is the concentration-dependent Born radius used to define $\Omega_{\text{ion}}$ in Figure 1 when $c_{\text{i}} \neq 0$. The Born radii $R_{\text{i}}^0$ in Table 1 are

![Table 1](image_url)

<table>
<thead>
<tr>
<th>Symbol</th>
<th>Meaning</th>
<th>Value</th>
<th>Unit</th>
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<tbody>
<tr>
<td>$k_B$</td>
<td>Boltzmann constant</td>
<td>$1.38 \times 10^{-23}$</td>
<td>J/K</td>
</tr>
<tr>
<td>$\tau$</td>
<td>Temperature</td>
<td>298.15</td>
<td>K</td>
</tr>
<tr>
<td>$e$</td>
<td>Proton charge</td>
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<td>C</td>
</tr>
<tr>
<td>$\varepsilon_0$</td>
<td>Permittivity of vacuum</td>
<td>$8.85 \times 10^{-14}$</td>
<td>F/cm</td>
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<tr>
<td>$\varepsilon_{\text{in}}, \varepsilon_w$</td>
<td>Dielectric constants</td>
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<td></td>
</tr>
<tr>
<td>$\delta_i$</td>
<td>Correlation length</td>
<td>$i = \text{Na}^+, \text{Ca}^{2+}, \text{Cl}^-$</td>
<td>A</td>
</tr>
<tr>
<td>$Q_{\text{w}}, Q_{\text{w}}^+$</td>
<td>Radii</td>
<td>0.95, 0.99</td>
<td>A</td>
</tr>
<tr>
<td>$A_{\text{Cl}^-}^+, A_{\text{Cl}^-}^0$</td>
<td>Radii</td>
<td>1.81, 1.4</td>
<td>A</td>
</tr>
<tr>
<td>$R_{\text{Na}^+}^0, R_{\text{Ca}^{2+}}^0, R_{\text{Cl}^-}^0$</td>
<td>Born radii in Eq. (12)</td>
<td>1.617, 1.706, 2.263</td>
<td>A</td>
</tr>
<tr>
<td>$\delta_{\text{Na}^+}, \delta_{\text{Ca}^{2+}}, \delta_{\text{Cl}^-}$</td>
<td>in Eq. (11)</td>
<td>4.2, 5.1, 3.8</td>
<td></td>
</tr>
<tr>
<td>$\Omega_{\text{i}}$</td>
<td>in Eq. (10)</td>
<td>18</td>
<td></td>
</tr>
</tbody>
</table>
cited from [2], which are computed from the experimental hydration Helmoltz free energies of these ions given in [17]. All values in Table 1 are either physical or experimental data except that of $\delta_i$, which is the only adjustable parameter in our model. All these values were kept fixed throughout calculations.

The PF results of Na+, Ca2+, and Cl− activity coefficients agree well with the experimental data [11] as shown in Figures 2 and 3 for NaCl and CaCl2 electrolytes, respectively, with various [NaCl] and [CaCl2] from 0 to 2.5 M. In Figure 4, we observe that the Debye–Hückel theory oversimplifies the Ca2+ activity coefficient to a straight line as frequently mentioned in physical chemistry texts [4] because the theory does not account for the steric and correlation effects of ions and water, let alone the atomic structure of the ion and its hydration shell as shown in Figure 1. Both PB and PF results in Figure 4 were obtained using the same atomic Fermi formula 10 for shell radii $R_{Sh}^i$ in $\Omega_{Sh}$ and the same concentration-dependent Born formula 12 for Born radii $R_{Born}^{i\text{ion}}$ in $\Omega_{i\text{ion}}$. Therefore, the only difference between PB and PF is in $\Omega_{\text{Solv}}$, where $l_e = S_{\text{trc}} = 0$ for PB and $l_e \neq 0$ and $S_{\text{trc}} \neq 0$ for PF. Note that these two formulas are not present in previous PB models. Figure 4 shows that the correlation and steric effects still play a significant role in the solvent domain $\Omega_{\text{Solv}}$ although the domain is $R_{Sh}^{\text{Ca}^{2+}} = 4.95 \, \text{Å}$ (not shown) away from the center of the Ca2+ ion. The ion and shell domains are the most crucial region to study ionic activities. For example, Fraenkel’s theory is entirely based on this region—the so-called smaller-ion shell region [1].

The PF model can provide more physical details near the solvated ion ([CaCl2], for example) in a strong electrolyte ([CaCl2] = 2 M) such as the dielectric function $\varepsilon(r)$ of varying permittivity (shown in Figure 5), variable water density $\rho_{\text{H2O}}(r)$ (in Figure 5),

![Figure 3](image3.png)

**Fig. 3.** Comparison of PF results with experimental data [11] on $i = \text{Ca}^{2+}$ and Cl− activity coefficients $\gamma_i$ in various [CaCl2] from 0 to 2 M.

![Figure 4](image4.png)

**Fig. 4.** Comparison of Poisson–Fermi (PF), Poisson–Boltzmann (PB), and Debye–Hückel (DH) results on $i = \text{Ca}^{2+}$ activity coefficients $\gamma_i$ in various [CaCl2] from 0 to 2 M.
Fig. 5. Dielectric $\tilde{\varepsilon}(r)$ and water density $C_{H\text{O}}(r)$ profiles near the solvated ion $\text{Ca}^{2+}$ with $[\text{CaCl}_2] = 2\,\text{M}$.

Fig. 6. $\text{Cl}^-$ concentration $C_{\text{Cl}^-}(r)$, electric potential $\phi^\text{PF}(r)$, and steric potential $\phi^\text{st}(r)$ profiles near the solvated ion $\text{Ca}^{2+}$ with $[\text{CaCl}_2] = 2\,\text{M}$.

concentration of counterion ($C_{\text{Cl}^-}(r)$ in Figure 6), electric potential ($\phi^\text{PF}(r)$ in Figure 6), and the steric potential ($\phi^\text{st}(r)$ in Figure 6). Note that the dielectric function $\tilde{\varepsilon}(r)$ is an output, not an input of the model. The steric effect is small because the configuration of particles (voids between particles) does not vary too much from the solvated region to the bulk region. However, the variation of mean-field water densities $C_{H\text{O}}(r)$ has a significant effect on the dielectrics in the hydration region as shown by the dielectric function $\tilde{\varepsilon}(r)$. The strong electric potential $\phi^\text{PF}(r)$ in the Born cavity $\Omega_{\text{ion}}$ and the water density $C_{\text{H\text{O}}}(r)$ in the hydration shell $\Omega_{\text{sh}}$ are the most important factors allowing the PF results to match the experimental data. PF theory deals well with the much more concentrated solutions in ion channels where void effects are important [9].

4. Conclusion

We have proposed a Poisson–Fermi model for studying activities of single ions in strong electrolyte solutions. The atomic structure of ionic cavity and hydration shells of a solvated ion is modeled by the Born theory and Fermi distribution using experimental data. The steric effect of ions and water of nonuniform sizes with interstitial voids and the correlation effect of ions are also considered in the model. Atomic structure and these two effects play a crucial role to match experimental activity coefficients that vary nonmonotonically with salt concentrations. The model contains one adjustable parameter, held at a single fixed value here.
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References


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Bob Eisenberg has just retired as Chairman of Molecular Biophysics at Rush University after thirty nine years. He was tutored at Harvard College by John Edsall in the early years of molecular biology and worked with John Pappenheimer and Steve Kuffler. He graduated summa cum laude after three years. His Ph.D. was with Paul Fatt in Bernard Katz’ Biophysics Department, University College London. Alan Hodgkin was external examiner, and Andrew Huxley, a continuing mentor. Bob studies biology, from molecules to muscles, using anything that helps: experiment, instrument design, simulation, and theory. He is the senior designer of the AxoPatch Clamp amplifier, used in thousands of laboratories to record picoamp currents through single channel molecules.