A Mathematical Model of Electrodiffusion and Osmotic Water flow Yoichiro Mori^{a,b}, Chun Liu^{b,c} and Bob Eisenberg^d

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Background and Overview

We propose a system of partial differential equations (PDE) that describe electrodiffusion and osmotic water flow. From a physical standpoint, this is a far-reaching generalization of the standard treatment of osmosis and electrodiffusion in irreversible thermodynamics to spatially extended systems^{1,2}. To the best of the authors' knowledge, this is the first model in which osmotic water flow and electrodiffusion with deformable and capacitance-carrying membranes have been treated in a mechanically and thermodynamically consistent fashion.

Systems in which both electrodiffusion and osmotic water flow are important abound in physiology³. These include brain ionic homeostasis, fluid secretion by epithelial systems, electrolyte regulation in the kidney, fluid circulation in ocular systems, gastric protection, water uptake by plants etc. We believe the proposed system will have wide-ranging applications in the description of such physiological systems.

Model

Consider biological tissue as a threedimensional region demarcated by the cell membrane Γ , separating the intraand extracellular spaces Ω_i and Ω_e . lonic concentrations c_k satisfy the following drift diffusion equation with the Poisson equation in Ω_i and Ω_e . We solve for **u** (fluid velocity), c_k , ϕ (electrostatic potential) in a selfconsistent fashion.



$$\frac{\partial c_k}{\partial t} + \mathbf{u} \cdot \nabla c_k = \nabla \cdot (c_k D_k \nabla \mu_k), \quad \mu_k = k_B T \ln \mathbf{u}$$
$$-\nabla \cdot (\epsilon \nabla \phi) = \sum_{k=1}^N q z_k c_k$$

The electrolyte fluid is assumed incompressible and **u** satisfies a fluid force balance equation (e.g. Stokes equation). On Γ , we impose boundary conditions on both sides of the membrane.

• Continuity of the electric flux density across and through the dielectric membrane, thus including membrane capacitance.

• Continuity of the ionic flux, where the ions pass through the membrane via ionic channels/transporters/pumps. Channel currents are driven by voltage and concentration gradients, and transporter/pump currents may be driven by energy supplied to the system (e.g. via ATP). • Slip boundary conditions for the fluid velocity, taking into account the transmembrane water flow. The membrane thus moves with the flow. • A stress-jump boundary condition across the membrane. Mechanical properties of the membrane are included.

 $a c_k + q z_k \phi$

Variational Structure and Thermodynamic Consistency

An important property of this model is that it satisfies the following free energy equality.

 $\frac{d}{dt}(G_S + E_{\text{elas}} + E_{\text{elec}}) = -I_p - J_p + J_a$

 G_S : entropic contribution to free energy, E_{elas} : elastic energy of membrane, E_{elec} : electrical energy (capacitive and bulk), I_n : dissipation due to electrodiffusion and viscous fluid flow, J_n : dissipation through passive transmembrane currents and transmembrane water flow, J_a : energy generation through active transmembrane currents *Proof*: Integrate by parts. Treatment of boundary terms is tricky due to the nonlocal nature of electrostatic interactions and the fact that the membrane is dynamic; a novel calculus identity is proved to overcome this difficulty.■

Significance/Implications:

• We recover van t'Hoff's law of osmotic pressure (Π =cRT where Π is the free energy of ions.

• The free energy identity makes it straightforward to identify equality of cross coefficients as dictated by irreversible thermodynamics^{1,2}. • Osmotic effects are typically slow, in which case electroneutrality is a very good assumption. The model can be formulated under this assumption without destroying the variational structure. ϕ becomes a

Lagrange multiplier.

• Assumption of spatial homogeneity leads to a system of classical ODEs used in the study of cell volume control⁴. In this context, the above serves as a hitherto unknown Lyapunov function⁵, giving us a powerful tool to study the stability of steady states.

Tissue Level Model

To describe tissue level physiological phenomena, it is important to develop macroscopic models that do not track the biophysical details at the cellular or subcellular spatial scales. Such a model can be obtained by taking a homogenization limit of the cellular model. This procedure parallels the derivation of the bidomain model in cardiac and lens electrophysiology from three dimensional cable theory^{6,7}. The tissue level model retains the variational structure of the cellular level model. We are currently working to apply this theory to the study of ocular fluid circulation⁸(cornea and lens) as well as to cortical spreading depression⁹ in the brain, a (patho)physiological phenomenon associated with massive redistribution of ionic concentrations across the neuronal and glial membranes.

osmotic pressure and $c=\sum c_k$) as the Legendre transform of the entropic

is shown below¹²:



Abstract:

We propose a system of partial differential equations describing electrodiffusion and osmotic water flow in a three-dimensional setting. We propose models at two spatial scales, the cellular and the tissue level, the latter, a homogenization limit of the former. A salient feature of these models is the presence of a variational structure. The model is a far-reaching generalization of the classical treatment of electrodiffusion and osmosis using irreversible thermodynamics. We expect that the models we propose here to have numerous applications in various physiological systems/phenomena, including epithelial fluid secretion, lens circulation, brain ionic homeostasis, etc.

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Related Computations

We are in the course of developing numerical methods to simulate our model, both at the cellular and tissue levels. Here, we present previous work in which electrodiffusion (without osmotic water flow) at the cellular level was fully simulated in a three-dimensional setting^{10,11}.

We study the propagation of cardiac action potential under reduced gapjunctional coupling. Due to the very narrow cleft space geometry between cardiac muscle cells, there is a possibility, especially, under reduced gap junctional coupling, that ionic concentration effects and field effects could become important. A sample computational example of such a simulation

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