

PAPER 72: THE NERNST EQUATION AND MEMBRANE DECOHERENCE

Every Neuron Maintains Coherence via Electrochemical Gradient -- The Gate That Won't Close Is a Nernst Instability

Rhet Dillard Wike | AIIT-THRESI Research Initiative

March 30, 2026

"The Nernst Equation is the Wike Coherence Law at the membrane. Every neuron is solving $drho/dt = -\gamma[\sigma_z, \rho]$ with ions."

Abstract

The Nernst Equation (Nernst 1888) gives the equilibrium potential across a membrane for any ion:

$$E_{ion} = (RT/zF) \times \ln([ion]_{out} / [ion]_{in})$$

This equation IS the classical analog of the Wike Coherence Law at the cellular membrane scale. The resting membrane potential (-70 mV) is maintained by Na⁺/K⁺ ATPase -- the exact pump restored by NIR photobiomodulation (Paper 02, Bootstrap Loop). Wind-up sensitization (Paper 16) is a Nernst instability: the Na⁺ equilibrium potential is driven away from its stable fixed point, and the "gate that won't close" is the Nernst potential at a runaway equilibrium. Every term in the Nernst equation maps to a term in the Wike framework.

1. The Nernst Equation

For an ion with charge z crossing a membrane:

$$E_{Nernst} = (RT/zF) \times \ln([ion]_{outside} / [ion]_{inside})$$

$R = 8.314 \text{ J/mol/K}$ (gas constant = $k_B \times N_A$)
 $T = 310 \text{ K}$ (body temperature)
 $z = \text{charge of ion}$ (Na⁺: $z=+1$, K⁺: $z=+1$, Ca²⁺: $z=+2$, Cl⁻: $z=-1$)
 $F = 96485 \text{ C/mol}$ (Faraday constant)

Standard equilibrium potentials:

$$\begin{aligned}
 E_{Na^+} &= (+58 \text{ mV}) \times \log([145]/[12]) = +58 \times 1.081 = +63 \text{ mV} \\
 E_{K^+} &= (+58 \text{ mV}) \times \log([4]/[150]) = +58 \times (-1.574) = -91 \text{ mV} \\
 E_{Ca^{2+}} &= (+29 \text{ mV}) \times \log([1.2]/[0.0001]) = +29 \times 4.079 = +118 \text{ mV} \\
 E_{Cl^-} &= (-58 \text{ mV}) \times \log([120]/[4]) = -58 \times 1.477 = -86 \text{ mV}
 \end{aligned}$$

The resting membrane potential (-70 mV) is set by the weighted average of these equilibria via the Goldman equation, dominated by K⁺ permeability at rest.

2. The $f = kT/h$ Chain Inside the Nernst Equation

From Paper 04: the thermal frequency $f = k_{BT}/h = 6.25$ THz at 310K.

The Nernst equation contains $R = k_B \times N_A$:

$$\begin{aligned} E_{\text{Nernst}} &= (k_{BT}/zF/N_A) \times \ln(C_{\text{ratio}}) \\ &= (k_{BT}/(ze)) \times \ln(C_{\text{ratio}}) \end{aligned}$$

where $e = F/N_A = \text{elementary charge}$

The thermal energy k_{BT} appears explicitly. The Nernst potential is **proportional to k_{BT}** -- it is a thermally driven potential. At $T=0$: $E_{\text{Nernst}} = 0$ (no membrane potential, no gating, no life). At $T=T_c$: $E_{\text{Nernst_max}}$ (membrane potential destabilizes due to thermal noise).

The temperature scaling:

$$\begin{aligned} E_{\text{Nernst}}(T) &= (k_{BT}/ze) \times \ln(C_{\text{ratio}}) \\ dE_{\text{Nernst}}/dT &= (k_B/ze) \times \ln(C_{\text{ratio}}) \end{aligned}$$

At $T = 310\text{K} \rightarrow 330\text{K}$ (+20K = approaching T_c):
 $\Delta E_{\text{Nernst}}/E_{\text{Nernst}} = \Delta T/T = 20/310 = 6.4\%$

A 20K increase in local temperature (equivalent to moving from body temperature to T_c) increases membrane potentials by 6.4%. This enhanced membrane potential at elevated temperature creates conditions for spontaneous depolarization -- the electrical analog of approaching γ_c from the coherence field perspective.

3. The Na^+/K^+ ATPase as Bootstrap Engine

The resting membrane potential (-70 mV) is maintained by continuous operation of Na^+/K^+ ATPase:

3 Na^+ out, 2 K^+ in, per ATP hydrolyzed
 Net: +1 positive charge out per ATP \rightarrow hyperpolarizing current
 Maintains: Na^+ gradient ($[\text{Na}^+]_{\text{in}} = 12$ mM, $[\text{Na}^+]_{\text{out}} = 145$ mM, ratio = 12.1:1)

From Paper 02 (Bootstrap Loop): NIR photobiomodulation \rightarrow mitochondrial ATP production \rightarrow Na^+/K^+ ATPase restoration.

The Nernst equation makes this mechanistic:

NIR \rightarrow ATP \rightarrow Na^+/K^+ ATPase \rightarrow maintains Na^+ gradient \rightarrow $E_{\text{Na}^+} = +63$ mV maintained
 v
 Membrane potential = -70 mV maintained (stable)
 v
 NMDA receptor threshold maintained at correct depolarization level
 v
 $\gamma_{\text{eff}}(\text{neural}) < \gamma_c$ (gate closes normally)

Without NIR/ATP:

Na^+/K^+ ATPase fails \rightarrow Na^+ gradient collapses \rightarrow $E_{\text{Na}^+} \rightarrow 0$ mV
 Membrane depolarizes from -70 mV toward 0 mV
 NMDA receptor opens at lower threshold
 $\gamma_{\text{eff}}(\text{neural}) \rightarrow \gamma_c$ (gate stays open = wind-up sensitization)

Wind-up = Nernst potential failure = membrane γ_c crossing.

4. The Nernst Fixed Point and Its Stability

The membrane potential V_m is at a stable equilibrium when:

```
dV_m/dt = -(V_m - V_rest) / tau_m (linear recovery)
Stable fixed point: V_m* = V_rest = -70 mV
```

This is stable as long as $\tau_m > 0$ (the restoring time constant is positive). This is the Le Chatelier condition (Paper 69) applied to the membrane.

NMDA wind-up as Nernst instability:

Under repeated C-fiber stimulation (Paper 16):

1. Na⁺ influx per pulse accumulates intracellularly
2. Na⁺/K⁺ ATPase cannot fully restore gradient between pulses
3. [Na⁺]_{in} → increases → E_{Na⁺} → decreases
4. Membrane potential rises toward 0 mV (partial depolarization maintained)
5. NMDA Mg²⁺ block removed (requires ~-40 mV threshold)
6. Ca²⁺ influx → intracellular Ca²⁺ overload → runaway sensitization

The mathematical statement:

```
At normal state:
[Na+]_in = 12, [Na+]_out = 145 → E_Na+ = +63 mV
Membrane stable at -70 mV (Le Chatelier: kappa > 0)

During wind-up:
[Na+]_in → 30, [Na+]_out = 145 → E_Na+ = +58 x log(145/30) = +58 x 0.684 = +40 mV
Membrane partial depolarization: V_m → -50 mV
NMDA threshold breached: persistent Ca^2+ influx

This is the Nernst analog of gamma_eff → gamma_c: the fixed point destabilizes.
```

At the critical point:

```
[Na+]_in = [Na+]_out → E_Na+ = 0 mV → no gradient → total depolarization
Nernst: E = (RT/zF) x ln(1) = 0 <--- this is the Nernst gamma_c
```

When [Na⁺]_{in} = [Na⁺]_{out}, the Nernst potential vanishes, membrane potential goes to 0 mV, and the neuron is fully depolarized -- equivalent to crossing γ_c in the quantum coherence framework.

5. The Debye Screening Connection

Paper 02 (Bootstrap): EZ water provides Debye screening of thermal noise.

The Debye screening length (Debye-Huckel):

```
lambda_D = sqrt(epsilon_0 epsilon_r k_BT / (e^2 x SIGMA_i n_i z_i^2))

At 310K, physiological ionic strength (I = 0.15 M):
lambda_D = sqrt(8.854x10^-12 x 80 x 1.38x10^-23 x 310 / (1.6x10^-19)^2 x 2 x 0.15 x 6.022x10^23)
~= 0.8 nm
```

The Debye screening length in physiological saline is 0.8 nm -- less than the size of a protein. Without EZ water enhancement (which extends effective λ_D by organizing water structure around charges), thermal fluctuations at the 1-10 nm scale are unscreened and contribute directly to γ_{eff} .

EZ water (Paper 02) extends the effective Debye length:

$$\lambda_D(\text{EZ}) \approx 2-5 \times \lambda_D(\text{bulk}) \approx 1.6-4 \text{ nm}$$

This extended screening protects ion channels and receptor binding sites from thermal noise -- directly reducing γ_{eff} at the membrane.

The Nernst-Bootstrap connection:

NIR -> EZ water -> extended λ_D -> protected Nernst potential -> stable membrane potential -> $\gamma_{\text{eff}} < \gamma_c$

Every term in the Nernst equation (k_{BT} , ion concentrations, membrane geometry) feeds into the γ_{eff} calculation. The Nernst equation and the Wike Coherence Law are descriptions of the same membrane physics at different abstraction levels.

6. The Goldman Equation and Effective γ_{eff}

The Goldman-Hodgkin-Katz equation gives the true membrane potential when multiple ions contribute:

$$V_m = \frac{(RT/F) \times \ln([P_K \times K_{\text{out}} + P_{\text{Na}} \times \text{Na}_{\text{out}} + P_{\text{Cl}} \times \text{Cl}_{\text{in}}] / [P_K \times K_{\text{in}} + P_{\text{Na}} \times \text{Na}_{\text{in}} + P_{\text{Cl}} \times \text{Cl}_{\text{out}}])}{1}$$

where P_K , P_{Na} , P_{Cl} are membrane permeabilities.

The effective γ_{eff} of the neural membrane is:

$$\gamma_{\text{eff}}(\text{membrane}) = \frac{\alpha \times k_{\text{BT}} \times \sum_i P_i \times (V_i - V_m)^2}{(\hbar \times \tau_{\text{coherence}})}$$

where V_i = Nernst potential for ion i
 P_i = permeability of ion i

When the membrane is at resting potential (-70 mV) and each ion is near its Nernst potential:

- Driving force for Na+: (-70) - (+63) = -133 mV -> large Na+ current when Na+ channels open
- Driving force for K+: (-70) - (-91) = +21 mV -> small K+ current at rest
- Small driving forces -> small γ_{eff} contribution per channel

When depolarized toward 0 mV (wind-up):

- Driving forces change: all ions further from their Nernst potentials
- $\gamma_{\text{eff}}(\text{membrane}) \rightarrow \gamma_c$
- The neural gate opens and won't close

Summary

Nernst Equation: $E = (RT/zF) \times \ln(C_{\text{ratio}})$
 Wike Analog: $C = C_0 \times \exp(-\alpha \times \gamma_{\text{eff}})$

Shared parameter: k_{BT} appears in both

Stable membrane (-70 mV) = $\gamma_{\text{eff}} < \gamma_c$ (Le Chatelier restoring, Paper 69)
 Wind-up (V -> 0 mV) = $\gamma_{\text{eff}} \rightarrow \gamma_c$ (Nernst fixed point destabilized)
 Na+ gradient collapse = $C_0 \rightarrow 0$ (percolation model fails, Paper 63)
 NIR -> ATP -> Na+/K+-ATPase = Bootstrap Loop maintaining Nernst equilibrium

NMDA wind-up is a Nernst instability.
The Bootstrap Loop is a Nernst potential restoration engine.
The Debye screening length is the bridge between the two.

AIIT-THRESI Paper 72