

- Initially since $a_e + b_e = a_i + b_i$, the system is electroneutral and there is no potential difference across the cell membrane.

- Cl⁻ gates open. Na⁺ ions will flow from the exterior to the interior in response to the difference in concentrations. This corresponds to a diffusive

flux $J_D = - D_a \frac{\partial a}{\partial x}$ $D_a =$ diffusion coefficient

- In doing so, this builds up a potential difference. The flux of ions due to a potential difference

is $J_V = - \frac{u_a z_a a}{|z_a|} \frac{\partial \phi}{\partial x}$ $u_a =$ ion mobility
 $z_a =$ ion valency (= +1 for Na⁺)

$\phi =$ electric potential.

- The system reaches equilibrium when these fluxes are balanced: $J_D = J_V$

$\Rightarrow D_a \frac{\partial a}{\partial x} = \frac{u_a z_a a}{|z_a|} \frac{\partial \phi}{\partial x}$

$$\text{Nernst } D_a = \frac{u_a RT}{z_a F}$$

(Einstein's relation)

R = universal gas constant 1.2

T = temperature

F = Faraday's constant

$$\text{So } \frac{u_a RT}{z_a F} \frac{\partial a}{\partial x} = \frac{u_a z_a a}{z_a F} \frac{\partial \phi}{\partial x}$$

$$\frac{RT}{z_a F} \frac{1}{a} \frac{\partial a}{\partial x} = \frac{\partial \phi}{\partial x} \Rightarrow \phi = \frac{RT}{z_a F} \log(a) + \text{constant}$$

$$\frac{RT}{z_a F} \log\left(\frac{a_e^*}{a_i^*}\right) = \phi_e - \phi_i = \Delta \phi, \text{ potential difference across the membrane}$$

a_e^* = concentration of Na^+ at the cell membrane on the exterior

a_i^* = concentration of Na^+ at the cell membrane on the interior.

Due to the competing electrical potential we will still have $a_e^* > a_i^*$

Now the K^+ gates open.

The initial diffusive flux is $-D_b \frac{\partial b}{\partial x}$.

There is also a flux due to the electric potential generated by the Na^+ ions. This is $\frac{u_b z_b b}{|z_b|} \frac{\partial \phi}{\partial x}$

where $\phi = \frac{RT}{z_b F} \log(a) + \text{constant}$ is the potential due

to the Na^+ ions found earlier.

So the total initial flux is $J = -D_b \frac{\partial b}{\partial x} + \frac{u_b z_b b}{|z_b|} \frac{\partial \phi}{\partial x}$

and $D_b = \frac{u_b RT}{|z_b| F}$ by Einstein's relation and $z_b = 1$

gives the required result.

2.)

$$\frac{dn}{dt} = \alpha(V)(1-n) - \beta(V)n$$

$$\frac{dn}{dt} = \alpha(V) - (\alpha(V) + \beta(V))n$$

$$\frac{1}{\alpha + \beta} \frac{dn}{dt} = \frac{\alpha}{\alpha + \beta} - n$$

$$\tau(V) \frac{dn}{dt} = n_{\infty}(V) - n \quad (1) \quad \text{where}$$

$$\tau(V) = \frac{1}{\alpha(V) + \beta(V)}$$

V represents the voltage.

$$n_{\infty}(V) = \frac{\alpha(V)}{\alpha(V) + \beta(V)}$$

n represents the fraction of open gates.

α represents the rate of gate opening.

β represents the rate of gate closing.

V = V(t) prescribed so

$$\tau(t) \frac{dn}{dt} = n_{\infty}(t) - n$$

$$\frac{dn}{dt} + \frac{1}{\tau(t)} n = \frac{n_{\infty}(t)}{\tau(t)}$$

$$\frac{d}{dt} \left(n \exp \int_0^t \frac{ds}{\tau(s)} \right) = \frac{n_{\infty}(t)}{\tau(t)} \exp \int_0^t \frac{ds}{\tau(s)}$$

$$n = \exp \left(- \int_0^t \frac{ds}{\tau(s)} \right) \int_0^t \frac{n_{\infty}(\bar{t})}{\tau(\bar{t})} \exp \int_0^{\bar{t}} \frac{ds}{\tau(s)} d\bar{t} + n(0)$$

If $\tau = 1$, and $n_{\infty} = \cos(t) + 1$ then we may solve the ODE ① to get

$$n(t) = \frac{1}{2}(2n_0 - 3)e^{-t} + \frac{1}{2}(2 + \cos(t) + \sin(t))$$

$$= (n_0 - \frac{3}{2})e^{-t} + 1 + \frac{1}{2\sqrt{2}}\cos(t - \frac{\pi}{4})$$

↑
out of phase with n_{∞} by $\frac{\pi}{4}$

If $\tau = t+1$, $n_{\infty} = \text{constant}$, n_{∞}^* say, then we may solve the ODE ① to get

$$n(t) = \frac{n_0 + n_{\infty}^* t}{1+t}$$

As $t \rightarrow \infty$

$$n(t) = (n_{\infty}^* + \frac{n_0}{t}) (1 + \frac{1}{t})^{-1}$$

$$\sim (n_{\infty}^* + \frac{n_0}{t}) (1 - \frac{1}{t})$$

$$\sim n_{\infty}^* + \frac{(n_0 - n_{\infty}^*)}{t}$$

so approach to equilibrium is algebraic ($\propto \frac{1}{t}$)

3) The Hodgkin-Huxley model is:

All of this is done in the lectures

capacitance of the membrane

$$C_m \frac{dV}{dt} + (I_i - I_{app}) = 0 \quad (1)$$

applied current

except for identifying the magnitude of the parameters

$$\tau(V) \frac{dn}{dt} = n_{\infty}(V) - n \quad (2)$$

n = potassium activation.

$$\tau_m(V) \frac{dm}{dt} = m_{\infty}(V) - m \quad (3)$$

m = sodium activation.

$$\tau_h(V) \frac{dh}{dt} = h_{\infty}(V) - h \quad (4)$$

h = sodium inactivation.

$$I_i = \underbrace{g_{Na} m^3 h (V - V_{Na})}_{Na^+ \text{ current}} + \underbrace{g_K n^4 (V - V_K)}_{K^+ \text{ current}} + \underbrace{g_L (V - V_L)}_{\text{leakage (mainly Cl}^- \text{ chloride ions)}} \quad (5)$$

$\tau(V), \tau_m(V), \tau_h(V), n_{\infty}(V), m_{\infty}(V), h_{\infty}(V), C_m, I_{app}, g_{Na}, g_K, g_L, V_{Na}, V_K, V_L$ are all known.

Assumptions : 1) $T_m \ll 1$.

$$\textcircled{3} \Rightarrow m \approx m_{\infty}(V).$$

$$2) T = T_h$$

$$3) n_{\infty} + h_{\infty} = \text{constant}, \bar{h} \text{ say}$$

$$2), (3), \textcircled{2} + \textcircled{4} \Rightarrow n + h = \bar{h}$$

Then, $\textcircled{1}$ and $\textcircled{5}$ gives

$$C_m \frac{dV}{dt} = I_{app} - \left[g_{Na} m^3 (\bar{h} - n) (V - V_{Na}) + g_K n^4 (V - V_K) + g_L (V - V_L) \right]$$

$$\tau_n(V) \frac{dn}{dt} = n_{\infty}(V) - n$$

Two equations for two unknowns : V and n .

Actually, $\textcircled{2} + \textcircled{4}$ gives $\tau \frac{d(n+h)}{dt} = \bar{h} - (n+h)$

$$\frac{d}{dt} \left[(n+h) \exp \left[\int_0^t \frac{dt'}{\tau(t')} \right] \right] = \bar{h} \exp \left[\int_0^t \frac{dt'}{\tau(t')} \right]$$

$$n+h = \bar{h} + A \exp \left[- \int_0^t \frac{dt'}{\tau(t')} \right]$$

$$\Rightarrow \bar{h} \text{ as } t \rightarrow \infty, \text{ or equals } \bar{h} \text{ if } n+h = \bar{h} \text{ at } t=0.$$

Non-dimensionalize:

$$V = \frac{V - V_{eq}}{V_{Na} - V_{eq}}, \quad t = \tau t'$$

$$\Rightarrow \frac{dn}{dt} = n_{\infty}(V) - n$$

$$\epsilon \frac{dV}{dt} = I^* - g(V, n)$$

$$g(V, n) = \gamma_K(V + V_K^+)n^4 + \gamma_L(V - V_L^+) - (1 - \nu)(\bar{h} - n)m^3(V)$$

$$\gamma_K = \frac{g_K}{g_{Na}}, \quad \gamma_L = \frac{g_L}{g_{Na}}, \quad V_K^+ = \frac{-V_K}{V_{Na}}, \quad V_L^+ = \frac{V_L}{V_{Na}}$$

Using the numbers given provides

$$\gamma_K \sim 0.3,$$

$$\gamma_L \sim 0.003,$$

$$V_K^+ \sim 0.1,$$

$$V_L^+ \sim 0.1$$

$$\epsilon \sim 1.6 \times 10^{-3}$$

We may exploit the fact that $\epsilon \ll 1$ to perform fast-slow phase plane analysis; $\gamma_L \ll 1$ just simplifies the algebraic equation we must solve, $g(V, n) = 0$, for the nullcline.

$$4) \quad \dot{v} = J^* + f(v) - w$$

$$\dot{w} = \gamma v - w \quad \gamma \ll 1$$

$$f = v(a-v)(v-1) \quad 0 < a < 1$$

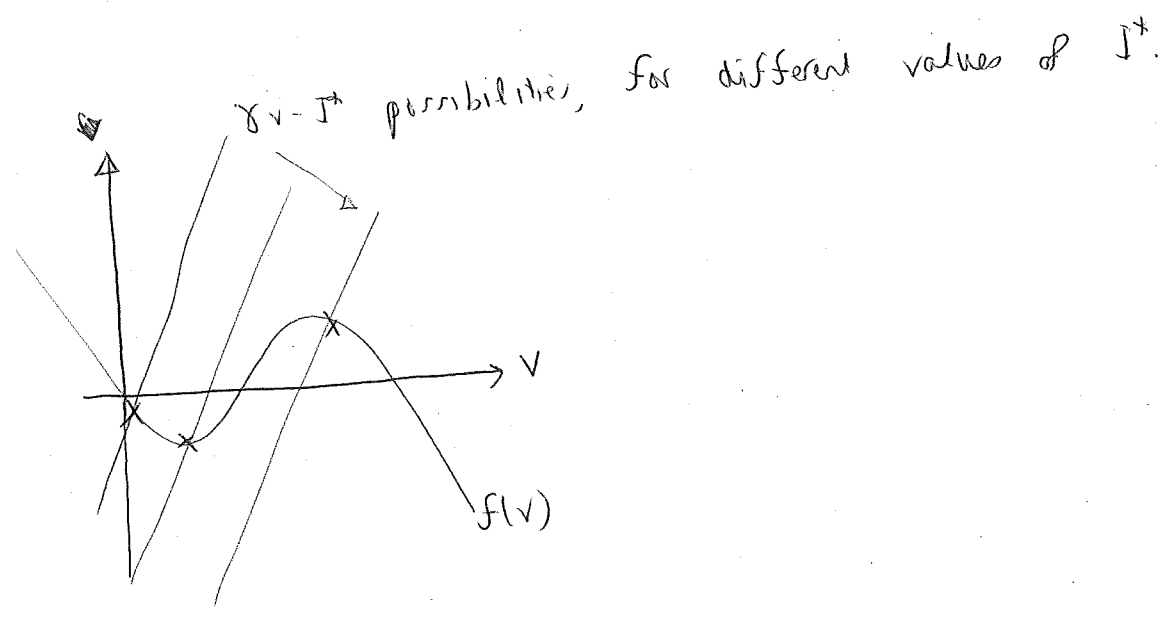
Steady states occur when $\dot{v} = \dot{w} = 0$

$$\Rightarrow f(v) - w + J^* = 0$$

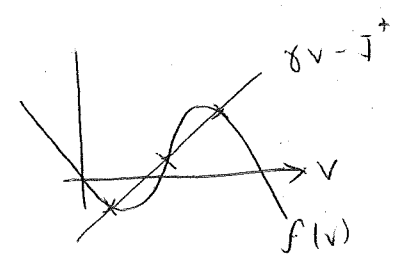
$$\gamma v = w$$

$$\Rightarrow v(a-v)(v-1) - \gamma v + J^* = 0$$

$$f(v) = \gamma v - J^*$$



For there to be a unique root, we need $f'(v) < \gamma$
 - otherwise we could have more than one root:



Now $f'(v) = \frac{d}{dv} \left[-v^3 + (a+1)v^2 + av \right]$

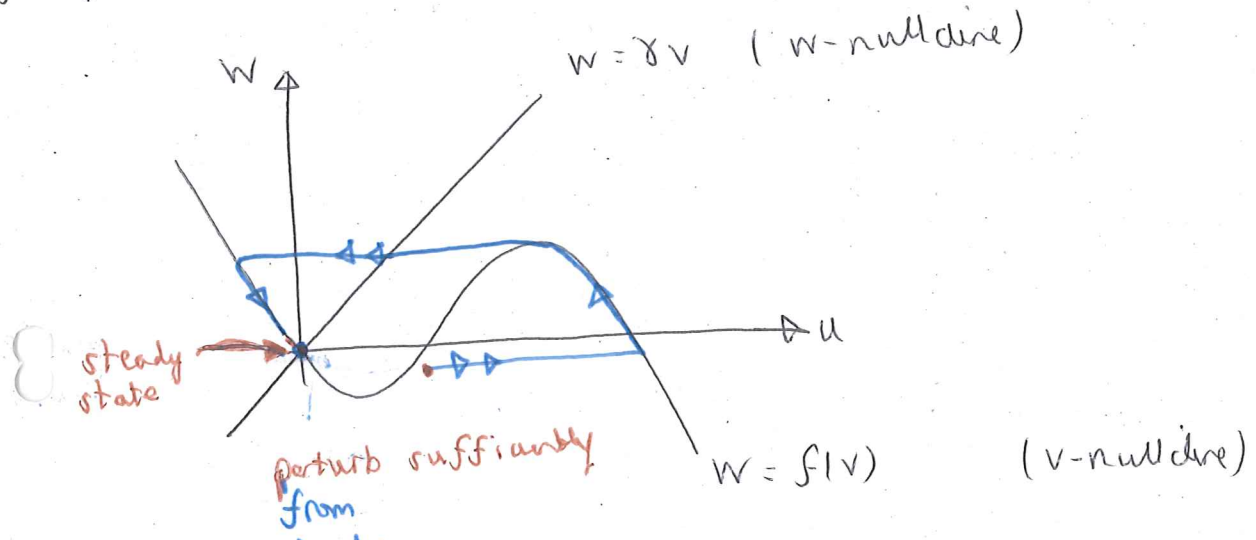
$= -3v^2 + 2v(a+1) + a$

$= -3 \left(v - \frac{a+1}{3} \right)^2 + \frac{1}{3}(a^2 - a + 1)$ (Completing the square)

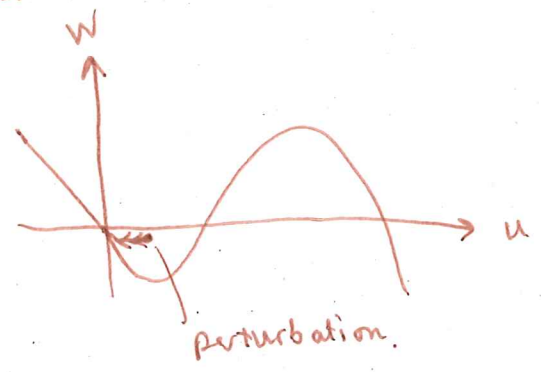
So $f'(v) < \frac{1}{3}(a^2 - a + 1)$

So we have a unique root if $\frac{1}{3}(a^2 - a + 1) < \delta$
as required.

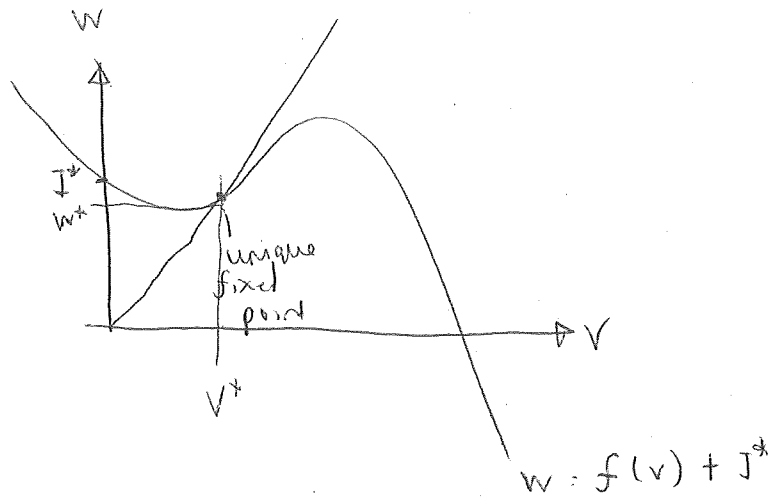
If $I^+ = a$ we have the phase plane:



If we don't provide enough of a perturbation then we do not make an excursion and we relax back to the equilibrium.



If $J^+ > 0$ we have the following phase plane:



Fixed point (v^+, w^+) satisfies
 ~~$f(v^+) - w^+ = 0$~~

$$J^+ + f(v^+) - w^+ = 0$$

$$\gamma v^+ - w^+ = 0$$

Linearize about this fixed point: $v = v^+ + \tilde{v}$
 $w = w^+ + \tilde{w}$

$$\Rightarrow \quad \varepsilon \dot{\tilde{v}} = f'(v^+) \tilde{v} - \tilde{w}$$

$$\dot{\tilde{w}} = \gamma \tilde{v} - \tilde{w}$$

$$\begin{pmatrix} \dot{\tilde{v}} \\ \dot{\tilde{w}} \end{pmatrix} = \underbrace{\begin{pmatrix} f'(v^+)/\varepsilon & -1/\varepsilon \\ \gamma & -1 \end{pmatrix}}_{\underline{M}} \begin{pmatrix} \tilde{v} \\ \tilde{w} \end{pmatrix}$$

The system is unstable iff $\text{tr}(M) > 0 \Rightarrow f'(v^+) > \varepsilon$
 and so if $f'(v^+) > \varepsilon$ then oscillations will occur.

Since $\varepsilon \ll 1$, oscillations will occur if $f'(v^*) \geq 0$

Thus, oscillations will occur if $v_- < v^* < v_+$ where

v_{\pm} are the roots of $f' = 0$.

$$\text{i.e. } v_{\pm} = \frac{a+1}{3} \pm \frac{1}{3} \sqrt{(a^2 - a + 1)}$$

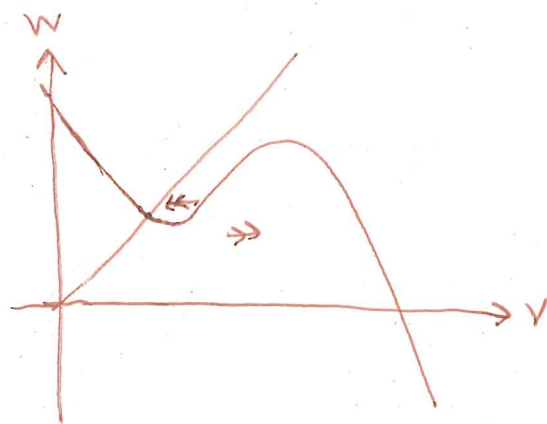
Thus, instability occurs for $I_- < J^* < I_+$ where

$$I_{\pm} = \gamma v_{\pm} - f(v_{\pm})$$

Substituting ① into ② gives (after algebra):

$$I_{\pm} = \frac{1}{3} \left[\gamma - \frac{2}{9} (a^2 - a + 1) \right] \left[(a+1) \pm (a^2 - a + 1)^{1/2} \right] + \frac{1}{9} a(a+1)$$

$f'(v^*) < 0$
Need to perturb beyond dip



$f'(v^*) > 0$
Infinitely small perturbation gives rise to an excursion.

