

Units: m μ n p f a z
 10^{-3} -6 -9 -12 -15 -18 -21 - - -

eg: mV μ A nA pA fF aF zJ
 PS PF

typical units for many cell data

(capacitance of 1 vesicle ~ 40 aF)

energy change to open MET $4zJ$

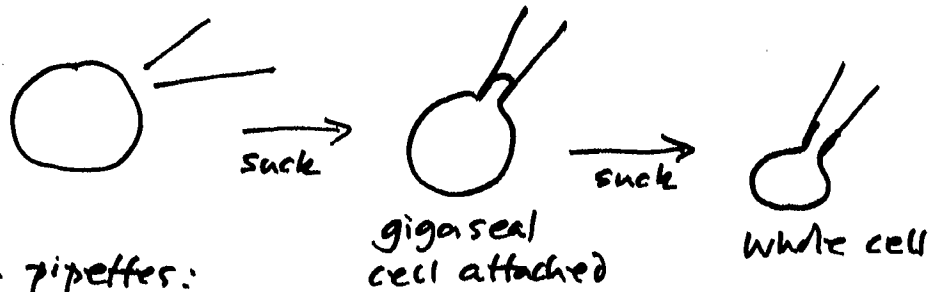
Basic

$V = iR$ or equivalently $i = gV$ OHM

R - resistance in ohms (Ω)

g - conductance in ohms⁻¹ = SIEMENS.

Patch clamping :

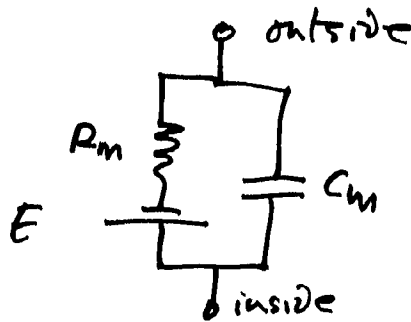


Advantages of patch pipettes:

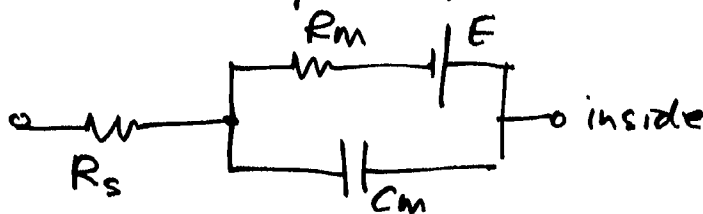
Low noise; better response time; high resolution

Equivalent Circuit of a cell

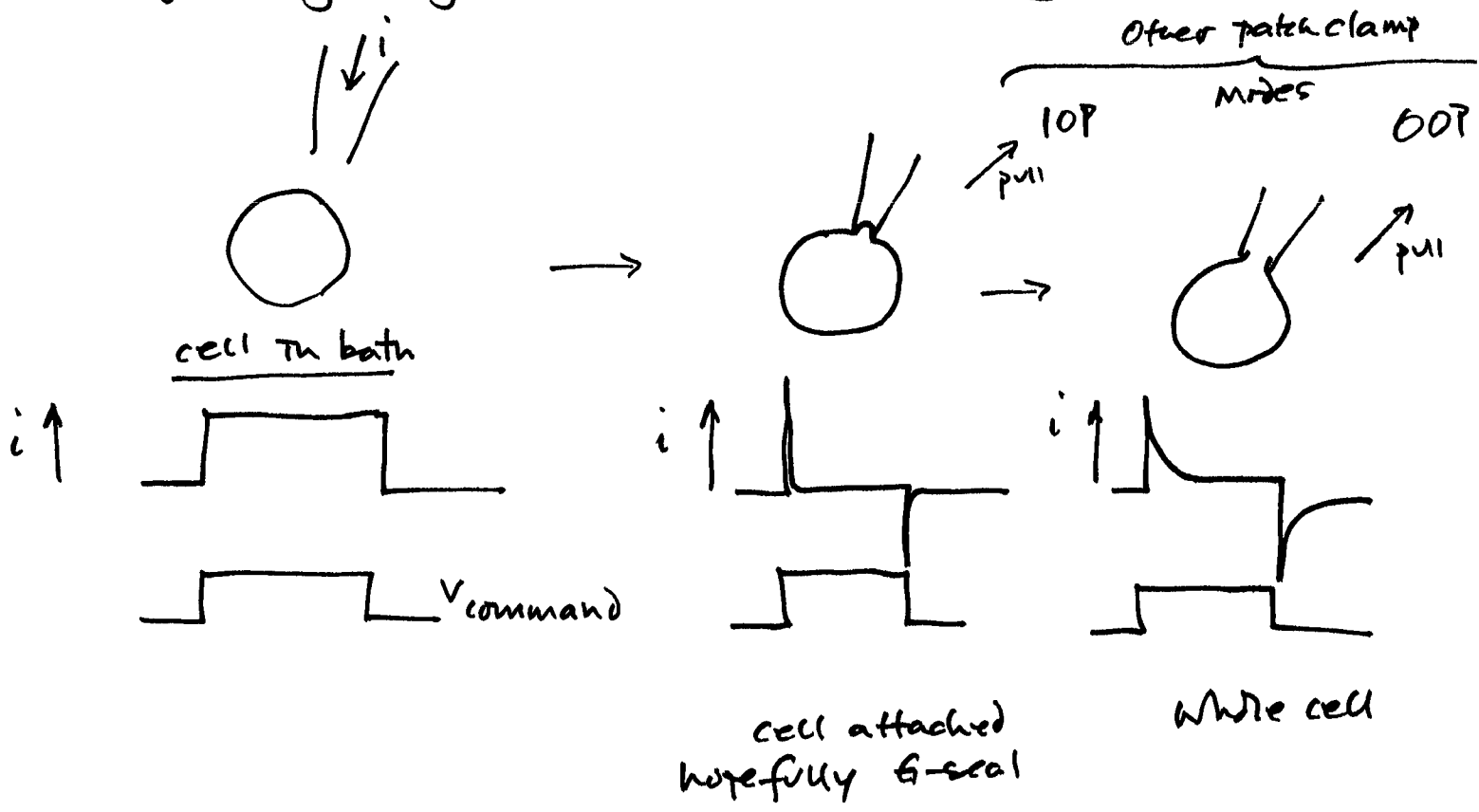
Membrane :



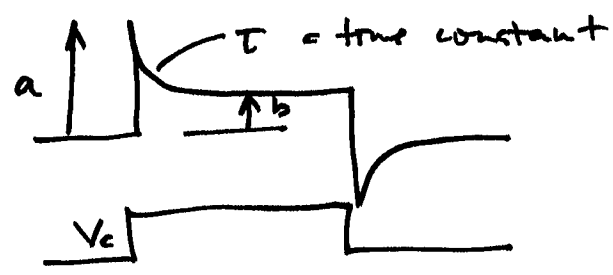
When recorded with a patch pipette circuit looks like



3 Steps to getting a whole cell recording



Various features of WC current trace tell you about params.

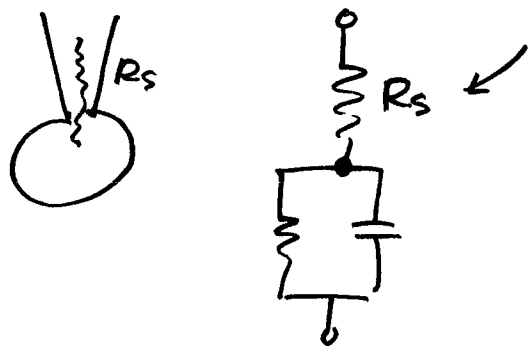


$$a = \frac{V_c}{R_s} ; b = \frac{V_c}{R_s + R_m} ; \tau = \frac{R_s R_m \cdot C_m}{R_s + R_m} \approx R_s C_m$$

Items to think about:

- Intracellular solution in pipette ; Extracellular solns.
- Stability and positioning of pipette ;
- Recording protocols ;
- Other simultaneous tasks eg imaging .
- eg deflecting hair bundle

Series resistance (or 'access' resistance)



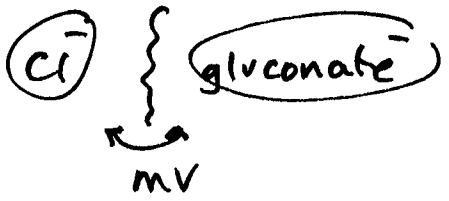
Current flowing through R_s leads to a voltage drop

- ⇒ potential at cell not what is dialled up
- ⇒ C_m not charged quickly enough
- ⇒ errors in current kinetics.

* Correct during expt + in analysis

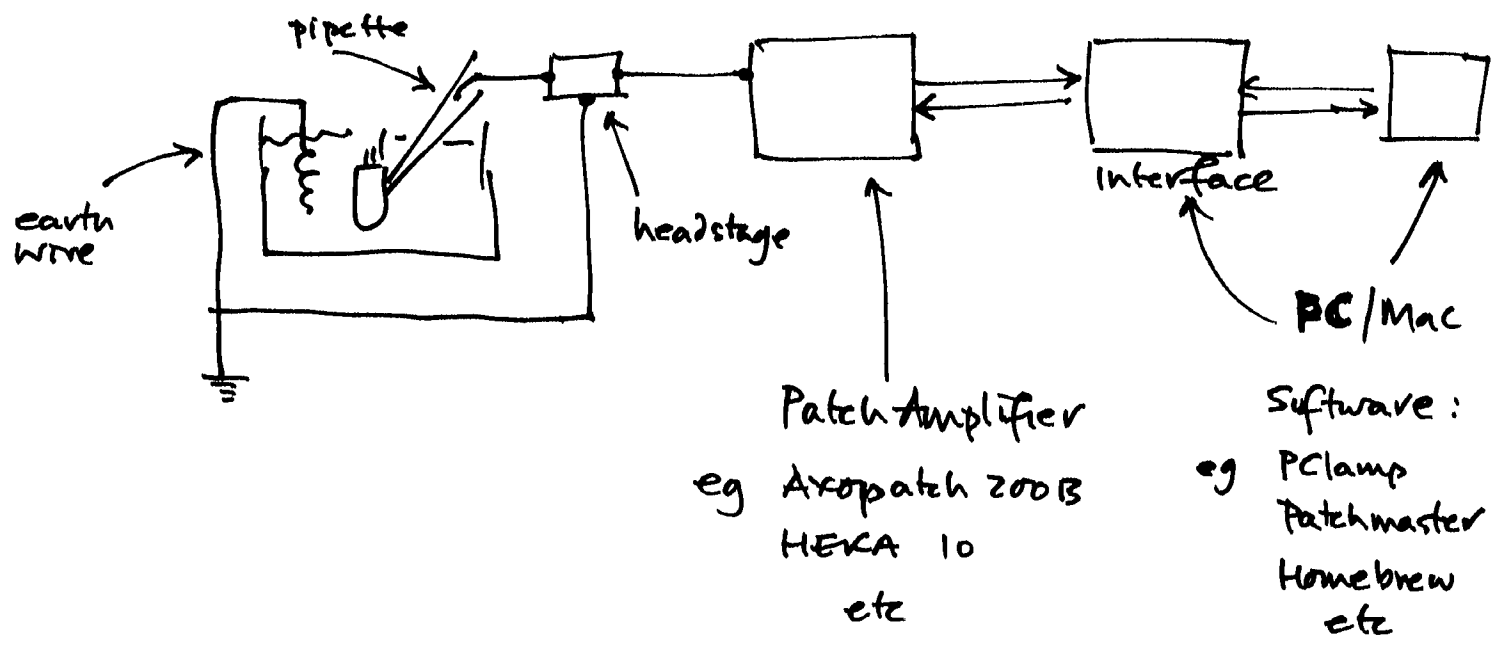
Junction potential.

Arises from eq interface between two different anionic solutions



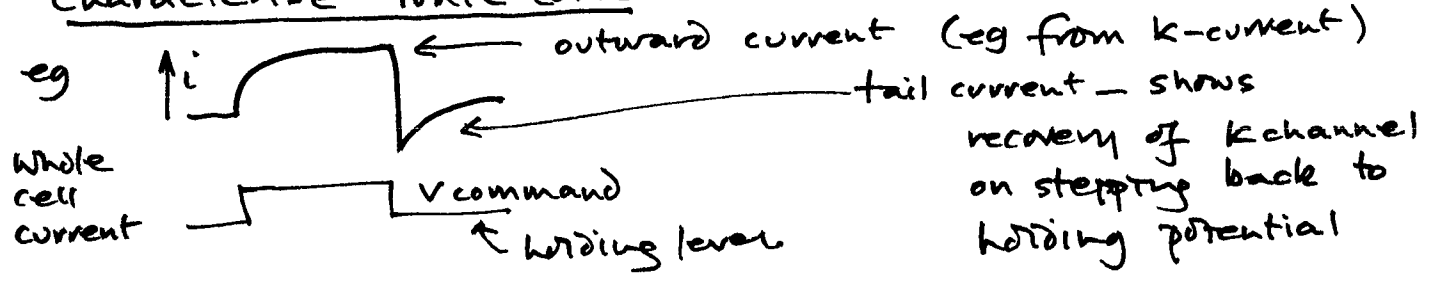
eg between solution and electrical connector wire

* Correct by using software estimates (eg in PClamp) by correctly zero-ing current before giga seal



Some things you can learn from EP recordings:

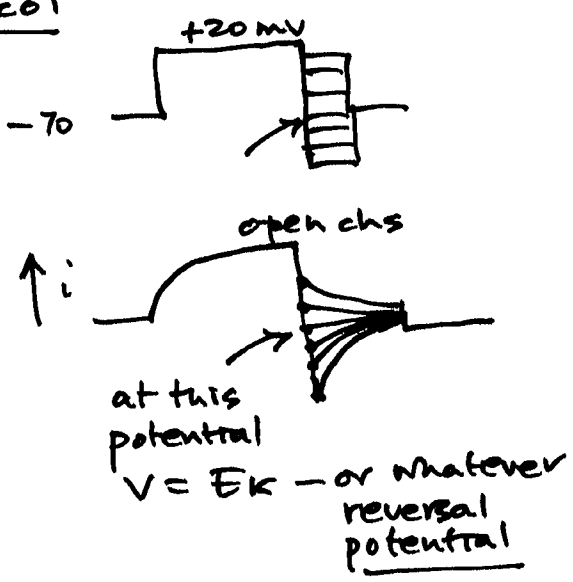
Characterise ionic currents



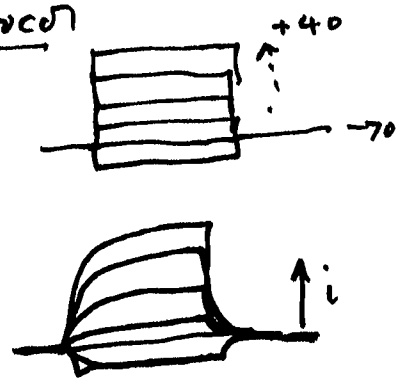
Tail current analysis

$$i_K = g_K(V) \cdot (V - E_K) \quad g_K(V) - \text{voltage activated conductance}$$

Protocol

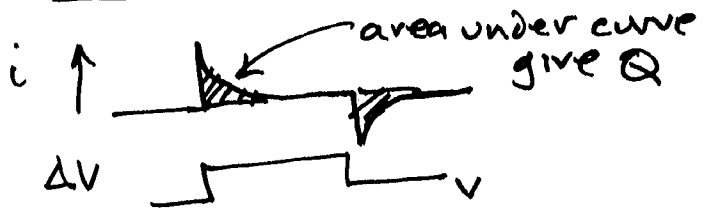


Protocol



measure $i \rightarrow g_K(V)$
the "activation"

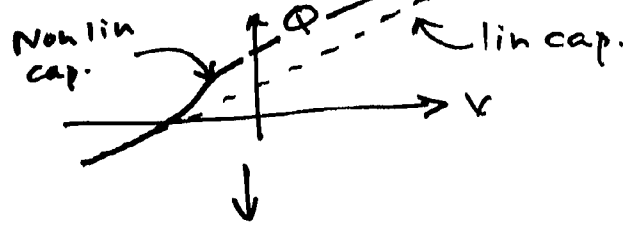
Characterise prestin EP



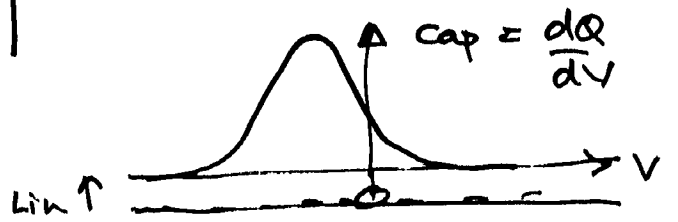
maths!

$$Q = \frac{\Delta V}{R_s} \int_0^{\infty} e^{-t/R_s C_m} dt = \Delta V \cdot C_m$$

In an QVC plot Q vs V



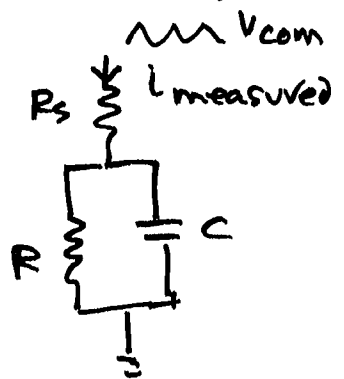
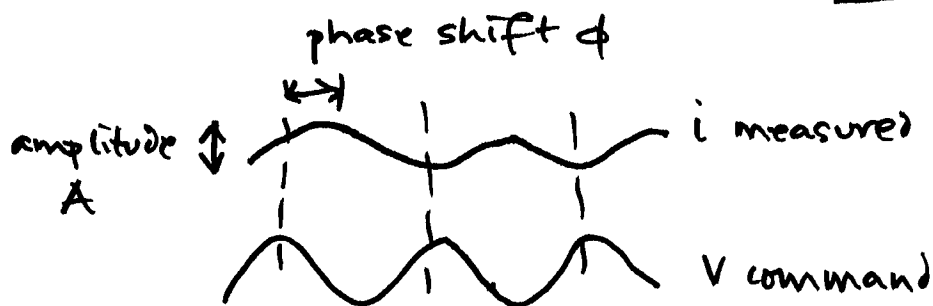
Non linear capacitance



To measure OHC motor function

1. Charge transients at different V
2. Use software to compute capacitance
3. Use a 'lock-in' amplifier to measure capacitance
4. Measure cell length

Principle of 3. Use sine wave commands not steps



Lock in amplifier gives A and ϕ — continuously
Arrange lock in amplifier so that it gives an output
 $\propto C_m$ on one channel
 $\propto I_{mess} \sim R_s$ on 2nd channel.

Can also be used to measure exocytosis, resolution
 $\sim 10fF \approx 300$ synaptic vesicles