

Gibbs Free Energy

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Gibbs Free Energy

- If we combine the two thermodynamics law;

$$\delta Q = dU + \delta W$$

$$\delta Q = TdS$$

$$TdS \geq \delta Q = dU + \delta W$$

Work done by the system; $\delta W = PdV + dW^*$

dW^* other parameters rather than PdV

$$\delta W^* \leq -(dU + PdV - TdS)$$

Gibbs free energy;

$$\delta W = PdV + Fdx + \Phi dq + \dots$$

dW^*

$$G = U + PV - TS$$

Gibbs Free Energy

$$\mathbf{G=U+PV-TS}$$

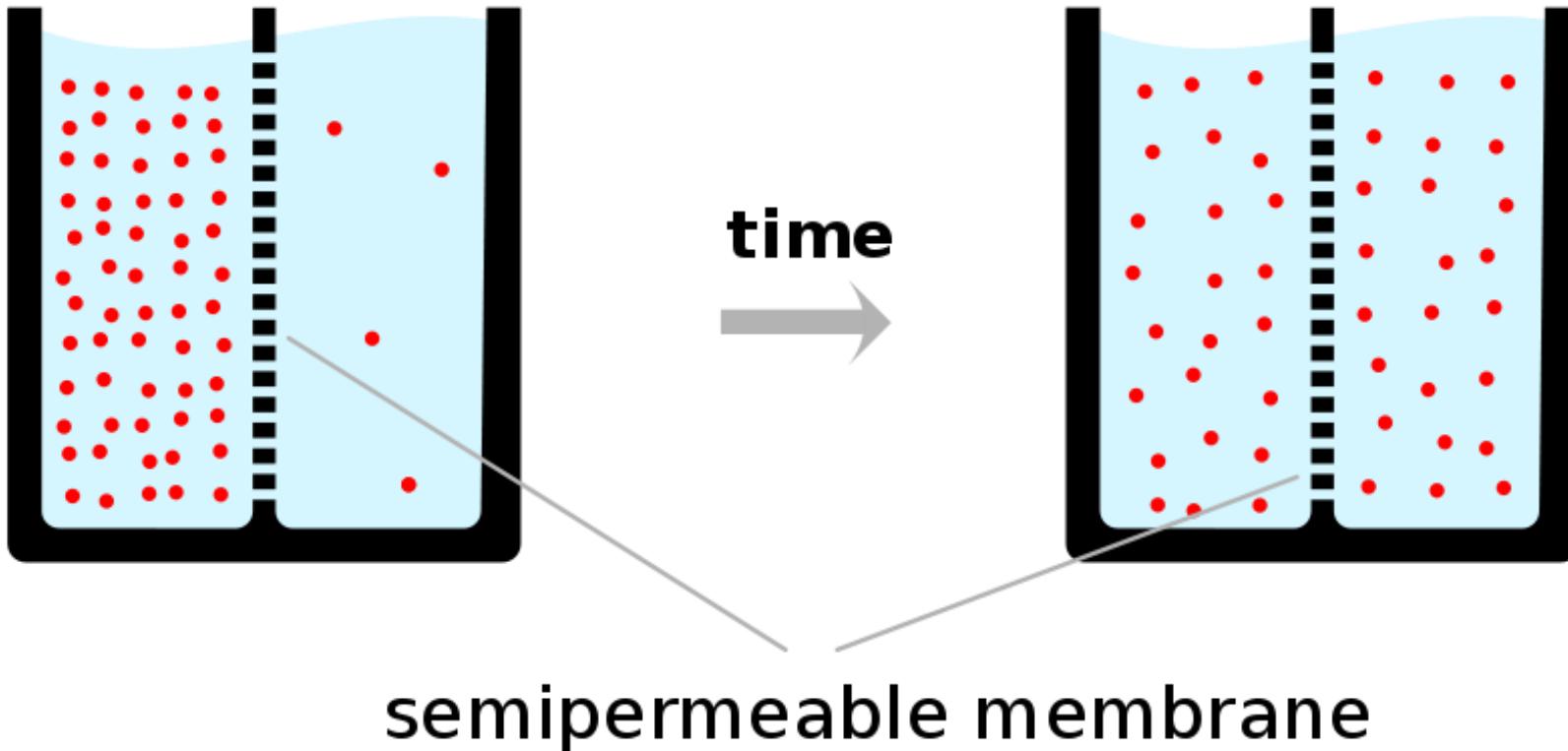
- $G_i = G_i^0 + RT \ln c_i + z_i FV + P_v i \dots \dots$

G_i^0 initial free energy

- **If the gibbs free energy is decreasing means that entropy is increasing**

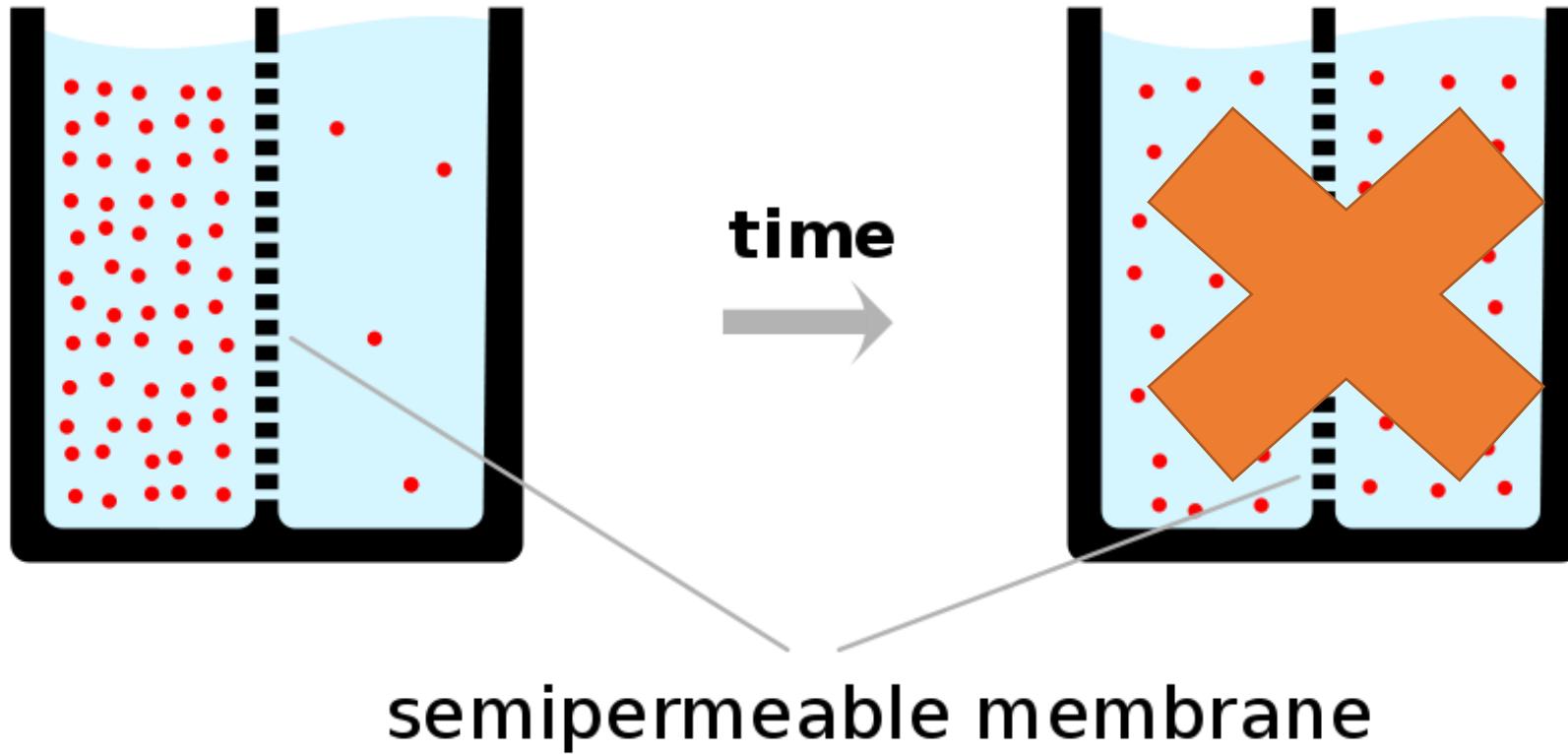
Nernst Equation:

$$G_i = G_i^0 + RT \ln c_i$$



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$$G_i = G_i^\circ + RT \ln c_i + z_i F V + P v_i + \dots \quad [\text{J/mol}]$$

$$G_i^{ic} = G_i^{dis}$$

$$RT \ln c_i^{ic} + z_i FV^{ic} + \dots = RT \ln c_i^{dis} + z_i FV^{dis} + \dots$$

Nernst Equation:

$$E_i = V^i - V^o = \frac{RT}{z_i F} \ln \frac{c_i^o}{c_i^i}$$

RT/F=25.67mV

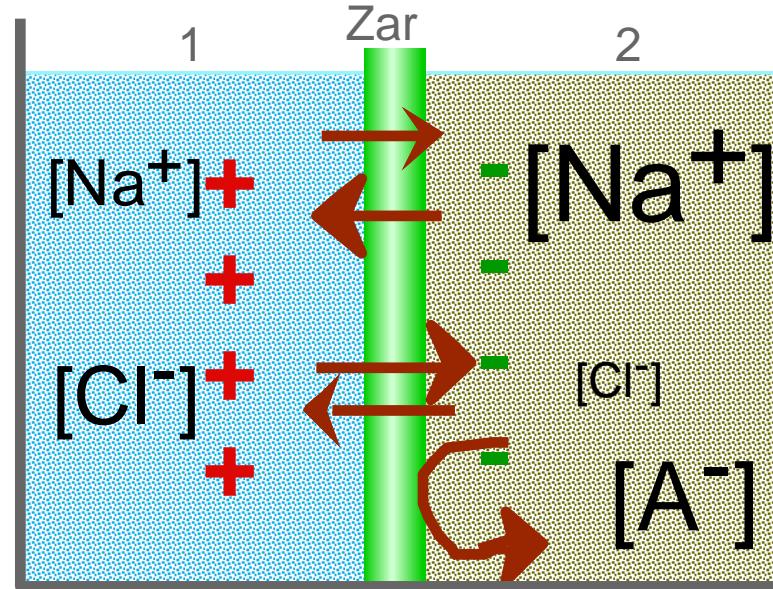
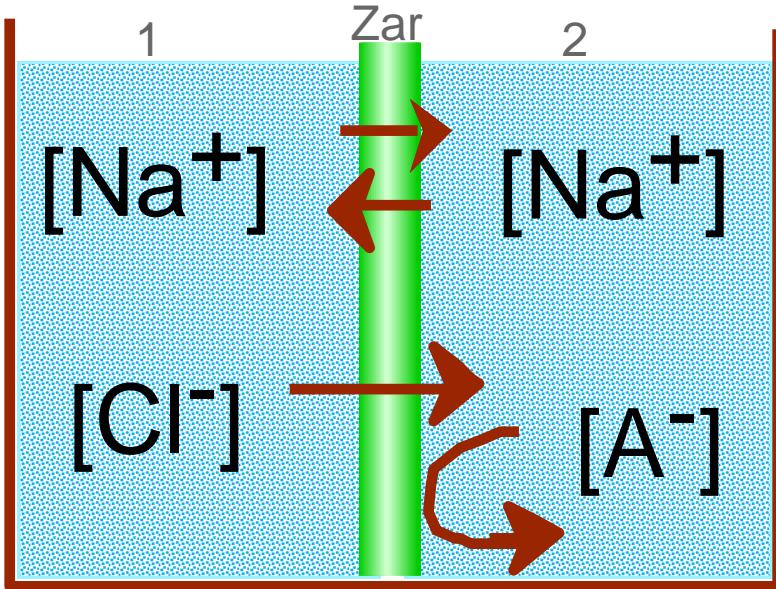
Z=Valence of the electron

E= Electrical potential

C_i=intracellular concentration of an ion

C^o =extracellular concentration of an ion

Gibbs-Donnan Equilibrium

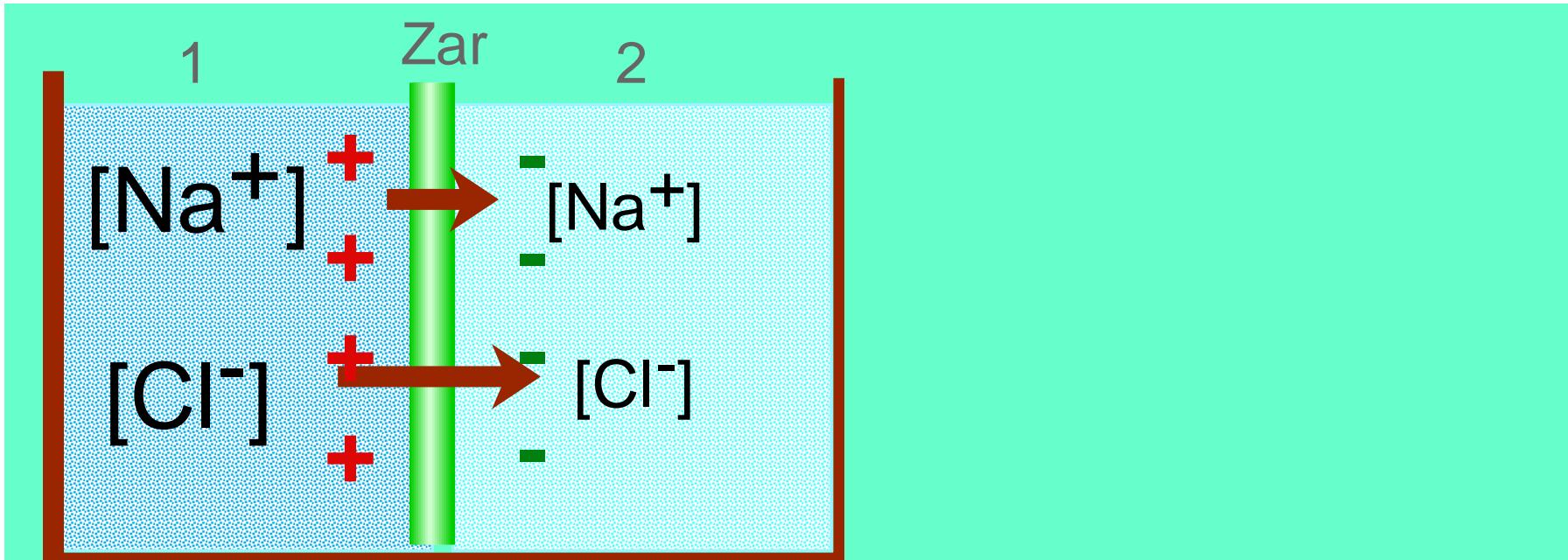


$$G_{\text{Na}^+}^\circ + RT \ln [\text{Na}^+]_1 + FV_1 = G_{\text{Na}^+}^\circ + RT \ln [\text{Na}^+]_2 + FV_2$$

$$G_{\text{Cl}^-}^\circ + RT \ln [\text{Cl}^-]_1 - FV_1 = G_{\text{Cl}^-}^\circ + RT \ln [\text{Cl}^-]_2 - FV_2$$

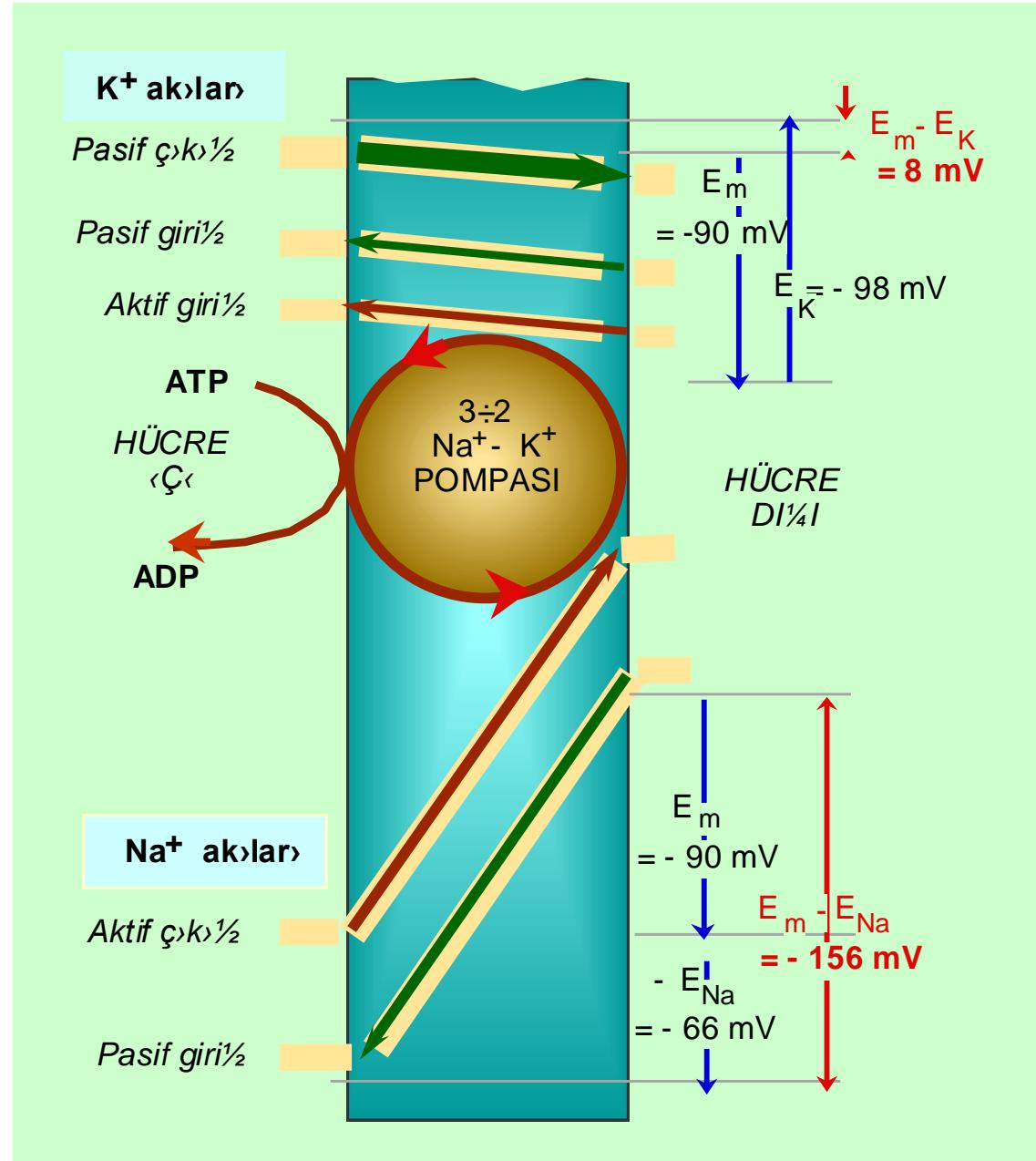
$$[\text{Na}^+]_1 = [\text{Cl}^-]_1 = c_1 \text{ ve } [\text{Na}^+]_2 = [\text{Cl}^-]_2 + [\text{A}^-]$$

Diffusion Potential

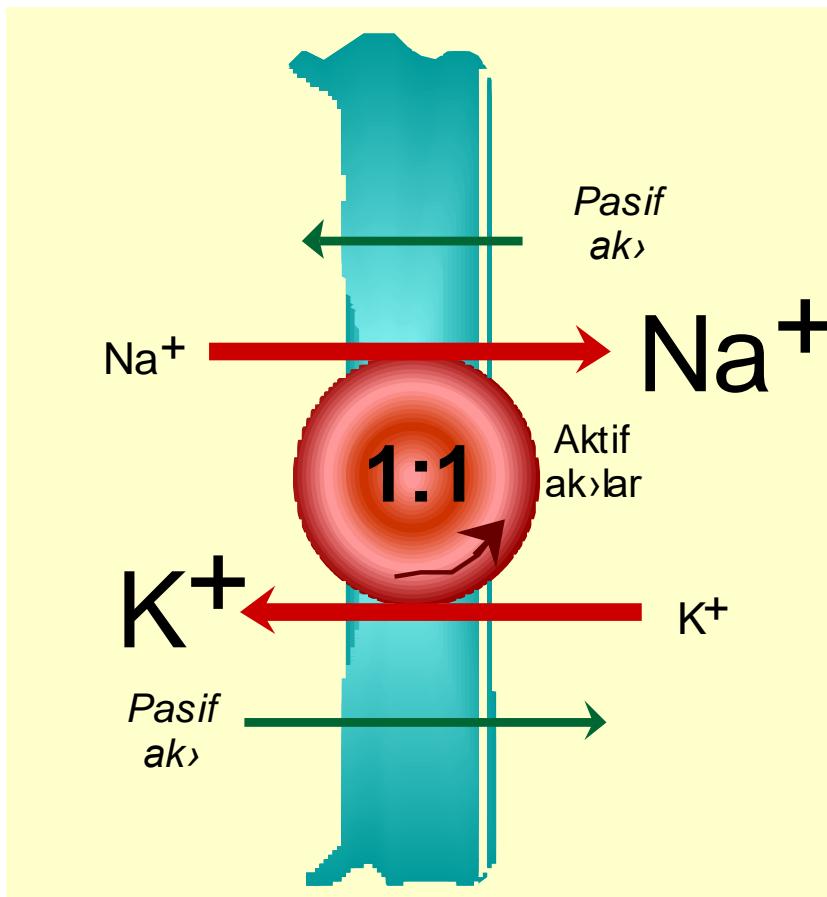


$$V = \frac{RT}{F} \frac{\mu_{Na} - \mu_{Cl}}{\mu_{Na} + \mu_{Cl}} \ln \frac{[c]_1}{[c]_2}$$

$$P_i = D_i / \delta = \mu_i RT / (z_i F \delta) \quad [m/s]$$



Gradient in cell membrane produced by pumps



Even if pump work as 1:1 ratio, it may still produce gradient

Goldman-Hodgkin-Katz Equation

The **Nernst equation** calculates the **equilibrium potential** (also referred to as the **Nernst potential**) for an ion based on the charge on the ion (i.e., its valence) and its concentration gradient across the membrane.

$$J = J_{Na} + J_K + J_{Cl}$$

$$E_m = \frac{RT}{F} \ln \frac{P_K [K^+]^{d/2} + P_{Na} [Na^+]^{d/2} + P_{Cl} [Cl^-]^{i\zeta}}{P_K [K^+]^{i\zeta} + P_{Na} [Na^+]^{i\zeta} + P_{Cl} [Cl^-]^{d/2}}$$

$$P_K \div P_{Na} \div P_{Cl} \approx 1 \div 0,04 \div 0,45$$

Resting membrane potential

For, $P_K \gg P_{Na}, P_{Cl}$

$$E_m \approx \frac{RT}{F} \ln \frac{[K^+]^{d/2}}{[K^+]^{i\zeta}} \rightarrow E_K$$

Equivalent electric circuit for the ion-selective membrane

