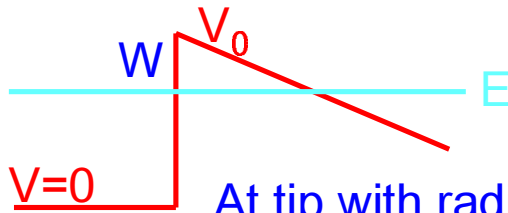
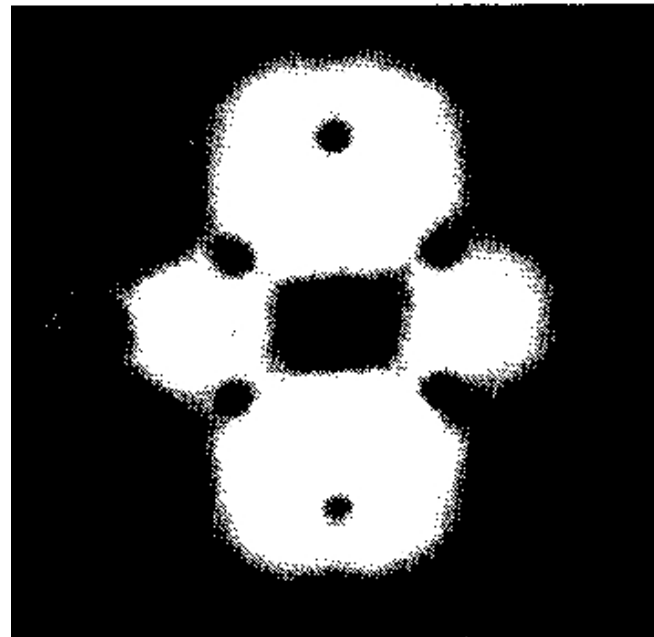
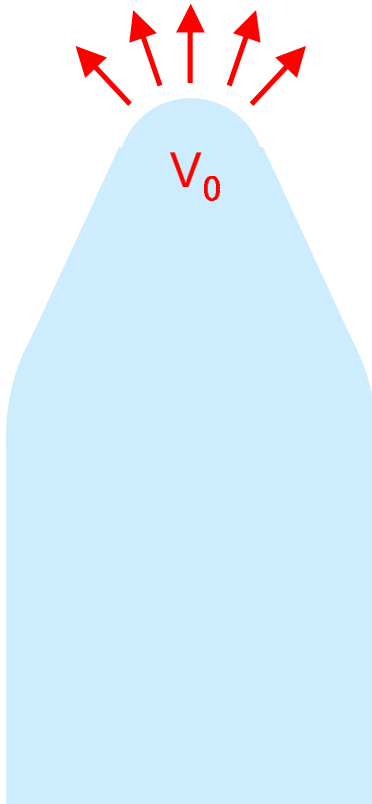


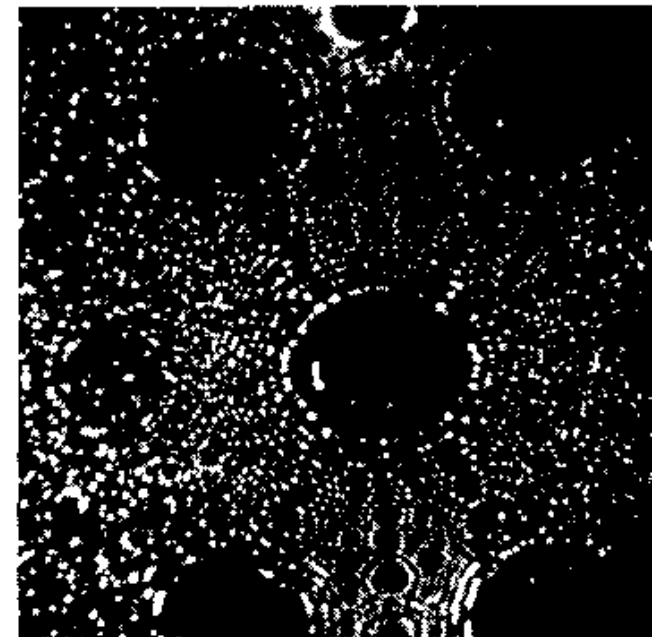
Field emission microscopy



At tip with radius $R = 500\text{nm}$ and $V_0 = 500\text{V}$ has $E_{\text{field}} = V/R = 1\text{GV/m}$



Electron field-emission
microscopic image



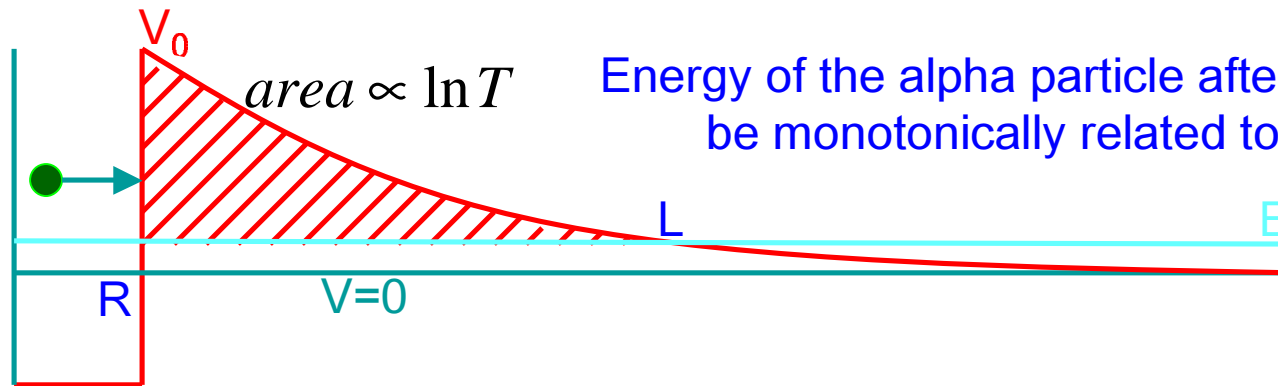
Helium ion field-emission
microscopic image
(He emission of a
previously applied Helium)



The alpha decay

04/11/2005

The transmission probability T for an alpha particle traveling from the inside towards the potential well that keeps the nucleus together determines the lifetime for alpha decay.



$$T \approx \exp\left[-2 \int_R^L \frac{\sqrt{2m[V(r)-E]}}{\hbar} dr\right]$$



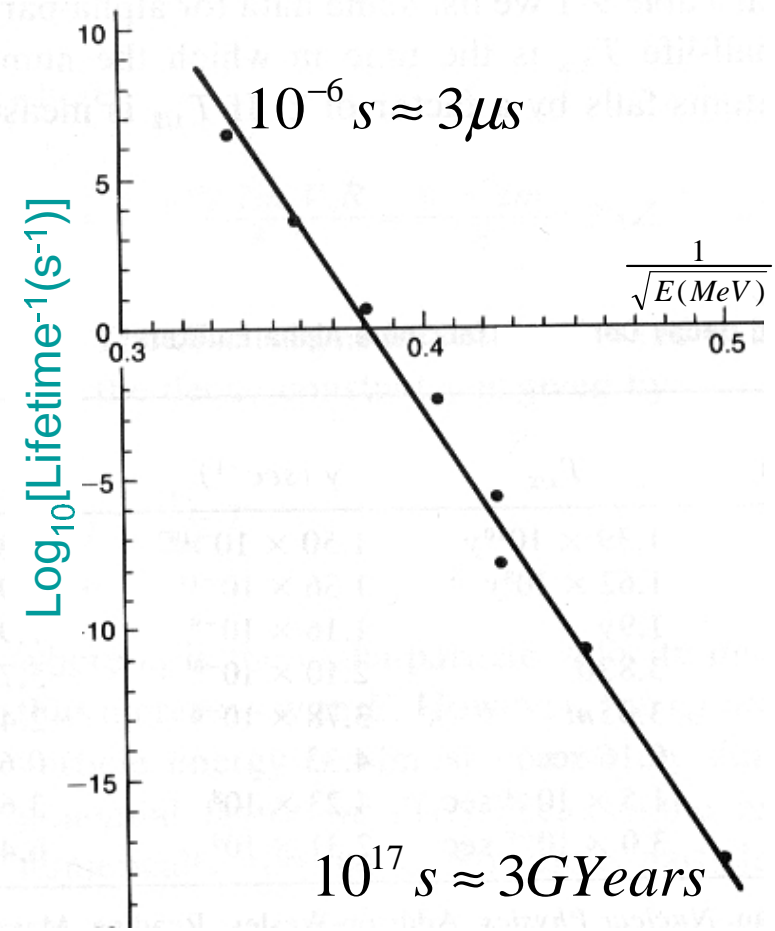
Scaling alpha decay times with Energy

The correct scaling of the alpha decay times with the alpha energies after the decay was one of the early successes of Quantum Mechanics, and of barrier penetration in 1928 (George Gamow).

$$\ln T \propto A - \frac{C}{\sqrt{E}}$$

$$C = \frac{\pi\sqrt{2m}}{\hbar} V_0 R = \frac{\pi\sqrt{2m}}{\hbar} 2(Z-2)e^2$$

$$\text{Lifetime} \propto T^{-1}$$



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The Schrödinger equation in three dimensions

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Review of one dimension:

$\Psi_n = A_n e^{i(k_n x_n - \omega t)}$
 $\Psi_{n+1} = A_{n+1} e^{i(k_{n+1} x_{n+1} - \omega t)}$
 $A_{n+1} = A_n e^{ik_n(x_n + \frac{\Delta}{2})} e^{-ik_{n+1}(x_{n+1} - \frac{\Delta}{2})}$

$$\Psi_{n+1}(x_{n+1} - \frac{\Delta}{2}) = \Psi_n(x_n + \frac{\Delta}{2})$$

$$\Psi_{n+1} = A_n e^{i[k_n(x_n + \frac{\Delta}{2}) - k_{n+1}(x_{n+1} - \frac{\Delta}{2}) + k_{n+1}x_{n+1}]} e^{-i\omega t} = \Psi_n e^{i[k_n \frac{\Delta}{2} + k_{n+1} \frac{\Delta}{2}]}$$

$$= \Psi_n e^{i \frac{k_{n+1} + k_n}{2} \Delta} \approx \Psi_n + ik_n \Psi_n \Delta \quad \rightarrow \quad \underline{\underline{\frac{\partial}{\partial x} \Psi(x, t) = ik \Psi}}$$

Conclusion: whenever $k \Psi$ needs to be computed, one can use $-i \frac{\partial}{\partial x} \Psi$

Dispersion relation $\hbar \omega = \frac{\hbar^2}{2m} k^2 + V(x)$ then leads to

$$i\hbar \frac{\partial}{\partial t} \Psi = -\frac{\hbar^2}{2m} \frac{\partial^2}{\partial x^2} \Psi + V(x) \Psi$$



From the dispersion relation to Schrödinger's equation

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The wave vector \vec{k} changes with position \vec{x} according to: $\frac{\hbar^2}{2m} \vec{k}^2 + V(\vec{x}) = E$
for a particle with fixed constant energy E , where ω does not change.

Time independent Schrödinger's equation

For a stationary state: $\Psi(\vec{x}, t) = \Phi(\vec{x})e^{-i\omega t} = \Phi(\vec{x})e^{-i\frac{E}{\hbar}t}$

$$\frac{\hbar^2}{2m} (k_x^2 + k_y^2 + k_z^2) \Phi + V(\vec{x}) \Phi = E \Phi$$



$$-\frac{\hbar^2}{2m} \left(\frac{\partial^2}{\partial x^2} + \frac{\partial^2}{\partial y^2} + \frac{\partial^2}{\partial z^2} \right) \Phi + V(\vec{x}) \Phi = E \Phi$$

Time dependent Schrödinger's equation

If Ψ_{ω_1} and Ψ_{ω_2} are solutions of for different energies E_1 and E_2 ,

then $\Psi = \Psi_{\omega_1} + \Psi_{\omega_2}$ is a solution of

$$-\frac{\hbar^2}{2m} \left(\frac{\partial^2}{\partial x^2} + \frac{\partial^2}{\partial y^2} + \frac{\partial^2}{\partial z^2} \right) \Psi(x, t) + V(x) \Psi(x, t) = i\hbar \frac{\partial}{\partial t} \Psi(x, t)$$



As in one dimension this holds for an arbitrary linear combination of stationary states.

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