

Confinement Induced Electron Capture

C. Martin^{1, a)} and R. Godes^{1, b)}

Authors' institution and/or address

(Dated: 14 March 2017)

We describe a Gedankenexperiment in which a bare proton can capture an electron due solely to confinement. We first briefly review the aspects around bare electron proton capture. We then provide a numerical solution of the Fermi VA-Theory for electron capture for an electron-proton pair confined in a classical box of size L . Interestingly, we find that the capture is most likely for $L=0.004-0.009$ Angstroms, well beyond the radius of the proton and the Compton wavelength of the electron. We also examine the implications for theoretical minimal power output for such a process.

PACS numbers: Valid PACS appear here

Keywords: Suggested keywords

I. BACKGROUND

We briefly review orbital electron capture and some considerations, like environmental effects on the rate, that motivate this study.

A. Orbital Electron Capture

In 1935, Yukawa proposed that a proton, bound in an atomic nucleus, could capture a low lying, bound atomic electron, transforming into a neutron, and releasing an electron neutrino.

$$p^+ + e^- \rightarrow n^0 + \nu^e$$

This may be called orbital electron capture, K-electron capture, or just electron capture (E.C.)

Electron capture usually occurs in unstable radioisotopes which lack the nuclear binding energy Q to decay by the more familiar Beta decay processes (β^- , β^+). The typical Q energies necessary are

$$Q_{\beta^+} \sim 2 - 4 MeV$$

$$Q_{\beta^-} \sim 0.5 - 2 (MeV)$$

$$Q_{E.C.} \sim 0.2 - 2.0 MeV$$

When $Q < 1.02 MeV$, which is twice the rest mass of an electron ($2m_e c^2$), a proton rich nucleus must decay by electron capture.

In particular, heavy elements may decay by E.C. and/or β^+ (positron emission) to a lower 'magic number' of stable nuclei, and by β^- decay to achieve a higher

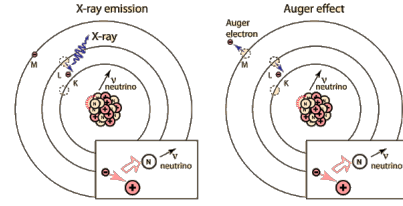
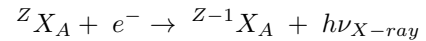


FIG. 1. orbital electron capture relaxation processes

magic number. E.C. is favored for high Z nuclei, but because of the energetic constraint, very light elements, such as 7Be , decay by primarily by E.C. Lacking any binding energy and/or internal nucleon structure, bare proton electron capture is not observed, and requires extreme, exotic conditions.

We observe electron capture by observing the resulting transmuted nuclei and/or the radiative relaxation processes. The captured electron is bound to the atom, and it is usually a K-shell electron, but may be L or higher. The nucleus may then absorb some energy, becoming excited, and then undergo internal conversion. During this, another higher lying, bound atomic electron is absorbed, and either an X-ray or Auger electron released [see Figure 1].

Because electron capture occurs in proton-rich nuclei, and, subsequently, releases a X-Ray photon, the reaction is also sometimes written as



(where Z is the total number of protons and neutrons, A the number protons, and $h\nu_{X-ray}$ is an X-ray photon)

Indeed, orbital electron capture is evidenced by high intensity x-rays and soft electrons. In 1938, Luis W. Alvarez observed the x-ray signature of orbital electron capture in activated Titanium¹. Since then, electron capture has been observed in about 150 radioactive isotopes.

^{a)} Also at Physics Department, XYZ University.

^{b)} Electronic mail: Second.Author@institution.edu.

1. Effect of Chemical Environment of Electron Capture rate

The lightest element that E.C. has been observed in is ${}^7\text{Be}^2$. In fact, there is so little energy that the competing β^+ positron emission process (described below) is prohibited, leading to a fairly long E.C. half-life of $\tau_{EC} \sim 50 \text{ days}$.

Being so light, and having such a large rate, electron capture in ${}^7\text{Be}$ can be slightly modified by both changing the chemical environment and/or the external pressure³⁻⁵. In particular, in 2004, Ohtsuki et. al. demonstrated a change of 0.83% by embedding Be in C-60 cages⁵.

How could such changes occur? The nuclear energy levels are in the keV to MeV region, and it is generally thought to be very difficult to impossible to effect. But the electron capture rate is proportional to electronic density at the surface of the nucleus—the *nuclear charge*. The electronic energy levels are in the eV range, so intense EM fields can alter the electronic structure and therefore slightly affect the E.C. rate.

B. The Weak Interaction and V-A theory

Electron capture is mediated by the Weak interaction, described most concisely by the Fermi V-A (Vector Axial) theory⁶⁻⁸. The V-A theory is a simple phenomenological approach, readily amenable to numerical calculations. While It is now understood in terms of ElectroWeak Unification and can be derived from the Standard Model, the original paper by Fermi, for which he won the 1938 Nobel Prize in Physics, was initially rejected by *Nature* because "It contained speculations too remote from reality to be of interest to the reader"⁹.

V-A theory is used to compute cross sections for scattering experiments and decay rates for electron capture for various atoms, even in different environments, chemical and otherwise. We can use machinery of the V-A theory to explore E.C. in a simple, idealized environment. To properly describe any reaction, however, we need to understand what reactions we can apply the theory to, and the other, potential competing reactions.

1. Electron Capture and other Weak processes

The Weak interaction describes several related processes within a single framework. *Neutron-rich* nuclei may become more stable as a result by undergoing one or more of the following:

- orbital electron capture $p^+ + e^- \rightarrow n^0 + \nu_e$
- positon emission (β^+ decay) $p^+ \rightarrow n^0 + e^+ + \nu_e$
- β decay $n^0 \rightarrow p^+ + e^- + \bar{\nu}_e$

There are also several related reactions, including

- reverse electron capture $n^0 + \nu_e \rightarrow p^+ + e^-$
- free neutron decay $n^0 \rightarrow p^+ + e^- + \bar{\nu}_e$
- inverse beta decay $p^+ + \bar{\nu}_e \rightarrow n^0 + e^+$

2. Neutron Decay

By detailed balance, reverse electron capture has the same rate as orbital electron capture—but is more favorable energetically. Indeed, inside the nucleus, the neutron is relatively stable. Free neutron decay has mean lifetime of $\tau = 881.5 \pm 1.5 \text{ sec}$, or about 15 minutes.

In contrast, orbital electron capture by a free proton is unspoken-of outside of a stellar environments. Even if the bare reaction could proceed, the reverse reaction would still dominate unless it is suppressed or is kinetically unfavorable.

3. Inverse Beta Decay

Electron capture is also sometimes called inverse β decay, but, here, we mean this to be the scattering of a proton and an electron anti-neutrino $\bar{\nu}_e$. It is characterized by emission of a positron e^+ .

4. Positron emission

In particular, any high energy relativistic process, we have to worry about positron emission. We noted above, however, that in ${}^7\text{Be}$, the competing positron decay reaction can not occur because there is not enough energy. Generally, this occurs at length scales below the Compton length of the electron, which, is smaller than we will need to consider.

5. Radiative Electron Capture

Also, in very rare cases, a gamma ray photon is emitted with the neutrino; this is called Radiative Electron Capture (REC)¹⁰⁻¹³. This can be thought of as a kind of Internal Bremsstrahlung (or so-called *braking*) radiation, caused by the electron accelerating toward the nucleus during capture, taking energy away from outbound neutrino¹⁴. It is traditionally been treated as a second order QED correction to the V-A theory¹². REC is 1000X less likely, but does occur. The resulting gamma rays are called soft because they do not exhibit sharp spectral lines. Recent, detailed rate calculations have elucidated the quantum mechanical details¹³.

C. Rate calculations

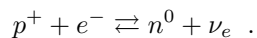
The electron capture rate can be computed using the Fermi VA-Theory^{7,8}. The most basic calculations require a only specifying the electronic wavefunctions(s), averaging over the possible electron-proton momenta, and numerically integrating over the outbound neutrino momentum. We describe this below.

The V-A theory is based on second order perturbation theory. It assumes an incoherent nuclear process, it is local, and that the interaction is phenomenological. It is treated as a simply a contact potential at the surface of the nucleus. Here, this means we need to compute the nuclear charge—the electron density on the surface of the nucleus $\|\psi_e(R=0)\|$, which we obtain from a classical particle-in-the-box wavefunction (below).

More complicated calculations are used for larger nuclei, second order processes, etc. They only require modifications to treat either atomic electronic structure of reactant and product atoms, and/or specific considerations for nuclear internal conversion and other second order processes.

1. Bare proton-electron capture

We frequently write electron capture as if it were simply proton-electron capture



At zero energy, proton electron capture is not possible because it violates energy-momentum conservation. Theoretically, a free proton can capture an electron from the continuum, but the interaction energy must be above threshold for neutron production. This is a huge amount of energy, although this happens regularly in accelerators, and, presumably, in stellar environments.

Observing proton electron capture outside of an accelerator, *on the desktop*, so to speak, would be an incredibly hard experiment because both final particles are neutral, and the neutrino is extremely weakly interacting.

2. Stellar Nucleosynthesis

Bare proton-electron capture is thought to occur in the early stages of the big bang and in stellar nucleosynthesis. It is thought to drive the formation of primordial elements, and to occur in the forming of neutron stars.

At very high temperatures, the proton electron collisions have sufficient energy to overcome the reaction barrier. For example, Bachall and coworkers have studied the electron capture rate in stellar media, and have computed the electron capture rate of ${}^7\text{Be}$ in the Sun^{15,16}. And it is believed that ionized Hydrogen captures an electron during the core collapse supernovae and in neutron stars [14]; in fact, it is thought to create stellar instability.

While we usually characterize a star by its temperature, these are also very dense systems, with $\rho \sim 10^6 \text{ g cm}^{-3}$. In contrast, the smallest star has density $\rho \sim 10^2 - 10^3 \text{ g cm}^{-3}$ [?]. As important, the reverse reaction is prohibited because, inside the dense neutron star, it is impossible to create a new electron; the Fermi sea is *full*.

So electron capture can occur by bare protons, but, presumably, only under extreme confinement (and with the reverse reaction is suppressed).

3. Metallic Hydrides

It has been suggested that bare proton-electron capture could actually arise metallic hydrides, such as PdH and NiH, as a result of some exotic, here-to-fore unrecognized physical mechanism. Widom and Larsen, in particular, have proposed that the effective Q value for bare proton E.C. would be lowered because the surface electrons may undergo mass renormalization due surface plasmon radiation, leading to high order radiative corrections to the Weak interaction pair production process¹⁷. One may define an effective mass \tilde{m} using

$$\tilde{m}c^2 \cong -\overline{p^\mu p_\mu} = mc^2 + \left(\frac{e}{c}\right)^2 + \overline{A^\mu A_\mu}$$

where A^μ is the gauge potential for the fluctuating surface plasmon radiation, and assuming $\overline{p^\mu A_\mu} = 0$

They argue that such a renormalization may be possible because the neutron scattering experiments on PdH show a collective proton oscillation that would be resonate with electronic surface plasmons.

Neutron scattering experiments also show that in PdH, the proton is confined such that the root-mean-square vibrational amplitude of the hydrogen atoms is be $0.25A$ (at $T = 293K$), which is anomalously small¹⁸.

II. CONFINEMENT INDUCED ELECTRON CAPTURE

We pose the following Gedankenexperiment: Suppose we confine a bare proton (and an electron) in a particle-in-a-box of volume L^3 . What box size L will 'induce' electron confinement?

We write this as

$$E_{box} + p^+ + e^- \rightarrow n^0 + \nu_e$$

where E_{box} is the *confinement energy*, which is induced by the box constraints.

1. Neutron post-reaction

To prevent the reverse reaction, we assume that the free neutron subsequently combines with another proton,

and gives us 2.2 GeV of energy in the process. This post-process contributes to the power output, and prevents the reverse reaction. Realistically, we expect this to happen at the maximum box size, at the energy threshold, where the outbound neutron has very low momentum and therefore a very small mean free path.

Still, for illustrative purposes, we will compute the power output, assuming this post-reaction, at all box sizes.

A. Compton length

The first obvious question is, should we use a classical or a relativistic box?

Most electron capture rate calculations use *ab initio* classical wavefunctions^{7,8}, perhaps with some relativistic corrections to the electronic Hamiltonian¹⁹.

We argue that we can safely use a classical box as long as $L_{min} \geq \frac{1}{2\pi} \lambda_e$, where λ_e is Compton wavelength of an electron²⁰⁻²². The Compton wavelength sets the scale, accounting for both quantum mechanics and special relativity.

$$\lambda_e = \frac{h}{m_e c} = \frac{e^2}{m_e c^2}$$

$$\lambda_e \approx 2.426 \times 10^{-12} \text{ m}$$

For an electron, the minimum L is on the order of 0.004 Angstrom

$$L_{min} \sim 0.004 \text{ \AA}$$

In any high energy, relativistic system, positrons can be produced; here it is through β^+ -decay. This generally occurs at or below the Compton length. We are seeking the maximum box size which can induce electron capture, and we assume that, at the max, positron emission will be very rare.

We also assume that the electron wavefunction does not change appreciably during the interaction, so that we may use a very simplified form for the cross section (σ) and rate (Γ). Again, this is reasonable for boxes $L \geq \lambda_e$.

1. Klein Paradox

Klein noted that a relativistic (Dirac) particle-in-a-box will *leak out* at box sizes near the Compton wavelength; this called the Klein paradox^{23,24}. And while this is usually taught as being simply particle-antiparticle creation, it has been suggested that the Klein paradox it can occur even at larger boxes sizes, and it is a general phenomena of confined relativistic particles. And recent experiments

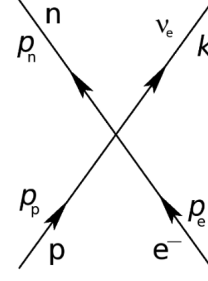


FIG. 2. EC particle production process

on Graphene have reopened the debate²⁵. Still, we will assume the traditional interpretation and that we can ignore the Klein paradox.

So we use a classical box, with minimum size $L_{min} = 0.004 \text{ \AA}$. We compute the maximum size below.

B. Theory

1. Particle production under the Weak interaction

Electron capture is mediated by the Weak Interaction, through the particle production process, given by the 4-point Interaction (see Figure 2). It requires *at least* 0.511 MeV energy to overcome the reaction barrier, which, here, is *provided* by the box.

$$0.511 \text{ MeV} + p^+ + e^- \rightarrow n^0 + \nu_e$$

We want to compute the rate of E.C. and the (minimum) power generated, as a function of the box size (L), using the full Weak Interaction Hamiltonian. To do this, we need to express the rate in terms of the differential cross section, and in a form suitable for numerical calculations.

C. Transition Rates and Power Calculations

Orbital electron capture and other Beta-decay processes follow first order kinetics, so the capture rate is described by a single number. To calculate the rate, we require a full relativistic, quantum mechanical treatment because the capture process involves both creating particles and the kinetic energy spectrum is of order $m_e c^2$.

The capture rate Γ_{EC} is given by multiplying the cross section σ_{EC} by the incident velocity \mathbf{v}_{ep}^{in} and electronic (box) density $|\Psi_{ep}(0)|^2$ of the (e^-, p^+) pair at the origin

$$\Gamma_{EC} = |\Psi_{ep}(0)|^2 \mathbf{v}_{ep}^{in} \sigma_{EC} .$$

We will compute the electronic density classically, using box wavefunctions, and, for each box size, determine

the incident velocity (really momentum). We then compute the cross section using full relativistic kinematics and Dirac spinors.

1. Cross Section

We write the differential cross section in the C.M. frame as

$$d\sigma_{EC} = \left(\frac{1}{2\pi}\right)^2 \frac{\sum_{fi} |\mathcal{M}_{fi}|^2}{16 |\mathbf{k} \cdot (E_n \mathbf{k} - k^0 \mathbf{p}_n)|} \frac{k^3 p_e d\Omega_k}{|\mathbf{p}_e \cdot (E_p \mathbf{p}_e - E_e \mathbf{p}_e)|}$$

where \mathcal{M}_{fi} is a matrix element of the weak interaction Hamiltonian, k represents the neutrino, and we use natural units ($\hbar = 1, c = 1$).

2. Rate

This gives the (differential) rate as

$$d\Gamma_{EC} = \left(\frac{1}{2\pi}\right)^2 \frac{\sum_{fi} |\mathcal{M}_{fi} [p_{ep} \rightarrow i\nabla]| \psi_{ep}(\mathbf{x})|_{\mathbf{x}=0}|^2}{16 E_p E_e |\mathbf{k} \cdot (E_n \mathbf{k} - k^0 \mathbf{p}_n)|} k^3 d\Omega_k$$

where we explicitly specify the electronic wavefunction.

We represent the (e^-, p^+) pair using a 3D box wavefunction, and obtain the Energies and 3-momenta from the relativistic kinematics. We then average over all 8 permutations of the incident velocities (really momenta \mathbf{p}_{ep}) for the 3D box, and integrate over outbound neutrino solid angle $d\Omega_k$ using numerical quadrature. The final rate is computed as, for each box size, as a function the kinematics, using

$$\Gamma_{EC} = \frac{1}{8} \sum_{\mathbf{p}_{pe}} \int_{d\Omega_k} d\Gamma_{EC}$$

where $|\mathbf{p}_{ep}|$ is given by the box size (described below).

3. Power

We estimate the excess power generated by the confined electron capture, resulting if/when the outbound neutron reacts with the environment. The power \mathcal{P} is

$$\mathcal{P} = \Gamma_{EC} * Q * \rho$$

where Q is the nuclear decay energy, or Q-value, and ρ is the density of confined elements. We choose the density to that of a typical material, of order Avogadro's number, $N_A \sim 6.02 \times 10^{23}$.

We estimate the power for both the bare proton-electron capture

$$\mathcal{P}_{pe} = \Gamma_{EC} * [(\mathbf{E}_e - m_e) + (\mathbf{E}_p - M_p)] * \rho$$

and the subsequent neutron post-reaction

$$\mathcal{P}_n = \Gamma_{EC} * [2.2 + (\mathbf{E}_n - M_n)] * \rho$$

We also define an excess power as

$$\mathcal{P}_{XS} = \mathcal{P}_{pe} - \mathcal{P}_n$$

which we argue is a good measure of the potential reactive power output of the confined E.C. process. We are interested in box lengths where the excess power is significant enough to drive the reaction.

4. Particle-in-the-box Wavefunctions

While most electron capture calculations assume a specific, bound, atomic electronic wavefunction(s), we perform a much simpler calculation; we treat the electron-proton pair as classical particle-in-a-box, and analyze the problem just above the Compton scale using the low order VA theory.

$$\psi_{ep}(\mathbf{x}) = \left(\frac{2}{L}\right)^{\frac{3}{2}} \cos\left(\frac{\pi x}{L}\right) \cos\left(\frac{\pi y}{L}\right) \cos\left(\frac{\pi z}{L}\right)$$

expressions for momentum.

Because the VA theory assumes an incoherent process, the electron, proton wavefunction is usually factored as an electron wavefunction, with a point-particle in the center

$$\psi_{ep}(\mathbf{x}) = \psi_p(0) \psi_e(\mathbf{x})$$

We only consider the ground state ψ_{ep}^0 wavefunction.

We note, in *ab initio* electronic structure calculations, it is now generally possible to treat the Hydrogen proton wavefunction explicitly, and to treat the electron-proton coupling at the level of Hartree Fock¹⁹. This has proved useful, for example, for describing isotope effects on electronic structure.

Here, we treat the confined electron, proton pair in the C.M. frame so that the 3-momentum of the electron and proton related

$$\mathbf{p}_e = \mathbf{p}_{pe}$$

$$\mathbf{p}_p = -\mathbf{p}_{pe}$$

and we consider all 8 permutations for a given box size L :

$$\pm \mathbf{p}_{pe}(1), \pm \mathbf{p}_{pe}(1), \pm \mathbf{p}_{pe}(3)$$

5. Relativistic Kinematics and Energetics

For consistency with the particle production process, we treat all kinematics and energetics relativistically. Given energy-momentum conservation

$$E^2 = m^2 + \mathbf{p}^2$$

and the 3D particle-in-the-box energy ground state energy

$$E_{gs} = \frac{3\pi^2}{2mL^2} ,$$

we use $E = \frac{\mathbf{p}^2}{2m}$ to write

$$E_e^2 = m_e^2 + 3 \left(\frac{\pi}{L} \right)^2 ,$$

and

$$E_p^2 = M_p^2 + 3 \left(\frac{\pi}{L} \right)^2 .$$

The threshold Kinetic energy in the center of momentum (C.M.) frame is given as

$$EK_{e_{min}} : K_e = E_e - m_e = \frac{(M_n - m_e + m_\nu)^2 - M_p^2}{2(M_n + m_\nu)}$$

which is approximately 781.6 KeV (or 783.1 KeV in the proton rest frame).

Of course, the proton rest frame is approximately the electron rest frame, but it should be mentioned that the in the electron rest frame, the threshold kinetic energy is 2000X greater. Therefore, it is assumed that the energy transfer to induce electron capture is in the C.M. frame of the (e,p) pair.

The minimum momentum is

$$\mathbf{p}_{min} = \sqrt{(K_e + m_e)^2 - m_e^2} .$$

For the final state, the neutrino (ν) kinetic energy is

$$K_\nu = k^0 - m_\nu = \frac{(E_p + E_e - m_\nu)^2 - M_n^2}{2(E_p + E_e)}$$

and the neutron (n) kinetic energy is

$$K_n = E_n - M_n = \frac{(E_p + E_e - M_n)^2 - m_\nu^2}{2(E_p + E_e)}$$

At this point, we can compute the relativistic energies for all particles, simply as a function of box length (L),

for a confined (e^-, p^+) pair. Of interest is the shape of the curve, and when the excess power output becomes favorable. To compute the exact rate and estimate the power, we need to evaluate the matrix element(s) of the Weak Interaction Hamiltonian. While most practitioners simply estimate this, for our purpose, we can simply compute them exactly.

[maybe show the curve, and state the maximum box size. or wait until later ?]

[say something about estimating the size of the interaction here, which should be enough for a baseline calculation]

D. Weak Interaction Hamiltonian and Matrix Elements

The Hamiltonian for the V-A theory for E.C. is^{7,8}

$$\mathcal{H}(x) = -\frac{G_F}{\sqrt{2}} [J^\mu(x) L_\mu^+(x) + h.c.]$$

where J^μ and L_μ are (in modern parlance) the Hadron and Lepton currents, resp., and are given by

$$J_\mu = \bar{u}_n \gamma_\mu (1 + \lambda \gamma_5) u_p$$

$$L_\mu = \bar{u}_\nu (1 - \gamma_5) u_e$$

where u_n, u_p, u_e, u_ν are Dirac (free-particle) wavefunctions.

G_F is the Universal Fermi Weak Coupling Constant, and $\lambda = -\frac{G_A}{G_V}$, which is determined by experiment (and subject to minor changes). G_V is the Axial-Vector Weak Coupling Constant, and G_A is the vector weak coupling constant. The most recent value is $G_V = 1.2767(16)$, and $G_A = 1^{26}$.

1. VA Matrix Elements

In our calculations, however, we use a slightly different, more modern convention for the matrix elements, multiplying through by G_V^{27} . The matrix elements \mathcal{M}_{fi} are then given by

$$\mathcal{M}_{fi} = \frac{G_F}{\sqrt{2}} \bar{u}(p_n, s_n) (G_V - G_A \gamma^5) \gamma^\mu u(p_p, s_p) \times \bar{u}(p_\nu, s_\nu) \gamma_\mu (1 - \gamma^5) u(p_e, s_e)$$

where $u(p, s)$ are Dirac 4-spinors, with the convention

$$u(p, s = 1) = \sqrt{E + m} \begin{bmatrix} 1 \\ 0 \\ \frac{p_3}{E+m} \\ \frac{p_1 + ip_2}{E+m} \end{bmatrix}$$

$$u(p, s=2) = \sqrt{E+m} \begin{bmatrix} 0 \\ 1 \\ \frac{p_1 - ip_2}{E+m} \\ \frac{p_3}{E+m} \end{bmatrix}$$

$$\langle \bar{u} | u \rangle = \bar{u}_0 u_0 - \bar{u}_1 u_1 - \bar{u}_2 u_2 - \bar{u}_3 u_3$$

and $\bar{u} = u^\dagger \gamma^0$. The γ are 4-component Gamma matrices, and $(\dots \gamma^\mu \dots \gamma_\nu \dots)$ is the Einstein summation convention.

Writing the matrix elements in terms of the currents gives

$$\mathbf{J}_{had} = \bar{u}(p_n, s_n)(G_V - G_A \gamma^5) \gamma^\mu u(p_p, s_p)$$

$$\mathbf{L}_{lep} = \bar{u}(p_\nu, s_\nu) \gamma_\mu (1 - \gamma^5) u(p_e, s_e)$$

[Express power in terms of real part of current]

Notice that when applying the VA theory this way, we assume that the electron-proton interaction is a contact potential, operating at the surface of the nucleus, and that the underlying quantum process is incoherent.

[how much more to describe here ? Specify the 4-norm / gamma metric tensor convention?]

2. Sum over all spin combinations

[[I have not yet decided how much of this section to include and work out. The idea is to look at the detailed spin combinations of the matrix elements, and present an analysis similar to the discussion in "Beta Decay for Pedestrians". this is relevant since , as we see below, even though we sum over 16 matrix element combinations, only 4 are non-negative in this very specific system. it is possible, in other environments, that forbidden transitions might be allowed.. I also wanted to discuss this to grok the psudecode.]]

We sum over the 16 allowed transitions in the V-A theory, as depicted in Figure[?]. We note, however, that due to symmetry, only 4 spin configurations contribute appreciably:

$$\bar{u}(p_n, 1) u(p_p, 1) \bar{u}(p_\nu, 2) u(p_e, 2)$$

$$\bar{u}(p_n, 2) u(p_p, 2) \bar{u}(p_\nu, 1) u(p_e, 1)$$

$$\bar{u}(p_n, 2) u(p_p, 1) \bar{u}(p_\nu, 2) u(p_e, 2)$$

$$\bar{u}(p_n, 1) u(p_p, 2) \bar{u}(p_\nu, 2) u(p_e, 1)$$

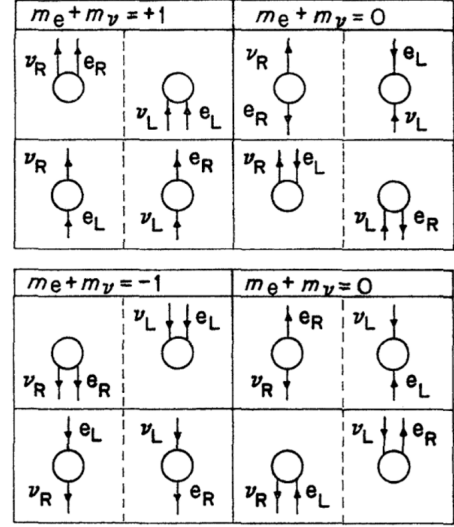


Fig. 1.1. The sixteen possible allowed transitions.

FIG. 3. Allowed spin transitions

E. Numerical Calculations

The full calculations sums of over all 8 possible incident momenta permutations $(\pm \mathbf{p}_{ep}(1), \pm \mathbf{p}_{ep}(2), \pm \mathbf{p}_{ep}(3))$, and the 16 allowed spin transitions (see figure below). By conservation of momenta, we can eliminate the neutron and only need to average over the outbound neutrino momenta solid angle $(d\Omega_k)$; we do this using an 8-point gaussian quadrature.

1. Power Calculations

[[Again, I have not worked this out in detail. In particular, we discuss the power above, so this is just for understanding the code, and I may leave this to the python notebook. It is necessary , however, to really grok all the pseudocode]]

$$\mathcal{F} = r_c \left(\frac{p_{ep}}{2\pi} \right)^3 \left(\frac{G_F^2 (E_\nu^2 - m_\nu^2)^3}{512\pi^2 E_p E_e \hbar c |\mathbf{k}(E_n p_\nu - E_\nu p_n)|} \right)$$

F. Results

Figure 4 presents the results of relativistic kinematics and excess power calculations, as a function of box length. For illustrative purposes, we include results box sizes even below the Compton length. The excess power ratio is positive for $L > 0.004\text{\AA}$, with a maximum of almost a factor of 3.

Algorithm 1 Rate and Power Calculations

<pre> 1: for $\mathbf{p}_{ep} \in [\mathbf{p}_{max}, \mathbf{p}_{min}]$ do 2: $Tran(\mathbf{p}_{ep}) = 0$ 3: $Power_n(\mathbf{p}_{ep}) = 0$ 4: for $w_x, w_\phi \in \mathbb{Q}[1, 8]$ do 5: for $s_n, s_p, s_\nu, s_e \in [0, 1]$ do 6: for $s_x, s_y, s_z \in [-1, 1]$ do 7: $p_n, p_p, p_\nu, p_e \leftarrow pekin(w_x, w_\phi, \mathbf{p}_{ep})$ 8: $\bar{u}(p_n, s_n), u(p_p, s_p), \bar{u}(p_\nu, s_\nu), u(p_e, s_e)$ 9: for $u \in [1, 4]$ do 10: $\mathbf{J}_{had} = \bar{u}(p_n, s_n) \cdot (G_V - G_A \gamma^5) \cdot \gamma^\mu \cdot u(p_p, s_p)$ 11: $\mathbf{L}_{lep} = \bar{u}(p_\nu, s_\nu) \cdot \gamma_u \cdot (1 - \gamma^5) \cdot u(p_e, s_e)$ 12: $\mathbb{C} = \langle \mathbf{L}_{lep}^\dagger, \mathbf{J}_{had} \rangle$ 13: $\mathcal{F}^2(\mathbf{p}_{ep}, p_p, p_e, p_n, p_\nu)$ 14: $Tran(\mathbf{p}_{ep}) += w_x w_\phi Re[\mathbb{C}^\dagger \mathbb{C}] \mathcal{F}^2$ 15: $Power_n(\mathbf{p}_{ep}) = Tran(\mathbf{p}_{ep}) * (2.2 + (E_n - M_n))$ </pre>	<pre> ▷ $\mathbf{p}_{ep} = \ \mathbf{p}_{ep}(L)\$ ▷ Transfer ▷ Neutron Power ▷ quadrature weights ▷ $s_{up}, s_{down} = 0, 1$ ▷ $\mathbf{p}_{ep} = \mathbf{p}_{ep}[s_x, s_y, s_z]$ ▷ 4 vectors \leftarrow rel. kinematics ▷ spinors by p, s-index ▷ 4-vector indices ▷ Hadronic current ▷ Leptonic current ▷ Current Amplitude ▷ Matrix Element Factor ▷ Integrated Rate ▷ Neutron Power estimate </pre>
--	---

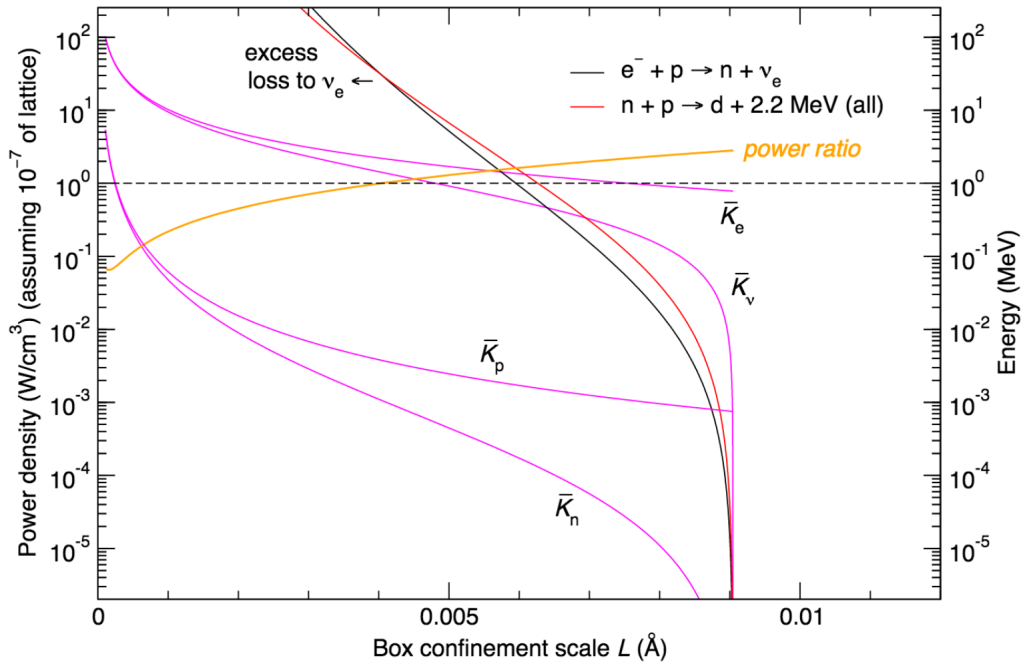


FIG. 4. Kinetic Energy and Power as a function of Box Length

ACKNOWLEDGMENTS

We wish to acknowledge the support of the Anthro-
pocene Institute . . .

Appendix A: Appendixes

To start the appendixes, use the `\appendix` command. This signals that all following section commands refer to appendixes instead of regular sections. Therefore, the `\appendix` command should be used only once—to set up

the section commands to act as appendixes. Thereafter normal section commands are used. The heading for a section can be left empty. For example,

```
\appendix
\section{}
```

will produce an appendix heading that says “APPENDIX A” and

```
\appendix
\section{Background}
```

will produce an appendix heading that says “APPENDIX

A: BACKGROUND” (note that the colon is set automatically).

If there is only one appendix, then the letter “A” should not appear. This is suppressed by using the star version of the appendix command (`\appendix*` in the place of `\appendix`).

Appendix B: A little more on appendixes

Observe that this appendix was started by using

`\section{A little more on appendixes}`

Note the equation number in an appendix:

$$E = mc^2. \quad (\text{B1})$$

1. app:subsecA subsection in an appendix

You can use a subsection or subsubsection in an appendix. Note the numbering: we are now in Appendix ??.

a. app:subsubsecA subsubsection in an appendix

Note the equation numbers in this appendix, produced with the subequations environment:

$$E = mc, \quad (\text{B2a})$$

$$E = mc^2, \quad (\text{B2b})$$

$$E \gtrsim mc^3. \quad (\text{B2c})$$

They turn out to be Eqs. (B2a), (B2b), and (B2c).

¹L. W. Alvarez, “The capture of orbital electrons by nuclei,” **54**, 486 (1938).

²M. F. L’Annunziata, *Radioactivity: Introduction and History and From the Quantum to Quarks* (Second Edition and Elsevier and, 2016).

³B. W. et al, “Change of the ${}^7\text{Be}$ electron capture half-life in metallic environments,” (2006).

⁴A. Ray, P. Das, S. K. Saha, A. Goswami, and A. De, “Enhanced orbital electron-capture nuclear decay rate in compact medium,” **679**, 106 (2009).

⁵T. Ohtsuki, H. Yuki, M. Muto, J. Kasagi, , and K. Ohnol, “Enhanced electron-capture decay rate of Be-7 encapsulated in c-60 cages,” **93(11)**, 112501 (2004).

⁶E. Fermi, “An attempt of a theory of beta radiation.1,” **88**, 161 (1934).

⁷W. Bambynek, H. Behrens, M. H. Chen, and R. L. Intemann, “Orbital electron capture by the nucleus,” **49**, 1 (1976).

⁸R. Bouchez and P. Deponimier, “Orbital electron capture by the nucleus,” **23**, 395 (1960).

⁹F. Close, “Small things and nothing,” (2013).

¹⁰R. J. Glauber and P. C. Martin, “Radiative capture of orbital electrons,” **104**, 158 (1956).

¹¹R. J. Glauber and P. C. Martin, “Relativistic theory of radiative orbital electron capture,” **109**, 1307 (1958).

¹²J. M. Jauch, “On the theory of inner bremsstrahlung in β decay,” (1951).

¹³K. Pachucki and U. Jentschura, “Radiative orbital electron capture by the atomic nucleus,” **75**, 055502 (2007).

¹⁴J. D. Jackson, *Classical Electrodynamics* (Third Edition and Wiley and, 1998).

¹⁵J. N. Bahcall, “Electron capture and nuclear matrix elements of ${}^7\text{Be}$,” **128**, 1297 (1962).

¹⁶A. V. Gruzinov and J. N. Bahcall, “The ${}^7\text{Be}$ electron capture rate in the sun,” **155**, 511 (1969).

¹⁷A. Widom and L. Larsen, “Theoretical standard model rates of proton to neutron conversions near metallic hydride surfaces,” (2007).

¹⁸G. A. F. Jr., A. I. Schindler, T. Tanaka, , and T. Morita, “Neutron diffraction study of temperature-dependent properties of palladium containing absorbed hydrogen,” *Phys. Rev* **137(A483)** (1965).

¹⁹C. H. Martin and M. C. Zerner, *Electronic Structure Calculations on Transition Metal Complexes: Ab Initio Models and Approximate Models* (Inorganic Electronic Structure and Spectroscopy and Volume I: Methodology and E. I. Solomon and A. B. P. Lever and Wiley, 2006).

²⁰P. Alberto, C. Fiolhais, and V. M. S. Gil, “Relativistic particle in a box,” **17**, 19 (1996).

²¹Garay and L. J, “Quantum gravity and minimum length,” **A10**, 145 (1995).

²²B. Sidharth, “Planck scale to compton scale,” (2006).

²³Bjorken and Drell, *Relativistic Quantum Mechanics* (McGraw-Hill, 1964).

²⁴N. Dombey and A. Calogeracos, “70 years of the klein paradox,” **315d** (1999).

²⁵M. I. Katsnelson¹, K. S. Novoselov², and A. K. Geim, “Chiral tunnelling and the klein paradox in graphene,” **2**, 620 (2006).

²⁶M. et. al, “Determination of the weak axial vector coupling from a measurement of the beta-asymmetry parameter a in neutron beta decay,” .

²⁷C. Itzykson and J. Zuber, “Quantum field theory,” (1985).