heavy fast charge light atomice; mass me mess m, speed v, charge Ze theory anchors

We've been boling of this problem M. We comprted

dE

1 (4TH 60)2 4 TT N Z 22 e 4

atoms electrons

vol every every len (bruin)

We registed the electron binding energy (\$\frac{1}{2} \text{ atomiz} \text{ wavefunction}) treating . It as a free point charge. This is whit the energy transfer is large compared to the quantum mechanizal binding energy - ie, we computed the ronization energy loss, assuming small deflections (bun = \frac{7}{2} \cdot \gamma\_{max}^2) and fast ollisions (bun = \frac{7}{2} \cdot \gamma\_{max}^2) and fast ollisions

As in our discussion of bremsstrahlung, Quinten mechanics can modify boun in some regimes.

Uncertainly principle: Ap 1x = ti/2. We compreted a classical momentum transfer Dp = 22e2 with classical impact parameter b. The classical appx requires (Ap)(b) = 27e2 > tr/2 Or 42ez 21. If this is villated, we should use a different estimate to get 6 min. The bean particle "sees" an electron coming at it long todinelly " frame | b = 8 mev e In order to- this moment on to make sense as "anostly long. todard," the spread in transverse momenta around zero needs to be smaller than it! (typical Ptromsverse) & (Vanev)

The quantum uncertainty in Procusurese 3 O(th/binn) if we demand that the et is localized transversely on scales of order binn (to be determined.)

So, we require themin < 8 me v o- bonn = to/8mer. We could do the same argument for the e rest frame and And bown = ty Sour & beam mass but she we assumed m> me the requirement burn = the ris the more stringent of the two, and we must take it. So, if ret is smell, we must switch from the classical boin = Ze /4TTEO Your to the quantum m= 1711co.e

2 my/

town

bump quantum

La typical

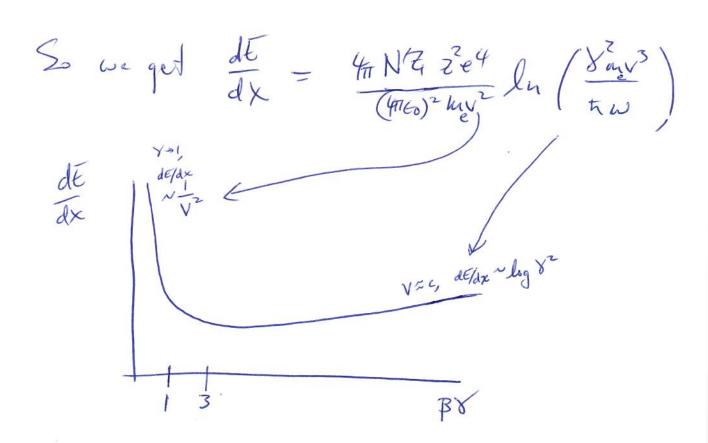
atomic

bregionny. boin = ti/mer. powers of & in this!

F The

(1) The transverse E field grows with 8. Therefore as 8 increases we can increase the impact parameter and Still get a high frequency kich

(2) energy transfer goes up, so bank ulf



This is useful in experiments: energy deposed / unit length is almost independent of energy for relativistic particles.

The rise at small v is why contaction >> brem for nouvelativists particles.

The stopping powers were tally O(1) (MeV/cm) x (1/9/13)

## Dens. Ly effects

So far we've ignored the fields of other atoms near the beam particle. In reality the polarization of the material in the vicinity of the beam will reduce the effectiveness of energy transfer.

lets see what happens of we put in a dielectric constant.

A long there ago, we wrote down the Lorentz garge wave equations for the scalar & vector potential!

$$\frac{1}{c^2} \frac{\partial^2 \varphi}{\partial t^2} - \nabla^2 \varphi = \frac{P}{40}$$

$$\frac{1}{c^2} \frac{\partial^2 \vec{A}}{\partial t^2} - \nabla^2 \vec{A} = \frac{P}{40}$$

These were invacuum. In a non-magnetic medium with dielectric constant 6, there are two changes:

(a) polaritation reduces the field

from a point charge 1/60 -> p/6

(b) phase velocity c-> 1/5=10. = c

The wave equalities become;

$$\frac{1}{\overline{c}^2} \frac{\partial^2 Y}{\partial t^2} - \nabla^2 Y = P/\epsilon$$

Let's define a 4D Fourier transform:

$$F(x,t) = \int_{-\infty}^{\infty} \frac{d^3k dw}{(2\pi)^4} e^{i(\vec{k}\cdot\vec{x}-\omega t)} \widetilde{F}(\vec{k},\omega)$$

Plugging in the transforms for gis, found A, we find

$$\left(-\frac{\omega^2}{\tilde{c}^2} + k^2\right) \tilde{\varphi}(k, \omega) = \frac{1}{\epsilon} \tilde{\rho}(k, \omega)$$

$$\left(-\frac{\omega^2}{c^2} + h^2\right) \tilde{A}(h, \omega) = \mu_0 \tilde{J}(h, \omega)$$

Now let assume our 
$$\approx$$
 const velocity point  $2e$  -charge beam particle  $s$  the source;  $p = 2ed(x-vt)$   $= \sqrt{p}$   $= \sqrt{2}ed(x-vt)$   $= \sqrt{2}e$   $= \sqrt{2}ed(x-vt)$   $= \sqrt{2}e$   $= \sqrt{2}ed(x-vt)$   $= \sqrt{2}$ 

(Why is  $\epsilon = \epsilon(\omega)$ ? Think of a single hormonically bound electron.  $\vec{X} + \vec{D} \cdot \vec{x} + \vec{W} \cdot \vec{X} = -\frac{e}{m} \vec{E}$  is its Equation of modon. The Fourier transform is algebraic of cambe  $\overline{X}(w) = \frac{-e_{/m} \overline{E}(w)}{(w_o^2 - w^2) + i \Gamma w}$ 

Now the polarization comes from svaning over electron displacements:  $\tilde{p} = -NZe \times \tilde{x}(\omega) = (e(\omega) - e_0) \tilde{e}(\omega)$ e/viit dipole moment/vnitvol linear dielectrice (assumption)

So E(w) = 60 + NZe2. Eo. (wis-w)-iw[

at least, in the hormonic model. It retkets resonance and absorption (17). Since the applied fields are the dependent and the electrons are bound, it makes sense that the response will have some frequency dependence beyond that of the applied fields.)

$$\vec{E} = -\vec{\nabla} \varphi - \vec{A}$$

$$\vec{B} = \vec{\nabla} \times \vec{A}$$

$$\Rightarrow \vec{E}(k,\omega) = -i\vec{k} \, \hat{\vec{\gamma}}(\vec{k},\omega) + i\omega \, \vec{A}(\vec{k},\omega)$$

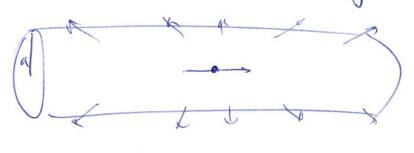
$$\vec{\vec{\beta}}(k,\omega) = + i\vec{k} \times \vec{A}(\vec{k},\omega)$$

and for the point charge,

$$\widetilde{E} = -i \, \widetilde{k} \, \widetilde{\varphi} + i \, \underline{w} \, \overline{v} \, \widetilde{\varphi}$$

$$\widetilde{B} = i \, \widetilde{k} \, \underline{v} \, \widetilde{v} \, \widetilde{\varphi}$$

Now let's relate these to the energy loss.



Integrably over all x at one time is equivalent to integrating over all time at 1 x since the particle 15 in constant relocity motion. So  $\left(\frac{dt}{dx}\right)_{b = a} = \frac{2\pi a}{\text{Mov}} \int_{-\infty}^{\infty} \overline{t_i} \, \overline{b_3} \, dt.$ Now convert to a frequency integral!  $\frac{-2\pi a}{m_0} \int_{\infty} dt \left( \int \frac{d^3h dw}{(2\pi)^4} e^{i(h_y a - wt)} \widetilde{\mathcal{E}}_i(\widetilde{h}, w) \right) \left( \int \frac{d^3h dw}{(2\pi)^4} e^{i(h_y a - wt)} \widetilde{\mathcal{B}}_3 \right)$ here we choose our fixed point to be (0, a, 0) for  $=\frac{-2\pi\alpha}{m\omega}\int\frac{d\omega}{2\pi\tau}\left(\int\frac{d^3k}{(2\pi)^3}e^{iky\alpha}\widetilde{E}_{,}\left(k,\omega\right)\right)\left(\int\frac{d^3k'}{(2\pi)^3}e^{ik'y\alpha}\widetilde{B}_{3}(k,-\omega)\right)$  $=\widetilde{\mathcal{B}}_{3}(-\omega)$ Shee I dw e int B fw) has to be real,

= I dw e int B3(-w) = I dw e int B3(w)

So B3(-W) = B \* (W)

Furthermore
$$\int_{-\infty}^{\infty} d\omega \, \widetilde{E}_{i}(\omega) \, \widetilde{B}_{3}^{*}(\omega) = \int_{\omega}^{\infty} \widetilde{E}_{i}(\omega) \, \widetilde{B}_{3}^{*}(\omega) + \int_{\omega}^{\infty} d\omega \, \widetilde{E}_{i}(\omega) \, \widetilde{B}_{3}^{*}(\omega) + \underbrace{\widetilde{E}_{i}(\omega)}_{i} \widetilde{B}_{3}^{*}(\omega) + \underbrace{\widetilde{E}_$$

So we need - 4th Re So dw E, Bs.

It's a bit of work to get E, (w) and Bs(w).

Jackson does it around 13, 30. Here's
the result! More Bessels!

$$\widetilde{E}_{1}(\omega) = \frac{-i2e\omega}{2\pi^{2}eV^{2}} \left(1 - V^{2}_{C^{2}}\right) K_{0}(\lambda a)$$

$$\widetilde{E}_{2}(\omega) = \frac{2e}{2\pi^{2}eV} \lambda K_{1}(\lambda a)$$

$$\widetilde{B}_{3}(\omega) = V \widetilde{E}_{2}$$

$$\lambda = \frac{\omega}{V} \sqrt{1 - V^{2}_{C^{2}}}$$

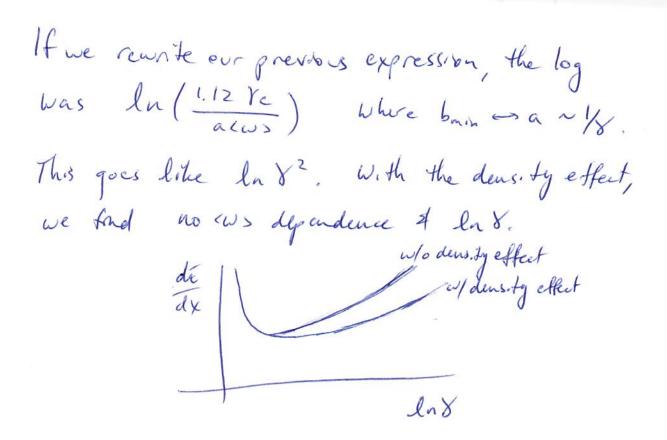
So  $\frac{dE}{dx}\Big|_{b>a} = -\frac{4\pi}{h_0} \operatorname{Re} \int_0^\infty \frac{d\omega}{2\pi} \left[ \left( \frac{-iZ^2 e^2 \omega}{k\pi^2 J^2 e^2 v^2 e^2} \right) \left( 1 - \frac{v^2 e^2}{L^2} \right) \left( \frac{1}{2} - \frac{v^2 e^2}{L^2} \right) \left( \frac{v^2 e^2}{L^2} \right) \left($ The integrand is imaginary if e is real and Be c1. Then the result vanishes. We need the complex (absorptive) part of the dielectric function. That turns out to correspond to optical frequencies, so that typically Hadeel when I is "very complex." Then we can expand the Besseli! Ko(x) s -ln(x/z) + ...

K, (x) = 1/x + ...

Then the integral can be done, using the harmonia model to- E. The result is (in the relativistic limit)

dk ) = (4TKo)2 4TINZe2(te)2 ln (1.12c a wp) W= 411NZe/m

The energy loss only depends on the electron density, not on atomiz details. Same to all meterals with Similar densities!



Experimentally, this screening effect is observed & curves rise like lux.