

# Approximation of Range in Materials as a Function of Incident Electron Energy

Gregory Wilson and JR Dennison

**Abstract**— A simple composite analytic expression has been developed to approximate the electron range in materials. The expression is applicable over more than six orders of magnitude in energy (<10 eV to >10 MeV) and range ( $10^{-9}$  m to  $10^{-2}$  m), with uncertainty of  $\leq 20\%$  for most conducting, semiconducting and insulating materials. This is accomplished by fitting data from two standard NIST databases [ESTAR for the higher energy range and the electron IMFP (inelastic mean free path) for the lower energies]. In turn, these data have been fit with well-established semi-empirical models for range and IMFP that are related to standard materials properties (e.g., density, atomic number, atomic weight, stoichiometry, band gap energy). A single free parameter, the effective number of valence electrons per atom  $N_v$ , is used to predict the range over the entire energy span.

**Index Terms**—range, inelastic mean free path, spacecraft charging

## I. INTRODUCTION

THE range,  $R$ , or maximum distance an electron of a given incident energy can penetrate through a material before all kinetic energy is lost and the electron comes to rest, is a common way to parameterize electron interactions with materials. The range is used in spacecraft charging calculations to predict the charge distribution of deposited electrons in materials and to model secondary and backscattered electron emission. It is also used to predict the distribution of energy deposited by incident electrons as they traverse a material; this distribution is further used to model radiation induced conductivity. It is therefore important for spacecraft charging models to have a realistic, reasonably accurate, and efficient expression to predict the approximate range of electron energies commonly encountered in space plasma fluxes, from  $\sim 10$  eV to  $\sim 10$  MeV. This expression needs to be readily implemented for a wide array of conducting, semiconducting and insulating spacecraft materials with a minimal number of fitting parameters.

Figure 1 offers dramatic visual evidence of the validity of the range of electrons in a material. The white line seen at the

center of the side view results from melting of the plastic target during discharge, as electrons deposited in a narrow distribution at a depth of  $R \approx 3$  mm by a monoenergetic  $\sim 1$  MeV electron beam are released.

## II. THEORY

The desired range expression can be developed by merging well known semi-empirical models for the interaction of electrons with materials in different energy regimes by employing the continuous-slowing-down approximation (CSDA). In the CSDA, the rate of energy loss (or total stopping power) at every position along the penetration path,  $\frac{dE}{dz}$ , is assumed constant; variations in energy-loss rate with energy,  $E$ , or penetration depth,  $z$ , are neglected. For a given incident energy,  $E_b$ , the CSDA range is obtained by integrating the reciprocal of the total stopping power with respect to energy over the full penetration depth such that  $E_b = \int_0^R \frac{dE}{dz} dz$  [1, 2].

For a constant energy-loss rate in the CSDA,  $\frac{dE}{dz} = E_b/R = E_m/\lambda_m = E_{mim}/\lambda_{min}$  is a constant. Here  $E_m$  is equal to mean energy lost per collision occurring at mean free path  $\lambda_m \equiv \lambda_{IMFP}(E_m)$ , and  $E_{mim}$  is the energy at the minimum in the inelastic mean free path curve at  $\lambda_{min} \equiv \lambda_{IMFP}(E_{min})$ . A reasonable approximation for  $E_m$  is the geometric mean of the plasmon energy and the bandgap energy,  $E_g$ , times an empirically determined factor of 2.8 [3]

$$E_m = 2.8 \left[ (E_p)^2 + (E_{gap})^2 \right]^{1/2}$$

where the plasmon energy is

$$E_p = \hbar(N_v q_e^2 / m_e \epsilon_0)^{1/2}$$

$N_v$  is the effective number of valence electrons per atom,  $\hbar$  is Planck's constant divided by  $2\pi$ ,  $\epsilon_0$  is the permittivity of free space, and  $q_e$  and  $m_e$  are the electron charge and rest mass, respectively.

Tabulated values of the electron ranges at high energies using the CSDA can be found in the NIST ESTAR database spanning incident energies from  $E_{HI} \sim 20$  keV up to  $\sim 1$  GeV [4]. The CSDA can also be applied to lower energy ranges. The NIST electron inelastic mean free path (IMFP) database [5] has tabulated values for the IMFP, which is closely related to the range as shown below, which are valid for energies from  $\sim 30$  eV to  $E_{LO} \sim 1$  keV. Thus, in order to create an

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Figure 1. Front (Left) and side (Right) views of a Lichtenberg discharge tree. The white line (Right) indicates the narrow distribution of deposited charge from a ~1 MeV electron beam at  $R \approx 3$  mm.

analytic expression for the full range of energies desired, the problem can be broken into three parts according to energy of the incident electron: a high energy range for  $E_b > E_{LO} \equiv 1$  keV; a mid-energy range for  $E_m < E_b < E_{LO}$ ; and a low energy range for energies  $E_b < E_m$ .

#### A. High Energy Range

A simple power law approximation for the range is used

$$R_{HE}(E) = bE^n \left\{ 1 + \frac{E/N_V}{m_e c^2} \right\}$$

where the term in brackets is a first order relativistic correction that becomes significant above  $\sim 10^5$  eV. Figure 2 shows the fit to tabulated data for Au from the ESTAR database, using both a simple power law and a power law with the first order relativistic correction. Figure 3 shows fits to the Au data using several range approximation formulas.

The stopping power exponent  $n$  is determined by requiring that the expressions for  $R_{HE}(E)$  matches the more accurate low energy (non-relativistic) Bethe-Joy range expression based on the Bethe stopping power formula [6] as extended by Tanuma [7] used in conjunction with the NIST ESTAR database [4] ,

$$R_{BJ}(E) = \frac{(4\pi\epsilon_0)^2 M_A}{2\pi q_e^2 Z_A \rho_m} \frac{E^2}{\ln \left( \sqrt{\frac{e}{2}} \left[ \frac{E/N_V}{E_m} + k \right] \right)}$$

at two energies  $E_{LO}$  and  $E_{HI}$ . Here  $M_A$  is the atomic weight,  $Z_A$  as the atomic number,  $\rho_m$  is the mass density, and  $k=0.8$  is a fixed empirical constant.  $E_{HI} \equiv 20$  keV is the lower energy at which data are available for all materials in the ESTAR database and  $E_{LO} \equiv 1$  keV is the upper energy at which data are

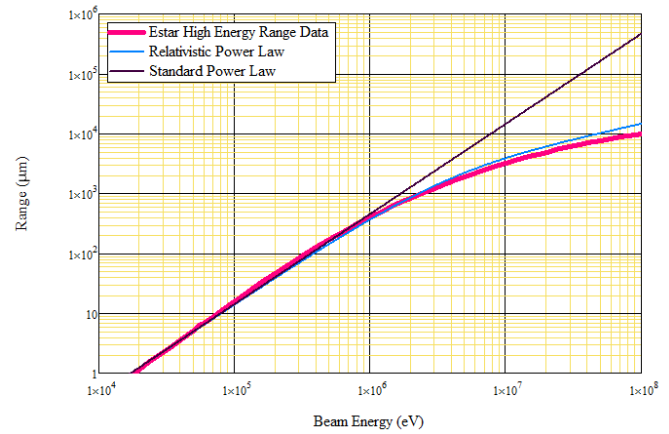


Figure 2. Comparison between the standard power law and the relativistic power law for Au. The relativistic power law allows approximations for energies up to 10 MeV with percent errors  $\sim 20\%$ .

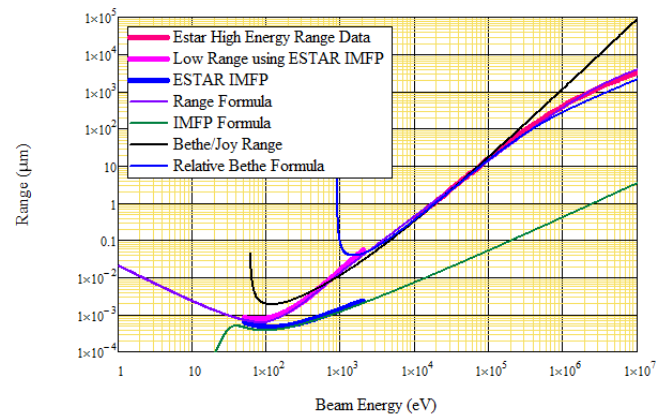


Figure 3. Comparison between several range approximations and the data from the ESTAR database for Au. The IMFP data for Au are also plotted along with the TPP-2M IMFP formula for  $\lambda_{IMFP}(E)$ .

available for all materials in the IMFP database. This leads to an expression for the stopping power exponent

$$n = \frac{\ln \left[ \frac{E_{LO} \ln \left\{ \sqrt{\frac{e}{2}} \left[ \frac{E_{HI}/N_V}{E_m} + k \right] \right\} (2m_e c^2 N_V + E_{HI}) (m_e c^2 N_V + E_{LO})^2}{E_{HI} \ln \left\{ \sqrt{\frac{e}{2}} \left[ \frac{E_{LO}/N_V}{E_m} + k \right] \right\} (2m_e c^2 N_V + E_{LO}) (m_e c^2 N_V + E_{HI})^2} \right]}{\ln[E_{LO}/E_{HI}]}$$

Note that the only free parameter in this expression is  $N_V$ , along with the electron mass and the fixed empirical constant  $k=0.8$ .

The high energy expression for  $R_{HE}(E)$  is normalized to the mid-energy expression at  $E_{LO}$ , as detailed below, by setting

$$b = \frac{E_{LO}^{1-n}}{E_m \left( 1 - \exp \left[ -\frac{E_{LO}}{E_m} \right] \right) \left\{ 1 - \left[ 1 + \left( \frac{E_{LO}/N_V}{E_m} \right)^2 \right]^{-1} \right\}}$$

Again, note that the only free parameter in this expression is  $N_V$ , along with  $E_m$  which is expressed above in terms of  $N_V$  and the band gap energy,  $E_{gap}$ .

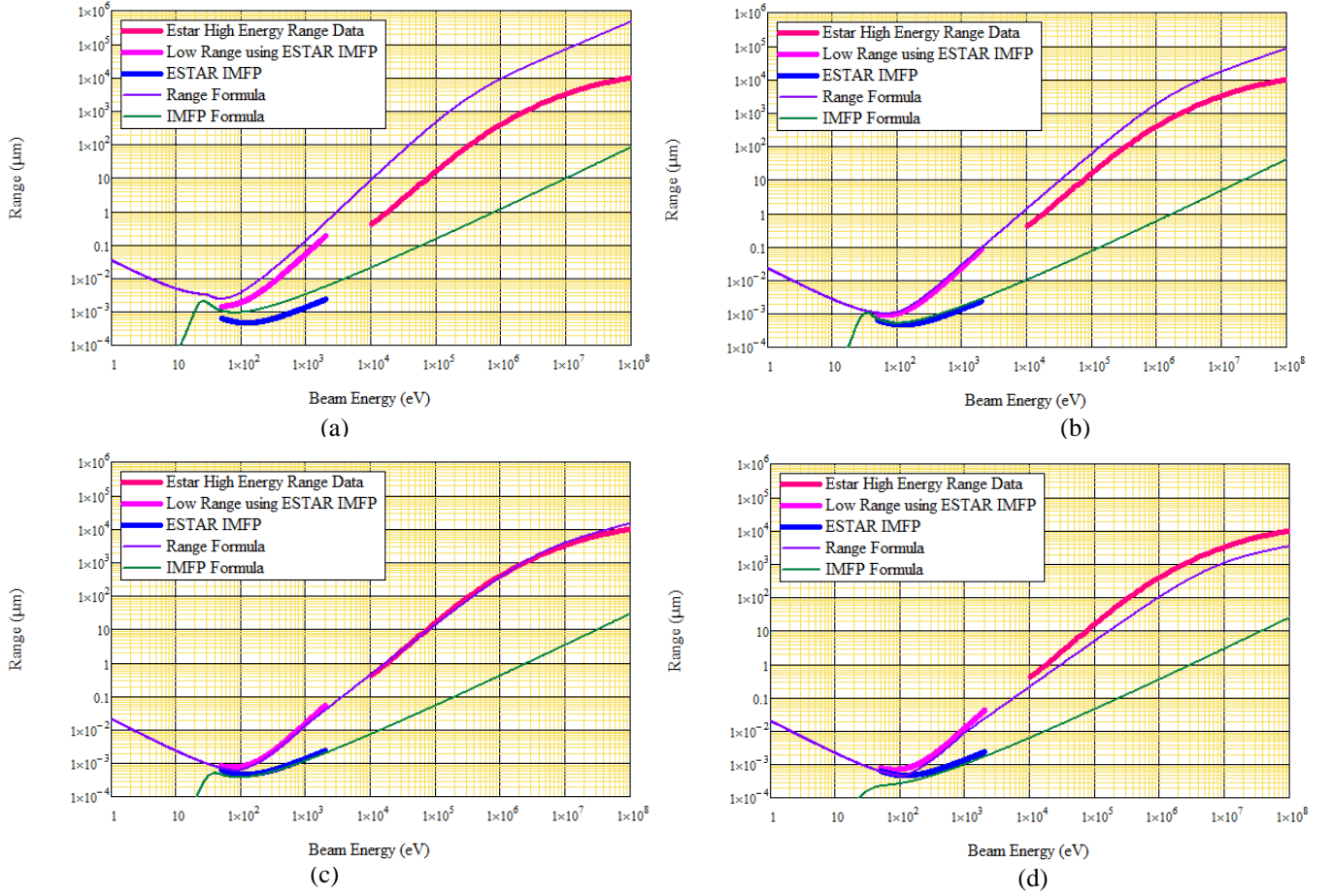


Figure 4. Graphs showing the variation of the range expression by changing the single fitting parameter  $N_v$ . For graphs (a) through (d),  $N_v = 1, 5, 12$  (best fit) and 20, respectively.

### B. Mid-Energy Range

Direct extrapolation of the range from the ESTAR data to lower energies is not valid for energies comparable to the atomic electronic structure, typically a few keV and below, because the discrete energy nature of the collisions becomes important. However, a simple extension of the CSDA to lower energies can relate the range to the electron IMFP, where

$$\frac{dE}{dz} = E_b/R(E_b) = \left[ \frac{E_m}{\lambda_{IMFP}(E_b)} \right] (1 - e^{-E_b/E_m})$$

Here the stopping power is again assumed equal to the total energy lost (incident energy,  $E_b$ ) divided by the total distance traveled (range,  $R$ ). This is set equal to the mean energy lost per collision,  $E_m$ , divided by the mean distance traveled per collision all times the probability that a collision occurs,  $(1 - e^{-R/\lambda_m}) = (1 - e^{-E_b/E_m})$ . Here, the energy dependence in the range is fully contained in the energy dependence of the mean free path. For  $E_b > E_m$ ,  $\lambda_{IMFP}(E_b)$  is assumed to be given by the TPP-2M formula used in conjunction with the NIST IMFP database [5]:

$$\lambda_{IMFP}(E) = EE_p^{-2} [\beta \ln(\gamma E) - CE^{-1} + DE^{-2}]^{-1}$$

where

$$\begin{aligned} \beta &\equiv -0.1 + 0.34E_m^{-1} + 0.069\rho_m^{0.1} \\ \gamma &\equiv 0.191\rho_m^{-\frac{1}{2}} \\ C &\equiv 1.97 - 0.91U \text{ with } U \text{ in eV} \\ D &\equiv 53.4 - 20.8U \\ U &\equiv \frac{N_v \rho_m}{M_A} \end{aligned}$$

Once again, by using the proposed equations of Tanuma contained in the TPP-2M model, the only free parameter in this expression is  $N_v$ , along with the materials constants  $E_{gap}$  (through  $E_m$ ) plus  $M_A$  and  $\rho_m$ . Because of the shallow core levels (generally with binding energies  $< 30$  eV) that may contribute significant intensity to the energy-loss function, there arises an ambiguity in the choice of the value of the number of valence electrons [8]. While  $E_{gap}$  may be considered an additional fitting parameter for insulators, its effect on  $R$  is minimal causing primarily a small vertical shift in the range curve.

### C. Low Energy Range

To calculate the range for  $E_b < E_m$ , we assume that the energy loss per collision of the low energy collisions is constant and equal to the mean excitation energy  $E_m$ , but that the probability that an electron undergoes one such inelastic collision falls off as  $R/\lambda_m (e^{-R/\lambda_m}) = E_b/E_m (e^{-E_b/E_m})$ , while for  $E_b < E_m$ , the IMFP is constant and equal to the IMFP

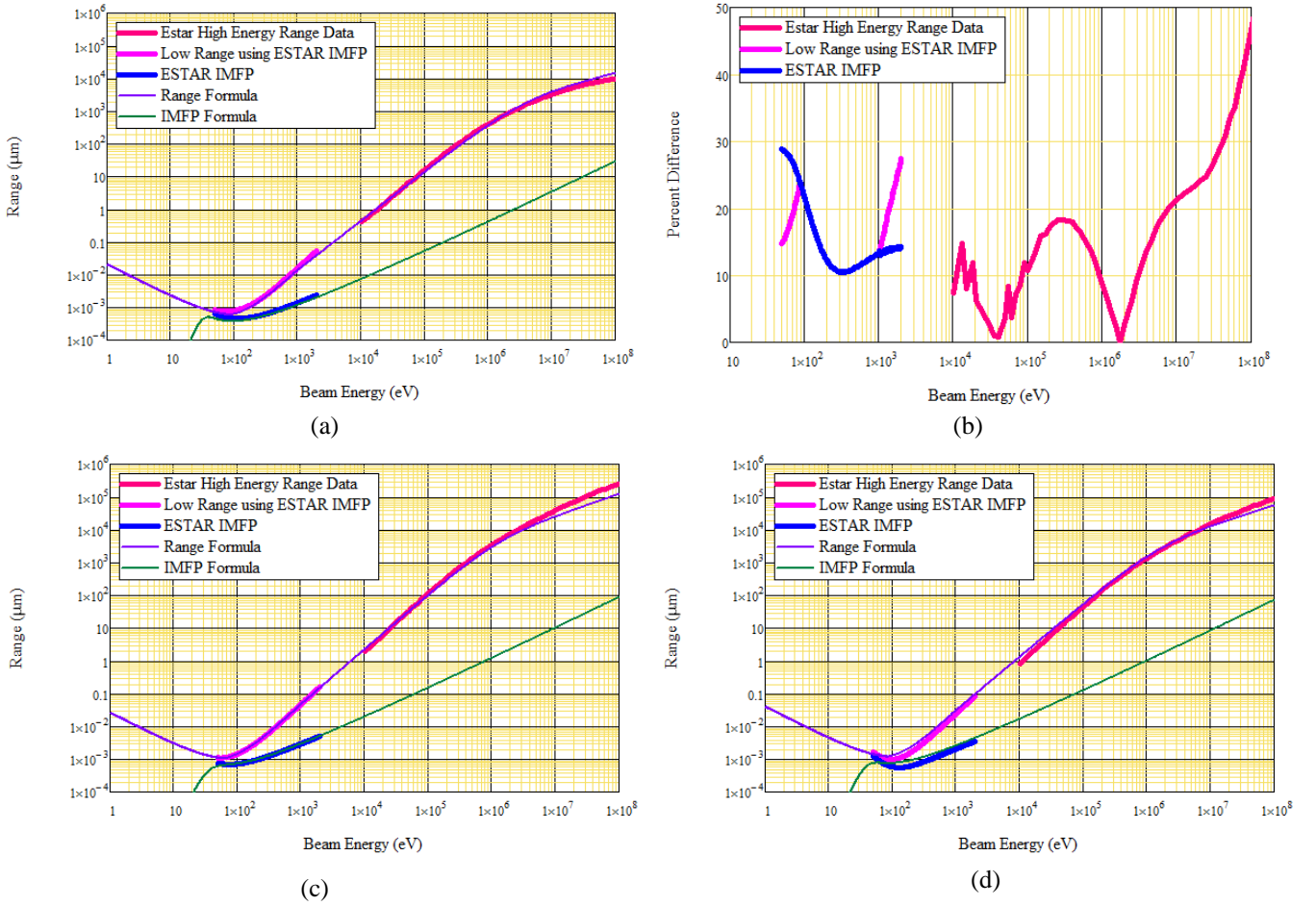


Fig. 5. (a) Comparison of the range formula for Au with  $N_V = 12.0$ . (b) Residual plot of Au range data in Fig 5. (c) Comparison of the range formula for Kapton with  $N_V=4.1$ . (d) Comparison of the range formula for  $Al_2O_3$  with  $N_V=5.0$ .

at the mean energy loss or  $\lambda_{IMFP}(E_m)=\lambda_m$ . This simple low energy approximation avoids the unusual asymptotic behavior exhibited by the TPP-2M expression at energies below  $E_m$  that is evident in the green curves in Fig. 4.

#### D. Composite Range Function

The final result is a continuous composite analytic approximation to the range with a single fitting parameter spanning from  $<10$  eV to  $>10$  MeV:

$$R(E; N_V) = \begin{cases} \left[ \frac{E}{E_m} \right] \frac{\lambda_{IMFP}(E_m)(1 - \exp[-1])}{\left(1 - \exp\left[-\frac{E}{E_m}\right]\right)^2} & ; E < E_m \\ \left[ \frac{E}{E_m} \right] \frac{\lambda_{IMFP}(E)}{1 - \exp\left[-\frac{E}{E_m}\right]} & ; E_m \leq E \leq E_{HI} \\ bE^n \left(1 - \left[1 + \frac{E/N_V}{m_e c^2}\right]^{-2}\right) & ; E > E_{HI} \end{cases}$$

Figures 5(a), 5(c) and 5(d) show fits to data for three prototypical materials: the conductor Au; the polymeric insulator polyimide (Kapton), and the insulating ceramic

$Al_2O_3$ . Table I lists the fitting parameter  $N_V$ , along with materials properties and derived values, for 14 typical spacecraft materials. The residual curve for the fit for Au is shown in Fig. 5(b).

### III. APPLICATIONS

The usefulness of an analytical approximation of the range to spacecraft applications can easily be demonstrated by considering expressions for the dose rate and the radiation induced conductivity; both expressions require an energy dependent range expression.

The dose rate is defined as the energy deposited by incident radiation per unit mass. The dose rate in the CSDA for a homogeneous material is proportional to the volume in which radiation energy is deposited, which is equal to the beam cross section times  $R$  [9]. Thus,

$$\dot{D} \equiv \frac{\partial D}{\partial t} = \frac{E_o J_{in}}{\rho_m R q_e} \propto R^{-1}$$

The dose rate for Au as a function of incident energy is shown in Fig. 6.

Table I. Materials Properties and Fitting Parameters

Material		Fitting Parameter $N_V$	Material Properties					Derived Values				
Name	Formula		$\rho_m$ (gm/cm <sup>3</sup> )	$Z_A$	$M_A$ (amu)	$E_{gap}$ (eV)	$n$	$b$ ( $\mu\text{m}/\text{eV}^{-n}$ )	$E_P$ (eV)	$E_m$ (eV)	$\lambda_{\min}$ (nm)	
Graphite	C	5.3	1.7	6	12.01	0.1	0.642	0.7143	24.87	69.6	0.793	
Amorphous C	C	4.0	2.0	6	12.01	0.1	0.676	0.3877	23.43	65.6	0.614	
Aluminum	Al	5.0	2.7	13	26.98	0.0	0.668	0.5075	20.31	56.9	0.467	
Silicon	Si	5.0	2.33	14	28.09	1.11	0.676	0.5422	18.49	51.9	0.438	
Copper	Cu	8.3	8.96	29	63.55	0.0	0.561	0.7821	31.06	87.0	0.422	
Germanium	Ge	9.8	5.32	32	72.64	0.66	0.571	1.355	24.32	68.1	0.477	
Silver	Ag	10.6	10.5	47	107.87	0.0	0.536	1.217	29.17	81.7	0.416	
Gold	Au	12.0	19.32	79	196.97	0.0	0.508	1.261	31.15	87.2	0.371	
Polyethylene	[C <sub>2</sub> H <sub>4</sub> ] <sub>n</sub>	2.5	0.94	2.65	4.64	2.9	0.727	0.2354	20.43	57.8	0.642	
Polyimide	[C <sub>22</sub> H <sub>10</sub> N <sub>2</sub> O <sub>5</sub> ] <sub>n</sub>	4.1	1.42	5.01	9.769	2.3	0.678	0.4582	22.165	62.4	0.652	
PTFE	[C <sub>2</sub> F <sub>4</sub> ] <sub>n</sub>	6.0	2.2	8.01	16.023	6	0.620	0.8794	26.06	78.9	0.865	
Aluminum Oxide	Al <sub>2</sub> O <sub>3</sub>	5.0	3.97	10	30.392	9.9	0.628	0.5188	28.33	84.0	0.746	
Silicon Dioxide	SiO <sub>2</sub>	5.0	2.32	9.98	19.99	8.9	0.653	0.6215	21.87	66.1	0.711	
Glass, Pyrex	doped SiO <sub>2</sub>	6.2	2.32	9.98	19.99	4	0.626	0.8150	24.36	69.1	0.656	

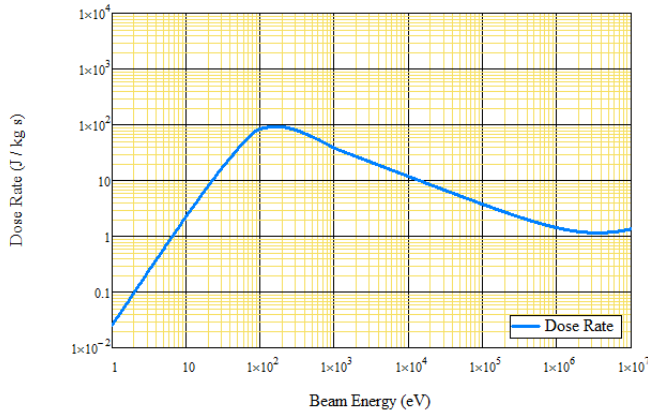


Figure 6. Dose rate as a function of energy in the CSDA for Au.

Radiation Induced Conductivity (RIC) is the enhanced conductivity that results from the energy deposited in this volume. In the CSDA

$$\sigma_{ric}(\dot{D}) = k_{ric} \dot{D}^\Delta \propto R^{-\Delta}$$

with  $1/2 < \Delta < 1$  [10]. Figure 7 shows the RIC for Kapton as a function of incident energy for three values of  $\Delta$ . Notice that both  $\dot{D}$  and  $\sigma_{RIC}$  exhibit energy dependent maxima as a consequence of the minimum in the range expression.

Secondary electron (SE) emission is another extension of the range which would be highly beneficial. In the CSDA, the SE yield can be expressed as

$$\delta_{SE}(E, N_V, \delta_{max}, E_{max}) = \frac{\delta_{max}}{1 - \exp[-R(E)/\lambda_{SE}]} \left\{ 1 - \exp\left[-\frac{R(E)}{\lambda_{SE}} \left(\frac{E}{E_{max}}\right)^\eta\right] \right\}$$

Work is underway to develop an expression for the SE yield in terms of the range expression developed here. The expression would have three independent free parameters;  $N_V$  and the

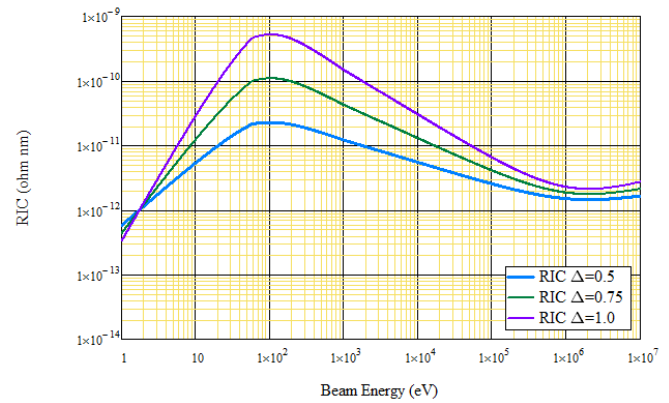


Figure 7. RIC as a function of energy in the CSDA for polyimide.

maximum SE yield  $\delta_{max}$  at energy  $E_{max}$ .

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