

where α is the molecular polarizability. The quadrupole energy is $V_q = -\frac{1}{2}eQ_{\text{mole}}r^{-3}(1-3\cos^2\theta)$, where Q_{mole} is the quadrupole moment of the molecule relative to the molecular axis and θ is the angle between the molecular axis and the line joining the center of the molecule with the ion. When the measured^{3,4} values of α and Q_{mole} for N_2 are inserted into the above expressions, it is seen that $|V_p| > |V_q|$ for the whole range of separations at which $|V_p|$ is of the order of kT . For pure polarization scattering the momentum transfer cross section averaged over all thermal energies is approximately equal to the cross section at the kinetic energy $(64/9\pi)kT$. Comparing the two interactions at the separation $r=3.8\text{A}$, $|V_p| = (64/9\pi)kT = 0.059\text{ ev}$ ($T=300^\circ\text{K}$) while $|V_q(\theta=\pi/2)| = 0.018\text{ ev}$ and $|V_q(\theta=0)| = 0.035\text{ ev}$.

An exact calculation of the momentum transfer cross section resulting from the interaction V_p+V_q is intractable because the potential is not spherically symmetric and translational kinetic energy is not conserved. However, in the case of a pure polarization interaction the Langevin formula⁵ gives the mobility of ions (single positive charge, atomic mass M) in nitrogen as $2.11(1+28/M)^{1/2}\text{ cm}^2/\text{volt sec}$. The inclusion in the Langevin potential model of a repulsive force at small separations would cause an increase in this calculated mobility by at most 15 percent, depending on the effective radius of the repulsion. This agrees reasonably well with the empirical result² of $2.04(1+28/M)^{1/2}\text{ cm}^2/\text{volt sec}$ and indicates that the quadrupole interaction is not of primary importance in determining the ionic mobility at room temperature.

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Electrical Conduction in Halide-Contaminated Ice

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THE anomalous electrical properties of halide-contaminated ice have been the subject of several papers by the authors.¹⁻⁴ Such ice, frozen according to the methods prescribed, exhibits unusual features: (1) it acts as a rectifier; (2) it possesses an extremely high dielectric constant for fields applied in the non-conductivity direction; and (3) the conductive and dielectric properties depend in a complicated way upon temperature, bias, and frequency. The general behavior with respect to the properties enumerated is such as to suggest electronic conduction. This interpretation was given further support by the fact that the passage of an electric current for long time intervals produced no marked change in the conductive behavior of the crystal. It appeared reasonable, therefore, to formulate an explanation based on the premise that the partially shielded proton is displaced under the stress of an applied electric field in such a manner as to increase considerably the ionic contribution to the hydrogen bonding between neighboring molecules. In spite of the relative ease of proton displacement considered essential, the proton appeared to be bound because of the high-energy barrier postulated for proton transfer (26 kcal/mole or even more⁵). Moreover, there is persistence of a sense of direction to the c axis under all conditions of applied field—a condition considered incompatible with the process of molecular rotation inherent in the transfer mechanism.

An experimental re-examination of the conduction process has revealed that the conduction of electricity in the forward direction through the contaminated ice results in the liberation of oxygen

and hydrogen at the positive and negative electrodes respectively. The amount of gas liberated is in quantitative agreement with that associated with the total charge transfer. Contrary to our earlier interpretations it now appears certain that the conduction process is ionic—by a proton transfer mechanism.

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Polarization of Nucleons Elastically Scattered from Nuclei*

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SEVERAL recent experiments¹⁻³ have shown that protons of energy 200-350 Mev scattered from nuclei are polarized. The elastically scattered protons have a polarization (~ 60 percent) that is somewhat larger than the inelastically scattered ones. Fermi⁴ has proposed an explanation of the polarization for elastic scattering in terms of a nuclear spin-orbit interaction potential similar to that assumed in the nuclear shell model. He used the Born approximation in his estimates. The purpose of this note is to investigate the polarization effects of a nuclear spin-orbit potential using the transparent nuclear model of Serber.⁵ That is, we add to the nuclear complex potential which is constant over a sphere of radius R , a term

$$-\hbar^{-1}U(r)\mathbf{L}\cdot\boldsymbol{\sigma}, \quad (1)$$

where \mathbf{L} and $\hbar\boldsymbol{\sigma}/2$ are the orbital and spin angular momenta of the proton. U is taken to be real since the absorption cross section of protons in nuclear matter is independent of its spin. The radial and energy dependence of U are unknown, as well as its variation with atomic number. We have assumed two specific forms for U :

$$U_1(r) = u_1 R \delta(r-R), \quad U_2(r) \cong u_2 (R/r)^2. \quad (2)$$

$U_1(r)$ is the one considered by Fermi and is suggested by the Thomas precession of a particle with spin under acceleration, which, in the transparent nuclear model, is concentrated at the boundary of the potential hole. By contrast, $U_2(r)$ is concentrated near the origin. [The exact form of U_2 , which was chosen for convenience in numerical computation, is implied in Eq. (8).]

The calculations are performed by a partial-wave analysis. The degree of polarization (as defined by Oxley *et al.*¹) is

$$P(\theta) = (|f-ig|^2 - |f+ig|^2) / (|f-ig|^2 + |f+ig|^2), \quad (3)$$

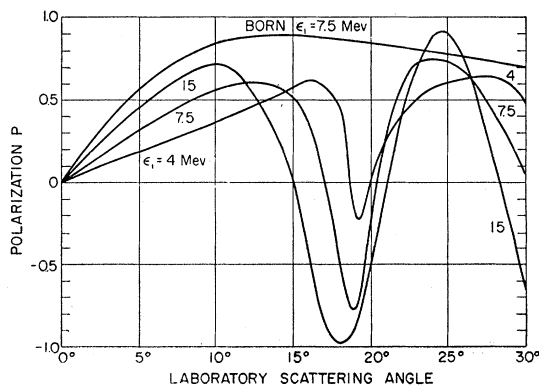


FIG. 1. Polarization P as a function of scattering angle θ for various ϵ_1 calculated from potential U_1 [Eq. (7)] for 316-Mev protons scattered by Be. The top curve gives the Born approximation values for $\epsilon_1 = 7.5$ Mev.