For the Lamb shift $(\delta E_{2S_2} - \delta E_{2P_2})$ for He⁺, Eq. (1) gives approximately 16 900 Mc/sec, and Eq. (3) gives about 13 880 Mc/sec, while the experimental value for this quantity as found by Lamb and Skinner³ is around 14 020 Mc/sec. This shows the correctness of the quantum-electrodynamical formula (3) in the approximation used and the nonvalidity of Eq. (1).

Some time ago, I proposed a phenomenologic theory of the Lamb shift,² in which self-interactions were completely "forbidden," and in which the Lamb shift was due to some assumed direct interaction between electron and proton and between the spins and the electromagnetic field. This theory led to Eq. (1) with Eq. (2) substituted. It is hard to see how this theory could ever be made to yield the additional term with $-2(\ln Z)\delta_{l,0}$ by any simple modification of the interactions proposed. Therefore the experimental evidence on He⁺ rules out the explanation of the Lamb shift by such direct interactions, and thus yields new evidence for the reality of the finite part of the quantum-electrodynamical self-interaction of the electron.

As regards the attempts to postulate the nonexistence of selfinteractions, this evidence shows that such a postulate would have to be modified in such a way as to leave at least such part of the self-interaction as corresponds to the finite part of the self-energy leading to the Lamb shift, thus removing the necessity of ad hoc assumed direct interactions. Other restrictions to be imposed on this postulate of nonexistence of part of the selfinteractions have been discussed by me earlier.⁴

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The Nuclear Magnetic Moment of Cr⁵³ and Sr⁸⁷

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ROM the hyperfine structure of the paramagnetic resonance absorption of a magnetically dilute single crystal, Bleany and Bowers¹ have observed the spin of Cr^{53} to be $\frac{3}{2}$ and the magnetic moment to be $|\mu| = (0.5 \pm 0.1)$ nm, the sign being undetermined. Using a nuclear induction spectrometer, we have observed the nuclear magnetic resonance of Cr⁵³ at a frequency of 2.40 Mc/sec in a field of 10 000 gauss. The nuclear induction signals have been observed in a 1.1-molar aqueous solution of Na₂CrO₄ in which the Cr has been isotopically enriched to 90 percent Cr⁵³. The signals have also been observed, with reduced amplitude, in a saturated solution of natural abundance Na₂CrO₄. The polarity of the Cr⁵³ nuclear induction signal is opposite to that of Na²³, indicating that Cr53 has a negative magnetic moment. In the same magnetic field the ratio of the nuclear resonance frequency of Cr⁵³ to that of D² in a sample of D₂O containing 1 molar of MnCl₂ has been observed to be $\nu(Cr^{53})/\nu(D^2) = 0.36820 \pm 0.00003$. From this we find for the magnetic moment of Cr53, without diamagnetic correction, $\mu(Cr^{53}) = -(0.47351 \pm 0.00007)$ nm, where we have used in this calculation the spin $\frac{3}{2}$ of Cr⁵³ as observed by Bleany and Bowers,¹ the ratio $\mu(D^2)/\mu(H) = 0.307015$ as given by Mack,² and $\mu(H) = 2.7925$ nm as determined by Bloch and Jeffries.³

From optical hyperfine structure measurements Heyden and Kopfermann⁴ have determined the spin of Sr^{87} to be 9/2 and the magnetic moment to be $\mu = -1.1$ nm. Using a saturated aqueous solution of SrBr₂, we have observed a nuclear induction signal at a frequency of 1.84 Mc/sec in a field of 10 000 gauss. We have enhanced the signal amplitude and also verified that this is indeed the magnetic resonance of Sr⁸⁷ by using a 3.1-molar aqueous solution of SrBr₂ in which the Sr had been enriched to 60-percent Sr⁸⁷. The polarity of the Sr⁸⁷ signal was observed to be opposite to that of Br⁷⁹, verifying that the magnetic moment of Sr⁸⁷ is negative. We have detected no "chemical shift"5 between the resonance frequencies of Sr⁸⁷ in SrBr₂ and SrI₂. Using the enriched sample,

we have observed that, in the same magnetic field, the ratio of the nuclear resonance frequency of Sr⁸⁷ to that of D² in D₂O containing 1-molar MnCl₂ is $\nu(\hat{Sr}^{87})/\nu(\hat{D}^2) = 0.28232 \pm 0.00003$. From this we find for the magnetic moment of Sr⁸⁷, without diamagnetic correction, $\nu(Sr^{87}) = -(1.0892 \pm 0.00015)$ nm, where we have used a spin of 9/2 for Sr⁸⁷ as measured by Heyden and Kopfermann⁴ and the values of $\mu(D^2)/\mu(H)$ and $\mu(H)$ as quoted above.

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Lower Limit for the Lifetime of the 665-kev Excited State of Mo⁹⁷†

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CCORDING to Goldhaber and Hill¹ the 665-kev gamma ray A emitted in the decay of Nb⁹⁷ is probably a magnetic dipole transition between states with orbitals $g_{7/2}$ and $d_{5/2}$. As the Nb⁹⁷ decay involves a rather energetic beta ray which can compensate for the gamma-ray recoil, Nb⁹⁷ seemed to be a favorable case for a lifetime measurement using resonance scattering.²

A measurement of the nuclear resonance scattering of a gamma ray is a determination of the width of the initial nuclear level involved in the transition. The resonance scattering technique is especially well suited for lifetimes of the order of 10⁻¹³ second and shorter (widths $\geq 6 \times 10^{-3}$ ev). This is just the region into which, according to Weisskopf's lifetime formula,3 the magnetic dipole transitions of $\approx mc^2$ energy should fall. The only M1 transition in this energy range whose lifetime is known is the 478-kev transition in Li7. From an observation of the Doppler broadening due to the motion of the Li^{7*} nucleus in the $B(n, \alpha)$ reaction, Elliott and Bell⁴ deduced a lifetime of 7×10^{-14} second, which is even somewhat shorter than expected from Weisskopf's formula. On the other hand, the lifetimes of several low-energy M1 transitions have been measured by Graham and Bell⁵ using delayed coincidence techniques. Most of these lifetimes were found to be \sim 100 times longer than predicted from Weisskopf's formula. If the 665-kev Mo⁹⁷ gamma ray belonged to this "slow" group, resonance scattering would not be observable with our present means; if, however, the 665-kev gamma ray followed Weisskopf's formula, a large nuclear resonance scattering effect could be expected.

Ten milligrams of ZrO₂ enriched⁶ in Zr⁹⁶ were bombarded in the Brookhaven pile and yielded a 0.3-millicurie source of Zr⁹⁷ and its daughter Nb97. The gamma rays from this source were scattered alternately from Mo and Zr scatterers. The scattered radiation was observed with a scintillation spectrometer which accepted only pulses corresponding to the 665-kev photoelectron line. As the difference in atomic number between Zr and Mo is small, Rayleigh scattering from the two scatterers was almost the same. The small difference in Rayleigh scattering was determined in a separate experiment using the 663-kev gamma ray from Cs¹³⁷.

After correcting for the difference in Rayleigh scattering the counting rates for the Nb⁹⁷ source with the Mo and Zr scatterers were identical, the experimental uncertainty being two percent.

If one assumes the contribution of the resonance scattering from the Mo scatterer to be two percent of the counting rate (i.e., equal to the uncertainty), and if one takes into account that only a small percentage of the gamma rays is emitted while the nucleus still has the momentum imparted to it by the beta ray,