

Higher-Precision Determination of the Fine-Structure Interval in the Ground State of Positronium, and the Fine-Structure Density Shift in Nitrogen*

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The fine-structure interval $\Delta\nu$ in the ground state of positronium has been remeasured with higher precision to obtain $\Delta\nu_{\text{expt}} = 203.396 \pm 0.005$ GHz (24 ppm). The current theoretical value is $\Delta\nu_{\text{theor}} = 203.4155$ GHz. The difference $\Delta\nu_{\text{expt}} - \Delta\nu_{\text{theor}} = -19 \pm 5$ MHz is about 4 standard deviations of the experimental error, indicating the need for calculations of higher-order radiative contributions. Values are given for both the linear and quadratic coefficients of the fine-structure density shift in N_2 .

The importance of a highly precise measurement of the fine-structure interval $\Delta\nu$ in the ground state of positronium for a test of quantum electrodynamics, and in particular of the Bethe-Salpeter equation for the bound-state lepton-antilepton system, is well known.^{1,2} The present Letter reports a value for $\Delta\nu$ with a substantially smaller error than the best previously quoted value.³ Significant comparison of this new precise value for $\Delta\nu$ with theory clearly requires the calculation of higher-order radiative contributions to $\Delta\nu$.⁴

The method of the present experiment is the measurement of the Zeeman transition frequency between the $M=0$ and $M=\pm 1$ magnetic sublevels of orthopositronium by a magnetic resonance technique, and is basically similar to that used in previous measurements.^{3,5,6} The transition frequency f_{01} in a magnetic field H is given by

$$f_{01} = \frac{1}{2} \Delta\nu [(1+x^2)^{1/2} - 1] \quad (1)$$

in which $x = 2\mu_B g' H / h \Delta\nu$. The quantity g' is given by

$$g' = g \left(1 - \frac{5}{24} \alpha^2\right) = g(1 - 11.1 \times 10^{-6}), \quad (2)$$

where g is the gyromagnetic ratio of the free electron (or minus that of the free positron),⁷ $g = 2(1.001\,159\,656\,7 \pm 0.000\,000\,003\,5)$.

The factor involving α^2 is a relativistic bound-state contribution.⁸⁻¹⁰ It was obtained using a Hamiltonian for positronium in an external magnetic field consisting of the Pauli Hamiltonian,¹¹ the Breit interaction,¹² and the virtual annihilation interaction.¹³ A transformation developed by Chraplyvy¹⁴ for two-body equations was used to decouple the large and small components. This approach is an extension of the Foldy-Wouthysen transformation¹⁵ for the Dirac equation. The α^2 term arises from the relativistic change of mass with velocity and from the spin-orbit and spin-

other-orbit interactions, but does not have a contribution from the virtual annihilation interaction. The α^2 factor was also derived¹⁶ from a two-particle Breit equation by the method of Perl and Hughes.¹⁷ Finally this factor can be obtained by the general derivation of Grotch and Hegstrom.¹⁸

Important improvements in apparatus and in experimental technique have enabled us to obtain the substantial improvement in accuracy. A schematic diagram of the experimental apparatus is shown in Fig. 1. A new electromagnet¹⁹ with a pole diameter of 38 cm and a pole gap of 10 cm was used to produce a field of about 7800 G homogeneous to ± 4 ppm over a volume of 7 cm³. The field was regulated to ± 0.5 ppm by an NMR stabilization system. The microwave system at 2.3 GHz was essentially that previously used,³ and the positron source was 2.5 mCi of Na^{22} . The stopping gas was ultrahigh purity N_2 , which allowed the use of sufficiently high input microwave

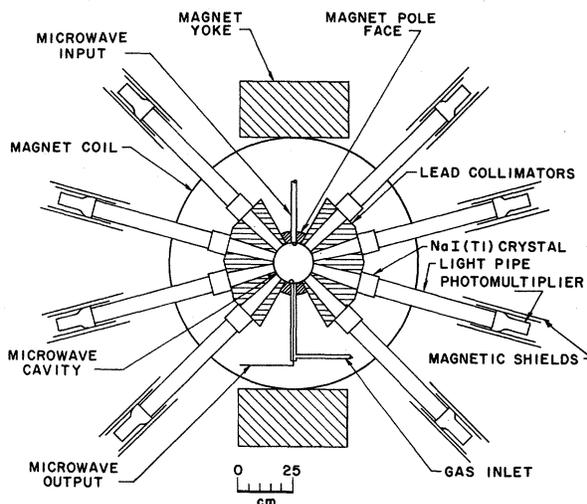


FIG. 1. Schematic diagram of experimental apparatus.

power at the lower pressures without electrical discharge. Eight NaI (Tl) γ -ray detectors were used to provide four coincidence detectors for two 0.51-MeV annihilation γ rays with a 0.2- μ sec time resolution and an overall increase in detection solid angle by a factor of 9 compared to an earlier experiment.³ The active volume for positronium observation of 7 cm³ is about twice that of the previous experiment. Data were collected with a multichannel pulse-height analyzer.

A resonance curve was obtained by varying the magnetic field H with fixed microwave frequency and power. Signal heights were 3% to 11% and linewidths (full width at half-maximum) were 30 to 47 G. A nonresonant magnetic-field-dependent background, which arises from the magnetic focusing of positrons and from the increase in the $2\gamma/3\gamma$ branching ratio with increase in field and which amounts to about 5% of the resonance signal, was subtracted from the raw data by using a representation of the background linear in H over the resonance region. The remaining signal was fitted by the theoretical line shape as described for a previous experiment.³ 54 resonance curves were obtained for N₂ pressures from 0.14 to 3.40 atm.

For each resonance curve a resonance microwave frequency f_{01} and central magnetic field H_0 is determined by the fitting procedure. Then with the use of Eq. (1) a value for $\Delta\nu$ is calculated. A plot of the values of $\Delta\nu$ versus N₂ density is shown in Fig. 2. The solid circles are the weighted means of all values of $\Delta\nu$ obtained at a given density, and the associated error bars are 1-standard-deviation errors, due predominantly to statistical counting errors. The data are fitted by the function

$$\Delta\nu(D) = \Delta\nu(0)(1 + aD + bD^2) \quad (3)$$

in which D is the gas density, and a and b are the linear and quadratic coefficients, respectively, of the fine-structure density shift. The results are given in Table I and indicate that not only a linear but also a quadratic density shift is pre-

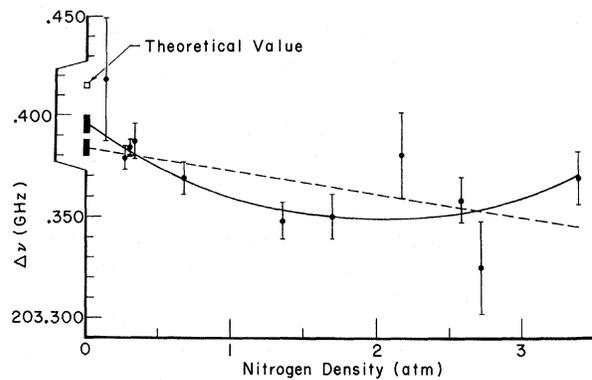


FIG. 2. Plot of $\Delta\nu$ versus N₂ gas density in units of atmospheres at 23°C. The solid and dashed lines are the quadratic and linear fits using Eq. (3), respectively. The solid bars show 1-standard-deviation errors for the fitted values $\Delta\nu(0)$.

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The principal result of our experiment is a value for $\Delta\nu$ of free positronium based on the fit of Eq. (3) to our data:

$$\Delta\nu_{\text{expt}} = 203.396 \pm 0.005 \text{ GHz (25 ppm)}, \quad (4)$$

in which the error is mainly due to a statistical counting error.²⁰ This result agrees with, and has 1/2.5 times the quoted error of, the best previously reported result³:

$$\Delta\nu_{\text{expt}} = 203.403 \pm 0.012 \text{ GHz (60 ppm)}.$$

The current theoretical value is⁴

$$\begin{aligned} \Delta\nu_{\text{theor}} &= \frac{1}{2} \alpha^2 c R_{\infty} \left[\frac{7}{3} - \frac{\alpha}{\pi} \left(\frac{32}{9} + 2 \ln 2 \right) - \frac{3}{2} \alpha^2 \ln \alpha \right] \\ &= 203.4155 \pm 0.0006 \text{ GHz (3.1 ppm)} \quad (5) \end{aligned}$$

using²¹

$$\alpha^{-1} = 137.036\,02 \pm 0.000\,21 \text{ (1.5 ppm)},$$

$$R_{\infty} = (1.097\,373\,1 \pm 0.000\,000\,1) \times 10^5 \text{ cm}^{-1} \text{ (0.1 ppm)},$$

TABLE I. Results of fits of the data $\Delta\nu(D)$ by Eq. (3) for N₂ gas.

	Linear fit	Quadratic fit
$\Delta\nu(0)$ (GHz)	203.384 ± 0.004	203.396 ± 0.005
a (10^{-5} atm^{-1} at 23°C)	-5.7 ± 1.4	-23 ± 6
b (10^{-5} atm^{-2} at 23°C)	0	$+5.7 \pm 1.8$
χ^2 per degree of freedom	17.1/9	6.8/8
$\Delta\nu_{\text{expt}} - \Delta\nu_{\text{theor}}$ (MHz)	-31 ± 4	-19 ± 5

and²²

$$c = (2.997\,924\,62 \pm 0.000\,000\,18) \times 10^{10} \text{ cm/sec} \\ (0.06 \text{ ppm}).$$

The indicated uncertainty in $\Delta\nu_{\text{theor}}$ is due to the uncertainty in the constants used in evaluating $\Delta\nu_{\text{theor}}$ from Eq. (5). The next uncalculated radiative correction term is of order $\alpha^4 R_\infty$ which is about 50 ppm.

The difference between the experimental and theoretical values for $\Delta\nu$ is

$$\Delta\nu_{\text{expt}} - \Delta\nu_{\text{theor}} = -0.019 \pm 0.005 \text{ GHz}, \quad (6)$$

where the uncertainty is the 1-standard-deviation experimental error. This difference is about 4 standard deviations of the experimental error. Clearly a more significant comparison of experiment and theory requires the calculation of the $\alpha^4 R_\infty$ radiative correction term.

The experimental values for the linear and quadratic coefficients of the fine-structure density shift are obtained from the quadratic fit of Table I. We note the large difference between the values of a determined with or without the allowance for a quadratic term. In earlier work with Ar gas, not enough data were obtained to test for the quadratic term.³ No quantitative theory has yet been given for the positronium fine-structure pressure shifts.

We believe that our present apparatus and technique, when supplemented with improved data acquisition equipment and a higher-intensity Na²² source, are capable of achieving an accuracy of 10 ppm in the determination of $\Delta\nu$, which would correspond to a choice of the resonance line center of $\frac{1}{1000}$ of its full width at half-maximum.

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