First Measurement of the Velocity of Slow Antihydrogen Atoms

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The speed of antihydrogen atoms is deduced from the fraction that passes through an oscillating electric field without ionizing. The weakly bound atoms used for this first demonstration travel about 20 times more rapidly than the average thermal speed of the antiprotons from which they form, if these are in thermal equilibrium with their 4.2 K container. The method should be applicable to much more deeply bound states, which may well be moving more slowly, and should aid the quest to lower the speed of the atoms as required if they are to be trapped for precise spectroscopy.

When the goal of producing “cold” antihydrogen (H̄) was laid out long ago [1], the objective was H̄ atoms that were cold enough to be confined in a neutral particle trap for precise spectroscopy and gravitation studies. This stringent definition of “cold” requires H̄ energies significantly below the 0.5 K depth of superconducting magnetic traps, when these are placed in the ~ 1 Tesla bias field needed to confine the antiprotons (̄p) and positrons (e⁺) for H̄ production.

Antihydrogen produced during the positron cooling of antiprotons [2] in a nested Penning trap [3] was called “cold” antihydrogen in reports of its observation [4–6]. However, no H̄ energy, velocity, or temperature was actually measured. The observed atoms were clearly cold compared to H̄ moving at nearly the speed of light [7,8]. Almost certainly the H̄ energy was less than the tens of eV well depths of the potential wells used to confine the ̄p and e⁺ from which the H̄ were formed. It was naturally hoped that the H̄ were in thermal equilibrium with the 4.2 K [5,6] or 15 K [4] temperature of the electrodes confining the ̄p and e⁺.

In this Letter we report the first measurement of the velocity of H̄ atoms. The change in H̄ transmission efficiency through an oscillating electric field is measured as a function of the field’s oscillation frequency. ATRAP’s background-free field ionization detection method [5] registers only H̄ that reach the detection well intact. Atoms moving slowly enough will never make it through the electric field without being ionized. Faster atoms are sometimes able to pass while the oscillating field is too weak to ionize them, depending upon the phase of the field. In this first demonstration we deduce that the most weakly bound H̄ that we detect have an energy that is about 200 meV, a speed that is about 20 times higher than an average thermal speed at 4.2 K. More deeply bound H̄ observed to survive a 360 V/cm electric field may move more slowly; this method should make it possible to check, though the measurements will take much more time than has been available so far. No attempt has yet been made to minimize the ̄p driving forces that bring ̄p and e⁺ into repeated contact [6].

The ATRAP apparatus is represented in Fig. 1, with a cross section of the crucial volume in Fig. 2(a). In preparation for H̄ production, typically 2 × 10⁵ ̄p from CERN’s Antiproton Decelerator are accumulated for this demonstration. The well-established techniques for slowing, trapping, cooling, and stacking [9,10] are now used in all H̄ experiments. The ̄p cool by collisions to equilibrium with trapped, 4.2 K electrons. (The electrons...
No potentials or fields are varied along this trajectory. The magnetic field axis to the normalization well \([\text{Fig. 2(b)}]\).

Atoms traveling in the \(\hat{z}\) direction will be similarly counted in the detection well \([\text{Fig. 2(d)}]\) if they are not ionized by the electric field they pass through on their way to this well. An atom will ionize if the magnitude of the axial electric field it encounters along its trajectory exceeds a value \(F\) which is directly related to the size of its internal orbit \([12]\). The field along an \(\bar{\text{H}}\) trajectory always has a static component \(F_{\text{dc}}(\rho, z)\), and we can add a time varying axial field \(F_{\text{ac}}(\rho, z) \cos (\omega t + \phi)\).

While we can only measure the number of \(\bar{\text{H}}\) atoms that travel into the very small solid angle of our detection well, a simple example gives clear evidence that driven \(\bar{\text{H}}\) production sends \(\bar{\text{H}}\) preferentially directed along the magnetic field axis, \(\pm \hat{z}\). From \(4.5 \times 10^6 e^+\) and \(2.9 \times 10^5 \bar{\text{p}}\) we detect \(7.6 \times 10^3 \bar{\text{H}}\) in the detection and normalization wells. Since the combined solid angle of the detection and normalization wells is less than \(4\pi/100\), the \(\bar{\text{H}}\) production cannot be isotropic since several times more \(\bar{\text{p}}\) would be required than the number available.

A measurement of the \(\bar{\text{H}}\) velocity is very simple in principle. With the oscillating field turned off, the ratio of \(\bar{\text{H}}\) atoms ionized in the detection and normalization wells is obtained (open square in Fig. 3). This point has no frequency associated with it, but is plotted on the same vertical scale as the other points on the graph. These come from measurements with the oscillating field turned on, for different values of oscillation frequency \(\omega/(2\pi)\) (solid points in Fig. 3). The fraction of \(\bar{\text{H}}\) atoms detected decreases as \(\omega\) increases. As \(\omega\) increases, fewer atoms travel quickly enough through the oscillating field to avoid ionization, and fewer atoms are thus detected. The \(\bar{\text{H}}\) atoms are produced under the same conditions for each measurement, and the count from the detection well is normalized as discussed above. Electric field gradients can change the speed of a highly excited and polarizable \(\bar{\text{H}}\) atom, but this is a small affect here since the preionizing field is relatively spatially uniform \([12]\).

The vertical scale for Fig. 3 is chosen so that the measurement with no oscillating field is consistent with 1, and so that the measured \(\omega \rightarrow 0\) limit of the oscillating field measurements is consistent with 0.62. The latter is the fraction of the time that the magnitude of the oscillating electric field, along the trajectory of an \(\bar{\text{H}}\) traveling to the detection well, is less than the maximum static field along this path. In this limit 0.62 of all the \(\bar{\text{H}}\) traveling to the detection well should thus not be ionized by the oscillating field.

To interpret our measurements quantitatively we consider the \(N(\rho, v, F)d\rho dv dF\) antihydrogen atoms produced at \(z = 0\) and \(t = 0\), at a radius between \(\rho\) and \(\rho + d\rho\), with an axial velocity between \(v\) and \(v + dv\), and which cool via synchrotron radiation to thermal equilibrium with the 4.2 K trap electrodes. The \(\bar{\text{p}}\) are then positioned in the side wells of a nested Penning trap \([\text{Fig. 2(b)}]\) by manipulating the potentials applied to the vertical stack of ring electrodes. A magnetic field, \(B\hat{z}\) with \(B = 5.2\) T, is parallel to trap axis. Typically \(4 \times 10^5 e^+\) are accumulated at the same time as the \(\bar{\text{p}}\) \([11]\). They are then positioned in the center of the nested Penning trap \([\text{Fig. 2(b)}]\), between the side wells for antiprotons. Like the electrons they cool to 4.2 K by radiating synchrotron radiation.

\(\bar{\text{H}}\) atoms are produced near the center of the nested Penning trap, at the location of electrode T7 as indicated by “\(\bar{\text{H}}\) ” in Fig. 2(a). \(\bar{\text{H}}\) atoms are produced when the \(\bar{\text{p}}\) are driven from one side well \([\text{Fig. 2(b)}]\), by a radiofrequency driving force, through the \(e^+\) \([6]\). The \(\bar{\text{p}}\) that do not form \(\bar{\text{H}}\) get more chances when they are driven back and forth from one well to the other. The drives give \(\bar{\text{p}}\) velocities along the axis of the trap, either in the direction of the normalization well or in the direction of the detection trap. (Likely such weakly driven \(\bar{\text{H}}\) production can produce slower atoms than those produced by injecting higher energy \(\bar{\text{p}}\) into a nested Penning trap for cooling \([4,5]\).) If an \(\bar{\text{H}}\) is formed before the \(e^+\) completely cool the \(\bar{\text{p}}\), then the \(\bar{\text{H}}\) will have a residual axial velocity that is larger than the average thermal velocity of the \(\bar{\text{p}}\) at the 4.2 K temperature of the \(e^+\) awaiting \(\bar{\text{H}}\) production.

Atoms traveling in the \(-\hat{z}\) direction travel along the magnetic field axis to the normalization well \([\text{Fig. 2(b)}]\). No potentials or fields are varied along this trajectory. The \(\bar{\text{H}}\) ionized by the strong electric fields in this well \([\text{Figs. 2(c) and 2(d)}]\) leave their \(\bar{\text{p}}\) in the well for back-
are in a state that will be ionized by an axial electric field between \( F \) and \( F + dF \).

When no time varying electric field is applied, the \( \bar{H} \) atoms on their way to the detection well experience an axial electric field \( F_{dc}(\rho, z = vt) \). Figure 2 shows \( F_{dc}(\rho, z) \) for \( \rho = 0 \) (c) and \( \rho = 3 \) mm (d), along with the corresponding values \( F_i(\rho = 0) = 34 \text{ V/cm} \) and \( F_i(\rho = 3 \text{ mm}) = 45 \text{ V/cm} \). The number of detected \( \bar{H} \) is thus

\[
N_0 = \int_0^{\rho_{\text{max}}} d\rho \int_0^{\infty} dv \int_{F_i(\rho)}^{F_{\text{max}}(\rho)} dF \mathcal{N}(\rho, v, F). \tag{1}
\]

Detected atoms are those that ionize between \( F_i(\rho) \) and \( F_{\text{max}}(\rho) \), the maximum values of \( F_{dc}(\rho, z) \) that they encounter before and within the detection well, respectively.

When the alternating axial electric field, \( F_{ac}(\rho, z) \times \cos(\omega t + \phi) \), is applied the \( \bar{H} \) atoms traveling towards the detection well encounter a net axial field

\[
F(\rho, z, \omega/v, \phi) = F_{dc}(\rho, z) + F_{ac}(\rho, z) \cos\left(\frac{\omega z}{v} + \phi\right), \tag{2}
\]

where we must later average over all phases \( \phi \). The number of detected \( \bar{H} \) is

\[
N = \int_0^{\rho_{\text{max}}} d\rho \int_0^{\infty} dv \int_{F_i(\rho, \omega/v, \phi)}^{F_{\text{max}}(\rho, \omega/v, \phi)} dF \mathcal{N}(\rho, v, F). \tag{3}
\]

The limit \( F_{\text{max}}(\rho, \omega/v, \phi) \) is the maximum axial field magnitude (i.e., the maximum \( |F(\rho, z, \omega/v, \phi)| \)) that an \( \bar{H} \) sees at any \( z \) before the detection well begins.

We make the simplest assumption—that the \( \bar{H} \) are produced uniformly out to a radius \( \rho_{\text{max}} = 3 \text{ mm} \)—approximately the measured extent of our \( e^+ \) plasma [13]. (Later we show that we are not very sensitive to this assumption.) We also assume the measured power law dependence \( N \sim F^{-3} \) [14]. Finally we assume that the \( \bar{H} \) have a velocity \( v_0 \hat{z} \), since we expect that driven \( \bar{H} \) production produces atoms traveling along the magnetic field direction. We would expect the same result for any velocity spread that is narrow compared to the average velocity. Thus

\[
N(\rho, v, F) \sim \frac{2\pi \rho}{\pi \rho_{\text{max}}^3} \delta(v - v_0) F^{-3}. \tag{4}
\]

The solid curves in Fig. 3 show the fraction of \( \bar{H} \) atoms that should be detected when the oscillating electric field is turned on, for various values of the \( \bar{H} \) kinetic energy \( 1/2 M v_0^2 \), where \( M \) is the \( \bar{H} \) mass.

An \( \bar{H} \) kinetic energy of 200 meV is a good fit to our measurements (2400 K in temperature units). This conclusion is rather insensitive to our assumptions. For example, if we had assumed that all the \( \bar{H} \) was produced on the central axis, then we would have concluded that the \( \bar{H} \) velocity was about 100 meV. If we had instead assumed that all the \( \bar{H} \) was produced 4 mm off axis, then we would have concluded that the \( \bar{H} \) velocity was 300 meV. A 200 meV \( \bar{H} \) velocity is much higher than the 0.3 meV average energy for a 4.2 K thermal distribution—the lowest possible average \( \bar{p} \) and \( \bar{H} \) energy. Even the 1 meV curve in Fig. 3 is far from consistent with our data.

An electric field gradient \( \partial F_z/\partial z \) exerts a force on a highly polarizable \( \bar{H} \)[12]

\[
f_z = \frac{e^2 \rho^3}{2r_e m_e c^2} \frac{\partial [F_z(z)^2]}{\partial z}. \tag{5}
\]

Here \( e \) is the proton charge, \( \rho \) is the radial size of the \( \bar{H} \) atom, \( r_e \) is the classical electron radius, and \( m_e c^2 \) is the electron rest energy. For fields nearly strong enough to ionize the atom the force increases beyond this value. We use the overestimates \( F_z = 100 \text{ V/cm} \) and a gradient \( \partial F_z/\partial z = 100 \text{ V/cm}^2 \) applied over 1 cm, together with the size \( \rho = 0.3 \mu m \) of the largest atom that survives \( F_{dc} = 40 \text{ V/cm} \). The resulting force would change the velocity of a 200 meV \( \bar{H} \) by less than a percent, justifying the neglect of the gradient force on a polarized \( \bar{H} \) in this work. For lower \( \bar{H} \) speeds more care must be taken.

For a \( \bar{p} \) traveling in the \( \hat{z} \) direction through the \( e^+ \) plasma, one \( \bar{p} \) speed that seems important is the one that equals the average axial speed of the \( e^+ \) that are going in the same direction as the \( \bar{p} \). For 4.2 K \( e^+ \) this corresponds to a \( \bar{p} \) energy of 210 meV. This is close to what we measure, likely by coincidence given the approximate character of the estimate. One might expect increased \( \bar{H} \) production at this \( \bar{p} \) energy, but this depends in a complicated way upon how quickly the \( \bar{p} \) are being cooled by the \( e^+ \). A recombination rate that depends upon the relative velocity of the \( \bar{p} \) and \( e^+ \) will become insensitive to \( \bar{p} \) energies below this value, since the relative velocity will be determined by the \( e^+ \) velocity.
Another important speed is

\[ v_p = n_e (\pi b^2) v_e L. \]  

(6)

This speed would allow just enough time in the \( L = 1 \text{ mm} \) thick plasma for there to be a deexcitation collision [15] between the \( e^+ \) initially picked up by the \( \pi \) and another \( e^+ \) in the plasma on average. The expected \( e^+ - e^+ \) collision rate should be of order \( n_e (\pi b^2) v_e \) where \( n_e = 1.6 \times 10^9 \text{ cm}^{-3} \) is the \( e^+ \) density and \( v_e \) is the average thermal speed for a \( e^+ \) at \( T_e = 4.2 \text{ K} \). The distance of closest approach \( b \) comes from equating the potential energy \( (4 \pi e_n)^{-1} e^2 / b \) and the thermal energy \( kT_e \). The corresponding \( \pi \) energy, and hence the \( \pi \) energy, is \( 460 \text{ meV} \), larger than we observe. The cross section used is only an estimate, of course, and the \( e^+ \) may be heated somewhat by the \( \pi \)’s. Notice that \( v_p \propto n_e L T_e^{-3/2} \), suggesting that a \( e^+ \) plasma with a lower density, a shorter length and a higher temperature will produce \( \pi \) with lower velocities.

The relatively high velocities that we observe may also be related to the fact that this first demonstration experiment measures only the speed of the most weakly bound \( \pi \) states, which may have had less cooling time. Because the number of \( \pi \) atoms detected goes down steeply as the strength of the ionization field \( F \) is increased, as \( F^{-3} \) [14], the \( \pi \) atoms that we detect are essentially all atoms which will ionize just above the maximum static electric field \( [F \pi (\rho)] \) in the region before the detection well.

More deeply bound \( \pi \) states may well have much lower velocities if these arise because of more collisions of the \( \pi \) with \( e^+ \) in the plasma. It should be possible to use this new technique to measure the velocity of more deeply bound states. The experiment is simple in principle. The deeply bound states can be selected by raising the maximum value \( F \pi (\rho) \) of the static electric field \( F_{dc} (\rho, z) \). The challenge in practice is that the number of deeply bound states observed goes down inversely as the cube of this field value, greatly increasing the time required to accumulate data.

In conclusion, a new method makes it possible to measure the speed of slow \( \pi \) atoms for the first time. For the most weakly bound states, our measurements fit well to a 200 meV \( \pi \) energy. This is close to the estimated 210 meV \( \pi \) energy below which the \( \pi \) formation rate no longer increases. It is lower than the 460 meV energy that is estimated to allow a deexcitation collision between the \( e^+ \) in an \( \pi \) and a \( e^+ \) in the surrounding plasma on average. It is higher than the lowest possible energy, the 0.3 meV average energy of a 4.2 K thermal distribution of \( \pi \).

The new method should reveal how the \( \pi \) velocity depends upon the strength of the driving force, the number and density of the \( e^+ \) and \( \pi \), and upon the binding energy or size of the \( \pi \) atoms. Slowly lowering the depth of the center well in the nested Penning trap may produce slower \( \pi \), as may interchanging the position of the \( e^+ \) and \( \pi \) in this trap. The first step towards devising ways to get the coldest possible \( \pi \) atoms, of course, is a method to measure the \( \pi \) speed, which is now available.

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