Recombination of H$_3^+$ and D$_3^+$ Ions with Electrons.

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Abstract. Flowing-afterglow measurements in decaying H$_3^+$ or D$_3^+$ plasmas suggest that de-ionization does not occur by simple binary recombination of a single ion species. We find that vibrational excitation of the ions fails to provide an explanation for the effect, contrary to an earlier suggestion. Instead, we suggest that collisional stabilization of H$_3^{**}$ Rydberg molecules by ambient electrons introduces an additional dependence on electron density. The proposed mechanism would permit plasma de-ionization to occur without the need for dissociative recombination by the mechanism of potential-surface crossings.

INTRODUCTION

It is usually taken for granted that typical plasma afterglow experiments yield the binary dissociative recombination coefficient for molecular ions and that, in principle, the measured values would be the same as those inferred from beam experiments. The assumption seems plausible since the capture of an electron into a repulsive state of the neutral is thought to induce rapid dissociation$^1$, leaving little time for third-body interactions. The argument, however, presupposes the existence of suitable dissociation channels. If such do not exist, then perhaps long-lived Rydberg molecules can be formed and third-body interactions may be more important. The experiments described here suggest that this situation may exist in plasmas containing H$_3^+$ and D$_3^+$ ions.

Theoretical arguments$^{2,3}$ indicate that the recombination

\[ H_3^+ + e^- \rightarrow H+H+H \]  
\[ \rightarrow H_2+H \]  

for vibrationally cold H$_3^+$ (v=0) ions by direct dissociative recombination should be slow due to a lack of suitable crossings between the ionic and neutral potential surfaces. However, vibrationally excited H$_3^+$ ions in v>3 are predicted to exhibit faster recombination. This expectation disagreed with the early microwave-afterglow experiments$^4,5$ which gave fairly "normal" recombination coefficients ($\sim 2\times 10^{-7}$ cm$^3$/s) for H$_3^+$ ions believed to be in v=0. This "H$_3^+$ dilemma" has existed for about 20 years.

The structure and energy levels of H$_3^+$ and H$_3^*$ are quite well known. A brief introduction and extensive references can be found in the "thumbnail sketch
of H₃⁺⁺ by Oka and Jagod ⁶. The importance of H₃⁺ in astrophysical environments is discussed in reviews by Dalgarno ⁷ and by Tennyson et al. ⁸. A fairly complete account of the history of dissociative recombination work has been given by Bates et al. ⁹.

One possible resolution of the H₃⁺ dilemma was offered by Adams et al. ¹⁰. They interpreted their observation of an anomalous decay of a flowing afterglow plasma by assuming the presence of two species of H₃⁺ ions, one of which recombined fairly fast while the other recombined slowly (α<2x10⁻⁸ cm³/s). By identifying the fast and slowly recombining ions as, respectively, v>3 or v=0 ions, theory and experiment could be reconciled. This solution, while being in conflict with the microwave afterglow results, seemed quite plausible until Amano ¹¹, ¹² showed that spectroscopically identified H₃⁺ ions in v=0 recombined readily with electrons (α~ 1.8 x 10⁻⁷ cm³/s). A repetition of the flowing-afterglow measurements by Canosa et al. ¹³, ¹⁴ gave no evidence for the existence of two distinct types of H₃⁺ ions.

Recombination cross sections for H₃⁺ have been measured repeatedly in the merged-beam apparatus at the University of Western Ontario ¹⁵, ¹⁶, ¹⁷, ¹⁸. The results showed that the cross section varied with conditions in the ion source and this was ascribed to vibrational excitation. In later work ¹⁶, the H₃⁺ vibrational state was deduced from the threshold energy for electron-ion dissociative excitation. The result indicated that v=0 ions recombined only slowly with electrons.

When ion storage rings became available, they were very quickly applied to H₃⁺ recombination. In these instruments, vibrationally excited ions have time to relax by radiative transitions (including the v₁=1 metastable level of H₃⁺ with a lifetime ¹⁹ of 1.2 s). Larsson et al. ²⁰ and Sundström et al. ²¹, using the CRYRING storage ring, obtained recombination cross sections for H₃⁺ (v=0) over the wide energy range from 0.0025 to 30 eV. The analysis of the experimental cross sections yielded a recombination coefficient of (1.15 ± 0.13) x10⁻⁷ cm³/s at an electron temperature of 300 K.

Since the storage-ring measurements also yielded fairly large recombination coefficients for v=0 ions, the problem of finding a recombination mechanism for v=0 ions became more acute. New mechanisms were proposed ²², ²³ that do not require crossings between ionic and neutral potential curves. It is possible that such mechanisms may play a part in the recombination of H₃⁺ ions, but no detailed calculations have been made.

Not all experimentalists were willing to accept a sizable recombination coefficient for H₃⁺ in v=0. Smith and Spanel ²⁴ repeated and refined the afterglow experiment of Adams et al.'s ¹⁰ and concluded that only H₃⁺ ions in v>3 recombined efficiently with electrons. Again, those authors found that the decay of H₃⁺ plasmas was anomalous in the sense that the recombination coefficient appeared to become smaller in the late afterglow. It was, in part, this work that motivated us to carry out an additional experiment. As will be seen, we largely reproduced Smith and Spanel's ²⁴ results that there is something peculiar about the decay of H₃⁺ plasmas but our interpretation of the experiment is quite different.
To avoid ambiguities, we will use the term "recombination coefficient" ($\alpha$) only for purely binary recombination. We will use the word "de-ionization coefficient" ($\beta$) to denote the quantity that is derived from plasma decay experiments. The two quantities may be equal, but this is not assumed from the onset.

A more complete description of the experiment, additional details of the analysis, and the interpretation will be given in a journal article.

**EXPERIMENTAL METHODS**

Our flow tube is a stainless-steel tube of diameter 6 cm and length 36 cm. The carrier gas was helium (at pressures near 1 Torr and temperatures near 295 K). Flow velocities ranged from 5000 cm/s to 6000 cm/s. Reagent gases are added to the flowing plasma through several movable and fixed inlets. A movable Langmuir probe serves to measure electron densities $n_e$ as a function of axial position $z$. A quadrupole ion mass spectrometer analyzes the ion composition at the downstream end of the flow tube.

**Plasma generation and ion conversion**

The techniques of recombination measurements in flow tubes are quite well known and will not be discussed here. The method consists of creating a well-characterized plasma that contains only the desired ion species and measuring the rate at which free electrons disappear. The creation of the $H_3^+$ plasma involves several steps:

1. The carrier gas (helium) is partially excited to the metastable state and (to a lesser degree) ionized by passing it through a microwave discharge.
2. Argon (-15 % of helium) is added downstream to convert metastable helium atoms to $Ar^+$ ions by Penning ionization. After this, the plasma contains mainly $Ar^+$ ions with a small admixture of $He^+$ and $He_2^+$ ions (< 0.1 % of $Ar^+$). The electron density at this point is typically $4 \times 10^{10}$ cm$^{-3}$.
3. To create $H_3^+$ ions, $H_2$ is added at densities from $10^{14}$ to $10^{15}$ cm$^{-3}$. The formation of $H_3^+$ via the reactions
   
   \[ Ar^+ + H_2 \rightarrow ArH^+ + H \quad (+1.53\text{eV}) \quad (2) \]
   \[ ArH^+ + H_2 \rightarrow Ar + H_3^+ \quad (+0.57\text{eV}) \quad (3) \]
   
   (see Radzig and Smirnov for thermochemical data) takes place very rapidly compared to recombination since the rate coefficients for reactions (2) and (3) are large ($\sim 10^{-9}$ cm$^3$/s).

**Vibrational excitation and quenching of the $H_3^+$ ions**

If the entire energy of 2.1 eV that is released in reactions (2) and (3) were to appear as vibrational excitation, which seems unlikely, then $H_3^+$ could be produced in vibrational states up to $v=5$. However, $H_3^+$ ions in $v>1$ would be destroyed rapidly by proton transfer to Ar,

\[ H_3^+ + Ar \rightarrow ArH^+ + H_2 \quad (4) \]
which is energetically possible for \( \text{H}_3^+ \) ions with internal energies above 0.57 eV. The non-reacting, surviving \( \text{H}_3^+ \) vibrational states are the ground state, the \( \nu_2=1 \) bending-mode vibration at 0.3126 eV, and the \( \nu_1=1 \) breathing mode vibration at 0.394 eV (see Lie and Frye \(^{29}\) or Oka and Jagod \(^6\)). At \([\text{Ar}]= 5 \times 10^{15} \text{ cm}^{-3}, \nu \geq 1 \) ions would be destroyed by proton transfer \((k \sim 10^{-9} \text{ cm}^3/\text{s})\) in less than 1 μsec, a time that is far shorter than the time scale of recombination (about 1 ms).

The radiative lifetimes of the \( \nu_1=1 \) and \( \nu_2=1 \) levels \((1.2 \text{ s} \text{ and } 4 \text{ ms}, \text{ respectively})\) \(^{19}\) are too long for radiative relaxation to be important. Quenching of \( \nu=1 \) ions in collisions with \( \text{H}_2 \) is more effective. Amano's \(^{30}\) absorption studies of the \( \nu_2 \) hot band indicate that the \( \nu_2=1 \) level is quenched by \( \text{H}_2 \) with a rate coefficient of approximately \( 3 \times 10^{-10} \text{ cm}^3/\text{s} \). This eliminates \( \nu_2=1 \) ions; they will be destroyed in 30 μs at \([\text{H}_2]= 1 \times 10^{14} \text{ cm}^{-3} \). On the other hand, there is evidence \(^{31,32}\) for slower quenching \((k \sim 10^{-12} \text{ cm}^3/\text{s})\) of other vibrational states. It seems very likely that the slowly-quenched ions are those in the \( \nu_1=1 \) level. A similar conclusion was reached by Bawendi et al. \(^{33}\) who studied the absorption of \( \text{H}_3^+ \) hot bands in \( \text{He}/\text{H}_2 \) discharges.

The foregoing considerations indicate that only \( \text{H}_3^+ (\nu_1=1) \) ions live long enough to undergo recombination. Smith and Spanel \(^{24}\) did not consider the destruction of the higher vibrational levels by proton transfer with argon and consequently believed that most \( \text{H}_3^+ \) ions would be in states \( \nu \geq 3 \). We do not agree with this conclusion; others have expressed the same criticism earlier \(^{13,14}\). This is an important point, since Smith and Spanel's proposed reconciliation of theory and experiment rests on the assumption that vibrationally excited ions dominate.

There is no \textit{a priori} reason to expect that \textit{para} and \textit{ortho} \( \text{H}_3^+ \) ions should recombine equally fast. According to Oka \(^{34}\), \textit{ortho}-\textit{para} equilibration by proton exchange with \( \text{H}_2 \) is rapid. We assume that the proton-transfer rate is sufficiently large \((k \sim 10^{-10} \text{ cm}^3/\text{s})\) would suffice at \([\text{H}_2]= 10^{14} \text{ cm}^{-3} \) to keep the para-to-ortho ratio at a constant value in the recombining plasma.

### DATA ANALYSIS

The most common method of deducing \( \beta \) from the slope of a graph of the reciprocal of the electron density \( n_e(z) \) vs. axial position \( z \) \((V \text{ being } \text{the plasma velocity})\),

\[
\frac{1}{n_e(z)} = \frac{1}{n_e(z=0)} + \frac{\beta z}{V}
\]

provides quick, but only only approximate values of a \( z \)-averaged de-ionization coefficient. Since we are looking for deviations from a simple decay law, we were forced to employ a more rigorous approach (for details see ref. 25): It consisted of solving numerically the continuity equation for the electron density \( n_e(r, z) \) in a flowing plasma with a parabolic velocity field. The computed solutions were then used to validate a computationally faster, approximate method that could be used for routine data analysis. The approximate method extracts a \( z \)-dependent de-ionization coefficient \( \beta(z) \) from the measured electron density \( n_e(z) \). This form of
EXPERIMENTAL RESULTS

40 decay curves $n_e(z)$ were recorded for $[Ar] = 4 \times 10^{15} \text{cm}^{-3}$ to $8 \times 10^{15} \text{cm}^{-3}$, and $[H_2] = 1 \times 10^{14} \text{cm}^{-3}$ to $1.7 \times 10^{15} \text{cm}^{-3}$. Even the simplest form of data analysis (based on Eq. (9)) showed that $\beta$ increased noticeably with increasing $H_2$ concentration (see Fig. 1). A similar, but less scattered graph was obtained when the average values of $\beta(z)$ in the first 4 cm were plotted as a function of $[H_2]$.

Figures 2 and 3 show examples of the electron density $n_e(z)$ and $\beta(z)$ for two different $H_2$ concentrations. It is apparent that $\beta$ is not constant but that it decreases with increasing $z$. This finding reproduces the observations of Smith and Spanel 24; the decay of the plasma clearly does not follow a simple recombination law.

A smaller set of data was taken for $D_3^+$ ions. The observations largely mirrored those for $H_3^+$. Again, the recombination coefficient increased with increasing $D_2$ concentration, but the $D_3^+$ de-ionization coefficients were significantly smaller (from $0.7 \times 10^{-7} \text{cm}^3/\text{s}$ to $1.3 \times 10^{-7} \text{cm}^3/\text{s}$ in the range $[D_2] = 1$ to $14 \times 10^{14} \text{cm}^{-3}$).

INTERPRETATION OF RESULTS

The interpretation should explain the enhancement of $\beta$ upon hydrogen addition, the decline of $\beta(z)$ with increasing position $z$, and if possible, it should elucidate the recombination mechanism. We attempted to explain the peculiar decay of the plasma by two different models. The first model assumes the presence of $H_3^+$ ions in two different states. The second model invokes an entirely different explanation in terms of Rydberg molecules.

Two - state model

The assumption that the $z$ dependence of $\beta$ is caused by the presence of $H_3^+$ ions in $v=0$ and in $v_1=1$ leads to

$$\beta(z) = \alpha_0 f_0(z) + \alpha_1 f_1(z)$$

where $\alpha_0$ and $\alpha_1$ denote the recombination coefficients of $v=0$ and $v_1=1$ ions, and $f_0$ and $f_1$ are their fractional concentrations. Quenching of $v_1=1$ ions to $v=0$ by $H_2$ (rate coefficient $k_q$) and differing recombination losses make the concentrations $z$-dependent. It is not difficult to construct a kinetic model that yields a fit to the measured $\beta(z)$ curve at $[H_2] = 1.3 \times 10^{14} \text{cm}^{-3}$. Figure (2) shows fits for $\alpha_0 = 2.2 \times 10^{-7} \text{cm}^3/\text{s}$, $\alpha_1 = 4 \times 10^{-8} \text{cm}^3/\text{s}$, $f_0 = 0.7$, and two quenching coefficients of $k_q = 1 \times 10^{-12} \text{cm}^3/\text{s}$ and $k_q = 1 \times 10^{-13} \text{cm}^3/\text{s}$. However, when the same values are
used to compute a fit to the data at \([H_2]= 1.7 \times 10^{15} \text{ cm}^{-3}\). The model (see Fig. 3) indicates that \(\beta(z)\) should now increase with \(z\). A decline of \(\beta(z)\) can be forced by reducing the quenching rate to \(k_q=1\times10^{-13} \text{ cm}^3/\text{s}\), but the model fails to reproduce the observation that \(\beta\) at \(z=0\) (only 1 cm from the reagent inlet) is larger at the larger \(H_2\) concentration. We eventually concluded that the two-state model does not provide a consistent explanation of the experimental findings.

**De-ionization by Rydberg molecule stabilization**

We now return to the subject that was mentioned in the Introduction, namely de-ionization mechanisms which involve interactions with third bodies. The \(\beta(z)\) data suggest that the de-ionization coefficient is larger at higher electron densities and higher \(H_2\) densities. Is it possible that these particles affect the mechanism of de-ionization directly, rather than changing the state of the recombing ions? At first sight, this does not seem to be a viable proposition. Electron-stabilized and neutral-stabilized recombination are known to be too slow (by several orders of magnitude\(^{35}\) to have an effect on recombination in low-density (\(n_e< 10^{11} \text{ cm}^{-3}\)) plasmas. Third-body effects are inefficient because electron capture requires energy transfer to a third particle during the brief electron-ion encounter.

In the following, we will argue that additional stabilization mechanisms exist in the case of molecular ions, especially those that do not undergo fast dissociative recombination. Theory indicates the \(H_3^+\) ions may fall into this category. Non-dissociating, long-lived Rydberg states may be formed in collisions of electrons with molecular ions. These can be stabilized not only by energy-removing collisions but also by angular-momentum changing (\(l\)-mixing and possibly \(m\)-mixing) collisions.

The following sequence may occur: In the first step, an electron of low angular momentum (\(s\) or possibly \(p\)) is captured into a ro-vibrational level of a Rydberg state of \(H_3\),

\[
e^{-} + H_3^+ \rightarrow H_3^{**} (v=1, n, s).
\]

The \(n=7\) state should be favored since excitation to the \(v_2=1\) state (0.3126 eV) in \(n=7, l=0\) is resonant at an electron energy of +0.0309 eV (1.2 kT at 300 K).

We postulate that in the second step the angular momentum of the Rydberg molecule is randomized in collisions with ambient electrons (\(l\)-mixing and possibly \(m\)-mixing). It is known from studies of dielectronic recombination\(^{36}\) and zero-electron-kinetic-energy (ZEKE) spectroscopy\(^{37}\) that redistribution of angular momenta of Rydberg electrons (\(l\)-mixing) due to electric microfields and collisions can increase predissociation and autoionization lifetimes by several orders of magnitude. The basic reason is that the wave functions of high-\(l\) states do not interact with the ion core. Since the lifetimes of some \(H_3^{**}\) autoionizing states are quite long (of the order of \(10^{-7}\) s)\(^{38,39}\) and the \(l\)-mixing efficiency of thermal electrons is extremely large (rate coefficients \(>10^{-3} \text{ cm}^3/\text{s}\))\(^{40}\), \(l\)-mixing should occur in less than \(10^{-7}\) s under the conditions of our experiment (\(n_e = 4 \times 10^{10} \text{ cm}^{-3}\)).
FIGURE 1. $\beta$-values in H$_3^+$ or D$_3^+$ plasmas (from the $1/n_e$ vs. z plots), as a function of $[\text{H}_3]$ (triangles) or $[\text{D}_3]$ (filled circles). Lines: Least-squares fit to the data. Dotted lines: 95% confidence limits the least-squares fit.

FIGURE 2. $\beta(z)$ values (filled squares) derived from $n_e(z)$ data (open circles) at $[\text{H}_3]=1.3\times10^{14}$ cm$^{-3}$. Lines through $\beta(z)$ points: Model calculations for $k_1=1\times10^{-12}$ cm$^3$/s (dashed) and $k_2=1\times10^{-13}$ cm$^3$/s (dotted). Arrow: Position of the H$_2$ gas inlet.

FIGURE 3. Same as Fig. 3, but for $[\text{H}_3]=1.7\times10^{13}$ cm$^{-3}$.

FIGURE 4. Comparison of observed values of $\beta(n_e)$ at $[\text{H}_3]=1.3\times10^{14}$ cm$^{-3}$ (circles) and $[\text{H}_3]=1.7\times10^{13}$ cm$^{-3}$ (squares) to the model calculations described in the text.
In the third step, the long-lived $H_3^{**}(n=7, v=1, l>1)$ can be returned to the autoionizing state (s, or p) by further collisions, they may be quenched to lower $n$-states by neutral atoms or electrons, they may undergo slow predissociation, radiate to a lower state, or react with an $H_2$ molecule. Some of the decay rates are known or can be estimated. Collisional energy loss (n-changing) with helium atoms is not important compared to radiative decay. The rate coefficient for n-changing collisions of electrons with $n=7$ Rydberg atoms (electron quenching) has been measured by Devos et al.\textsuperscript{43} to be $10^{-5}$ cm$^3$/s. Radiative decay rates for $n=7$ hydrogenic states are close to $10^6$ s$^{-1}$ for s-states, $2x10^7$ s$^{-1}$ for p-states, and about $2x10^6$ s$^{-1}$ for a statistical mixture of l-values up to $l=6$.\textsuperscript{42} The reaction with $H_2$

$$H_3^{**} + H_2 \rightarrow H_2 + H_3^* \tag{8}$$

provides the needed enhancement of the loss rate of electrons in the presence of $H_2$. It will compete with radiative decay if it occurs on a time scale of $<$1μs. At $[H_2] = 1x10^{15}$ cm$^{-3}$, this requires a rate coefficient of $1x10^{-9}$cm$^3$/s for reaction (8), which is not unreasonable.

The set of reaction equations is easily solved to obtain an effective de-ionization coefficient. The capture and autoionization step

$$H_3^+ + e^- \rightarrow H_3^{**}(n=7, s,p) \tag{9}$$

is described by a capture coefficient $k_c$ and an autoionization rate $v_a$. Stabilization by l-mixing due to electrons

$$H_3^{**}(n=7, s,p)+ e^- \rightarrow H_3^{**}(n=7, l>1)+ e^- \tag{10}$$

and its reverse are described by rate coefficients $k_s$ and $k_i$, respectively. The following destruction reactions for $H_3^{**}$ were included:

- $H_3^{**}(n=7) \rightarrow H_3^*(n<7) + hv$ (radiation, rate $v_{rad}$) \textsuperscript{(11a)}
- $H_3^{**} \rightarrow H_2 + H$ (or 3H) (predissociation, rate $v_p$) \textsuperscript{(11b)}
- $H_3^{**}(n=7) + e^- \rightarrow H_3^*(n<7) + e^-$ (rate coefficient $k_{qe}$) \textsuperscript{(11c)}
- $H_3^{**} + H_2 \rightarrow H_2 + H_3$ (low n) (rate coefficient $k_H$). \textsuperscript{(11d)}

To allow for the possibility that the rate coefficients for $H_3^{**}$ in the autoionizing states (s, p) differ from those in higher l-states, we treat their destruction separately and attach the subscripts "0" and "l" to the rate coefficients. After some algebra, the effective de-ionization coefficient can be put in the form

$$
\beta = \frac{1}{n_e} \frac{dn_e}{dt} = \frac{k_c}{1 + \frac{n_e k_s v_i}{v_0} + \frac{n_e k_i v_i}{v_a (n_e k_i + v_i) v_a}} \tag{12}
$$

where $v_0 = v_{rad,0} + v_{p,0} + k_{qe} n_e + k_H [H_2]$ and $v_l = v_{rad,l} + v_{p,l} + k_{qe} n_e + k_H [H_2]$.\textsuperscript{42}
It is possible to find a set of plausible rate coefficients so that Eq. (12) gives a reasonable fit to the data. Fig. 4 shows a comparison to experimental values of $\beta (n_e)$ for the following set of rate coefficients:

$$
k_c = 4.4 \times 10^{-7} \text{ cm}^3/\text{s}, \quad v_{a} = 1.3 \times 10^8 \text{ s}^{-1}, \quad k_s = 1.25 \times 10^{-2} \text{ cm}^3/\text{s}
$$

$$
k_i = 3.1 \times 10^{-4} \text{ cm}^3/\text{s}, \quad v_{\text{rad},0} = v_{\text{rad},1} = 1.1 \times 10^6 \text{ s}^{-1}
$$

$$
v_{p,0} = v_{p,1} = 0, \quad k_H = 4.5 \times 10^{-9} \text{ cm}^3/\text{s}, \quad k_{qe} = 1 \times 10^{-5} \text{ cm}^3/\text{s}
$$

We do not claim that this set of model parameters is unique or that the agreement with the data is sufficient proof for the validity of the model. More detailed studies would be required to prove that this model is valid. It is interesting, however, that the fitting parameters are quite close to those that were estimated earlier. If this model is correct, it would solve the theoretical dilemma of having to account for fast "binary recombination" of electrons with H$_3^+$ ions since this process would not be needed to explain the observed plasma de-ionization.

Our model is compatible with the most recent merged-beam observations of Mitchell and coworkers$^{43}$. They found that the apparent H$_3^+$ recombination cross sections decreased by about a factor of five when a "deflection field" in the post-collision region was increased in magnitude. The effect is ascribed to field-induced re-ionization of autoionizing H$_3^{**}$ molecules that were formed in electron-H$_3^+$ collisions. The role of electric fields in such experiments is not yet perfectly understood, but the observations demonstrate that long-lived H$_3^{**}$ states can be produced in such collisions. The nature of these states and the reasons for their unexpectedly long life-time, however, are not at all clear.

CONCLUSIONS

We reach the tentative conclusion that the recombination of H$_3^+$ plasmas does not exclusively occur by binary recombination. If this is true, then the results of afterglow measurements may be compatible with the theoretical expectation that binary recombination of H$_3^+$ ions in $v=0$ should be rather slow.

The "true" binary recombination coefficient of H$_3^+$ is needed for some applications, e.g. for models of the interstellar medium. It is conceivable that the recombination cross sections measured in merged-beam or ion-storage ring experiments are close to the true values, but the effects of residual electric fields in such experiments need to be examined more closely.

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