Balmer Alpha Emission and Hydrogen Atom
Energy in Ion-Source Discharges

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Abstract

The structure of the hydrogen Balmer alpha line emission profiles from three types of neutral beam injector ion source plasmas (filling pressure ~ 10 mTorr, electron density $\sim 1-2 \times 10^{12} \text{ cm}^{-3}$, electron temperature $\sim 2-4 \text{ eV}$) is studied with the aid of a simple model for the neutral particle balance and H_α emission. A large fraction of the H_α is produced by dissociative excitation of H_2 and dissociative recombination of H_2^+ , while the remainder is produced by excitation of H atoms, most of which have energies that are close to the characteristic H_2 dissociation energies. The H_α linewidth is thus insensitive to the discharge operating conditions and equals $\sim 0.27 \text{ \AA}$ when only slow ($\sim 0.3 \text{ eV}$) dissociatively excited atoms are present or $\sim 0.35 \text{ \AA}$ when fast ($> 1 \text{ eV}$) atoms, apparently also produced in dissociation reactions, are present as well.

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INTRODUCTION

In this paper we present a qualitative account of discharge phenomena based on an interpretation of the Doppler-broadened profiles of hydrogen Balmer alpha (H_α , 6563Å) emission from some ion source plasmas. Measurements of H_α light were made on three types of ion sources: a magnetic multipole line-cusp "bucket" source, a duopigatron ion source with a conventional oxide cathode, and the same duopigatron source with an incandescent tungsten thermal predissociator as a cathode. The data were obtained with a Fabry-Perot interferometer (FPI) whose high resolution made accurate analysis of the H_α profile shapes convenient. A model for the excitation of H_α in these discharges is developed in some detail. Wall recombination of H atoms, nonthermal fast electron populations, and various sources of H_α light and H atoms are taken into account. A clear distinction was found in some of the observed line profiles between light produced by dissociative excitation of molecules (and electron-impact excitation of atoms with energies close to the characteristic dissociation energies) and that produced by excitation of cold (wall temperature) background atoms. We believe that this is the first time slow dissociation product atoms (with most probable energies of about 0.27 eV) have been conclusively identified as such in a hydrogen discharge. The comparative invariance of the H_α linewidths from the ion source plasmas is explained by the presence of these atoms. Our results on the H_α line profile yielded information on other physical properties of the discharges, including the density and energy distribution of the neutral atom population and the possible occurrence of hot nonthermal electrons. This analysis should be of interest in the general spectroscopy of low density hydrogen plasmas and in the development of neutral beam injectors for thermonuclear fusion research.

I. EXPERIMENT

A. The ion sources. The injection of high energy neutral beams is the principal auxiliary heating scheme in large magnetically-confined fusion experiments.¹ The largest neutral beam injectors in use today employ ion sources with discharge currents of 1 kA or more to produce hydrogen or deuterium ion currents of up to 100 A. These ion sources have a uniform discharge plasma over a large volume and the beams extracted from them contain a high fraction of atomic ions (H^+) compared with molecular ions (H_2^+ and H_3^+). The partially ionized multicomponent plasmas in these ion sources are not well understood to a great extent because of the large number of competing collision processes within the discharge volume and at the vessel walls.

The sources examined in these experiments were a 15-cm-diam line-cusp bucket (modified MacKenzie) source,² a 7-cm-diam duopigatron source with a conventional oxide cathode,³ and the same 7-cm-diam duopigatron source with a hot tungsten predissociator as a cathode.⁴ The diameters specified here refer to the diameters of the beams extracted from the sources (and to the approximate diameters of the discharge plasmas). We shall call these sources the bucket source, the conventional duopigatron source, and the modified duopigatron source, respectively.

The bucket source² consists of a cylindrical anode chamber with arrays of long bar magnets running parallel to its axis and forming a multiple-magnetic-cusp, a number of tungsten filaments as a cathode, and an extraction structure consisting of three multiaperture grids with an extraction area of diameter 5 cm. The first grid (the "plasma grid") is in contact with the ion source plasma and is at the cathode potential. In our experiments, the FPI viewed

across the 25-cm-diam chamber through a 2.5-cm-diam port about 10 cm from the plasma grid. The arc discharge current was varied over 42-84 A and the arc voltage was about 80 V. The nominal flow-through filling pressure was 5-20 mTorr.

The duopigatron ion source³ consists of a cathode inside an intermediate electrode (connected to the positive side of the arc power supply through a 1 k Ω resistor), two anodes (No. 1 with a 0.03 Ω and No. 2 with a 0.01 Ω resistor to the positive side of the arc supply), and a 7-cm-diam plasma grid with multiple apertures (connected to the positive side of the arc supply through 200 Ω). The duopigatron discharge consists of two discharges separated by a double sheath near the intermediate electrode aperture: a glow discharge (duoplasmatron configuration) inside the intermediate electrode serves as a plasma feed for a Penning ion gauge (PIG) discharge in the extended anode region with magnetic cusps. We observed the H_{α} light from the later region a few cm above the plasma grid. Duopigatron discharges with two different types of cathode were examined: a conventional oxide filament³ and an incandescent tungsten tube through which the supply gas is fed and thereby thermally dissociated.⁴ A hot tungsten tube has been used previously as a cathode and as a dissociator, together with thin tungsten sheet liners on the inside walls, in order to enhance the fraction of atomic ions in the discharge.⁴ A somewhat wider range of filling pressures and arc currents was examined on the duopigatron sources than on the bucket source.

The plasma parameters for all three sources have the following measured values:²⁻⁴ electron density $n_e = 1-2 \times 10^{12} \text{ cm}^{-3}$; electron temperature $T_e = 2-4 \text{ eV}$; and molecular density $n_{H_2} \sim 10^{14} \text{ cm}^{-3}$. The magnetic field in the bucket source is $< 100 \text{ G}$ at the edge of the discharge and $< 5 \text{ G}$ in the central (10-cm-diam) region, while that in the duopigatron falls from about 200 G in

the intermediate electrode region to about 20 G at the plasma grid. The plasma sizes are 7-20 cm, roughly equal to the extracted beam diameters, and the copper vacuum vessels have inside dimensions of 10-25 cm. The gas and plasma parameters may be regarded as reasonably uniform over the discharge. Certainly n_e is uniform (that is one of the requirements for a large area source) and T_e is roughly so according to probe measurements. The densities of neutral molecules and atoms are close to uniform because their mean free paths for loss by electron collisions are much greater than the vessel dimensions.⁵ Charged particle species measurements on the ion beams extracted from these sources indicate that the ion population in the discharge is close to 50% H^+ , 30% H_2^+ , and 20% H_3^+ ($n_e = n_{H^+} + n_{H_2^+} + n_{H_3^+}$).

B. The Fabry-Perot interferometer. A Burleigh Instruments RC-10 piezoelectrically scanning FPI was used as a spectrum analyzer. The interferometer consists of two partially transmitting mirrors (reflectivity 99% for wavelengths of 6000-7000 Å) mounted parallel to one another. When the FPI cavity is illuminated by a beam of collimated monochromatic light it will transmit the light if the optical distance between the reflecting surfaces is an integral number of wavelengths, i.e., the condition for constructive interference is satisfied.⁶ The range of wavelengths that can be seen through the interferometer in the same order without overlapping into other orders is known as the free spectral range (FSR) and is given by $\Delta\lambda_{FSR} = \lambda^2/2d$, where d is the mirror separation and λ is the wavelength of the light (all dimensions in the same units). The smallest obtainable width (full width half maximum, FWHM) of the transmission peaks from the interferometer is called the minimum resolving bandwidth $\Delta\lambda_R$. The instrument function of the FPI is characterized by the finesse F which is defined as $F = \Delta\lambda_{FSR}/\Delta\lambda_R$. The finesse is determined

by numerous factors, including the reflectivity, flatness, and parallelism of the mirrors, the focal length and aperture of the collimating system, and mechanical vibrations. In our experiments the finesse was measured by using a helium-neon laser as a narrow linewidth source ($\lambda = 6328 \text{ \AA}$ with an overall linewidth of $\leq 0.02 \text{ \AA}$) and optimized by adjusting the mirrors to obtain maximum transmission of the laser beam. The distance between the mirrors and, therefore, the wavelength of peak transmission are varied periodically by applying a ramp voltage to piezoelectric transducers mounted on one of the mirror holders. The scan period, which includes from two to four interference orders, was varied between 20 and 200 msec, and the retrace lasted 15 msec. The rapid linear scan made it possible to observe the time variation of the light during a single arc pulse.

After passing through the FPI (5.0-cm-diam mirrors), the light from the plasma was focussed by a collimating lens through a pinhole and onto a light guide. The light guide was directed through an interference filter onto a photomultiplier tube. The interference filter blocked out background light from the plasma and the filaments. The photomultiplier output could be smoothed by an RC filter, but electrically unfiltered data were taken during each run to ensure that the RC filter did not distort the profiles.

C. Data. The top portion of Fig. 1 shows the location of the viewing port for observing the duopigatron ion source. (The lower portion shows the arrangements for optical studies of extracted neutral beams. They are the subject of a previous report.⁷) The optical system was aligned and the FPI tuned by passing a He-Ne laser through a viewing port located opposite the 1.25-cm-diam observation port and backwards along the optical train. No data were taken while the high-voltage beam-extraction system was operating.

Figure 2 shows oscilloscope traces of data from the conventional duopigatron discharge: (a) the arc current and the electrically unfiltered photomultiplier response to the H_{α} light from the plasma with three interference orders during the 200 msec arc pulse; (b) the RC filtered photomultiplier signal (triggered at the beginning of the arc) and a ten times magnified view of a single spike near the middle of the discharge; and, (c) the response of the entire optical system to a He-Ne laser beam and a ten times magnified trace of a single spike. In Fig. 2c the short and long scribe marks correspond, respectively, to the FWHM of the magnified spike and to the FSR of the interferometer at the laser wavelength. This shows that the finesse $F \sim 60$, so that the instrument resolution is $\Delta\lambda_R = 0.035 \text{ \AA}$ for the measured plate separation $d = 0.10 \text{ cm}$.

The signal from the right hand side of the expanded trace of Fig. 2b is plotted in Fig. 3 (curve A) as a function of the wavelength shift $\Delta\lambda$ ($\text{\AA}$) from the profile peak. The bars indicate the range of signals obtained from a large number of discharges from this source under various operating conditions. The small bumps roughly $0.1 \text{ \AA}$ from the peak on both sides are a reproducible feature of all the H_{α} spectra taken on the conventional duopigatron source. This feature was not evident in the profiles from the bucket source, although it could have been lost in the noise. Except for the modified duopigatron source (see Fig.4), all the plasmas yielded fairly symmetric H_{α} profiles with a FWHM of $0.27 \pm 0.02 \text{ \AA}$ under all conditions and at all times during the $< 200 \text{ msec}$ discharges. Furthermore, these H_{α} profiles showed no discernible enhancement in the wings (to within 4% of the integrated line intensity) above a "Gaussian fit" profile with the same half-width.

An asymmetric H_α spectrum with the long wavelength side higher was observed on the modified duopigatron source. The plot in Fig. 4 was constructed from a series of five H_α traces from this source under various conditions. The discharge operating conditions (arc current and voltage, filling pressure) were the same as for the conventional duopigatron source. The intensity of the H_α light, however, is far higher in the wings ($\Delta\lambda > 0.3 \text{ \AA}$) than in the other cases. The line had approximately the same (within a factor of 2-3) total intensity as, and a larger FWHM ($\sim 0.35 \text{ \AA}$) than, the profile of Fig. 3 (curve A).

II. H_α EMISSION PROCESSES IN LOW-TEMPERATURE DISCHARGES

A. Sources of H_α . Neutral particle energies. In these discharges several processes may produce excited hydrogen atoms (H^*). We are concerned with excitation to the $n = 3$ state and emission on the $n = 3 \rightarrow 2$ transition (H_α). The most probable excitation processes are: dissociative excitation or the formation of excited atoms during dissociation of H_2 molecules by electrons (cross section denoted by σ_{e2})⁸

dissociative recombination of H_2^+ ions with electrons to yield an excited product atom ($k\sigma_{dr2}$, where $k < 1$ is the fraction of the dissociative recombination reactions which yields a given state or line)⁹

and, electron-impact excitation of H atoms (σ_{e1})¹⁰

Radiative electron-ion recombination ($e + H^+ \rightarrow H^*$) is negligible.¹¹ It is not known whether excited atoms are produced during formation of H_3^+ in the reaction ($\sigma_{H_3^+}$)⁵

the major process by which H_2^+ is lost in the ion source discharges and an important source of H atoms (product H atoms could be excited only if the H_3^+ is formed in an electronically excited state). We shall neglect it and the dissociation reactions of H_3^+ as sources of H_α , but this should have little effect on our conclusions.

Excited atoms formed by electron impact dissociation of H_2 belong to one of two energy groups:^{12,13} slow, with energies $E \sim 0.3$ eV (formed in states with $n > 2$ by electrons with energies $E_e \gtrsim 15$ eV) and fast, with energies $E \sim 2-8$ eV, the customary Franck-Condon energy (formed by electrons with $E_e \gtrsim 28$ eV). In low-temperature plasmas with $T_e \lesssim 4$ eV, most dissociation product atoms will belong to the slow group because of the relatively small number of electrons with energies sufficient to produce fast excited atoms. The kinetic energies of the atoms produced in reactions of H_2^+ have not been measured, but H_α spectral profiles from tokamaks suggest that the kinetic energies of atoms formed in dissociative excitation of H_2^+ ($e + H_2^+ \rightarrow H^+ + H + e$) are similar to those formed in dissociative excitation of H_2 .¹⁴

Furthermore, from the potential energy curves for H_2 and H_2^+ , it is clear that most of the atoms formed by dissociative recombination of H_2^+ in low-temperature plasmas will belong to the slow group.

The mean free path for elastic collisions between a newly formed 0.3 eV H atom and the cold (~ 0.025 eV) background neutrals (H and H_2) is ~ 2 cm.¹⁵ The mean free path for a ground state 0.3 eV atom to be excited by electron impact to the $n = 3$ state in the ion source plasmas exceeds 10^4 cm, which is much greater than the vessel size or the elastic collision mean free path. Thus, unless they are completely absorbed at the walls, many of the H atoms that are eventually excited to the $n = 3$ level will have lost energy in collisions with other gas particles and the walls. The H_α transition occurs much more rapidly than any other collision process (the emission mean free path is < 1 mm). Therefore, the H_α produced by dissociation of H_2 or H_2^+ will come from emitting atoms with the characteristic dissociation energies (~ 0.3 or ~ 3 eV), while that produced by excitation of ground state H atoms will come from emitters that have a spread of energies ranging from the dissociation energy to the thermal energy of the neutral gas in the vacuum vessel. We shall distinguish these three sources of H_α and call them, respectively, slow molecular H_α , fast molecular H_α , and atomic H_α .

B. Model for neutral particle balance and H_α production. In order to analyze the production of H_α light we begin with an estimate of the relative populations of atoms and molecules in the ion source discharges ($T_e = 2-4$ eV, $n_e = 10^{12}$ cm⁻³, $n_{H_2} = 10^{14}$ cm⁻³, and $n_{H^+} : n_{H_2} : n_{H_3} = 50:30:20$). The following reactions produce most of the H atoms in moderate density thermal discharges with $T_e < 4$ eV:⁵ dissociation of H_2 (σ_d)

dissociative recombination of H_2^+ (σ_{dr2})

H_3^+ formation ($\sigma_{H_3^+}$), reaction (4); and, dissociative recombination of H_3^+ (σ_{dr3})

In these discharges almost all of the H_2^+ , formed by electron-impact ionization of H_2 (σ_{ion})

is lost inside the plasma before it reaches the walls since the time for loss in reactions (4) and (6) is much shorter than the time for H_2^+ to reach the walls from the center of the discharge. Furthermore, most of the H_2^+ will yield H atoms, rather than protons,⁵ either directly [reactions (4) and (6)] or indirectly [reaction (4) followed by reaction (7)]. Therefore, the rate of production of H atoms in the discharge is given approximately by twice the sum of the rates of dissociation and ionization of H_2 :

$$2 n_e n_{H_2} [\langle \sigma_d v_e \rangle + \langle \sigma_{ion} v_e \rangle] \text{ atoms/cm}^3 \text{ sec} , \quad (9)$$

where $\langle \sigma_x v \rangle$ denotes the rate coefficient for reaction x.

Following Gabovich's treatment¹⁶ of the particle balance in ion sources, we equate the rate of production of H atoms to the rate of recombination of H at the walls:

$$2n_e n_{H_2} (\langle \sigma_d v_e \rangle + \langle \sigma_{ion} v_e \rangle) V = \frac{1}{4} n_H \bar{v}_H \gamma S, \quad (10)$$

where V is the volume of the plasma, S is the surface area of the vacuum vessel, $\bar{v}_H$ is the mean velocity of the hydrogen atoms in the discharge (2.5×10^5 cm/sec for the wall temperature of $\sim 290^\circ\text{K}$ and 7×10^5 cm/sec for the most probable dissociation energy), and γ is the fraction of neutral atoms incident on the wall that undergo recombination to form molecules (more precisely, here γ represents the sticking probability for H atoms). Here we assume that all ions recombine at the walls to form molecules and that the plasma parameters are uniform. In the case of a sphere of radius R (chosen for simplicity of analysis) filled with plasma, this yields

$$\frac{n_H}{n_{H_2}} = \frac{8 R n_e (\langle \sigma_d v_e \rangle + \langle \sigma_{ion} v_e \rangle)}{3 \gamma \bar{v}_H}, \quad (11)$$

for the ratio of atom and molecule densities in the plasma.

The ratio of the atomic H_α emissivity ϵ_a (reaction 3) to the molecular H_α emissivity ϵ_m (reactions 1 and 2) is given by⁹

$$\frac{\epsilon_a}{\epsilon_m} = \frac{n_e n_H \langle \sigma_{e1} v_e \rangle}{n_e n_{H_2} \langle \sigma_{e2} v_e \rangle + 0.044 n_e n_{H_2} \langle \sigma_{dr2} v_e \rangle} = \frac{n_H \langle \sigma_{e1} v_e \rangle}{n_{H_2} \langle \sigma_{em} v_e \rangle}. \quad (12)$$

Here¹⁷

$$\langle \sigma_{em} v_e \rangle = \langle \sigma_{e2} v_e \rangle + 0.044 \frac{n_{H_2^+}}{n_{H_2}} \langle \sigma_{dr2} v_e \rangle \approx \langle \sigma_{e2} v_e \rangle + \frac{n_e}{n_{H_2}} \langle \sigma_{ion} v_e \rangle \quad (13)$$

is an effective cross section for H_α emission from molecular dissociation that includes the formation of H_2^+ from H_2 ($\langle \sigma_{dr2} v_e \rangle \sim 3 \times 10^{-8} \text{ cm}^3/\text{sec}$ for $T_e = 2-4 \text{ eV}$).⁵ Therefore, for a sphere of radius R ,

$$\frac{\epsilon_a}{\epsilon_m} = \frac{8 R n_e \{ \langle \sigma_d v_e \rangle + \langle \sigma_{ion} v_e \rangle \} \langle \sigma_{e1} v_e \rangle}{3 \gamma \bar{v}_H \langle \sigma_{em} v_e \rangle} \quad (14)$$

We note that the two contributions to $\langle \sigma_{em} v_e \rangle$ are roughly equal and that in Eq. (9) the second part contributes less than 20% over the entire temperature range of interest.^{5,8,9}

Equations (11) and (14) have a reciprocal dependence on the surface recombination coefficient γ . With copper, Wood and Wise measured $\gamma \leq 0.14$ for a filament probe and $\gamma = 0.19$ for an evaporated film.¹⁸ The copper walls of the ion sources are probably coated with contaminants, are rough on a microscopic scale, and do not qualify as high purity materials. Thus, it is difficult to specify a precise value of γ , but it is probably higher than these values. A qualitative theoretical discussion by Ali-Khan et al.,¹⁹ indicates that the sticking probability at metal surfaces for H atoms with energies below a few tenths of an electron volt approaches unity. Evidently, the larger γ is, the smaller the number of H atoms in the discharge and the

greater the relative amount of molecular H_2 . γ and $\bar{v}_H$ are related in these plasmas. This can be seen qualitatively as follows. At low pressures, for $\gamma \sim 1$, the H atoms will experience few collisions with background gas particles before being lost at the walls, so that the H atoms in the gas will have energies close to the characteristic dissociation energy, 0.3 eV. Conversely, if $\gamma \ll 1$, the atoms will be slowed down both at the walls and in gas-phase collisions, so the kinetic energy of most of them will approach the background gas temperature and n_H will become very large [cf. Eq. (11)]. Since $\gamma \sim 0.2$, we may expect the atoms to have an "intermediate" nonthermal energy distribution.

In accordance with this discussion of the surface recombination coefficient, let us assume that the atoms in the plasma belong to two groups, hot and cold, with energies of approximately 0.3 and 0.025 eV, respectively (we may then write $n_H = n_{Hhot} + n_{Hcold}$). In the spectral data for the duopigatron source (see Section III. A), because of its narrow Doppler width the cold atomic H_α can be isolated from the remainder of the H_α profile which is roughly half molecular H_2 . Thus, the observations provide a lower limit to the ratio of the intensity of emission from the cold H atoms (ϵ_{ac}) to the molecular emission, ϵ_{ac}/ϵ_m (ϵ_{ah} , the emissivity from hot H atoms cannot be separated optically from ϵ_m). As we shall see, this estimate provides a starting point for estimating the populations of the two groups of atoms. The ratio ϵ_{ac}/ϵ_m is evidently also given by $\epsilon_{ac}/\epsilon_m = [n_e n_{Hcold} \langle \sigma_{e1} v_e \rangle] / [n_e n_{H2} \langle \sigma_{em} v_e \rangle]$. Hence, for known T_e and n_{H2} , we have

$$n_{Hcold} = \left(\frac{\epsilon_{ac}}{\epsilon_m} \right) n_{H2} \frac{\langle \sigma_{em} v_e \rangle}{\langle \sigma_{e1} v_e \rangle}. \quad (15)$$

An analogous expression exists for $n_{\text{Hhot}} (\epsilon_{\text{ac}} + \epsilon_{\text{ah}})$.

If a monoenergetic group of hot nonthermal electrons constitutes a fraction η of the total electron population, then the effect of such a group can be included in Eqs. (11) and (14) by replacing the rates $\langle \sigma v_e \rangle$ by $(1 - \eta) \langle \sigma v_e \rangle + \eta \sigma_f v_f$, where σ_f is the cross section for an electron with energy E_f and v_f is the velocity of an electron with energy E_f . In an ion source, the primary electrons emitted from the cathode are accelerated across the cathode fall (cathode-anode potential ~ 80 eV). A fraction of these electrons may reach the observation volume without having lost all their energy in collisions, although this has not previously been confirmed experimentally in these ion sources.

The fraction $\epsilon_{\text{mf}}/\epsilon_{\text{m}}$ of the molecular H_a emissivity (ϵ_{m}) in the fast molecular component (ϵ_{mf}) is very sensitive to the presence of fast electrons in a low temperature plasma. For $E_e > 28$ eV, roughly equal amounts of the dissociatively excited $\text{H}(n=3)$ atoms are in the fast and slow groups, while for $16 < E_e < 28$ eV, they are all in the slow group.¹³ If a small nonthermal group with $E_f < 28$ eV is present it will slightly reduce $\epsilon_{\text{mf}}/\epsilon_{\text{m}}$, while if $E_f > 28$ eV the effect may be calculated as follows:

$$\frac{\epsilon_{\text{mf}}}{\epsilon_{\text{m}}} \approx \frac{1}{2} \frac{\eta \sigma_{\text{emf}} v_f + (1 - \eta) \sigma_{\text{em}}(28) v_e(28) \phi(28, T_e)}{\eta \sigma_{\text{emf}} v_f + (1 - \eta) \langle \sigma_{\text{em}} v_e \rangle} \quad (16)$$

Here $\phi(E, T_e) \sim 1.1(E/T_e)^{1/2} \exp(-E/T_e)$ is the fraction of an electron population at temperature T_e with kinetic energy above $E (>> kT_e)$, σ_{emf} denotes the cross section for molecular H_a emission for an electron energy of E_f [the appropriate generalization of Eq. (13) must include dissociative excitation of

H_2^+ , $e + H_2^+ + H^+ + H + e$ at high electron energies]⁵, and $\sigma_{em}(28)$ and $v_e(28)$ denote the quantities for $E_e = 28$ eV [the onset energy for fast $H(n=3)$]. While this expression is extremely sensitive to the presence of a small group of nonthermal hot electrons, ϵ_a/ϵ_m and n_H/n_{H_2} are comparatively insensitive. The effect of nonthermal electrons is more pronounced for low T_e plasmas because the fraction of fast electrons in the Maxwellian distribution is small.

C. Results of the model for the duopigatron source. Table 1 lists some sample calculations for the duopigatron source using the expressions derived in the preceding section. The radius of the ion source plasma is $R \sim 3.5$ cm, $n_e = 10^{12}$ cm⁻³, and $n_{H_2} \sim 10^{14}$ cm⁻³. As mentioned previously, a major difficulty in the use of this model is the lack of reliable data on γ and $\bar{v}_H$. Later, in interpreting our data, we shall consider two cases: (a) $\gamma \sim 1$ with $\bar{v}_H = 7 \times 10^5$ cm/sec, and (b) $\gamma = 0.2-0.3$ with $\bar{v}_H = 4-5 \times 10^5$ cm/sec (i.e., $\gamma\bar{v}_H = 1-7 \times 10^5$ cm/sec). The table lists the following quantities for thermal plasmas with $T_e = 2, 3$, and 4 eV in the absence of fast electrons ($\eta = 0$) and in the presence of 2% and 5% groups of monoenergetic electrons with energies $E_f = 25$ and 40 eV: n_H/n_{H_2} from Eq. (11); ϵ_a/ϵ_m from Eq. (14); $n_{H_{cold}} n_{H_2}^{-1} (\epsilon_{ac}/\epsilon_m)^{-1}$ from Eq. (15); and, ϵ_{mF}/ϵ_m from Eq. (16) for $E_f = 40$ eV (which is above the threshold for production of fast excited atoms in the dissociative excitation of H_2) and from a similar equation for $E_f = 25$ eV (below the threshold). The estimated errors in the calculations are $\lesssim 40\%$.

From the calculations given in the table we can notice a number of effects: (i) As T_e becomes higher, n_H/n_{H_2} increases and thus the relative amount of atomic H_α emission (ϵ_a/ϵ_m) rises as well. (ii) The fraction of fast molecular H_α (ϵ_{mF}/ϵ_m) increases with T_e for $E_f < 28$ eV. (iii) The extreme sensitivity of ϵ_{mF}/ϵ_m to the presence of a hot electron component with

$E_f > 28$ eV is evident in the table. A separate calculation shows that for $E_f = 40$ eV, $\epsilon_{mF}/\epsilon_m > 0.1$ for $\eta > 0.005$ when $T_e < 4$ eV. The presence of a hot electron component with $E_f < 28$ eV, on the other hand, reduces the relative amount of emission from the fast dissociation products for fixed T_e . (iv) The amount of atomic H_α relative to molecular H_α , ϵ_a/ϵ_m , exhibits a (weak) minimum at some fraction $\eta = \eta_{\min}$. This ratio, however, is comparatively insensitive to T_e when T_e is 3 eV or more. (v) One may expect to see substantial amounts of molecular H_α compared to atomic H_α in ion source discharges of this type for fixed n_e with combinations of the following characteristics: low bulk T_e , large wall recombination (sticking) coefficients, and very low pressures. Other molecular reactions besides reactions (1) and (2) may, of course, add to the molecular H_α contribution.

III. INTERPRETATION OF THE SPECTRAL DATA

A. Doppler broadening. The principal cause of the H_α broadening observed in our experiments is Doppler broadening. Other effects are relatively small:²⁰ the instrument width was 0.035 Å in the data of Fig. 3 (curve A), and Stark broadening and Zeeman splitting cannot exceed 0.02 and 0.01 Å respectively. The measured H_α linewidth was found to be constant at 0.27 ± 0.02 Å for the bucket and conventional duopigatron sources under all operating conditions. (We defer discussion of the modified duopigatron and its wider H_α profile until the end of the next section.) This suggests that the width is determined principally by molecular processes which yield atoms of fixed energies. Emitting atoms with an isotropic kinetic energy distribution $f_E(E)$ yield a Doppler broadened spectral profile whose shape $I(\Delta\lambda)$ is given by

$$I(\Delta\lambda) \propto \int_{E(\Delta\lambda)}^{\infty} E^{-1/2} f_E(E) dE, \quad (17)$$

where $E(\Delta\lambda) = mc^2\Delta\lambda^2/(2\lambda_0^2)$, $\Delta\lambda$ is the wavelength shift relative to the proper wavelength λ_0 , m is the mass of an atom, and c is the speed of light. Hydrogen atoms with a temperature T_H (eV) have Doppler-broadened 6563 Å H_α fine structure components with a Gaussian shape of FWHM $0.504T_H^{1/2}$ (Å). The kinetic energy distributions of both the fast and slow dissociation product atoms from H_2 , however, differ somewhat from Maxwellian.¹² In particular, the slow group is deficient at both high and low energies compared to a Maxwellian energy distribution having the same most probable energy, although this difference would probably not be observable with our apparatus.

Figure 3 shows: (A) the measured H_α profile from the conventional duopigatron source; (B) a calculated H_α spectrum using Eq. (17) for the case of two equally intense (as the observed profile A is symmetric) fine-structure components located 0.14 Å apart²¹ for atoms having the energy distribution of slow dissociation product $H(2s)$ atoms measured by Misakian and Zorn¹²; (C) as curve B except for Maxwellian atoms with temperature $T_H = 0.23$ eV; (D) the difference between curves A and C; and, (E) the calculated H_α spectrum for Maxwellian atoms with $T_H = 0.025$ eV and relative fine structure intensities of 1, 0.3, and 1 at $\Delta\lambda = -0.07$, 0, and 0.07 Å, respectively, combined with a Gaussian²² instrument function with an FWHM of 0.036 Å. Curves C and B are normalized to curve A at $\Delta\lambda = 0.14$ and 0.15 Å, respectively. Considering the rather large errors in Misakian and Zorn's measurements at low energies¹² (i.e., in curve B for $\Delta\lambda < 0.1$ Å), one can approximate the H_α spectrum (B)

from the slow dissociation product atoms rather well by curve C, which is based on a Maxwellian distribution of atoms. For $\Delta\lambda > 0.15 \text{ \AA}$, curves A, B, and C are in good agreement.

B. Energies of H atoms and fast nonthermal electrons in the discharges. The measured H_{α} linewidth of $0.27 \pm 0.02 \text{ \AA}$ is slightly less than that reported in spectroscopic observations of the excited $H(n=3)$ atoms formed during dissociative excitation of hydrogen gas by an electron beam,¹³ which was $0.31 \text{ \AA}$ (when the fine-structure correction is recombined with the derived Doppler width). Our experimental H_{α} profile (curve A of Fig. 3) is also somewhat narrower than that (curve B) derived for the measured $H(2s)$ kinetic energy distribution. The small symmetric bumps at $\Delta\lambda \approx \pm 0.1 \text{ \AA}$ in the traces from the duopigatron source appear to be a consequence of H_{α} emission from cold background atoms which have been excited by electron impact. Thus, as shown in Fig. 3, the measured H_{α} spectrum is the sum of contributions from slow dissociation product atoms (plus some amount from "re-excitation" of fresh dissociation product atoms that have not been slowed down, i.e., from n_{Hhot} , that is indistinguishable in terms of its Doppler width from the slow molecular H_{α}) and from thermalized background atoms; that is, curve A of Fig. 3 is the sum of curves C ($\approx$ curve B) and D. The closeness of the calculated spectrum (E) for thermal atoms with $T_0 = 0.025 \text{ eV}$ (290°K) to curve D supports this division of the emitting atom population into hot and cold groups (~ 0.3 and 0.025 eV). This part (D) of the spectrum causes the small bumps at $\Delta\lambda \sim \pm 0.1 \text{ \AA}$ and causes a slight narrowing of the spectrum compared to that obtained for dissociative excitation alone (curve B). Some narrowing may also be caused by the fact that some of the atoms that are "re-excited" will have lost some energy, but not enough to "put" them in the cold atom group. (The difference between curves A and B for $\Delta\lambda > 0.15 \text{ eV}$ suggests this, but the experimental errors in both preclude a firm conclusion.)

The area under curve C is roughly five times that under curve D; thus, $\epsilon_{ac}/\epsilon_m > 0.2$. For thermal plasmas with $T_e = 2, 3$, and 4 eV, Eq. (15) shows that $n_{Hcold} > 0.8, 1.8$, and $2.2 \times 10^{12} \text{ cm}^{-3}$, respectively ($n_{H_2} = 10^{14} \text{ cm}^{-3}$). From the table it is clear that model case (a) ($\gamma \bar{v}_H = 7 \times 10^5 \text{ cm/sec}$), besides implying that there should be no cold atoms, yields a value of n_H that is below these values of n_{Hcold} for $T_e < 4$ eV in a thermal plasma. We thus choose, still somewhat arbitrarily, case (b) ($\gamma \bar{v}_H = 1 \times 10^5 \text{ cm/sec}$) to obtain a crude estimate of the atom density and energy distribution, and we use the probe data and beam species measurements to set $T_e = 3$ or 4 eV. According to the table, a fast electron component that constitutes at most a few percent of an electron population with $T_e > 3$ eV has little effect on n_H/n_{H_2} or ϵ_a/ϵ_m . Thus, for $n_{H_2} = 10^{14} \text{ cm}^{-3}$, we have $n_H = 1.0 - 2.7 \times 10^{13} \text{ cm}^{-3}$ (corresponding to $T_e = 3-4$ eV). The corresponding $\epsilon_a/\epsilon_m = 1.1-2.5$, so that approximately 57-35% of the intensity under curve C of Fig. 3 is molecular H_2 and the rest is hot (~ 0.3 eV) atomic H_a . The actual value of ϵ_{ac}/ϵ_m is thus about 0.4 (.35-.57), so that $n_{Hcold} \sim 4 \times 10^{12} \text{ cm}^{-3}$, while $n_{Hhot} \sim 0.6-2 \times 10^{13} \text{ cm}^{-3}$. This qualitative relationship will hold for other reasonable choices of $\gamma \bar{v}_H$. It therefore appears (despite the lack of precise data on the surface recombination coefficient) that hot ($E \sim 0.3$ eV) atoms are more numerous than cold (~ 0.025 eV) in the duopigatron ion source; i.e., most atoms are not slowed down to the background (wall) temperature before they are lost. Since the bucket source also yields H_a profiles that are slightly narrower than those obtained from molecular H_2 alone, atomic H_a is probably superimposed on the molecular H_2 in this discharge as well. There the bumps corresponding to cold atomic H_a may be lost in the noise because of a smaller cold atom population.

In the modified duopigatron source the operating conditions were the same as in the conventional one, but the H_α spectrum obtained from it (Fig. 4) was strikingly different (cf. Fig. 3, curve A). Since this profile extends beyond $\Delta\lambda = 0.4 \text{ \AA}$, it is clear that atoms with velocities in excess of $1.8 \times 10^6 \text{ cm/sec}$ ($E = 1.5 \text{ eV}$) exist in this plasma. The asymmetry of this profile, unlike that of Fig. 3 (cf. Ref. 22), is consistent with the relative fine structure intensities reported in Ref. 21. The H_α spectra from the modified source can be resolved into two components of similar intensity: one with a width close to that observed in the conventional duopigatron (slow molecular + "hot," 0.3 eV, atomic H_α) and the other with a width $\gtrsim 0.7 \text{ \AA}$ (produced by emitting atoms with energies over 1 eV). The second component is not as wide as the fast molecular H_α spectrum observed in electron beam bombardment of H_2 gas,¹³ presumably because a significant portion of this component of the ion source profile is produced by excitation of fast dissociation product atoms that have been slowed down in repeated collisions with the walls²³ and the background gas. There are two possible mechanisms by which atoms with energies above 1.5 eV may appear in the discharge. The first involves Coulomb collisional heating of H_2^+ ions by H^+ in the discharge to temperatures on the order of 1-2 eV, followed by dissociation of the hot H_2^+ by low temperature electrons. (The H_2 , unheated, would provide the narrower H_α component.) However, since the known plasma parameters (n_e , T_e) are the same as in the conventional source, there is little reason to expect the H^+ to be significantly hotter than in the modified source. (This mechanism could be important in a plasma with the required T_{H^+} and a much lower n_{H_2} .) The second mechanism is for a small population of fast electrons with energies above the threshold for production of fast dissociation product atoms to enhance the fast molecular H_α intensity (see the Table).²⁴ A fraction $\eta \gtrsim 0.005$ is

sufficient to produce the effect observed here. (A small difference in the energy distributions of the hot nonthermal electron populations in the two sources would cause no noticeable difference in the other plasma parameters.) The appearance of a slightly hotter electron population in the modified source might be a result of some change induced by the additional dissociation and heating of H_2 by the hot tungsten (e.g., a pressure change) or simply a result of geometrical changes inside the intermediate electrode which permit hotter electrons to move deeper into the PIG region (e.g., greater length of the cathode). The modified duopigatron results are of particular interest because they appear, despite the obvious immediate differences in the H_α spectrum, to confirm our earlier conclusions about the mechanisms by which H_α light and H atoms are produced and about the magnitude of the surface recombination (sticking) coefficient.

IV. CONCLUSION

A model for the production of H_α in discharges with $T_e = 2-4$ eV, $n_e = 1-2 \times 10^{12} \text{ cm}^{-3}$, and $n_{H_2} \sim 10^{14} \text{ cm}^{-3}$ has been used to interpret the H_α line emission from three types of neutral beam injector ion sources. The relative amounts of H_α emission from the slow ($E \sim 0.3$ eV) and fast ($E \sim 3$ eV) atoms formed by electron-impact dissociative excitation of H_2 and dissociative recombination of H_2^+ and from the electron-impact excitation of cold ($\sim 290^\circ\text{K}$) and hot (~ 0.3 eV) background H atoms have been calculated for a duopigatron source. A rough analysis of the data from the conventional duopigatron source shows that $n_H = 1-2.5 \times 10^{13} \text{ cm}^{-3}$, of which about $4 \times 10^{12} \text{ cm}^{-3}$ are at the wall temperature and $6-20 \times 10^{12} \text{ cm}^{-3}$ are at energies close to the slow dissociation product energy (0.3 eV). 25-45% of the total intensity of the H_α

light from the plasma is produced by electron-impact dissociation of H_2 and H_2^+ (this fraction may be greater if other reactions of H_2^+ and H_3^+ yield H_α). The H_α spectral profile is sensitive to the presence of a fast nonthermal electron group and to the energy dependence of the surface recombination (sticking) coefficient for H atoms. In the conventional oxide cathode duopigatron and in the bucket source, all the H_α emission appears to come from slow dissociation product atoms (some are slowed down), while in the modified duopigatron source a substantial fraction of the emitting atoms have energies above 1 eV. These atoms are apparently fast product atoms from the dissociation of H_2 and the most probable reason for their appearance is the presence of a small amount (0.5% of the total electron population is sufficient) of nonthermal electrons with energies in excess of the threshold for production of fast dissociation product atoms.

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²²The FPI actually has an Airy function response⁶ but a Gaussian was used here for ease of calculation. The central fine structure peak reported in Ref. 21 has a relative intensity of about 0.15, but here it has been enhanced to improve the fit between curves D and E. This difference from Ref. 21 and, especially, the apparent symmetry of curve A of Fig. 3 suggest differences in the excitation conditions and fine structure intensities.

²³According to Ref. 19, γ for metals decreases with increasing atom energy over the range $0.1 \text{ eV} < E < 100 \text{ eV}$. Thus, because fast dissociation product atoms should undergo more wall reflections than slow atoms before they are lost, and will lose energy during this time, this variation in γ with E may enhance the relative amount of fast ($E \sim 3 \text{ eV}$) compared to slow ($E \sim 0.3 \text{ eV}$) atoms in the discharge and cause the "re-excitation" atomic H_α profile to have a ">1 eV" component that is more intense than the ".3 eV" component. Such a variation in γ may also partially explain the small amount of cold (0.025 eV) atoms seen in the conventional discharge since atoms with $E < .1 \text{ eV}$ have a much higher sticking probability than those with $E \sim .3 \text{ eV}$. On the other hand, since the measurements of Ref. 18 were presumably made with cold atoms (in equilibrium with the walls) it is doubtful that for our ion sources γ comes as close to unity at low E as suggested in Ref. 19.

²⁴Unexcited fast dissociation product atoms can be produced by electrons with $E_e > 18 \text{ eV}$ and since, as we have seen, "atomic" H_α is significant, the actual energy of the responsible fast electrons may be substantially lower than the nominal 40 eV of the Table. However, a slight difference in the fast electron populations of the two duopigatron sources remains a plausible explanation of the differences in their H_α spectra. Note also that the two mechanisms proposed here for the fast atoms are related to the extent that ionization of H atoms is the origin of the H^+ ions in the plasma.

Table 1

$T_e = 2 \text{ eV}$	$\eta = 0.02$		$\eta = 0.05$		
	$n = 0$	$E_f = 25 \text{ eV}$	40 eV	$E_f = 25 \text{ eV}$	40 eV
$n_H/n_{H_2} \times \gamma \tilde{\nu}_H$	2.4×10^3	9.5×10^3	1.2×10^4	2.0×10^4	2.4×10^4
$\epsilon_a/\epsilon_m \times \gamma \tilde{\nu}_H$	6.5×10^4	6.2×10^4	5×10^4	1.2×10^5	1.0×10^5
$n_{Hcold} n_{H_2}^{-1} (\epsilon_m/\epsilon_{ac})$	0.04	0.15	0.23	0.17	0.25
ϵ_{mf}/ϵ_m	2×10^{-3}	6×10^{-5}	0.48	4×10^{-5}	0.50
$T_e = 3 \text{ eV}$					
$n_H/n_{H_2} \times \gamma \tilde{\nu}_H$	1.0×10^4	1.7×10^4	1.9×10^4	2.7×10^4	3.2×10^4
$\epsilon_a/\epsilon_m \times \gamma \tilde{\nu}_H$	1.1×10^5	1.4×10^5	1.2×10^5	1.9×10^5	1.5×10^5
$n_{Hcold} n_{H_2}^{-1} (\epsilon_m/\epsilon_{ac})$	0.09	0.12	0.17	0.14	0.21
ϵ_{mf}/ϵ_m	1×10^{-2}	4×10^{-3}	0.36	2×10^{-3}	0.43
$T_e = 4 \text{ eV}$					
$n_H/n_{H_2} \times \gamma \tilde{\nu}_H$	2.7×10^4	3.5×10^4	3.5×10^4	4.5×10^4	5×10^4
$\epsilon_a/\epsilon_m \times \gamma \tilde{\nu}_H$	2.5×10^5	3.0×10^5	2.7×10^5	3.2×10^5	2.7×10^5
$n_{Hcold} n_{H_2}^{-1} (\epsilon_m/\epsilon_{ac})$	0.11	0.14	0.14	0.13	0.18
ϵ_{mf}/ϵ_m	3×10^{-2}	2×10^{-2}	0.22	1×10^{-2}	0.32

Calculated values of the ratios of the atom density to the molecule density (n_H/n_{H_2}) and of the atomic H_α emissivity to the molecular H_α emissivity (ϵ_a/ϵ_m), of the cold (wall temperature) atom density (n_{Hcold}), and of the fraction of molecular H_2 produced by fast dissociation product atoms (ϵ_{mf}/ϵ_m) for a deuterium plasma with $R = 3.5 \text{ cm}$, $n_e = 1 \times 10^{12} \text{ cm}^{-3}$, and $n_{H_2} \sim 10^{14} \text{ cm}^{-3}$ for fast electron fractions $\eta = 0, 0.02$, and 0.05 and fast electron energies $E_f = 25$ and 40 eV . The expected value of $\gamma \tilde{\nu}_H$ is in the range $1 \times 10^5 - 7 \times 10^5 \text{ cm/sec}$.

FIGURE CAPTIONS

- Fig. 1. A sketch of the duopigatron ion source and the beam chamber. The top portion shows the arrangement for FPI measurements on the 7-cm-diam source plasmas, and the bottom, that for observations⁷ of extracted beams.
- Fig. 2. FPI data from the 7-cm-diam duopigatron source with a conventional oxide cathode: (a) the arc current and unfiltered H_{α} photomultiplier output; (b) the filtered photomultiplier output with a 10X magnified trace of one spike; and, (c) the FPI response to a He-Ne laser, illustrating the resolution of the FPI.
- Fig. 3. (A) The signal from the right-hand side of the expanded trace of Fig. 2b; (B) the H_{α} spectrum calculated from the kinetic energy distribution of slow $H(2s)$ dissociation product atoms^{1,2} normalized to curve A at $0.15\text{\AA}$; (C) the H_{α} spectrum calculated for atoms with a temperature of 0.23 eV normalized to curve A at $0.14\text{ \AA}$; (D) the difference between curves A and C; (E) the H_{α} spectrum calculated for atoms with a temperature of 0.025 eV combined with a Gaussian instrument function of width $0.036\text{ \AA}$.
- Fig. 4. The H_{α} spectrum obtained from the modified duopigatron source with a hot tungsten predissociator. The bars indicate the range of signals from a series of five discharges.

Fig. 1

Fig. 2

Fig. 4