

# Impurities, temperature and density in a miniature electrostatic plasma and current source

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**Abstract.** We have spectroscopically investigated the Sterling Scientific miniature electrostatic plasma source—a plasma gun. This gun is a clean source of high-density ( $10^{19}$ – $10^{20}$  m<sup>-3</sup>), low-temperature (5–15 eV) plasma. A key result of our investigation is that molybdenum from the gun electrodes is largely trapped in the internal gun discharge; only a small amount escapes in the plasma flowing out of the gun. In addition, the gun plasma parameters actually improve (even lower impurity contamination and higher ion temperature) when up to 1 kA of electron current is extracted from the gun via the application of an external bias. This improvement occurs because the internal gun anode no longer acts as the current return for the internal gun discharge. The gun plasma is a virtual plasma electrode capable of sourcing an electron emission current density of 1 kA cm<sup>-2</sup>. The high emission current, small size (3–4 cm diameter), and low impurity generation make this gun attractive for a variety of fusion and plasma technology applications.

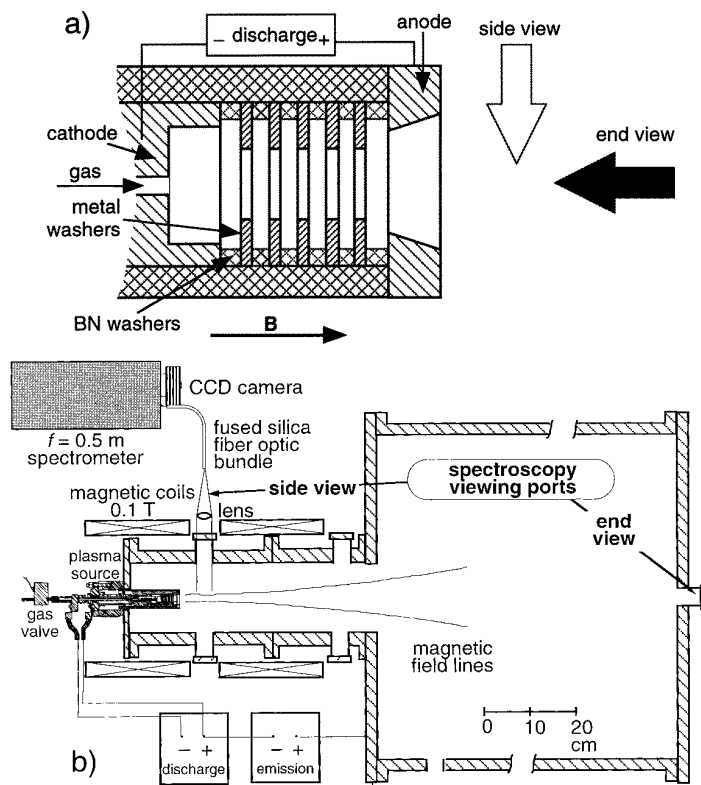
## 1. Introduction

The Sterling Scientific plasma gun is capable of producing a clean, high-density ( $10^{19}$ – $10^{20}$  m<sup>-3</sup>), low-temperature ( $T_i \approx T_e \approx 5$ –15 eV) plasma for pulse lengths of at least 40 ms. With the addition of a bias electric field, it is a high-current emission source (1 kA) at high current density (1 kA cm<sup>-2</sup>). A schematic diagram of a typical source assembly is shown in figure 1(a). The heart of the source is a set of electrodes used to form a small, cylindrical arc discharge plasma column. The principal electrodes are the anode and cathode. In addition to these, a stack of washers is used to define the arc channel between the anode and cathode. The washer stack is composed of alternating insulating and (electrically isolated) metal washers. Washers are used to minimize the plasma contact with material surfaces while providing a partial conducting boundary to help stabilize the plasma column. (The insulating washers have larger inner diameter than the metal washers and do not significantly contact plasma.) The source is very reproducible and reliable with metal washers, whereas it can become erratic with only insulating washers. This design represents the evolution of earlier designs using the same basic concepts [1, 2].

Spectroscopic investigation of the gun plasma had two major purposes. The first was to identify the major impurities and determine their behaviour, both within the

internal gun discharge and in the plasma flowing out of the gun. The second was to determine ion and neutral temperatures ( $T_i$  and  $T_n$ ) and ion density ( $n_i$ ) via Doppler and Stark broadening respectively. A key result is the observation that there appears to be only a very small amount of molybdenum (from the gun electrodes) in the hydrogen plasma flowing out of the gun when it is optimally operated. In addition, the gun plasma parameters actually improve (lower impurity contamination and higher  $T_i$ ) when current is extracted from the gun. These results confirm the hypothesis that the gun plasma acts as a ‘virtual electrode’ during current extraction.

The spectrometer used for this work was specifically designed and built to diagnose the gun plasma [3]. It is fibre optically coupled and equipped with a computer controlled CCD detector. This instrument is capable of both high resolution ( $\Delta\lambda \leq 0.015$  nm) for line broadening measurements and large bandpass (110 nm with  $\Delta\lambda \sim 0.3$  nm) for impurity surveys. Most of the spectroscopic data presented below were taken from one of two chordal views of the plasma (figure 1). The ‘end view’ was aligned with the axis of the gun discharge and viewed it directly through the hole in the anode. The ‘side view’ was perpendicular to the axis of the gun discharge and viewed the plasma flowing out of the gun, about 5 cm past the gun anode.



**Figure 1.** Overall view of the (a) plasma source and (b) test chamber, including placement of the side and end views used for spectroscopic measurements.

## 2. Impurity survey

### 2.1. Bright emission lines

Table 1 is a catalogue of the bright ultraviolet and visible emission lines identified in the variety of gun plasmas that were produced. Figure 2 shows typical spectra from 200 to 300 nm and 300 to 400 nm. The low- $Z$  impurities boron, carbon, and nitrogen come mostly from the boron nitride spacer washers inside the gun, while the metal impurities come from the internal gun electrodes and metal spacer washers. The bright lines observed all came from neutral to doubly ionized impurity ions. More highly ionized species did not appear to be present; the bright triplets of B IV at 282 nm and C V at 227 nm were notably absent even though the electron temperature (5 to 20 eV, depending on gun operation) was high enough to produce such ions. We speculate that particle confinement time in the gun plasma is not long enough to allow the impurities to reach the more highly ionized states.

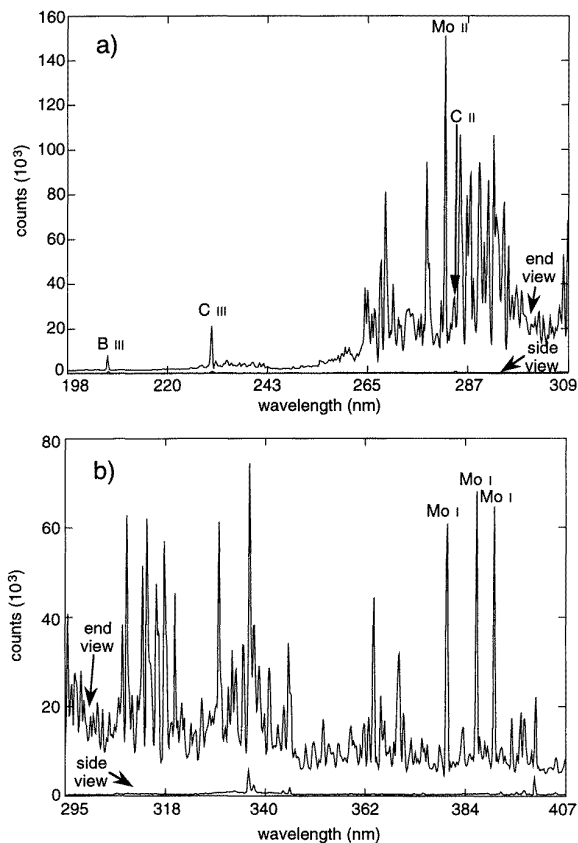
The majority of gun operations has been with molybdenum electrodes and washers; it was operated with copper electrodes and washers for a short time to quantify how deleterious the increased sputtering coefficient of copper was to gun operation. Most of the data presented in this section were taken by integrating (with the CCD detector) the line emission over the entire gun discharge. A single monochromator with a photomultiplier tube was also used to measure the time variation of a few specific lines during a gun discharge.

**Table 1.** Bright UV and visible emission lines in gun plasmas.

Species	Bright lines (nm)
H I	434.047, 486.133, 656.3
B II	345.129
B III	206.578, 206.723
C II	283.671, 283.760, 514.516
C III	229.687
N II	500.515, 567.956
Cu I	324.754, 327.396
Cu II	224.700
Mo I	379.825, 386.411, 390.296
Mo II	281.615, 281.744
Mo III	233.093

### 2.2. Molybdenum electrode guns

Several different molybdenum electrode gun configurations were examined spectroscopically, but no large difference in the behaviour of impurities was observed. Instead, the most striking result is the substantial difference in character between the spectra recorded with the end view from those recorded with the side view (figure 2). All of the impurity features are greatly suppressed in the side view compared to the end view. In particular, note that the Mo I and Mo II lines are not even visible on the side view spectra in figure 2, while the same lines are clearly apparent in the end view. Some of this difference is due to the fact that the end viewing chord integrates over the emission from



**Figure 2.** Comparison of relative impurity levels in the internal gun plasma (end view) and plasma flowing out of the gun (side view). (a) Spectrum from 200 to 300 nm, (b) from 300 to 400 nm.

a much larger volume of plasma than the side viewing chord. The plasma inside the gun seen by the end view is also more dense than that seen by the side view. The magnitude of these effects can be roughly estimated by comparing the ratios of the  $H\beta$ , B III, C II, and C III line emission from the side view to that from the end view. These ratios are typically about 10%, implying that Mo line emission seen by the side view could be reasonably expected to be about 10% of that seen by the end view. In fact, the Mo I emission is  $\leq 0.5\%$  of that seen by the end view, while the upper bound on the Mo II emission ratio is even smaller ( $\leq 0.2\%$ ). There are at least two non-exclusive explanations for this behaviour. First, sputtering of molybdenum may be more severe at the gun cathode than the anode because the cathode suffers ion bombardment while the anode draws only electron current. Thus the plasma surrounding the cathode (which is clearly seen by the spectrometer in the end view) may be more heavily contaminated since it is close to the source of molybdenum. Second, the molybdenum in the internal gun plasma may be partly 'trapped' so that it does not flow out of the gun with the external plasma stream. This trapping at the cathode may be simply that most of the molybdenum atoms sputtered off the cathode quickly ionize and fall back onto the cathode.

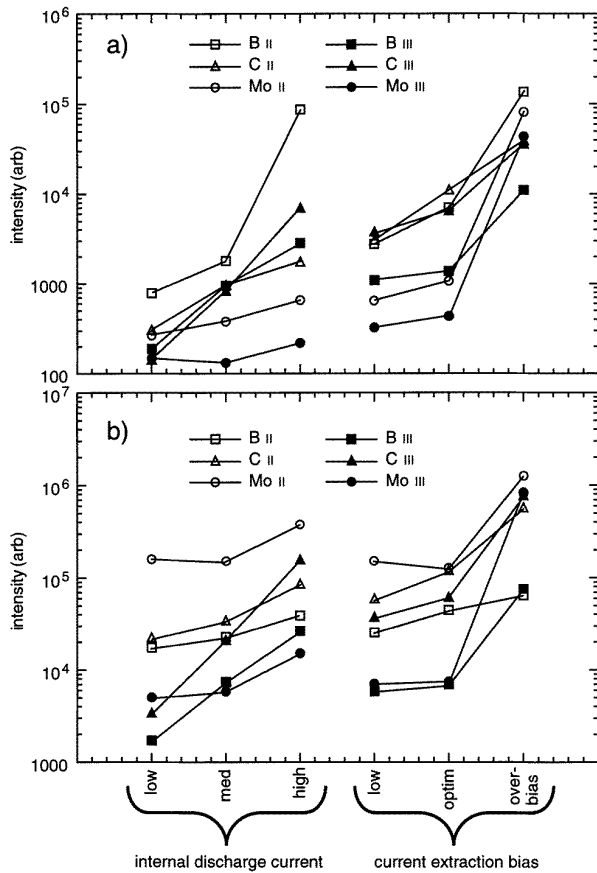
Operation of the plasma gun can be degraded if too much current is forced to flow in the internal discharge

and/or if too little hydrogen gas is supplied to fuel this discharge. When the gun is operated in these conditions, the overall level of impurity line emission increases and can become very large if the discharge is severely undergassed or if the internal current density greatly exceeds  $1 \text{ kA cm}^{-2}$  (this may be the current density limit for this type of gun).

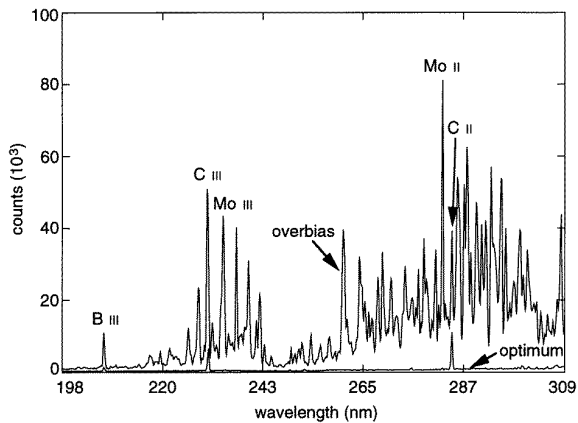
When the plasma gun is operated as an electron current source by application of a voltage bias between the gun and some distant current collector (e.g., a vacuum vessel wall), the plasma inside the gun acts as a virtual electrode. This type of operation does not raise impurity contamination of the plasma and may actually lower impurity contamination below that achieved during operation of the gun as a simple plasma source. Figure 3 contains a comparison of impurity line emission intensity obtained during 'biased' operation (electron current being extracted) and 'unbiased' operation. The first three sets of datapoints on the left side of the graphs are from unbiased operation of the gun; the optimum discharge current of about 1000 A lies between the point at medium current point at 820 A and the high-current point at 1120 A. Note that impurity contamination of the plasma begins to rise significantly at the high discharge current. The second set of three datapoints on the right side of the graphs is from biased operation of the gun, with the internal discharge current at the optimum of about 1000 A. In general, impurity levels decrease or remain unchanged at the low (40 V, 550 A) and optimum (85 V, 900 A) bias settings, but dramatically increase at the overbias point (115 V, 1030 A). Figure 4 clearly illustrates the change in character of the impurity spectra between optimum and overbiased cases. At optimum bias settings and below, the plasma in the internal discharge acts as a 'virtual electrode'; the plasma is the source of the electron emission current extracted from the gun by the external bias. Beyond the optimum bias point the electron emission from the internal gun plasma saturates and all additional emission current comes directly from the metal electrodes in the gun (thus the large increase in the molybdenum lines as metal is sputtered from the surface). We speculate that the decrease in impurity line emission upon the application of bias is caused by the removal of the gun anode from the current return path. All of the electron current from the internal discharge is drawn out of the gun by the bias and returned through the vacuum vessel walls. This minimizes the interaction of the gun plasma with the anode surface and reduces impurity emission.

### 2.3. Time variation of specific impurity lines

Examination of the time variation of specific impurity lines with a monochromator and photomultiplier tube reveals why 'overbiasing' the gun beyond the plasma electron emission saturation point causes such a dramatic increase in impurity contamination of the gun plasma. Figure 5 shows Mo II line intensities for several gun discharges (Mo I and Mo II are similar), as recorded from the side view. The Mo II line intensity is very similar for the 'no bias' and 'optimum bias' discharges, but exhibits dramatically different behaviour during the 'overbiased' discharge. Upon termination of the internal gun current

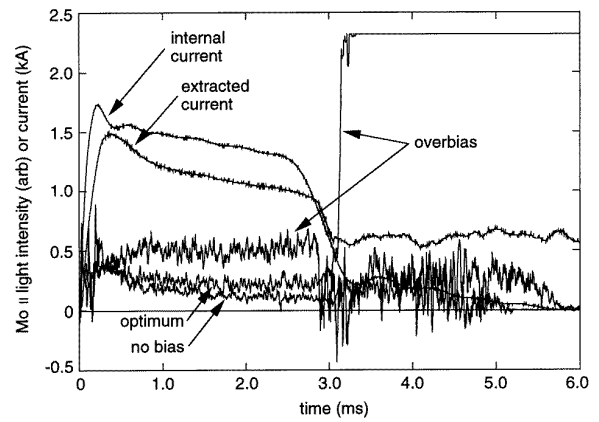


**Figure 3.** Comparison of impurity line emission intensity while varying internal gun discharge current and current extraction bias. (a) Side view and (b) end view.



**Figure 4.** Comparison of relative impurity levels with gun at optimum current extraction bias level (lower trace) and in 'overbias' condition (upper trace). These data are from the side view position.

(which occurs at 3 ms and is controlled by a pulse forming network), the Mo II line intensity immediately saturates the detector during the overbiased case. Substantial 'extracted current' also continues to flow beyond 3 ms. This does not occur in the unbiased or optimally biased cases. It seems reasonable to conclude that the extracted current that continues to flow after the internal gun discharge during



**Figure 5.** Comparison of the Mo II light intensity for no current extraction bias on the plasma gun, optimum bias, and overbias. Also shown is the internal gun discharge current and extracted current for the overbias case.

the overbiased case is being drawn directly from the molybdenum anode of the plasma gun, rather than from the virtual electrode formed by the plasma inside the gun. In fact, this situation clearly illustrates the advantage of a plasma virtual electrode over a real metal electrode. The virtual electrode provides clean emission of the high-density electron current, whereas surface erosion of a metal electrode contaminates the plasma and surrounding surfaces and eventually results in electrode failure when sufficient erosion has occurred.

As the current extraction bias is raised, it may be possible to sensitively determine the point at which overbias is about to occur by spectroscopically monitoring the metal contamination of the plasma flowing out of the gun. Note from figure 5 that the Mo II emission intensity during the gun discharge (0 to 3 ms) is larger for the overbias case than for optimum or no bias. This implies that some fraction of the extracted current is being drawn from the molybdenum anode even during the gun discharge, a situation that should be avoided since it leads to current extraction from the metal anode following the termination of the internal gun discharge. We have found that this problem can be somewhat mitigated by active control of the current extraction bias. If this bias is turned off immediately after the termination of the internal gun current, then large amounts of current will not be extracted from the molybdenum gun anode. However, such control does not prevent extraction of the fraction of the current that is drawn from the molybdenum anode during the gun discharge; this can only be remedied by lowering the extraction bias to the optimum level.

## 2.4. Copper electrode gun

The molybdenum electrodes and washers in the plasma gun were replaced with copper equivalents for a short operational period. Copper differs from molybdenum in two important ways: (1) the thermal conductivity is higher and (2) the surface sputtering threshold is lower. Higher thermal conductivity of the electrodes and washers is advantageous in that the surface heat load generated by

operation of the internal gun discharge can be drawn away and dissipated more effectively. However, an increase in erosion of the surface inevitably leads to increased impurity contamination of the plasma. Copper lines were observed to be bright in both the end and side views. Especially striking was the high relative brightness of these lines viewed from the side compared to the end. This is unlike the behaviour of the molybdenum lines shown in figure 2, where the Mo I and Mo II lines are so weak in the side view that they are almost invisible. It is likely that the electron current drawn to the copper gun anode erodes significantly more metal than the same current drawn to a molybdenum anode. These copper ions are not trapped in the same way that metal ions are trapped at the cathode, but instead flow out of the gun with the plasma stream.

Visual inspection of the copper electrodes and washers following a total of approximately 100 3 ms gun pulses revealed that substantial erosion of all plasma facing surfaces had taken place. The same inspection of molybdenum components after thousands of gun pulses reveals minimal surface damage and erosion. We have concluded that the gun electrodes and washers must be made from a refractory metal with a low sputtering coefficient.

### 3. Emission line broadening

#### 3.1. Thermal and Stark broadening

The two emission line broadening mechanisms useful for diagnosing the gun plasma are thermal and Stark broadening [4]. Thermal broadening arises from the Doppler shift caused by thermal particle motion, a Maxwellian velocity distribution giving rise to a Gaussian line profile. The B III line at 206.578 nm was used to measure ion temperature in the gun plasma because it is intrinsically bright, the line profile is dominated by thermal broadening, and it is isolated from any interfering spectral lines. In addition, the B III ions, since they were the highest ionization state of boron observed, should be reflective of conditions in the highest-temperature region of the gun plasma. The impurity ion–majority ion equilibration time for this high-density ( $10^{20} \text{ m}^{-3}$ ), low-temperature (10 eV) plasma is short, approximately 1  $\mu\text{s}$ . Even given what must be a very short particle confinement time in the gun plasma, the impurity ions should be near equilibrium to the majority species.

Stark broadening, also called pressure or collisional broadening, arises from the influence (via electric field) of nearby ions upon the emitting atom. Most useful for our purpose of estimating ion density of the gun plasma was a measurement of the Stark broadening of  $\text{H}\beta$  at 486.133 nm because the FWHM of this profile is approximately equal to  $0.040n_i^{2/3}$ , with ion density in  $10^{20} \text{ m}^{-3}$  [4]. Although not strictly correct, for our purposes we approximated the Stark broadening of this line with a Lorentzian profile. Since the effects of thermal and Stark broadening are combined and neither was dominant for the  $\text{H}\beta$  line emitted from the gun plasma, the data were fitted with a Voigt function. Furthermore, the  $\text{H}\beta$  emission is not spatially localized but

originates from the entire volume of the gun plasma and thus represents some sort of spatial average.

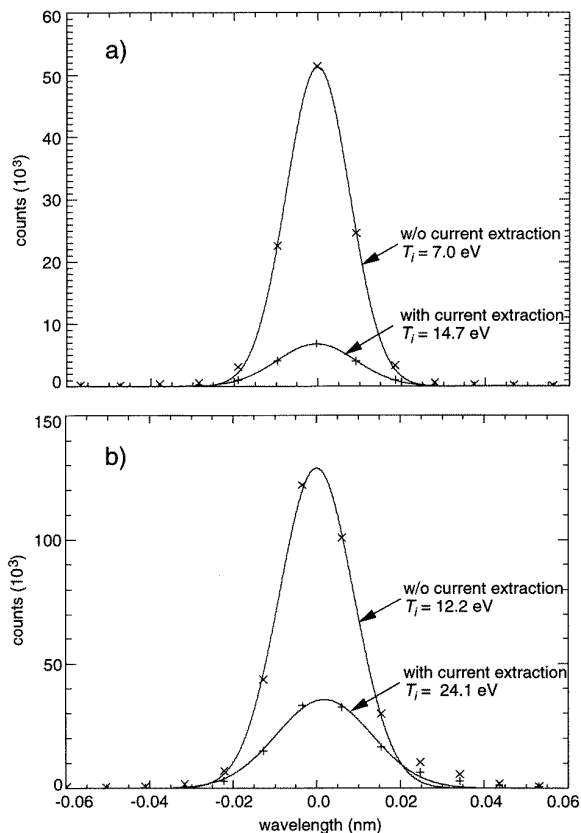
Two operational details should be mentioned. First, the B III line brightness was enhanced for this dataset by fitting the gun with a boron nitride spacer washer with a smaller inner diameter (similar to the inner diameter of the molybdenum washers, see figure 1(a)). This did not appear to affect operation of the gun. Second, the B III line profile was recorded by the spectrometer during the latter part of the gun discharge, skipping the first millisecond in order to avoid recording behaviour during the period in which the boron was reaching ionization equilibrium. The  $\text{H}\beta$  line emission was recorded over the entire 3 ms gun discharge, but care was taken to stop collecting light at the end of the discharge since the afterglow plasma that persisted 3 ms after the gun discharge produced a substantial amount of  $\text{H}\beta$  light.

#### 3.2. Ion temperature and density measurements

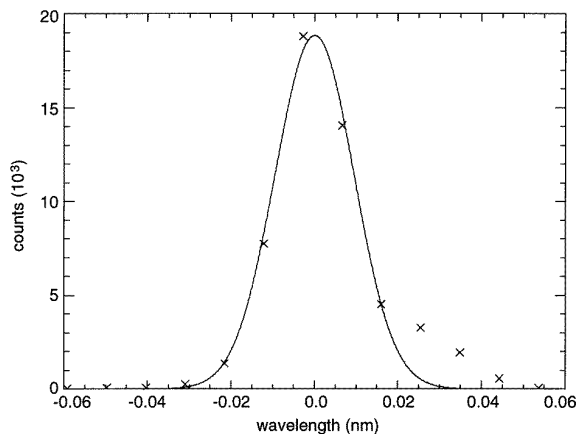
Two important observations summarize the B III measurements: upon application of the current extraction bias, the ion temperature doubles and the line brightness drops significantly. The current extraction bias seems to both heat the gun plasma and reduce the impurity level. The results from measurements of the broadening of the  $\text{H}\beta$  are not as easy to summarize. In general, the ion density is quite high, in the range of  $10^{20} \text{ m}^{-3}$ . This is in agreement with probe measurements of the plasma density [2]. The hydrogen neutral atom temperature is in the order of 1 eV.

Specific ion temperature measurements are illustrated in figure 6. The end view shows  $T_{i\parallel}$  going from 12 to 24 eV upon the application of bias, while  $T_{i\perp}$  from the side view goes from 7 to 15 eV. In both cases the line brightness recorded by the spectrometer dropped significantly upon the application of bias. This decrease does not appear to be simply an ionization to B IV, as the characteristic lines of that ion did not appear upon the application of bias. Some of the difference may be due to density changes in the plasma, but most likely it is simply a further indication that current extraction bias reduces the impurity content of the gun plasma. Note that in figure 6 (and in figures 7 and 8) the ordinate is in photoelectron counts; thus each data point shown on the graph represents the actual number of counts recorded in a spectrometer channel. Since the number of counts recorded in each channel is large, the statistical (Poisson) error bars are smaller than the cross used to mark the data points. Resulting absolute uncertainties for the ion temperature measurements are less than a few tenths of an eV.

Note that there appears to be a statistically significant enhancement of the light level in the spectrometer channels on the long-wavelength side of the end view Doppler broadened spectrum (figure 6(b)). This feature was always present and was most apparent in spectra taken in an ‘offset’ end view (figure 7) in which the spectrometer was in the end view location, but was shifted very slightly so as not to view the plasma in the internal gun discharge. We speculate that this may be a manifestation of directed flow of ions (with about 10 eV of kinetic energy) away from the spectrometer



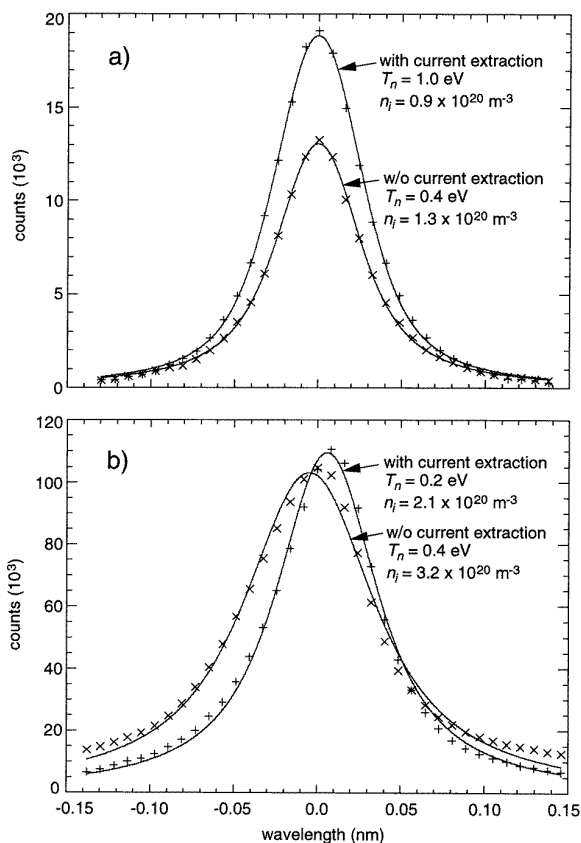
**Figure 6.** Comparison of the Doppler broadened profiles of the 206.58 nm B III line from the (a) side view and (b) end view.



**Figure 7.** The Doppler broadened profile of the 206.58 nm B III line from the 'offset' end view,  $T_i = 14.7$  eV. Note the clear presence of a non-Maxwellian feature on the long-wavelength side of the spectrum.

and toward the gun, but we do not understand the source or the mechanism.

Figure 8 presents specific measurements of the  $H\beta$  line profile. In the side view of the plasma from the gun,  $T_{n\perp}$  increases and  $n_i$  decreases when bias is applied. Both fits in figure 8(a) are quite good, so the values of ion density quoted on the figure are probably accurate to within a factor of two. In the end view, both  $T_{n\parallel}$  and  $n_i$  appear to decrease



**Figure 8.** Comparison of the Voigt profiles of the  $H\beta$  line from the (a) side and (b) end views, with and without current extraction bias.

when bias is applied to extract current, although changes are small and the fits are not as good, possibly because this view involves substantial spatial averaging.

#### 4. Summary

The major impurities in the gun plasma are boron, carbon, and nitrogen from the boron nitride washers and molybdenum from the metal electrodes. The plasma flowing out of the gun is largely uncontaminated by molybdenum; it appears to be trapped in the internal discharge. Typical plasma parameters are ion temperatures of 10 to 20 eV with ion densities in the order of  $10^{20} \text{ m}^{-3}$ . When a voltage bias is applied between the gun and some distant current collector, the plasma inside the gun acts as a virtual electrode, sourcing up to 1 kA of electron current. Application of current extraction bias removes the gun anode from the current return path and has two positive effects: the ion temperature doubles and the line brightness drops, indicative of ohmic heating and reduced impurity content.

#### Acknowledgments

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## References

- [1] Dimov G I and Roslyakov G V 1974 *Sov. Pribori. Tech. Exp.* **1** 29
- [2] Fiksel G, Almagri A F, Craig D, Iida M, Prager S C and Sarff S C 1996 *Plasma Sources Sci. Technol.* **5** 78
- Fiksel G 1991 *PhD Thesis* University of Wisconsin—Madison
- [3] Den Hartog D J and Holly D J 1997 *Rev. Sci. Instrum.* **68** 1036
- [4] Hutchinson I H 1987 *Principles of Plasma Diagnostics* (Cambridge: Cambridge University Press) pp 216–22