



# Volume and surface effects in Cs-free regimes in NIO1



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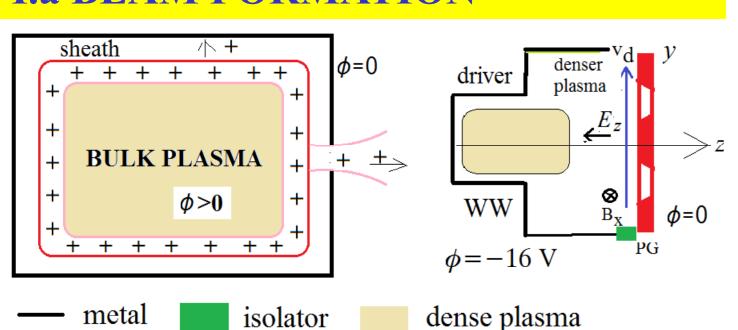
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Abstract: NIO1 (Negative Ion Optimization phase 1) is a compact multiaperture radiofrequency H- ion source whose design was optimized for sustainable prolonged beam on target (BOT) operation; installation economy implied a drastic scaling as respect to fusion device Dsources. The latter, in a consistent view for energy production request a beam on tokamak (BOT) span of 20 years, that is 6 108 s. Even if Cesium improves H- production as well known, also Cs-free regimes (and intermediate regimes) well deserve some development effort, in view of avoiding long term contamination of the accelerator and for use as cleaning procedure. Data collected by NIO1 in a true Cs-free regime (before 2020) are thus very important, and need a through statistical analysis, with special attention to the technique of gas conditioning that was discovered in NIO1 and to the issues concerned with long term operation. Gas conditioning macroscopically proves the importance of surface effects, even when the final production of H- happens in the source volume. As regards to the Electron Cyclotron Resonance Ion Sources (ECRIS), exchange of ideas and concepts, such as 'electron starvation' and biased disks, liners and wall coatings, is discussed. H- ion sources differ in terms of surface to volume ratio (over 10<sup>2</sup> m<sup>-1</sup> in matrix ion sources), practically achievable BOT (from 10<sup>4</sup> s to 10<sup>6</sup> s per year) and working frequency (from GHz in the ECRIS case down to 1 MHz) is reviewed. Gas mixing, conditioning and surface material perspectives are envisioned.

#### I. INTRODUCTION

Negative ions (eg. H<sup>-</sup>) production and extraction include much more complex systems ([12]) than positive ions do. Fusion applications add the major challenges: large current feasible only with multiaperture source; 2) CW operation time longer than hours; source life beam on target time BOT  $= 6 10^8 s (20 years)$ 

#### I.a BEAM FORMATION



As well known [Tonks, 1929] a positive ion plasma (say in a metal box) develops a positive ambipolar potential (normal sheath), which propels ions to walls. Piercing a hole in the box is enough for extraction. On the contrary, H- beams need: 1) formation of a so called inverted sheath near plasma grid (PG), vith one V<sub>b</sub> or more bias voltages [see Variale, Cavenago, poster 21]; 2) a magnetic filter field Bf.

#### I.b H<sup>-</sup> production

H<sub>2</sub> dissociation and H+ production (by impact ionization) needs over 15 eV electrons, that is electron temperature  $T_e > 4$  eV (driver region). This plasma will destroy H-, so a colder plasma (extraction region) near PG is obtained with filter Bf to store H-; filter Bf indeed

**Dissociative** attachment (DA) volume production:

e (cold) + 
$$H2* \rightarrow H- + H$$

H2\* = vibrationally excited molecule, also produced by >15 eV electrons. Charge exchange on cesiated wall:

H (fast) + Cs + e (wall) 
$$\rightarrow$$
 H- + Cs + wall

In Cs-based regimes, wall importance is obvious. In NIO1 true Cs-free regimes were studied until 2019; still we find wall effects, mainly gas conditioning.

Some symbols: p<sub>s</sub> source pressure; p<sub>2</sub> vessel pressure, P<sub>k</sub> forward rf power, I<sub>pg</sub> current in plasma grid PG electrode, j<sub>i</sub> H<sup>-</sup> current density, j<sub>e</sub> e<sup>-</sup> current density

# II. SET-UP and development

NIO1 has a modular setup, and achieved Hbeam runs longer than 10<sup>4</sup> s, trying to open worldwide a new window in fusion researches on source transient and stability effect. Achieved BOT  $< 10^6$  s/year limited by support.

#### II.a Overview

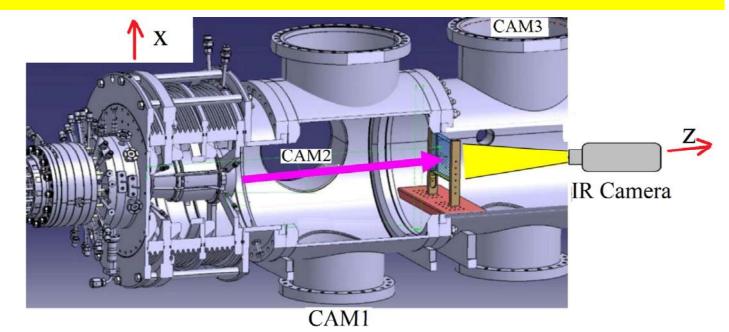
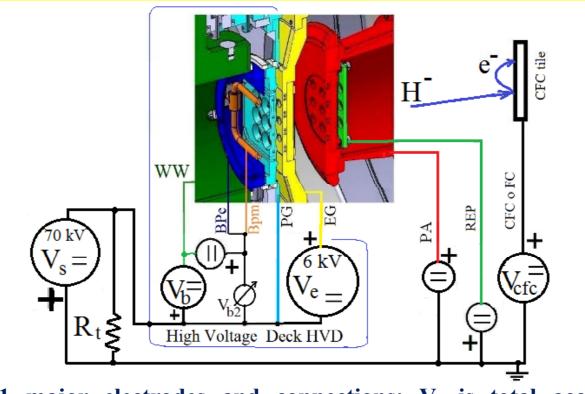


Figure 2. Overall 3D cut view, showing part of NIO1 accelerating electrodes; note CAM1, CAM2 and CAM3 placement; CFC tile recently moved after CAM3 position

# II.b Main connections



Nio1 major electrodes and connections; V<sub>s</sub> is total acceleration voltage,  $V_e$  the extraction voltage,  $V_h$  the main bias,  $V_{efc} = CFC$ polarization voltage (absolute values). Corrected ion current is  $I_a = I_s - (V_s/R_t)$ power supply current  $I_s$ 

### **II.c** Transients

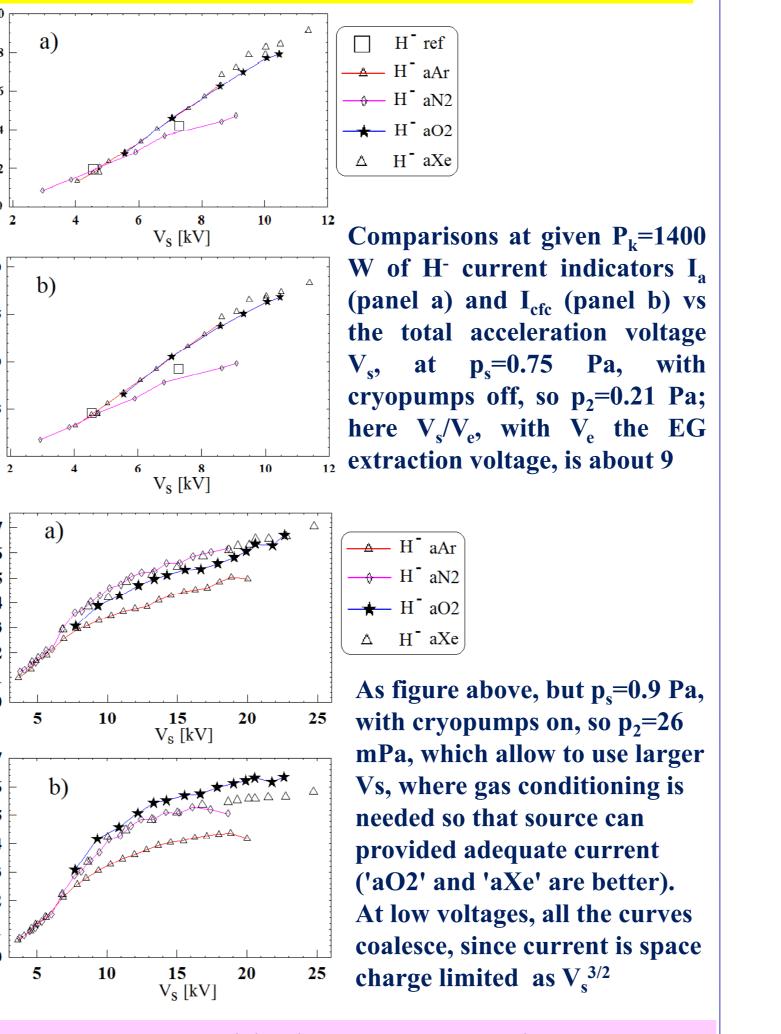
**Transients (seen from** <u>-</u> 10\*I<sub>cfc</sub>, case (a) 2017) are a spontaneous change of extracted ion current without major power supplies change. As a first remedies to get back to good current we develop: 1) improved rf window air cooling; 2) change of p<sub>s</sub>. So we observed anti-correlation

Fig 4: Anti-correlation of  $I_{cfc}$  and  $V_{pmt}$  during transients, for several fixed conditions:(a) at fixed control parameter as  $P_k=1.2$ kW,  $I_{ng}\approx 0$ ,  $p_s=0.75$  Pa, beam voltage  $V_s=4$  kV, extraction voltage  $V_p=0.5$  kV, compare [5]; (b):  $P_k=1.3$  kW,  $p_s=0.9$  Pa, filter current  $_{ng}$ =400 A with configuration 'f3' [13];  $V_s$ =4 kV and  $V_e$ =0.45 kV for better stability). Curve 'a' multiplied by 10; typical error 5 %

# III. Systematic experiments

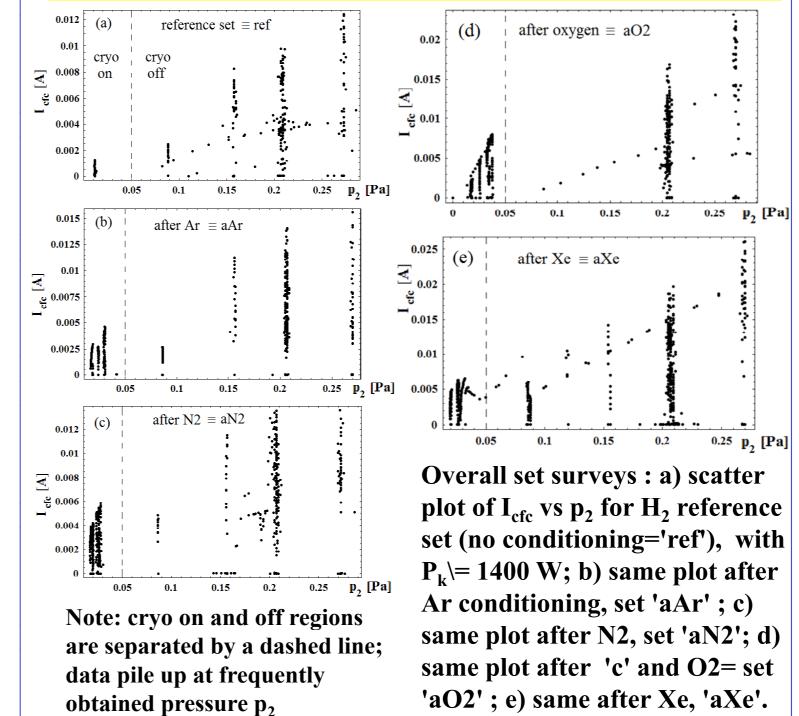
NIO1 support gas was O<sub>2</sub> or air and then H<sub>2</sub> To suppress them, at last we try gas conditioning: one day O2, then H2 again for a week, repeating. Gas conditioning with O<sub>2</sub> was very successful[13]; here we add results with selected other gases, as N<sub>2</sub> (suggested for comparison to air), Ar (used in many rf sources), Xe (mass similar to Cs). For busy schedule, Kr and Ne were unfortunately omitted. To cancel previous conditioning, we run source for several days, result for H- of last two days are the set 'ref'. Then we run with Ar for one day; results for H- in following two operation days are set 'aAr' (after Ar in brief). We repeated with N2 calling the H- result the set 'aN2'. Similarly for 'aO2' and 'aXe'.

## III.a Comparison at selected pressures



All tested gas conditionings show some improvement

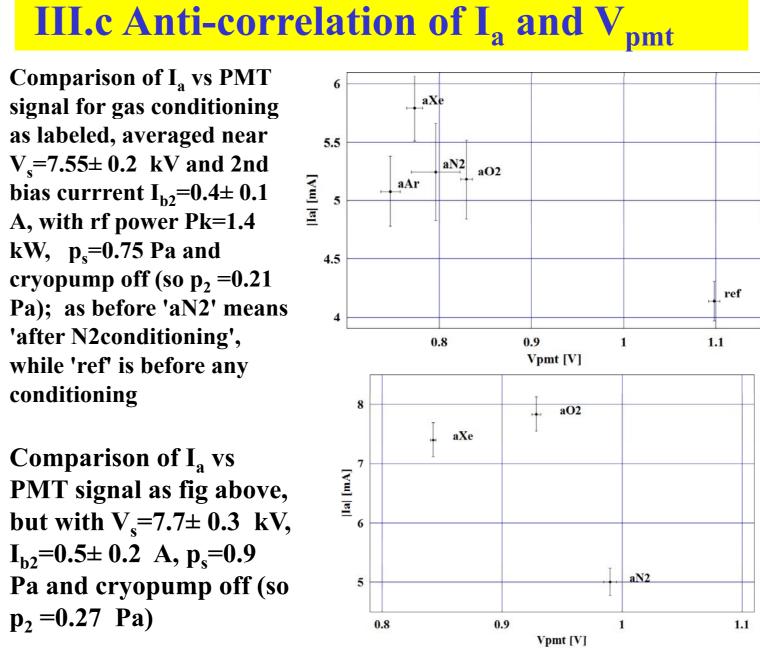
(2015). In 2017 we see undesired transients.



isolators soft iron

— Mo liners —— currents

III.b Overall comparisons



 $I_a$  vs  $V_{pmt}$  anti-correlation is an outstanding feature (explained in talk 7).

Efficacy of gas conditioning is a robust proof that surface effect matters also in Cs- free regimes

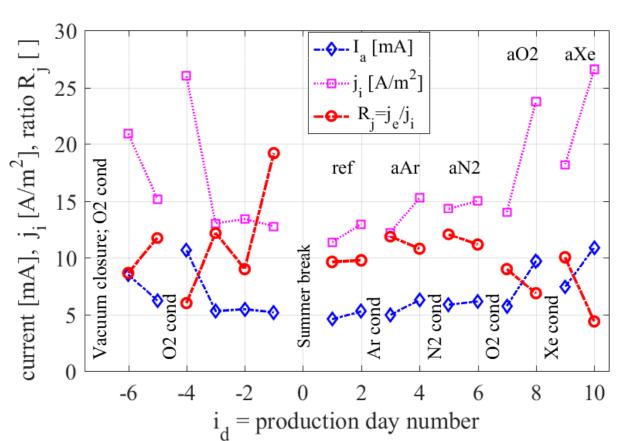


Figure 1: a) View of

plate (BP), plasma grid

permanent magnet (PM)

installation.

extensive

Summary of typical daily results for I<sub>a</sub>, j<sub>i</sub> and R<sub>i</sub> vs index i<sub>d</sub>, which enumerates the H- production days, eg. set 'ref' is  $i_d$ = 1,2, set after argon 'aAr' is [3,4], set 'aXe' id= 9,10; days  $i_d$ <0 conditioning ('cond') days; typical 2\σ error is 10 %.

## III.d Beams (see also poster P30)

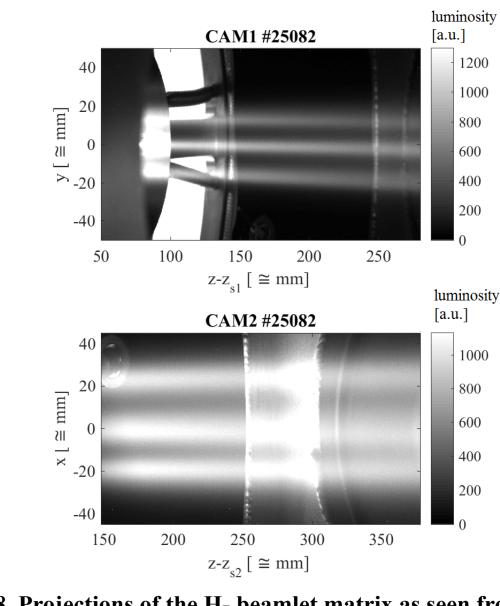
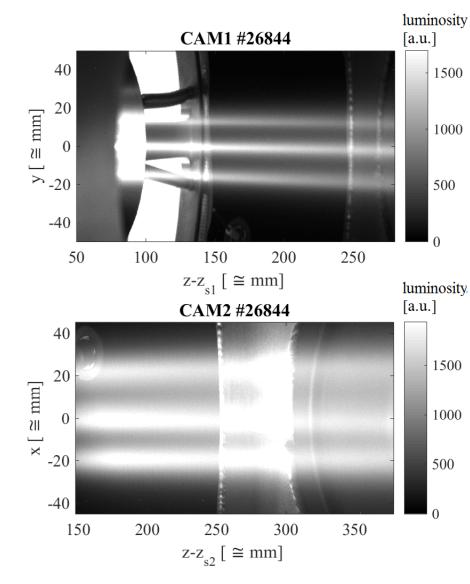


Figure 8. Projections of the H- beamlet matrix as seen from two lateral cameras; one example after Nitrogen conditioning 'aN2', with  $V_s=5.4$  kV (limiting  $I_a$  to 2.5 mA),  $V_s/V_e\approx 10$ ,  $p_s=0.9$  Pa, and  $I_{n\sigma}$ =400 A; labels show camera name and dataset index dsn.



As Figure 8, but with  $I_a=4.1$  mA, requiring  $V_e=0.75$  kV and  $V_s=8.1$ kV for reasonable optics; an example after Xenon conditioning 'aXe'

#### V. Conclusion and perspectives

The H<sup>-</sup> production can be improved in Cs-free regime, in a repeated way, with a gas conditioning technique, which also cancels the undesired transient fluctuation noted in previous works. Some fluid model is well in progress [see talk ID 7 this conf.]; analogies with gas mixing and other effects in ECRIS[7 19, 20] are noteworthy. In perspective, gas conditioning with Ar or Xe can be helpful in Cs-based regimes, or after Cs-based operation, both for cleaning[11] and for lowering the extraction plasma temperature  $T_e^a$ ; and it may be a valuable integrative method for H- ion sources.

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