

Boron erosion on a tungsten substrate using the TOMAS ICWC antenna

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1 Introduction

1.1 Boronization

The interior of a fusion reactor is extremely hostile with rapid temperature and pressure swings, as such a refractory material needs to be chosen for plasma facing components (PFC), tungsten is increasingly favoured over other materials due to its unique combination of properties such as low erosion rate, low tritium retention and resistance to heat-induced stress [1][2]. But there are two big drawbacks to using tungsten as a PFC: As it has a very high Z-value, if it does get sputtered, high amounts of radiation loss may occur due to brehmstrahlung [3] and second it's a very bad oxygen getterer which is the most deleterious of all impurities encountered in a fusion device. We can get the "best of both worlds" by coating our tungsten with a small (<100nm) layer of a low Z-material. Previously Beryllium was considered the main candidate but due to it's high toxicity and difficulty of handling the now most favored candidate is boron. Boronization (the deposition of a thin film of boron) has a proven track record of causing better confinement times and ELM (Edge Localized Mode) control [4][5][6][7]. There is however, at the time of writing, no direct research to the rapidity of Boron erosion under wall conditioning techniques such as ICWC and the outgassing efficiency of such a technique, that gap is what this paper attempts to fill.

1.2 Classification of PFMs

The properties of interest for PFMs may be quantisized as follows:

Performance measure	Physical Quantity	Definition
Thermal conductivity and strength	Thermal merit [8]	$\Delta_{th} = \frac{K\sigma_y}{\alpha\mathbb{E}(1-\nu)}$
ELM resilience	Disruption merit [8]	$\Delta_d = \frac{\sigma_u(C_p\rho K)^{1/2}}{\alpha\mathbb{E}}$
Impurity capturing/gettering	Gettering merit	$\Delta_{get} = \sum_i^{oxygen,\dots} \int_E T_i(E)\mathcal{F}_i(E)dE$
Tritium capturing/retention	Gettering anti-merit	$\bar{\Delta}_T = \int_E T_T(E)\mathcal{F}_T(E)dE$
Erosion probability and impact on the plasma	Impurity anti-merit	$\bar{\Delta}_{imp} = Z_{PFM}^2 \sum_j^{species} \int_E Y_j(E)\mathcal{F}_j(E)dE$

Where K is the thermal conductivity, σ_y is the yield strength, α the thermal expansion coefficient, \mathbb{E} Young's modulus, ν Poisson's ratio, $T_i(E)$ the chance of trapping particle species i incoming at energy E , $Y_i(E)$ the yield of particle species i incoming at energy E and $\mathcal{F}_i(E)$ the flux (incoming particles/s)

of particle species i at energy E . The last quantity was derived from the sputtering rate/erosion rate of PFM particles due to species i (outgoing particles/second):

$$\mathcal{S}_i = \int_E Y_i(E) \mathcal{F}_i(E) dE \quad (1)$$

and the Bremsstrahlung relation

$$\mathcal{P}[W] = \frac{Z_{PFM}^2}{C} n_{PFM} n_e \sqrt{T_{eV}} \quad (2)$$

I.e:

$$\mathcal{P}[W] = \left[Z_{PFM}^2 \sum_i \int_E Y_i(E) \mathcal{F}_i(E) dE \right] \cdot t \cdot \frac{n_e \sqrt{T_e}}{C} \quad (3)$$

$$\triangleq \bar{\Delta}_{imp} \frac{n_e \sqrt{T_{eV}}}{C} t \quad (4)$$

With t the time (as the amount of impurities grows with time, so does the radiated power/second), grouping the material-specific parameters as $\bar{\Delta}_{imp}$ captures the PFM impurity property (note that this is still plasma-dependent and thus machine-dependent). Merits should be maximized whilst anti-merits should be minimized. As tungsten is the go-to material for future fusion reactors, it may be used to norm the merits to the *reduced* versions for easier comparison:

$$\mathfrak{d}^{PFM} := \frac{\Delta^{PFM}}{\Delta^W} \quad \text{and} \quad \mathfrak{d}^{PFM} := \frac{\bar{\Delta}^W}{\bar{\Delta}^{PFM}} \quad (5)$$

These normed merits (henceforth called *normits*) scale differently and each have a different implementation on confinement (i.e different weight) as such a function of the form

$$\mathfrak{d}_{tot} := f(\mathfrak{d}_{th}, \mathfrak{d}_d, \mathfrak{d}_{get}, \mathfrak{d}_T, \mathfrak{d}_{imp}) \quad (6)$$

is not straightforward to deduce, instead, for now, the focus will be on the individual normits. These normits allow ease of comparison between different PFMs and straightforward extrapolation between material test facilities (such as TOMAS) and fusion test setups/fusion reactors (such as ITER and DEMO). This paper has two goals, one is to determine the boronized tungsten impurity normit \mathfrak{d}_{imp}^{B-W} experimentally, this will be done in the TOMAS reactor, the other is to extrapolate boron erosion rates to ITER-like reactors as to estimate the time needed inbetween boron coatings. The TOMAS reactor is capable of IC, EC, combined IC+EC and glow discharge, all for different gas species (H, He, D, Ar) and pressures. As \mathfrak{d}_{imp} is plasma-specific, the conducted experiments will try to find relations of the form $\mathfrak{d}(\text{species}, \text{power})$ for IC, EC and IC+EC. Here the power relations should give the possibility to extrapolate to different machines.

1.3 Experimental setup

One may analyse boron-tungsten erosion by using neutral and ion beams, mimicking actual IC, EC and IC+EC heated fusion devices, the obvious advantage being that the experiment is simpler with easily chosen and controlled parameters. But the disadvantage is that co-operative effects, which may occur only when a plasma is present, will not be observed [9]. The TOMAS machine [10] will be used as it is capable of forming a toroidal plasma with pure ICRH discharge, thus introducing these co-operative effects. It is equipped with a monopole ICWC antenna which will be set to operate at 50MHz, closely mimicking the 53MHz that will likely be used at ITER. The mask containing the samples (see figure 1) is mounted on a movable arm which we'll position on top of the vessel. We'll modify the pressure in such a way that the neutrals pressure (i.e the observed pressure during the discharge) is 10^{-5} mbar, closely mimicking what was the case in JET and will likely be the case in ITER [11]. The exposures will be deuterium at different power levels (1.5kW, 3.5kW and 5.5kW), the thickness of the boron layer will be tracked by using ellipsometry before and after on all samples and IBA+SEM (creating a hole using IBA and scanning thickness using SEM) post mortem and on one control sample.

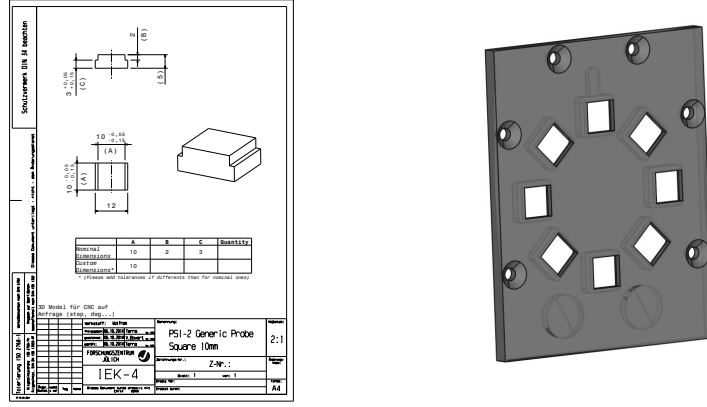


Figure 1: The mask (right figure) is made out of tungsten and can contain 8 samples (left figure), the 10x10mm top of the samples are coated with boron and attached in the mask such that it is facing towards the plasma in the vessel

2 Erosion estimation using neutrals

2.1 Explanation of the theory

The erosion rates should be easy to measure, as such it should be short enough to leave some boron intact but not too short as to have a statistically significant reduction in thickness. To this end an estimation of the sputtering rate will be made prior to the discharge and the time under plasma will be chosen accordingly. As the TOMAS machine's RFEA (Retarding Field Energy Analyzer) is at the moment of writing unable to measure the ion distribution, the only usable proxy to estimate the ideal time under load is the high energy neutrals measured using the ToF-NPA (Time of Flight Neutral Particle Analyzer) [?] with some factor to account for the ions.

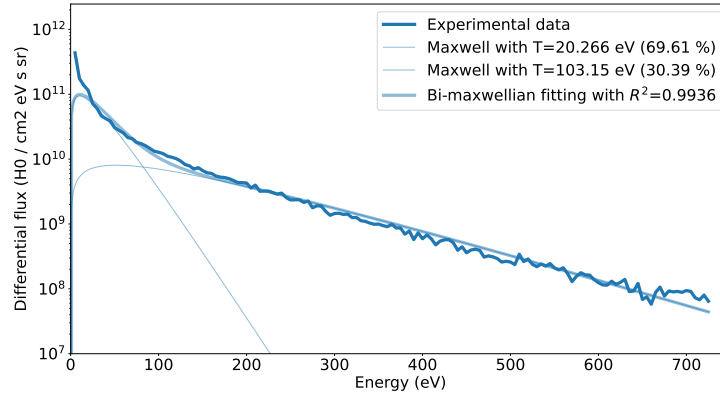


Figure 2: Our assumption behind the bi-maxwellian distribution is based on the characteristic that IC heating causes such a distribution in ions [12] combined with the knowledge of rate coefficients for atomic charge exchange which is dominant at the observed kinetic energies [9], i.e the bi-maxwellian ions "transfer" their energies to the neutrals causing the same bi-maxwellian to be visible in the neutral energies. Note that the R^2 parameter of the fitting on the graph is defined with the logarithmic difference to better emphasize the shape.

An example measurement made with the NPA is shown in figure 2. As the data is expressed as differential flux, it lends itself to be easily manipulated into other quantities. One such manipulation is using the following formula to transform it into an erosion rate (m/s):

$$S = \frac{2\pi}{N} \int_E Y(E) \partial \mathcal{F}(E) dE \quad (7)$$

With $Y(E)$ the yield of the neutrals on the material (i.e how many atoms of the material get sputtered for every incoming neutral), $\partial \mathcal{F}(E)$ the experimental differential flux as previously shown and N the number density. The validity of this formula is quite straightforward to see as:

- The NPA covers a certain solid angle, this has been accounted for as can be seen in the unit of the example experiment (/sr meaning per steradian), we can get an approximation for the average flux in the vessel by assuming homogeneity and thus multiplying this data by 2π steradians.
- The differential flux of particles in an energy bin surrounding E causes sputtering, to get this sputtering rate we multiply by the yield Y at that energy as it's defined to be the outgoing atoms per incoming, we integrate over all energies to get the full contribution.
- The number density of the target N dictates how the amount of outgoing atoms relates to the decrease in thickness, this can be seen from the units.

In the analysis software equation 7 is used in the discrete form, summing over the energy bins:

$$S = \frac{2\pi}{N} \sum_{E_0}^{E_{\max}} Y(E) \partial \mathcal{F}(E) \Delta E \quad (8)$$

Where we may get the yield $Y(E)$ using the software RustBCA[13], assuming ions to behave the same as neutrals (as the samples aren't charged, this should give a negligible discrepancy) with parameters from Wolfgang Eckstein's book [14], assuming perpendicular impingement¹.

2.2 Calculated estimates

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¹note that assuming a distribution such as $\cos(\theta)^2$ doesn't change the result by more than 10%

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