Fuel retention and carbon deposition on beryllium marker tiles from JET tokamak main chamber limiters investigated by ion beam analysis

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Abstract

JET tokamak with the ITER-like wall is operated with arrays of castellated beryllium limiters in the main chamber. In several locations Be marker tiles were installed for erosion-deposition studies. The castellation sides and the plasma-facing surfaces (PFS) of Be marker tiles from three different locations of the JET main chamber, from the experimental campaigns 2011-12 (ILW-1) and 2013-14 (ILW-2), were analysed employing ²H and ³He micro-beams in order to determine carbon impurity deposition and deuterium retention. The deposited carbon and deuterium amount on the castellation sides (up to 1.5 mm deep into the groove) was assessed with respect to the ion/electron drift direction. Both the carbon and deuterium amount on the investigated castellation sides either stays constant or reduces with depth from the edge of the PFS. No systematic difference is observed in the carbon deposition or deuterium retention on the different castellation sides of each sample with respect to the ion/electron drift direction. Carbon and deuterium content is found to be lower on the PFS than on surfaces in the gaps of castellation for the majority of the samples. The carbon amount is, in general, higher than the deuterium one. No systematic correlation between the carbon and the deuterium amount has been observed.

Keywords: JET tokamak, beryllium, carbon deposition, deuterium retention, ion beam analysis

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1. Introduction

The choice of appropriate plasma-facing materials (PFMs) is an issue of great importance as the plasma-wall interaction (PWI) affects both the lifetime of wall materials and plasma performance. Beryllium (Be) and tungsten (W) are the materials for plasma facing components (PFC) in ITER [1]. Beryllium has been chosen due to its low atomic number which limits plasma dilution and energy radiation losses. This metal has high thermal conductivity (~200 Wm⁻¹K⁻¹), low fuel retention in comparison to carbon (C) which reacts chemically with H isotopes [2] and it is an efficient oxygen getter thus reducing oxygen impurities and helps to keep the effective atomic number, Z_{eff}, in the vessel at low levels. A detailed overview of beryllium as PFC is presented in [3].

Until 2009 the Joint European Torus (JET) at the Culham Science Centre, the largest tokamak in the world, was operated with carbon as the main PFM (JET-C) [4]. Very high fuel inventories were measured because the presence of carbon is decisive for fuel retention by co-deposition [5]. This called for a large-scale test of a metal wall. Since 2011 JET has metallic PFC, called the ITER-like wall (JET-ILW) [6]. Beryllium is in the main chamber (castellated limiters and Be coatings on the inner wall cladding) and tungsten in the divertor: the load bearing plate in the base made of bulk metal, while W-coated carbon fiber composites tiles are in other locations. Three experimental campaigns were performed in 2011-2016 with deuterium fuelling: 2011-12 (ILW-1), 2013-14 (ILW-2) and 2015-16 (ILW-3) with input energy of 150, 201 and 245 GJ, respectively. The overview has been presented in [7], while detailed works have dealt with erosion, material deposition and deuterium (D) retention on the surface of the divertor [8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18] and on the main chamber [19, 20, 21, 22, 23, 24, 25, 26]. In addition, material deposition and fuel retention on surfaces located in the gaps of the castellated Be limiters have been examined [27, 28, 29]. It has also been consistently shown that fuel inventory with ILW was reduced by one order of magnitude in comparison to JET-C [30, 31]. The main source of carbon in JET was eliminated, but still the knowledge of the carbon content in co-deposits and the carbon-deuterium correlation is crucial for the detailed assessment of factors influencing the retention in JET-ILW and, by this, for improved predictions for ITER. Two are the sources of carbon after the JET transformation to a fully metallic device, a) the carbon residuals from the previous wall and b) the carbon fiber composite (CFC) tiles coated by tungsten in the divertor [32].

The relation between the co-deposition and the deuterium retention in JET-ILW had been investigated in detail on W-coated divertor tiles. Works [10, 12, 14, 24] report that the retention increases with the increase of the material (Be and C) deposition. The works [13, 15, 16, 17] claim that the increase of the D retention is due to Be deposition, while [8, 18] report that the C deposition increases the D retention. The C-D correlation in Be has also been investigated, on laboratory-prepared samples. Anderl et al. found the retention in pure Be to be lower than in C-coated beryllium [33]. Also Guseva et al [34] concluded that carbon impurities on the Be surface enhance the D retention. On the other hand, C. Porosnicu et al. irradiated different beryllium-carbon relative concentration with deuterium ions and found that lower carbon concentration retained higher deuterium content [35]. Thus, it is not clear from the literature whether the residual deuterium retention still exists as a result of the C-D chemistry or whether D is integrated into deposits irrespective of C.

Accelerator-based ion beam analysis (IBA) is the most efficient set of methods in surface studies of wall materials [36]. Among them, a ³He-based nuclear reaction analysis (NRA) has been the most frequently used for the simultaneous determination of D, Be, C in carbon-wall machines. In the case of JET-ILW, the C quantification on Be surfaces is not possible with the $^{12}\text{C}(^3\text{He},p_0)^{14}\text{N}$ reaction due to its superposition with the $^9\text{Be}(^3\text{He},p_3)^{11}\text{B}$ one, as detailed in [36]. The remaining options are in: (i) proton scattering via $^{12}\text{C}(p,p)^{12}\text{C}$ [37, 38] with the sensitivity at the level of $1x10^{17}$ ^{12}C cm $^{-2}$, (ii) heavy ion elastic recoil detection analysis (HIERDA) with high sensitivity (below $1x10^{15}$ ^{12}C cm $^{-2}$) but the information depth limited to less than 1 µm, (iii) NRA using a ^2H beam via $^{12}\text{C}(^2\text{H},p)^{13}\text{C}$. The latter method (with sensitivity of $1x10^{15}$ ^{12}C cm $^{-2}$) was applied for carbon studies in this paper.

A study of the carbon deposition on plasma-facing surfaces (PFS) and inside the castellation grooves of JET-ILW limiters was performed earlier with a standard ²H milli-beam [29]. In the current work we investigate whether carbon deposition on the castellation sides is affected by the orientation of the castellation side with respect to the ion or electron drift direction. Moreover, a ²H micro-beam has been applied to examine the morphology of carbon deposition. Deuterium retention is investigated by the use of a ³He micro-beam on both the PFS and the castellation sides. Whether carbon or deuterium are co-deposited with beryllium or not cannot be verified with the present study. The overall aim is to quantify carbon and deuterium on the castellation and plasma-facing surfaces of the JET-ILW beryllium marker tiles and investigate if carbon plays a role in the retention of deuterium in the deposits.

2. Materials and Experimental Details

Samples from different marker tiles of the main chamber and after the first and the second experimental campaigns were investigated: one sample from the upper Dump Plate (DP, 2B(C)2), two from the mid-plane of Outer Poloidal Limiter (OPL, 4D14) and four from the Inner Wall Guard Limiter (IWGL, 2XR10) (Figure 1). These tiles are castellated in order to reduce the eddy currents and thermal stresses [39], and therefore, the castellation sides are free to interact with the plasma. Moreover, the samples from the marker tiles have a nickel interlayer between the top beryllium layer and the bulk beryllium [40]. After the cut, one of the castellation sides was marked for reference reasons. The configuration of the tiles and the samples as well as the labelling of the castellation sides based on the ion/electron drift direction of ILW1 IWGL outer (27) sample, as an example, are presented schematically in Figure 2.

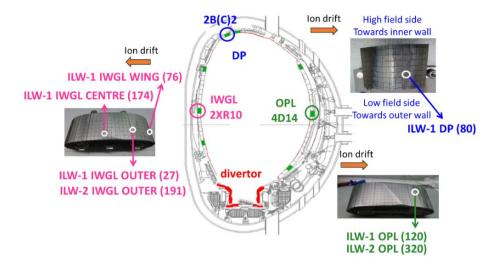


Figure 1: Position of the investigated Tiles and samples inside the main chamber of the JET tokamak.

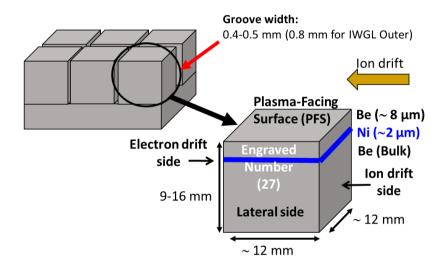


Figure 2: Schematic of the castellation configuration of the beryllium tiles and the configuration of the ILW1 IWGL outer (27) sample. The castellation sides are labelled based on the ion/electron drift direction.

The 2 H micro- and milli- beam measurements were performed using the 5.5 MV TN11 HV Tandem Accelerator at NCSR "Demokritos", in Athens, Greece. The beam energy was 1.35 MeV and a silicon surface barrier (SSB) detector with depletion depth of 1000 μ m was placed at an angle of 170° with respect to the beam axis. A kapton foil of 12.5 μ m was positioned in front of the detector in order to separate the peak 12 C(d,p₀) 13 C used for the carbon quantification from the peaks of alpha particles emitted via 9 Be(d,a₀) 7 Li and 9 Be(d,a₁) 7 Li reactions. The chamber was kept under vacuum (10 $^{-6}$ mbar). The beam spot of the microbeam had a diameter smaller than 100 μ m and the current was around 100 pA. The mapping

area was 1.5 x 1.5 mm² and the resolution 64 x 64 pixels. The data acquisition and the mapping was performed using the OMDAQ2007 software and appropriate hardware [41].

The ³He measurements were carried out at Ruder Boskovic Institute, in Zagreb, Croatia. The 3 4 ³He beam was accelerated by the 6 MV tandem Van de Graaff accelerator and 1.0 MV Tandetron accelerator. The beam energies varied between 2 and 3 MeV, and the mapping 5 areas were either $1 \times 1 \text{ mm}^2$ or $300 \times 300 \text{ }\mu\text{m}^2$. A Partially Depleted Silicon Surface Barrier 6 detector (PDSSB) with depletion depth of 2000 µm, with nominal active area of 300 mm² 7 collimated to 230 mm² was used and placed at an angle of 135° ± 19° with respect to the 8 9 beam axis. The distance between the target and the detector was approximately 2.5 cm, 10 which corresponds to a solid angle of 0.462 sr. Additionally, a Mylar foil of 120 μm thickness 11 was placed before the detector in order to absorb the alpha particles produced via 9 Be(3 He,a) 8 Be reactions and to detect only the 9 Be(3 He,p) 11 B and 2 H(3 He,p $_{0}$) 3 He peaks. A 12 chopper was used in order to estimate the collection charge of the measurements. The data 13 acquisition of the ³He beam was performed using the in-house developed software package 14 SPECTOR [42] and the hardware based on Xilinx Virtex 6 FPGAs; for more technical details 15 16 see [43].

- 17 It is noted that the investigated area from the castellation sides is at the entrance of the gap, 18 ranging from 0.3 to 1.5 mm into the gap, since according to M. Rubel et al. [28] the majority 19 of the deuterium retention is restricted in this area.
- The quantitative analysis of all the NRA spectra was performed with the SIMNRA software [44]. For the ³He micro-beam NRA measurements and the deuterium quantification, the V. Kh. Alimov et al [45] cross section data for the ²H(³He,p₀)⁴He reaction and the N. P. Barradas et al [46] one for the ⁹Be(³He,p_{0,1})¹¹B reactions were used. For the ²H beam NRA measurements and the carbon quantification, the evaluated cross section data from SigmaCalc archive [47] and P. Tsavalas et al [48] for the ¹²C(²H,p₀)¹³C and ⁹Be(²H,p₀)¹⁰Be reactions, respectively, were used.

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3. Results and discussion

3.1 Carbon deposition

The carbon content was measured on ion and/or electron drift sides, as well as on some lateral castellation sides, employing a 2 H beam. Figure 3 presents a representative NRA spectrum together with the simulated spectrum from the lateral side of the sample 27 (ILW1 IWGL outer) using 2 H micro-beam. In this figure, the energy range with the peaks corresponding to the nuclear reactions 12 C(2 H,p₀) 13 C and 9 Be(2 H,p₀) 10 Be is shown. The quantitative results of the determined carbon and deuterium amounts from a) the PFS and b) the castellation sides up to a maximum depth of 1.5 mm from the edge of the PFS are presented in Figure 4. The absence of C or D content for some castellation sides (C content on the ion drift side of ILW1 IWGL outer (27) and D content on the ion drift side ILW2 OPL (320), the electron drift side of ILW1 DP (80), and the lateral sides of ILW1 IWGL outer (27), ILW2 IWGL outer (191) and ILW1 centre (174)) is due to the lack of experimental data. The deposited carbon amount on the PFS is from our previous study [29]. It is observed that the orientation of the castellation side does not, in general, affect significantly carbon deposition. Only for sample 27 from the ILW1 IWGL outer, almost one order of magnitude higher carbon amount ((59±4) × 10^{17} at/cm 2) is found for the lateral side compared to the

electron drift one. On all the other castellation sides carbon deposition ranges from (3.2 \pm 0.2) \times 10¹⁷ at/cm² to (19 \pm 1) \times 10¹⁷ at/cm², with the ILW1 Dump Plate (80) presenting systematically the higher carbon deposition on its castellation sides and the IWGL 2XR10 Centre (174) the lowest. A trend of higher carbon amount on the castellation sides than that on the PFS is observed.

In Figure 5 the carbon mapping of the lateral side of all samples is depicted, with the exception of the ILW1 IWGL centre (174) sample, for which the ion drift side is presented. The PFS of the samples is at the top of the mapping and it is defined by a white line. On the ILW1 DP (80), some carbon agglomerates with diameter of about 150 μ m have been formed over the whole side. On the ILW1 OPL (120), we observe a slight decrease of carbon with the depth. On the ILW2 OPL (320), carbon agglomerates with diameter in the range 100-200 μ m are observed near the PFS of the sample. On ILW1 IWGL outer (27), at a depth of about 800 μ m from the PFS, a stripe rich in carbon, having a width of about 500 μ m, has been formed. On ILW2 IWGL outer (191), the amount of carbon decreases as a function of depth. On the ILW1 IWGL centre (174), 400 μ m from the PFS, there is a thin stripe, having a width of about 200 μ m, depleted of carbon. On the IWL1 IWGL wing (76), a drastic decrease of the carbon content with depth is observed.

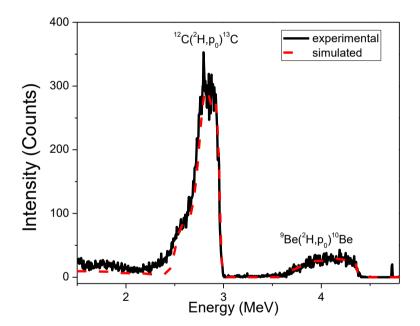


Figure 3: The measured (solid black line) and simulation (dash red line) NRA spectrum of ILW-1 IWGL Outer (27) castellation side using a ²H micro-beam.

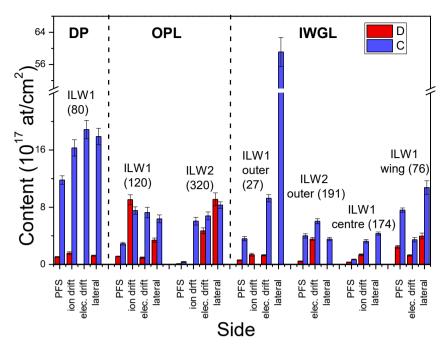
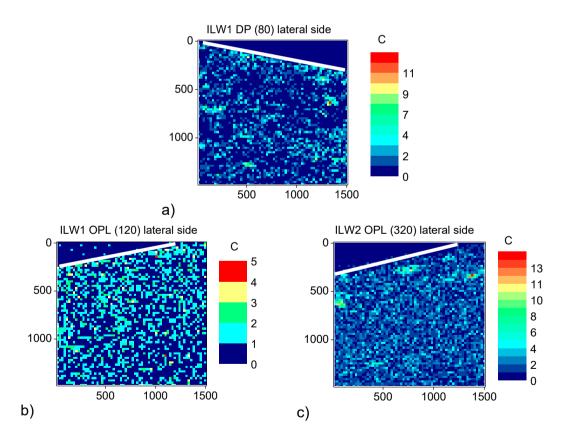


Figure 4: Deuterium and carbon content of the plasma-facing surface (PFS) and the castellation sides as measured by ³He and ²H beams, respectively.



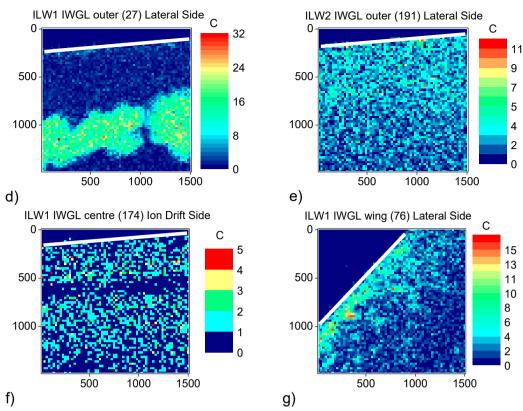


Figure 5: Mapping of the deposited carbon on the lateral side (a, b, c, d, e, g) and the ion drift side of the ILW1 IWGL centre (174) (f). The white line defines the edge of the plasmafacing surface. The unit of the axes is μ m.

3.2 Deuterium retention

The PFS and at least one castellation side of all the samples were measured using a ³He micro-beam. Figure 6 depicts representative experimental and simulated spectra of the ion drift side from the ILW1 Dump Plate (80) employing a ³He micro-beam. The determined deuterium content using a ³He beam is presented in Figure 4.

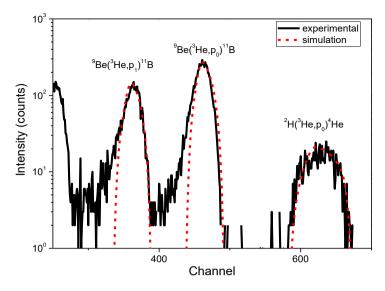


Figure 6: The experimental (black solid line) and the simulated (red dashed line) NRA spectra of the ion drift side of sample 80 from the ILW1 Dump Plate.

The deuterium content on the PFS of the samples is found to vary more than one order of magnitude, ranging from $(0.090\pm0.003)\times10^{17}$ at/cm² (sample 320 from ILW2 OPL) to $(2.5\pm0.2)\times10^{17}$ at/cm² (sample 76 from IWGL 2XR10 wing). The deuterium amount determined in the current work on the PFS of ILW1 DP ((1.05 ±0.05) \times 10¹⁷ at/cm²) is in reasonable agreement with that reported in [24] ((3.4 ±1.2) \times 10¹⁷ at/cm²). Additionally, integrating the mean D content (1.2 \times 10¹⁷ at cm²) of the different areas over the whole ILW1 IWGL tile, we observe that the total D content, 3.51 \times 10¹⁹ at, is half of the corresponding value (6.76 \times 10¹⁹ at) reported in [24].

On the castellation sides, the variation of the fuel retention between the various locations is reduced with the deuterium content ranging between $(0.96\pm0.10)\times10^{17}$ at/cm² (ILW1 OPL (120) electron drift side) and $(9.1\pm0.9)\times10^{17}$ at/cm² (ILW1 OPL (120) ion drift side and ILW2 OPL (320) lateral side). In general the castellation sides retain higher amounts of deuterium than the PFS.

From the first to the second campaign, the deuterium content on the PFS decreased; for the OPL from $(1.13\pm0.04) \times 10^{17}$ at/cm² to $(0.090\pm0.003) \times 10^{17}$ at/cm² and for IWGL outer from $(0.61\pm0.04) \times 10^{17}$ at/cm² to $(0.45\pm0.05) \times 10^{17}$ at/cm². On the contrary, deuterium amount on the castellation sides during ILW2 increases with respect to ILW1; for OPL from $(0.95\pm0.10) \times 10^{17}$ at/cm² to $(4.7\pm0.4) \times 10^{17}$ at/cm² (electron drift side) and from $(3.4\pm0.3) \times 10^{17}$ at/cm² to $(9.1\pm0.9) \times 10^{17}$ at/cm² (lateral side); and for IWGL outer from $(1.4\pm0.2) \times 10^{17}$ at/cm² to $(3.8\pm0.2) \times 10^{17}$ at/cm² (electron drift side).

Subsequently, the mean values of the deuterium content on the castellation sides are compared with those reported in [25]. There is agreement that the ILW1 DP castellation sides present the lowest deuterium retention with $(1.4\pm0.3)\times10^{17}$ at/cm² found in the current work and $<10^{17}$ at/cm² reported in [25]. For the ILW1 OPL castellation side, the value found in the current work $((4.5\pm2.4)\times10^{17}$ at/cm²) is very close with that reported in [25] $(\sim6\times10^{17}$ at/cm²). For the ILW1 IWGL, we find lower deuterium amount $((1.8\pm0.5)\times10^{17}$ at/cm²) than the low limit of the range reported in [25] $((7-20)\times10^{17}$ at/cm²).

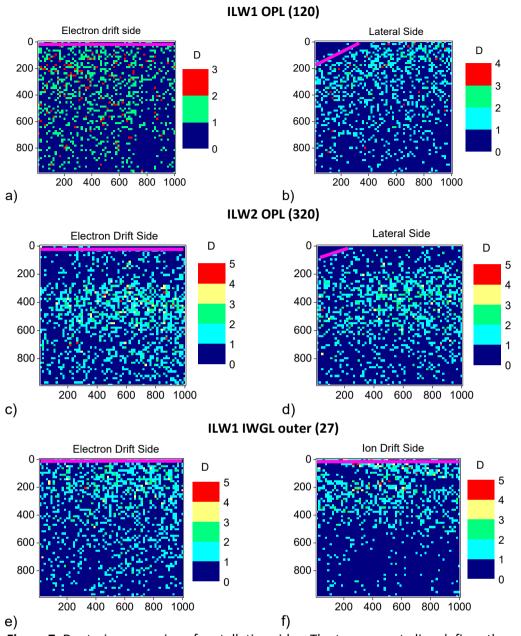


Figure 7: Deuterium mapping of castellation sides. The top magenta line defines the edge of the plasma-facing surface. The unit of the axes is μm .

Figure 7 depicts deuterium mappings of some of the castellation sides for IWL1 OPL (120), ILW2 OPL (320) and ILW1 IWGL outer (27), as determined with the 3 He micro-beam. The deuterium distribution on the PFS is homogeneous for all samples and therefore deuterium mappings of the PFS are not presented. On the other hand, deuterium is reduced with depth on the castellation sides of all samples apart from the ILW2 OPL (320) (Figure 7c and d) where a deuterium stripe of about 400 μ m width, 200 μ m from the PFS edge, is observed for both castellation sides. The deuterium distribution is similar on the castellation sides of ILW1 OPL (120) (Figure 7a and b). On the ion drift side of the IWL1 IWGL outer (27) (Figure 7f) the deuterium is reduced with depth more abruptly than on electron drift one (Figure 7e).

3.3 Deuterium retention versus carbon deposition

In this section we discuss possible correlation between deuterium retention and carbon deposition. From Figure 4, we conclude that high carbon amount is not necessarily accompanied by high deuterium content.

In Figure 8 the deuterium over carbon ratio (D/C) is presented for the PFS and the castellation sides. For the castellation sides the average content of C and D has been used. The D/C ratio ranges from 0.08 to 1.17 with DP presenting the smallest ratio (<0.1) and the castellation sides of OPL after the second campaign the highest one (\sim 1). Similar D/C ratios for the PFS and the castellation sides are observed during ILW1 campaign, whereas during ILW2 campaign the ratio is larger on the castellation sides compared to that of the PFS, being in the range of 3.8 - 6.6.

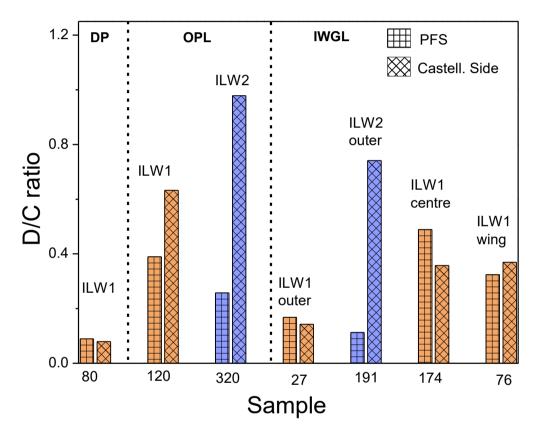
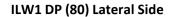
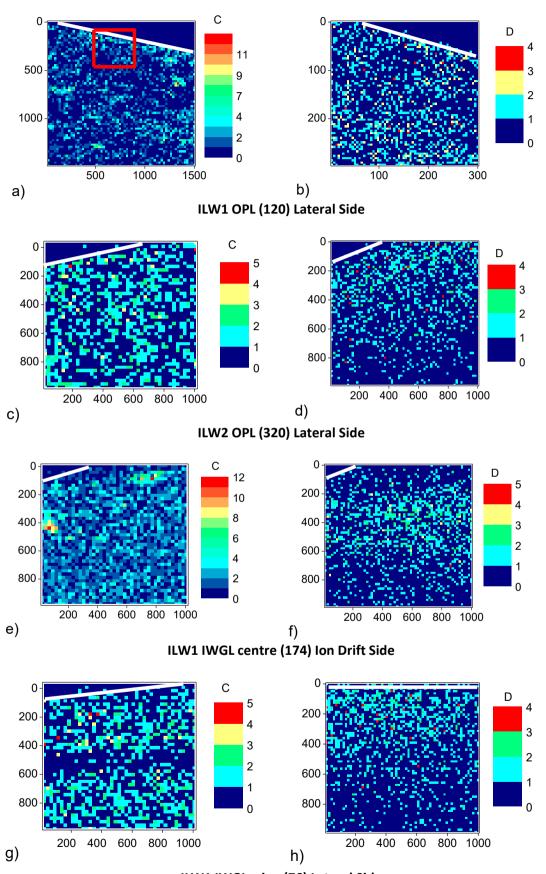


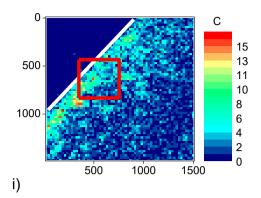
Figure 8: The deuterium over carbon (D/C) ratio of the plasma-facing surface (PFS) and the castellation. For the castellation the average of the measured castellation sides has been used.

Next we discuss the carbon and deuterium mappings from the same castellation side. Figure 9 depicts representative carbon and deuterium mappings of three castellation sides. The white line defines the edge of the plasma-facing surface. In Figure 9j the edge of the castellation side is not well defined, because the castellation side was not perfectly aligned with respect to the ion beam so signal from the PFS was also detected.





ILW1 IWGL wing (76) Lateral Side



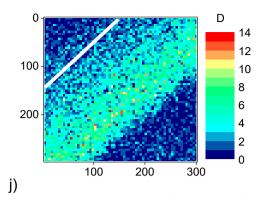


Figure 9: Carbon and deuterium mapping of the same castellation side of ILW1 DP (80) (a and b), ILW1 OPL (120) (c and d), ILW2 OPL (320) (e and f), ILW1 IWGL centre (174) (g and h) and ILW1 IWGL wing (76) (i and j). The area of deuterium mapping for b) and f) corresponds to the red square of the corresponding carbon mapping. The white line defines the edge of the plasma-facing surface. The unit of the axes is μm.

On ILW1 DP (80) lateral side (Figure 9a and b), carbon and deuterium have similar homogeneous distributions all over the mapped area. On the ILW1 OPL (120) lateral side the carbon distribution is nearly homogeneous (Figure 9c) while the deuterium decreases with depth (Figure 9d). On the ILW2 OPL (320) lateral side the stripe rich in deuterium (Figure 9f) is not observed on the carbon mapping (Figure 9e). On the ILW1 IWGL centre (174) ion drift side, there is a zone depleted of carbon (Figure 9g), while the amount of deuterium decreases smoothly with depth (Figure 9h). On ILW1 IWGL wing (76) lateral side, a similar stripe with high amount of carbon and deuterium is detected near the PFS (Figure 9i and j).

4. Summary and Conclusions

The PFS and castellation sides of samples from different beryllium marker tiles of the main chamber and after different campaigns of the JET tokamak were investigated employing ion beam analysis using ²H and ³He beams in order to assess carbon deposition and deuterium retention, respectively, their spatial distribution, their content with respect to the ion/electron drift direction and the correlation between them.

The carbon maps show that, in general, the carbon amount on the investigated castellation sides either stays constant or reduces with depth from the edge of the PFS. No systematic difference is observed in the carbon deposition on the different castellation sides of each sample with respect to the ion/electron drift direction.

Concerning deuterium, the PFS of the majority of the samples has retained less amount than that retained on the castellation sides. From the first to the second campaign the deuterium amount of the PFS decreases, while on the castellation sides it increases. The deuterium distribution on the PFS is homogeneous while on the castellation sides it decreases with depth for the large majority of the samples. Additionally, the carbon amount is, in general, higher than the deuterium one. No systematic correlation between the carbon and the deuterium amount has been observed.

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