

GA-A25482

**¹³C-TRACER EXPERIMENTS IN DIII-D PRELIMINARY IN
THERMAL OXIDATION EXPERIMENTS TO UNDERSTAND
TRITIUM RECOVERY IN DIII-D, JET, C-MOD, AND MAST**

by

**P.C. STANGEBY, S.L. ALLEN, N. BEKRIS, N.H. BROOKS, K. CHRISTIE, C. CHROBAK,
J.P. COAD, G.F. COUNSELL, J.W. DAVIS, J.D. ELDER, M.E. FENSTERMACHER,
M. GROTH, A.A. HAASZ, J. LIKONEN, B. LIPSCHULTZ, A.G. McLEAN, V. PHILIPPS,
G.D. PORTER, D.L. RUDAKOV, J. SHEA, W.R. WAMPLER, J.G. WATKINS,
W.P. WEST, AND D.G. WHYTE**

JUNE 2006



DISCLAIMER

This report was prepared as an account of work sponsored by an agency of the United States Government. Neither the United States Government nor any agency thereof, nor any of their employees, makes any warranty, express or implied, or assumes any legal liability or responsibility for the accuracy, completeness, or usefulness of any information, apparatus, product, or process disclosed, or represents that its use would not infringe privately owned rights. Reference herein to any specific commercial product, process, or service by trade name, trademark, manufacturer, or otherwise, does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States Government or any agency thereof. The views and opinions of authors expressed herein do not necessarily state or reflect those of the United States Government or any agency thereof.

¹³C-TRACER EXPERIMENTS IN DIII-D PRELIMINARY IN THERMAL OXIDATION EXPERIMENTS TO UNDERSTAND TRITIUM RECOVERY IN DIII-D, JET, C-MOD, AND MAST

by

P.C. STANGEBY,* S.L. ALLEN,[†] N. BEKRIS,[‡] N.H. BROOKS, K. CHRISTIE,[¶] C. CHROBAK,[¶]
J.P. COAD,[§] G.F. COUNSELL,[#] J.W. DAVIS,* J.D. ELDER,* M.E. FENSTERMACHER,[†]
M. GROTH,[†] A.A. HAASZ,* J. LIKONEN,[△] B. LIPSCHULTZ,[∞] A.G. McLEAN,* V. PHILIPPS,[◇]
G.D. PORTER,[†] D.L. RUDAKOV, J. SHEA,[¶] W.R. WAMPLER,[♦] J.G. WATKINS,[♦]
W.P. WEST, AND D.G. WHYTE[¶]

This is a preprint of a paper to be presented at the 33rd EPS Conf. on Plasma Physics, June 19-23, 2006, Roma, Italy, and to be published in the *Proceedings*.

*University of Toronto Institute for Aerospace Studies, Toronto, Canada

[†]Lawrence Livermore National Laboratory, Livermore, California

[‡]Forschungszentrum Karlsruhe, Eggenstein-Leopoldshafen, Germany

[¶]University of Wisconsin-Madison, Madison, Wisconsin

[§]EFDA-JET, Culham Science Center, Abingdon, United Kingdom

[#]EURATOM/UKAEA, Culham Science Center, Abingdon, United Kingdom

[△]EURATOM-Tekes, Helsinki, Finland

[∞]Massachusetts Institute of Technology, Cambridge, Massachusetts

[◇]Forschungszentrum Jülich GmbH EURATOM-Association, Jülich, Germany

[○]University of California-San Diego, La Jolla, California.

[♦]Sandia National Laboratories, Albuquerque, New Mexico.

Work supported by
the U.S. Department of Energy
under W-7405-ENG-48, DE-FC02-01ER54698, DE-FG02-04ER54762,
DE-FG02-04ER54758, and DE-AC04-94AL85000

GENERAL ATOMICS PROJECT 30200
JUNE 2006



¹³C-Tracer Experiments in DIII-D Preliminary to Thermal Oxidation Experiments to Understand Tritium Recovery in DIII-D, JET, C-Mod, and MAST

P.C. Stangeby¹, S.L. Allen², N. Bekris³, N.H. Brooks⁴, K. Christie⁵, C. Chrobak⁵, J.P. Coad⁶, G.F. Counsell⁷, J.W. Davis¹, J.D. Elder¹, M.E. Fenstermacher², M. Groth², A.A. Haasz¹, J. Likonen⁸, B. Lipschultz⁹, A.G. McLean¹, V. Philipps¹⁰, G.D. Porter², D.L. Rudakov¹¹, J. Shea⁵, W.R. Wampler¹², J.G. Watkins¹², W.P. West⁴, and D.G. Whyte⁵

¹*University of Toronto Institute for Aerospace Studies, Toronto, Canada*

²*Lawrence Livermore National Laboratory, Livermore, California 94550, USA*

³*Forschungszentrum Karlsruhe, Eggenstein-Leopoldshafen, Germany*

⁴*General Atomics, P.O. Box 85608, San Diego, California 92186-5608, USA*

⁵*University of Wisconsin-Madison, Madison, Wisconsin, USA*

⁶*EFDA-JET, Culham Science Center, Abingdon, United Kingdom*

⁷*EURATOM/UKAEA, Culham Science Center, Abingdon, United Kingdom*

⁸*EURATOM-Tekes, Helsinki, Finland*

⁹*Massachusetts Institute of Technology, Cambridge, Massachusetts, USA*

¹⁰*Forschungszentrum Jülich GmbH EURATOM-Association, Jülich, Germany*

¹¹*University of California-San Diego, La Jolla, California, USA*

¹²*Sandia National Laboratories, Albuquerque, New Mexico, USA*

Retention of tritium in carbon co-deposits is a serious concern for ITER. Developing a reliable in-situ removal method of the co-deposited tritium would allow the use of carbon plasma-facing components which have proven reliable in high heat flux conditions and compatible with high performance plasmas. Thermal oxidation is a potential solution, capable of reaching even hidden locations [1]. It is necessary to establish the least severe conditions to achieve adequate tritium recovery, minimizing damage and reconditioning time. The first step in this multi-machine project is ¹³C-tracer experiments in DIII-D, JET, C-Mod and MAST. In DIII-D and JET, ¹³CH₄ has been (and in C-Mod and MAST, will be) injected toroidally symmetrically, facilitating quantification and interpretation of the results. Tiles have been removed, analyzed for ¹³C content and will next be evaluated in a thermal oxidation test facility in Toronto with regard to the ability of different severities of oxidation exposure to remove the different types of (known and measured) ¹³C co-deposit. Removal of D/T from B on Mo tiles from C-Mod will also be tested. OEDGE interpretive code analysis of the ¹³C deposition patterns is used to generate the understanding needed to apply findings to ITER. First results are reported here for the ¹³C injection experiments in DIII-D.

¹³CH₄ was puffed through the upper pumping plenum of DIII-D (pump off) into lower single-null (LSN), neutral beam heated (6.5 MW), high density, detached, ELMy H-mode discharges [2]. The puff was toroidally symmetric and at a rate which did not significantly perturb the local plasma conditions. The puff rate averaged 18.8 torr l/s lasting 2 s during each of 17 repeat discharges. Immediately after the experiment, DIII-D was vented and a

total of 77 tiles were removed. The ¹³C content of 48 of the tiles was measured using nuclear reaction analysis (NRA) and proton-induced gamma-ray emission (PIGE). 37% of the ¹³C was found on front faces of tiles in the inner divertor region and the private flux zone, PFZ, Fig. 1, and ~10% near the puff. Fig. 1 also shows results from an earlier, similar experiment which used low density L-mode discharges where in total half as much ¹³C was injected [3]. The most significant difference in the ¹³C deposition pattern was that in the detached H-mode experiment there was substantial deposition on the PFZ wall. The NRA measurements were made at Sandia National Laboratories using the ¹³C(³He,p)¹⁵N reaction with an analysis beam of 2.5 MeV ³He [3]. The ¹³C deposits were also measured on a sub-set of the tiles at the University of Madison-Wisconsin, using the more sensitive PIGE technique [4], Fig. 1. Protons have a resonant nuclear reaction with ¹³C at 1748 keV, very narrow in energy ~0.075 keV. The excited nucleus emits a 9.2 MeV gamma ray, which is detected by a scintillator. By scanning the initial beam energy a highly resolved depth profile of ¹³C can be measured. PIGE measures the concentration of ¹³C versus depth with a higher depth resolution than NRA and hence lower detection limit for ¹³C deposited on graphite of about 10¹⁵ atoms/cm², compared with ~2 × 10¹⁶ atoms/cm² for NRA.

In the PIGE analysis, the depth profile of the ¹³C count rate peaks at the surface, and has a Gaussian “tail” which levels off at a depth ~0.5 microns into the tile. The Gaussian shape is fit to the raw data using a nonlinear least-squares fitting routine. Three parameters are fit at each analysis point: the surface ¹³C concentration (i.e. count rate, shown in Fig. 1), the 1/e half width of the Gaussian (i.e. the depth of the ¹³C) and the offset count rate which represents the “deep” natural ¹³C. From these parameters, plotted in Fig. 2, one can deduce important aspects of the ¹³C deposition in the divertor. First, the isotope enrichment at the surface is typically a factor of ~5-20. Since the natural ¹³C abundance is 1.1%, this indicates that the surface still has >80% ¹²C deposition. This is significant as it indicates that the ¹³C injection did not appreciably perturb the intrinsic deposition process, i.e. that the experiment was genuinely a *tracer* experiment, providing information on the natural co-deposition process. Second, the typical depth scale of the ¹³C enrichment is ~0.1-0.2 microns, indicating that the intrinsic deposition rate was ~1-4 nm/s over the ~50-100 s of total injection time. The constant ¹³C count rate seen deep into the sample is a result of the intrinsic 1.1% ¹³C.

Almost half the injected ¹³C is accounted for on the front faces of 7 tile-rows in the divertor and 3 tile-rows near the injection point. Although the inner wall tiles have not been measured yet using PIGE, the detection of ~40% (with large uncertainties) of the injected ¹³C at that location in the earlier L-mode ¹³C experiment in DIII-D suggests that much of the remaining ¹³C in the current H-mode experiment may be accounted for on the surfaces of tiles. It is hypothesized that the ¹³C-deposits in the divertor were not significantly disturbed by ongoing plasma exposure, including ELMs, although ELM-related changes in divertor carbon emissions remain to be explained. Any disturbance, if present, certainly did not cause the sort of massive migration of co-deposits to remote regions that occurred in the JET DTE1 experiment [5] and it appears that the ¹³C that entered the divertor remained on the divertor tiles. The JET results were for an entire campaign that included a variety of discharge conditions and configurations, in contrast with the single (detached) condition used here. That the co-deposits seem to stay approximately where they are initially formed — at least

for detached divertor conditions – could have important implications for tritium retention, namely the absence of migration of the tritium to remote locations that may be less accessible for tritium recovery. Deposition in tile gaps remains to be measured. The results from the outer baffle ring tile No. 24, (poloidal position >310 cm) contrast sharply with the other divertor (floor) tiles: the surface ¹³C is that expected for natural carbon, indicating that enriched ¹³C fluence was probably never present to these surfaces, since deposition and re-erosion would still leave some signs of an equilibrium ¹³C layer near the surface, particularly at locations toward the outer walls. This implies that transport of the ¹³C occurred essentially entirely via the inner SOL, resulting in a more concentrated co-deposition pattern than might be expected if significant quantities of the ¹³C were transported along both inner and outer SOL, or entered the confined plasma.

The interpretive Onion-Skin-Model Eirene Divimp edge (OEDGE) code was able to approximately reproduce the measured ¹³C deposition pattern, by making two key assumptions: (a) the existence of a fast parallel flow along the SOL toward the inner divertor, (b) a radial pinch of 10-20 m/s (in +R-direction, acting in the inner SOL, above the X-point) [6]. The existence of neither of these transport features has been predicted by standard edge codes, but fast parallel flows toward the inside have been directly measured on a number of tokamaks, and Kirnev [7] assumed the existence of a similar radial pinch in his EDGE2D modeling of JET, in order to reproduce the fast parallel SOL flow, where the pinch was needed to close the particle recirculation loop. The detailed breakup kinetics of the ¹³CH₄ was included in the OEDGE modeling and showed that the ¹³C-ions are produced too far out in the SOL to be able to replicate the observed deposition; some form of radial pinch is needed to move the ¹³C-ions toward the separatrix for deposition to occur where measured. Figure 3 illustrates the effect of assuming different radial pinches in the OEDGE analysis. With no radial pinch, too much of the deposition occurs well to the inside of the inner strike point. A radial pinch >20 m/s, on the other hand, pushes too much of the deposition outward and onto the PFZ wall. A radial pinch of ~10 m/s approximately replicates the measured deposition pattern of the ¹³C. Figure 4 shows that both a parallel flow and a radial pinch are required in order to replicate anything at all similar to the deposition pattern measured. The OEDGE modeling showed that most of the ¹³C deposited as neutrals formed by volume recombination in the cold, dense detached plasma in the inner divertor and the PFZ.

ITER operation is based on divertor detachment to control peak heat loads. DIII-D ¹³C-tracer experiments in detached H-mode suggests that it might be possible to arrange for a significant amount of the tritium co-deposition to occur on the PFZ wall. This could be advantageous since that surface need not possess significant heat-removal capability and so could be engineered to permit its being heated to high temperatures. If hot enough, carbon deposition will occur without co-deposition trapping of tritium. Intermittent heating to release the tritium, with or without simultaneous thermal oxidation, could be achieved in various ways, including flash heating by mitigated disruptions, since this surface is in direct view of the main plasma. Deposition of carbon as neutrals will also create “soft” co-deposits which release H/D/T more easily than “hard” deposits formed by ionic deposition, and are more readily removed by thermal oxidation [1]. For detached divertor conditions, co-deposits may

stay where first formed — much of it on relatively accessible locations — and not experience migration of the tritium by erosion-redeposition to inaccessible regions.

This work was supported in part by the US Department of Energy under W-7405-ENG-48, DE-FC02-04ER54698, DE-FG03-96ER54373, DE-FG02-04ER54758, and DE-AC04-94AL85000 and by a Collaborative Research Opportunities Grant from the Natural Sciences and Engineering Research Council of Canada.

- ¹G. Federici, C.H. Skinner, J.N. Brooks, et al, Nucl. Fusion **41** (2001) 1967.
²W.R. Wampler, A.G. McLean, S.L. et al., 17th Int. Conf. on Plasma Surface Interactions, Hefei, China (2006).
³W.R. Wampler, S.L. Allen, A.G. McLean, et al., J. Nucl. Mater. **337-339** (2005) 134.
⁴W.G. Whyte, Proc. of 5th ITPA Divertor and Scrape-Off Layer Group Meeting, Lisbon (2004).
⁵M. Keilhacker, M.L. Watkins, JET Team, J. Nucl. Mater. **266-269** (1999) 1.
⁶J.D. Elder, A.G. McLean, P.C. Stangeby, et al., *ibid* Ref. 1.
⁷Kirnev, et al., J. Nucl. Mater. **337-339** (2005) 271.

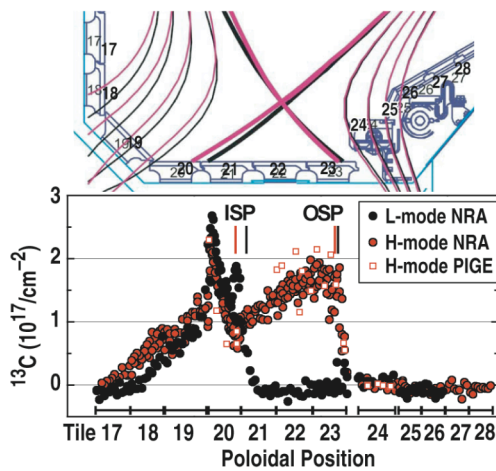


Fig 1. ¹³C measured in divertor by NRA and PIGE.

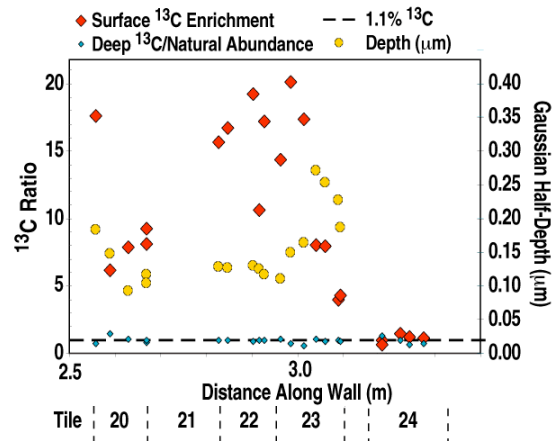


Fig. 2. Fitting parameters from PIGE analysis.

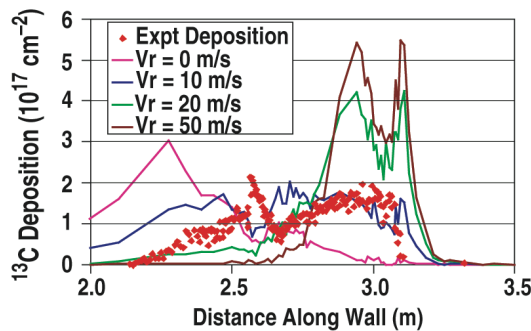


Fig. 3. OEDGE code analysis showing the influence of a radial pinch on the ¹³C deposition pattern. Fast parallel plasma flow along the inner SOL has also been included here, at M = 0.3 (Fig. 4). Modeling assumes time-invariant plasma conditions; see [4].

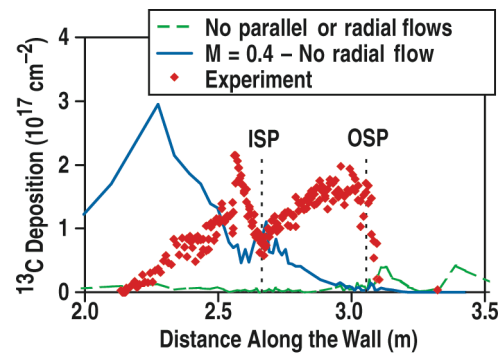


Fig. 4. OEDGE code analysis showing the dependence of deposition on the assumption of a radial pinch and a fast parallel flow in the inner SOL. For no parallel flow or radial pinch, the code-simulated deposition pattern is completely unlike that measured.