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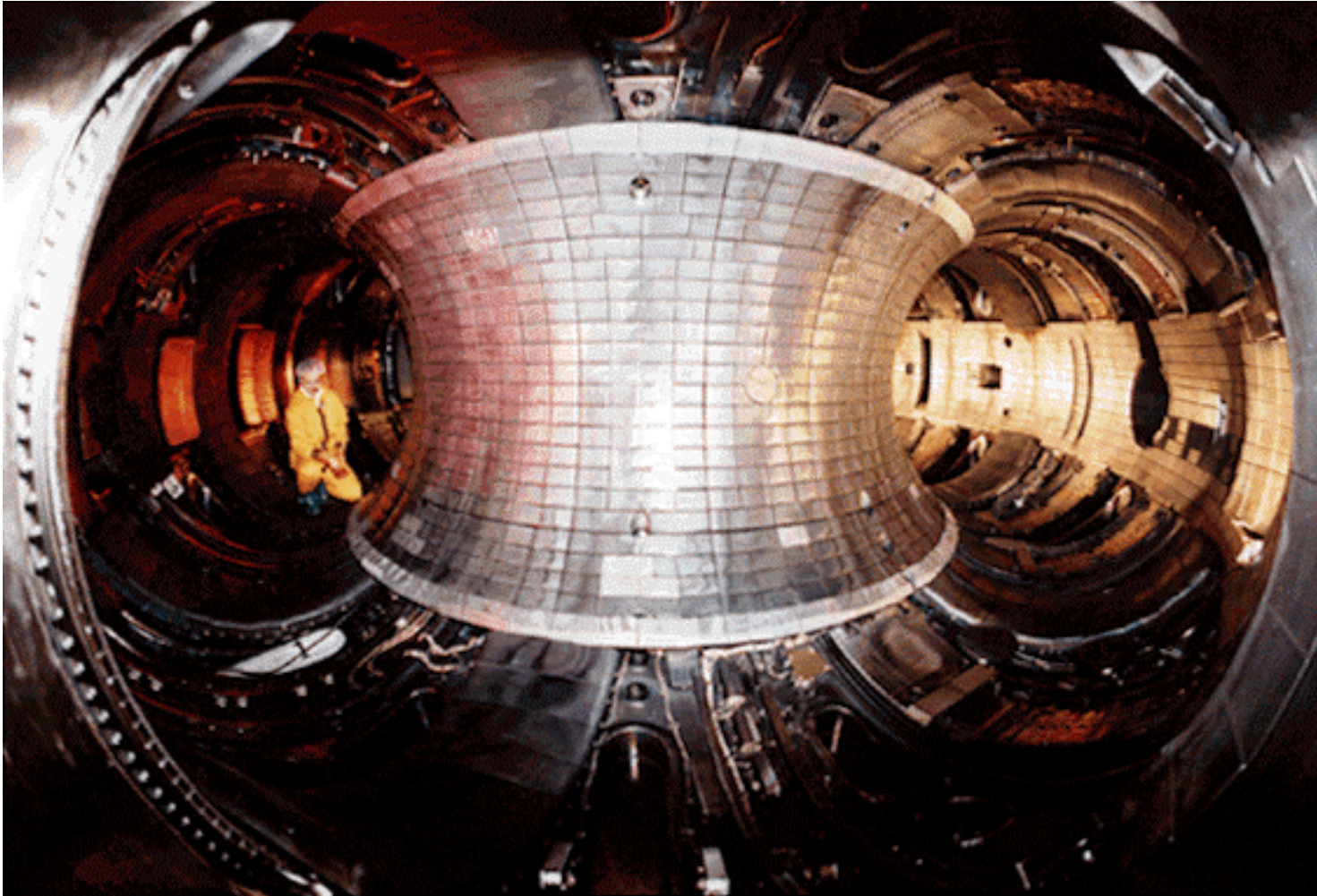
# *Tritium deposition patterns in TFTR*

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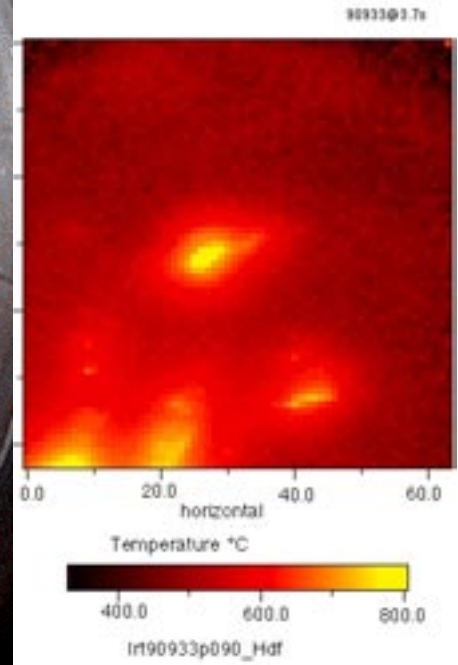
Presented by C. H. Skinner with key contributions from  
Charles Gentile, A Carpe, L Ciebiera, G Guttadora, S Langish (PPPL)  
John Hogan (ORNL)  
Mark Paffett, Robert Reiswig, Scott Willms, (LANL)  
Nicolas Bekris (FZK Karlsruhe), Paul Coad(JET),  
Tetsuo Tanabe, Kazuyoshi Sugiyama (Nagoya University).

*4th ITPA Meeting on SOL and Divertor Physics Topical Group  
13-16 January 2003, Naka, JAERI*





Limiter  
Temperature  
@ 28 MW NBI



- TFTR was a limiter machine - no divertor.
- Operated with tritium Nov '93 - April '97.
- Net deposition on VV walls (not erosion).
- Walls heated only by plasma (limiter hotspots reached  $\approx 800$  C).

Low density, high temperature edge

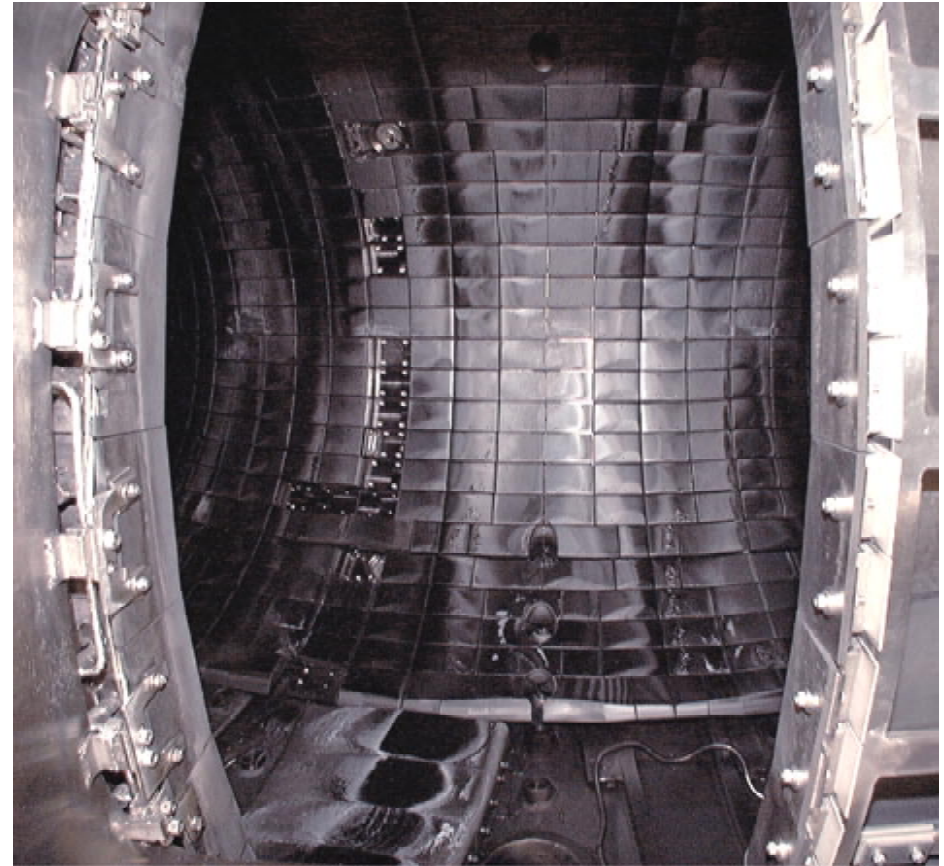
	TFTR SOL (TRANSP/DEGAS)	JET divertor (EDGE2D)
Ne	$0.1 \text{ e}^{19} - 1 \text{ e}^{19} \text{ m}^{-3}$	$\approx 10 \text{ e}^{19} \text{ m}^{-3}$
Te	200 - 600eV	<30 eV

# *Tritium deposition patterns in TFTR*

- Codeposition of tritium with eroded carbon both inboard & outboard
- Dust and debris observed
- Tiles, coupons, dust samples retrieved for analysis
- Tritium spatial distribution consistent with modeling...



Debris and dust on TFTR vessel floor

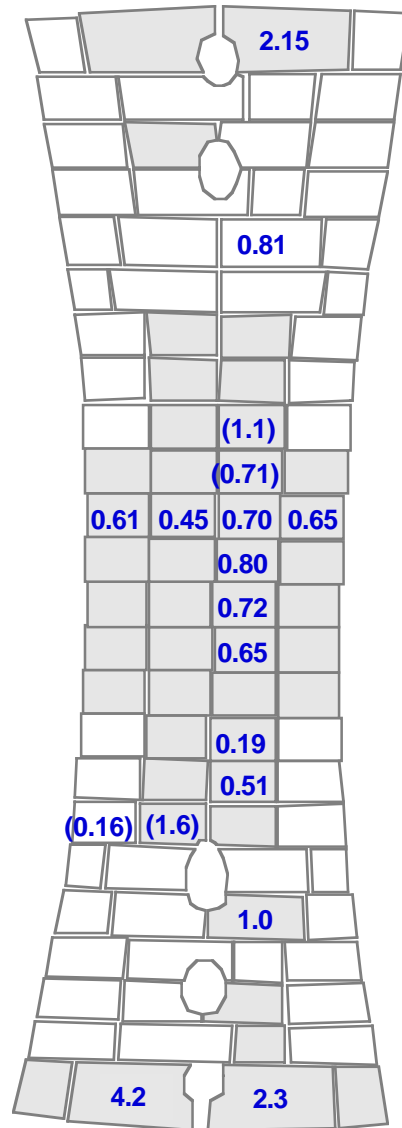


Co-deposition, flaking, deposits inside vessel.

# Tritium co-deposition on bumper limiter.



99E0014-04



Diagonal pattern on inner limiter segments due to geometry of 'scalloped' shape and connection length of field lines (Brooks et al., )

After plasma operations tritium in TFTR was located on inner limiter ( 0.2 g), and outer wall (0.36 g).

Highest concentrations were at top and bottom of limiter.

Numbers represent T (Ci) released by bakeout in air at 500 C for 1 hour.

9600 Ci = 1 g T

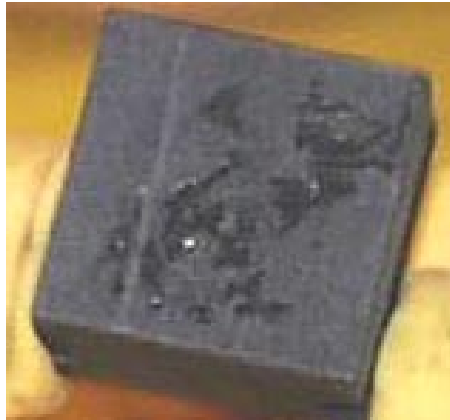
# Tritium on outboard side of vessel

From bakeout:

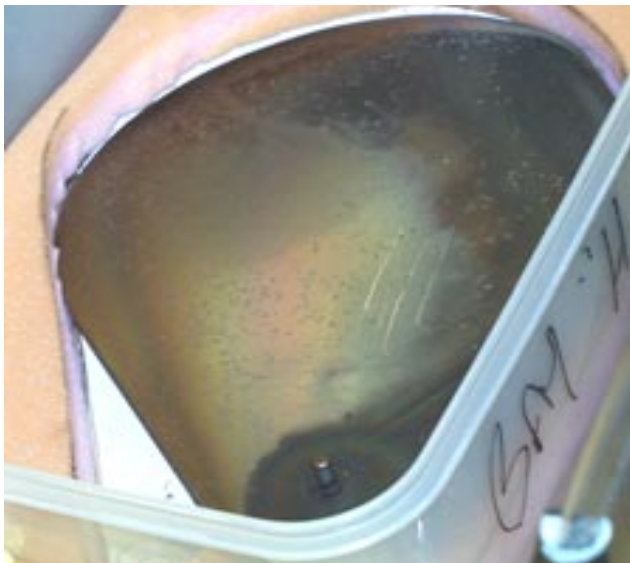
Bay H midplane graphite coupon: 24 Ci/m<sup>2</sup>

Bay N bottom graphite coupon: 65 Ci/m<sup>2</sup>

Bay P midplane graphite coupon: 16 Ci/m<sup>2</sup>

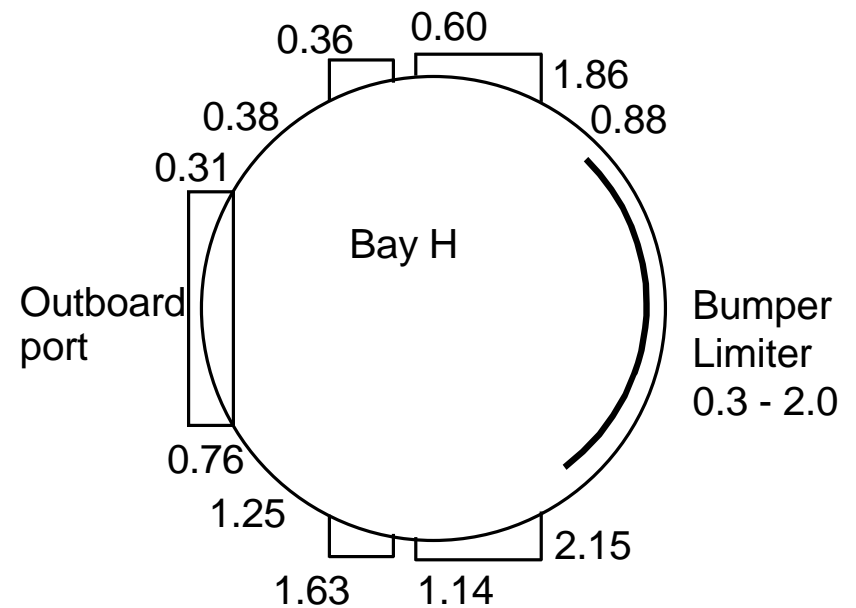


Bay H shutter (stainless steel) 9 Ci/m<sup>2</sup>



Tritium in top micron of surface ( $\beta$  range) measured by open wall ion chamber on vessel wall

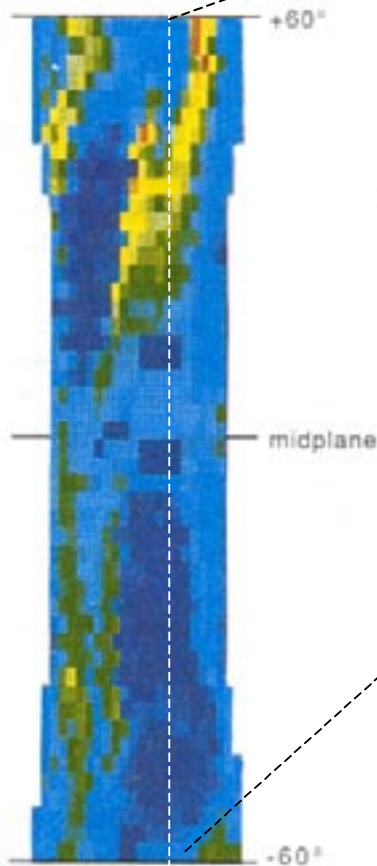
Maximum closest to limiter and on bottom of vessel.



Units Ci/m<sup>2</sup> [9,600 Ci/g T]

# Nuclear reaction analysis of deposited deuterium in DD phase

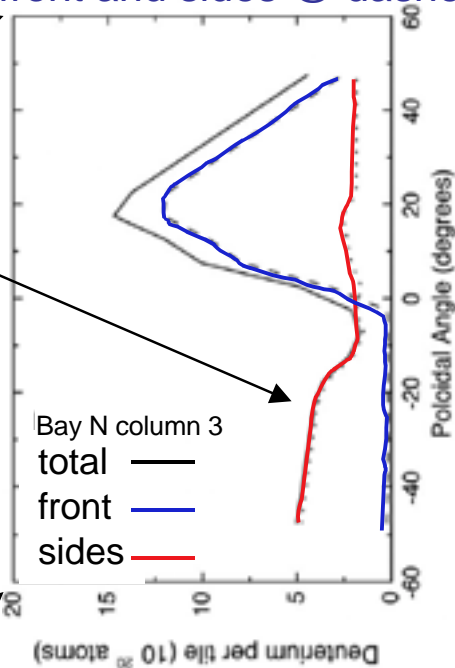
Metals deposition on plasma facing surface of bumper limiter (Beta backscattering)



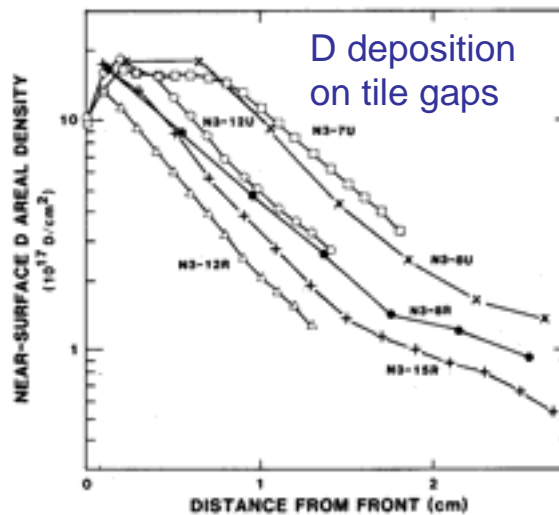
*Early demonstration of material migration in tokamaks*

D on tile front and sides @ dashed line.

Erosion areas have more D in gaps between tiles



Bay N column 3  
total —  
front —  
sides —



D deposition on tile gaps

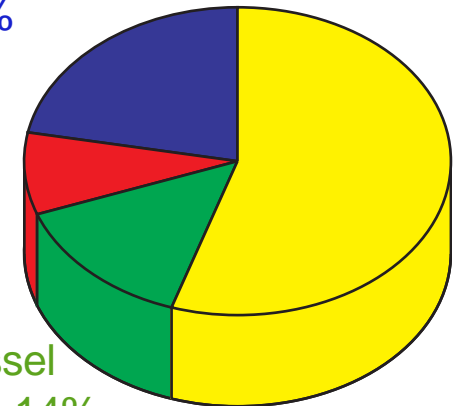
## Summary of DD phase

Plasma facing tile surface 22%

Gaps between tiles 9%

Vessel wall 14%

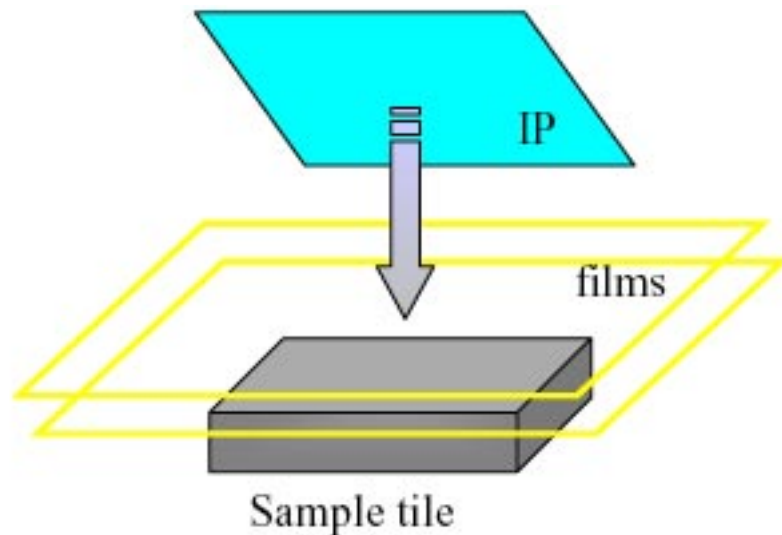
D not retained 55%



W. Wampler et al., J. Vac. Sci. Technol, A6 (1988) 2111,  
B E. Mills et al J. Nucl. Mater, 162-164 (1989) 343.

*Historical note: In 15 years we have not demonstrated an ITER relevant way to remove tritium in a tokamak!*

# High resolution T mapping



Photograph of CFC tile KC12

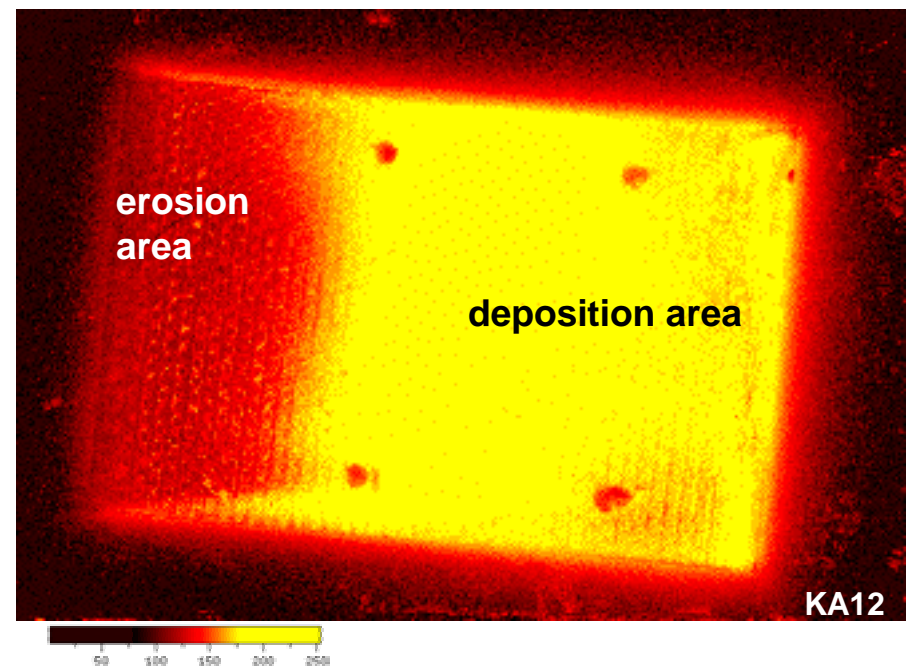


Imaging plate: tritium and  $^{60}\text{Co}$  on TFTR CFC tile

Fuji Imaging Plate(IP) provides high resolution contact image of tile radioactivity

1 hour exposure, IP remains behind contamination barrier of two  $1.2\ \mu\text{m}$  films placed in contact with tile.

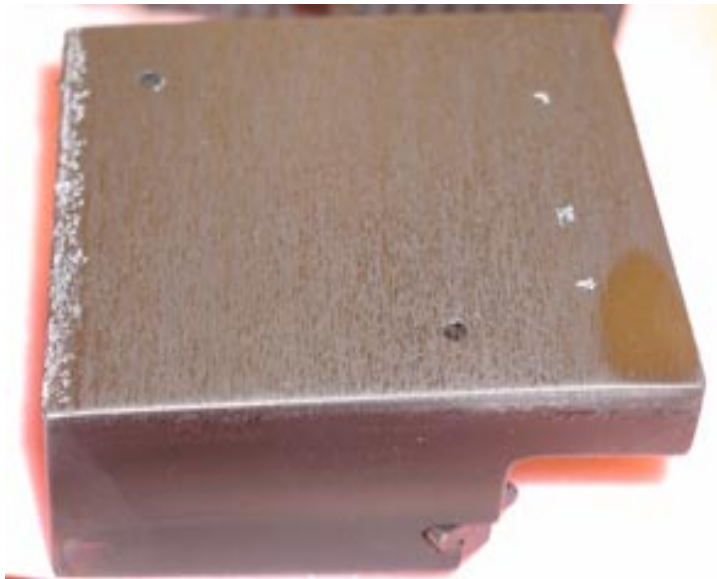
Image read out by laser  
(photo stimulated luminescence of BaFBr doped  $\text{Cu}^{2+}$ ).



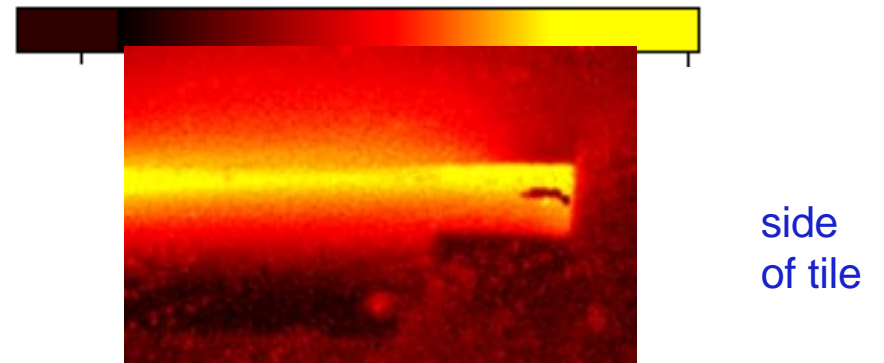
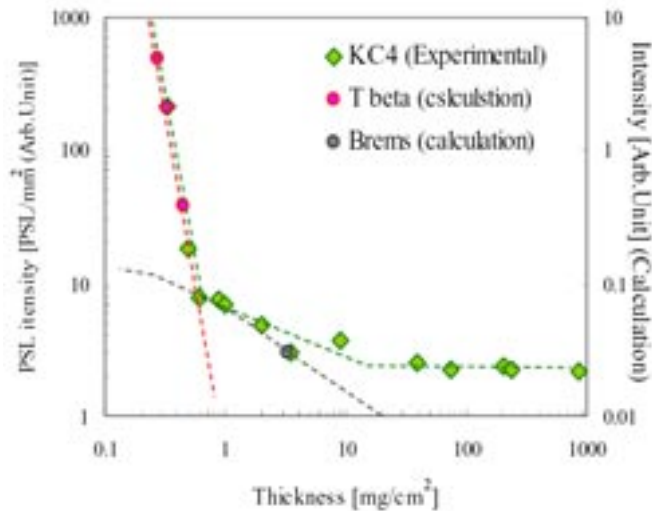
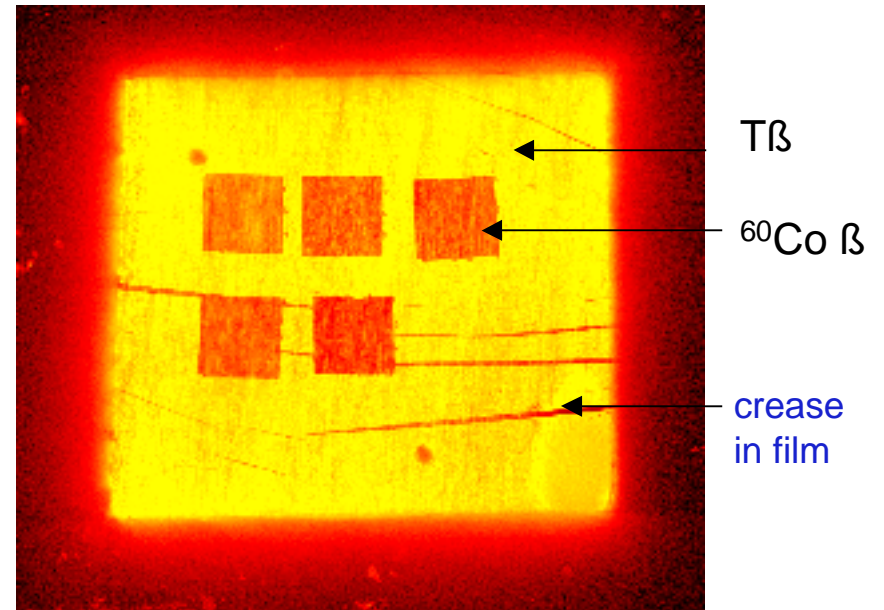
# Tritium on graphite tile in deposition area

Photograph of graphite tile KC21 from deposition area.

Image of tritium and  $^{60}\text{Co}$  (from stainless steel debris) on plasma facing surface and side of tile KC21



Additional film and Al square filters to identify 5keV betas from tritium and 318keV betas from  $^{60}\text{Co}$ . [ $^{60}\text{Co}$  and  $^{57}\text{Co}$  also identified by gamma spectroscopy.]



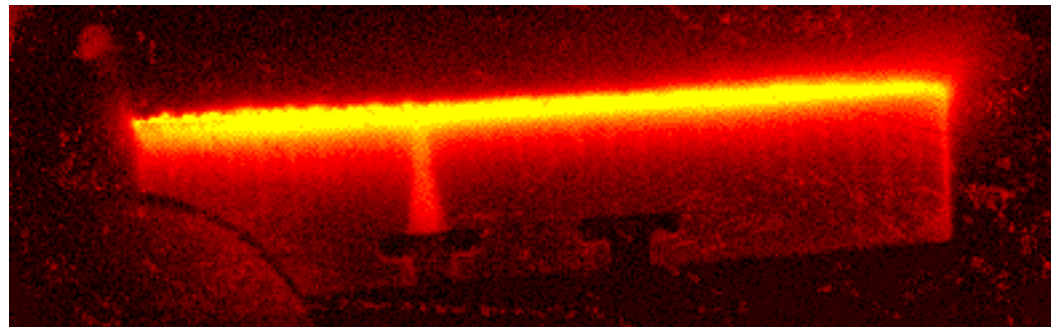
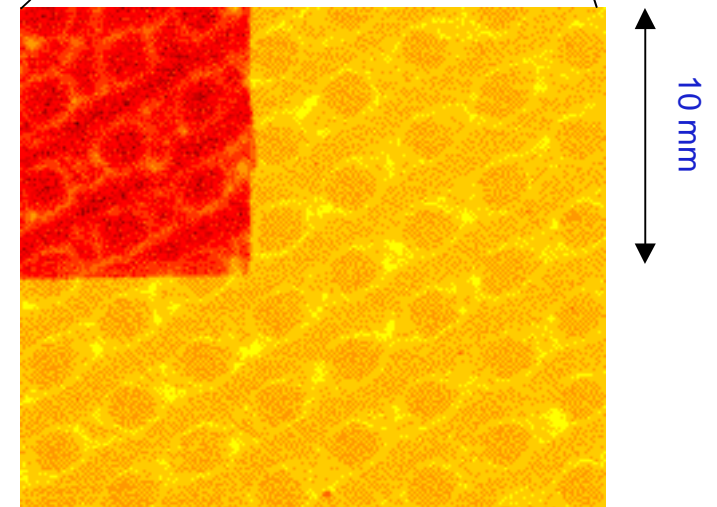
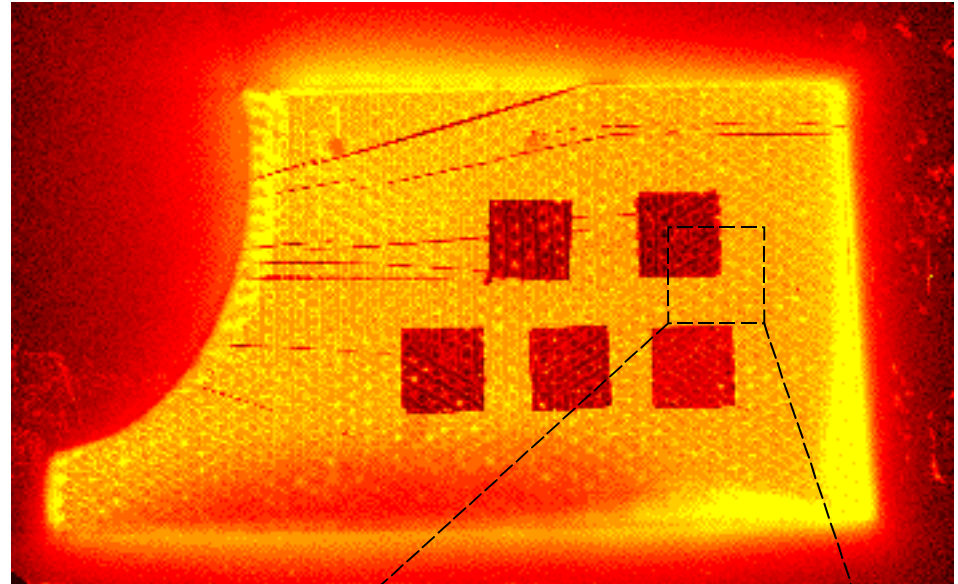


# Tritium on CFC tile in erosion area (KC3)

Photo of KC3



Imaging plate: Tritium &  $^{60}\text{Co}$

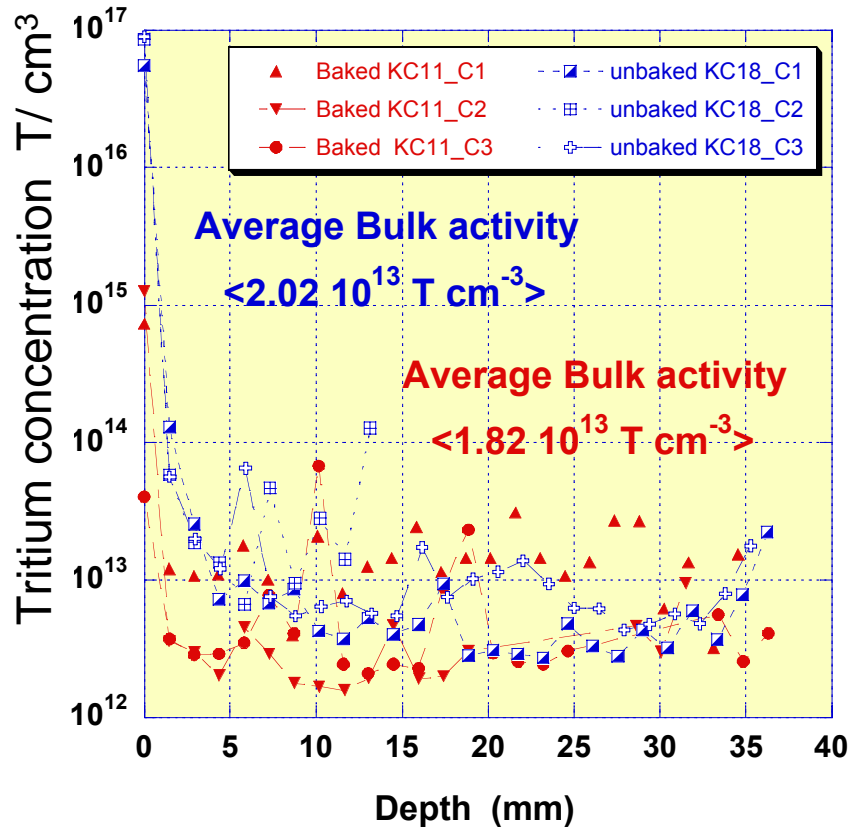


T +  $^{60}\text{Co}$  on tile side

Note tritium concentrated on fiber boundaries

# Tritium found in tile bulk

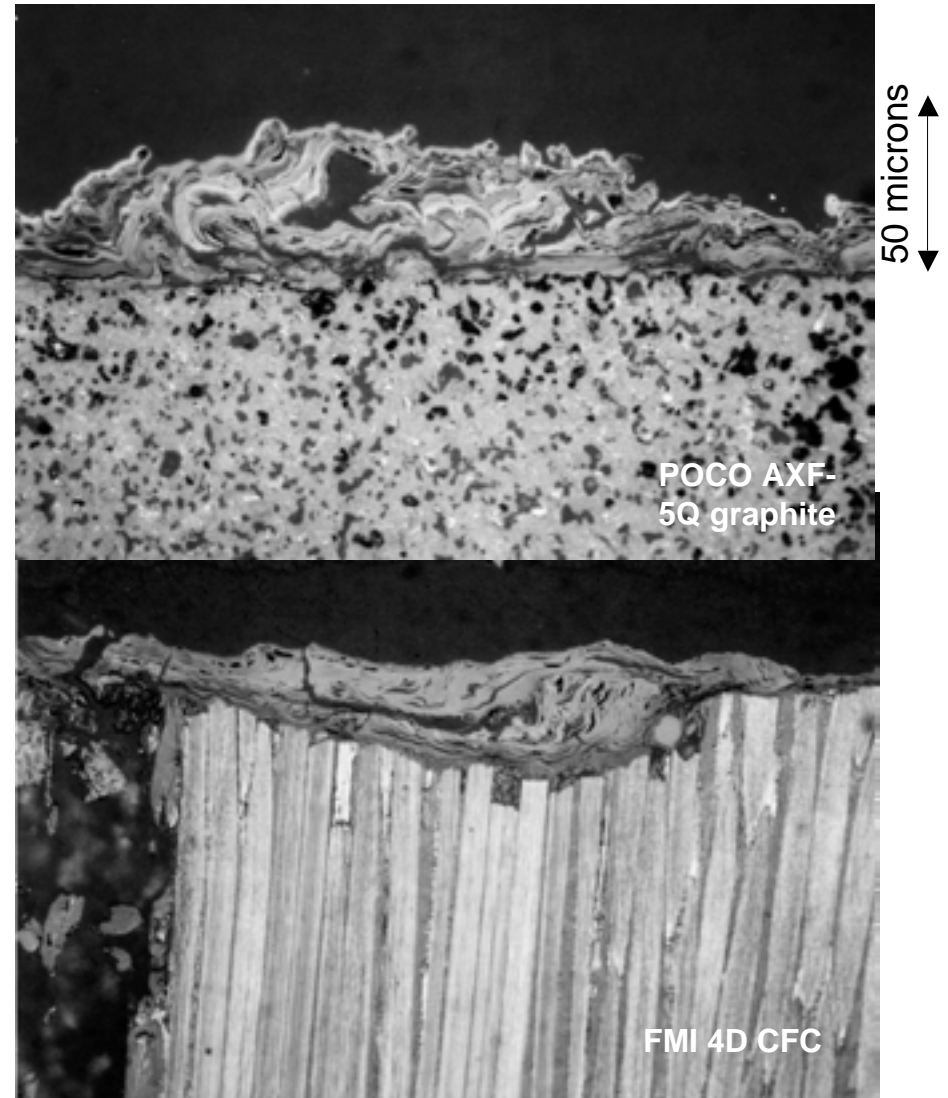
Tritium depth profile comparison for the CFC tiles KC18 (unbaked) and KC11 (baked)



Core sampling shows 0.5 - 1% of tritium has diffused to bulk of tile and is not released by baking

Nicolas Bekris FZK

Cross Section of TFTR co-deposit.



M Paffett, R Reiswig, S Willms, LANL

# Modeling of C production and Tritium retention in TFTR

BBQ code by John Hogan describes:

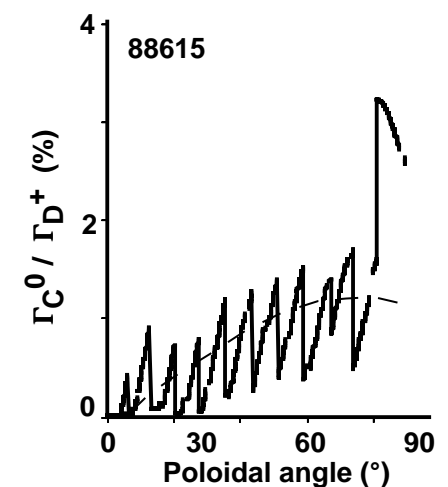
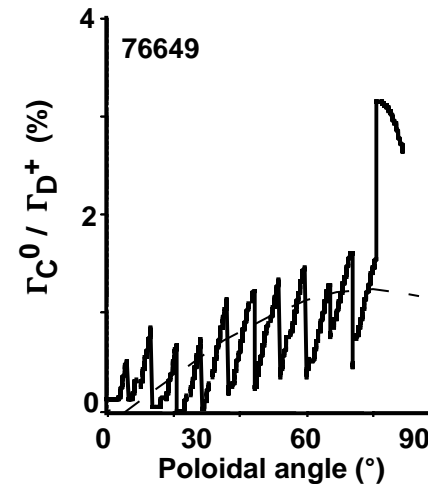
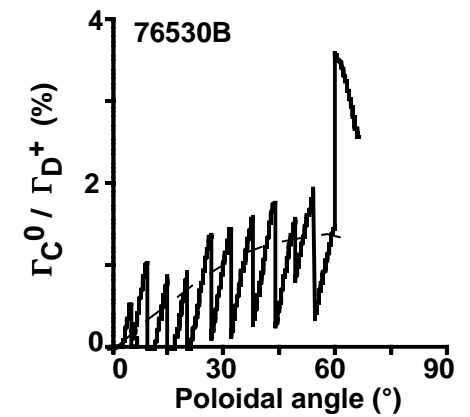
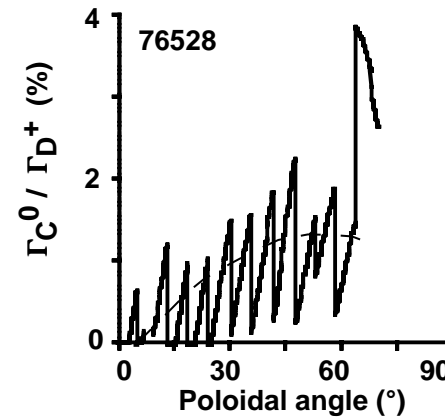
3D space, 3D velocity test particle Monte Carlo code for emitted C impurities from physical, chemical sputtering and radiation-enhanced sublimation (RES)

Parallel, perpendicular diffusion, electrostatic fields, friction with SOL flow, atomic/molecular physics (includes Erhardt-Langer database for CD4 breakup)

Combines detailed TFTR Bumper Limiter geometry (CAD) with impurity SOL transport and redeposition

Extrapolate carbon erosion from selected representative discharges  
H-isotope/C ratio in co-deposits  
approximately 0.2 (NRA) – estimate retention....

→ **Modeling can account for order of magnitude of retention**

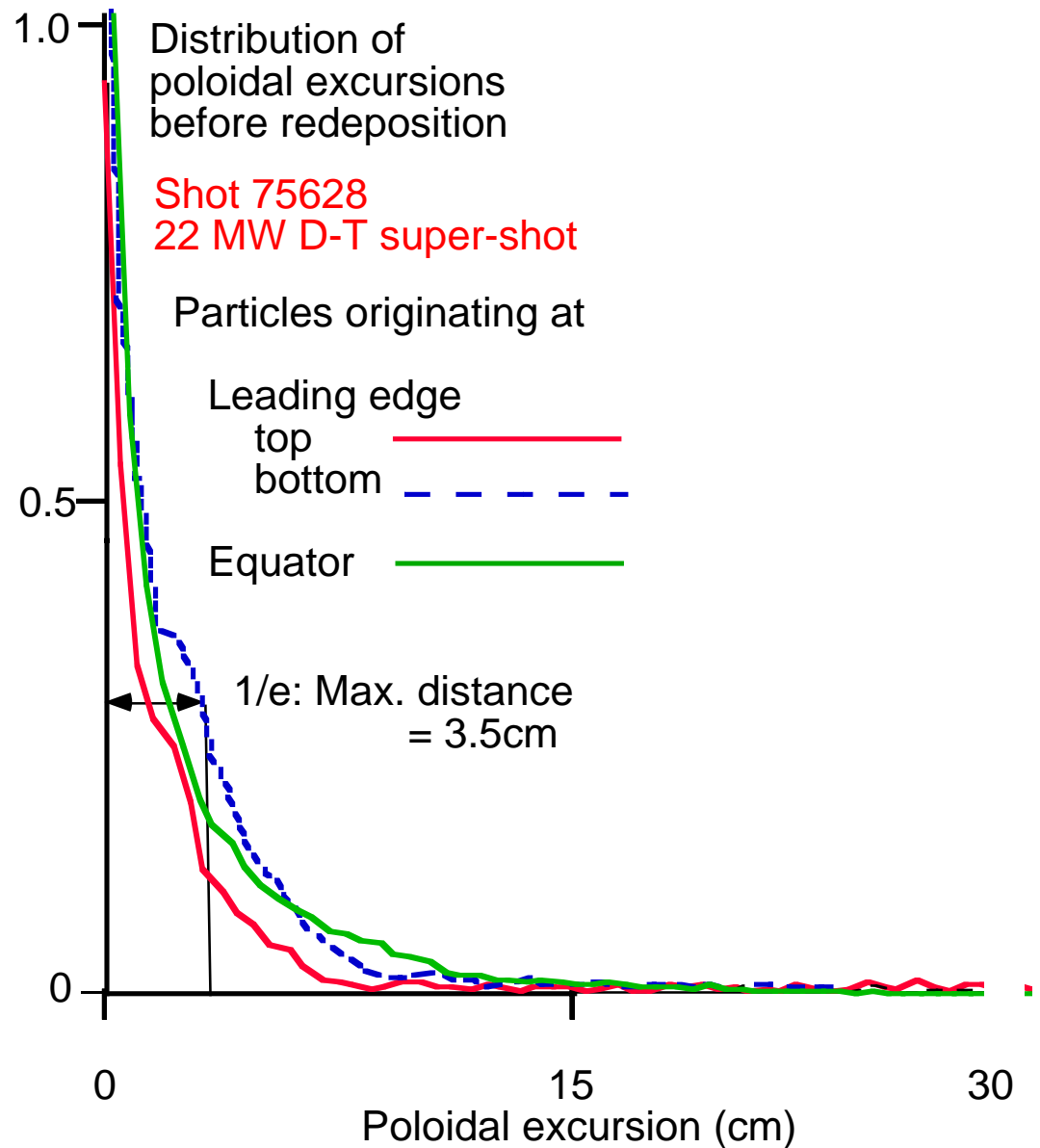


Local effective sputtering yield distribution on bumper limiter  
(emitted impurity flux / incident D+ flux) for 4 representative discharges.

John Hogan ORNL

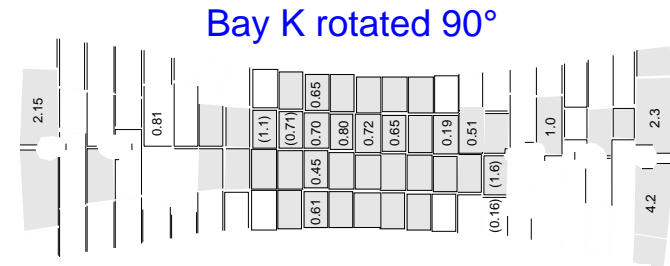
## Carbon + H-isotopes codeposited close to erosion point

- BBQ shows strong localization of D+ flux at top/bottom leading edges of TFTR limiter.
- Data consistent with considerable number of TFTR discharges with large ( $\approx 10\text{cm}$ ) radial decay length of the D+ flux due to inner wall recycling and flux amplification.
- Flight of sputtered carbon tracked in radial, poloidal and toroidal dimensions.
- Higher effective sputtering yield at high latitudes and prompt local redeposition leads to high codeposition in these areas.
- Significant concentrations of T predicted on upper and lower leading edges of limiter.

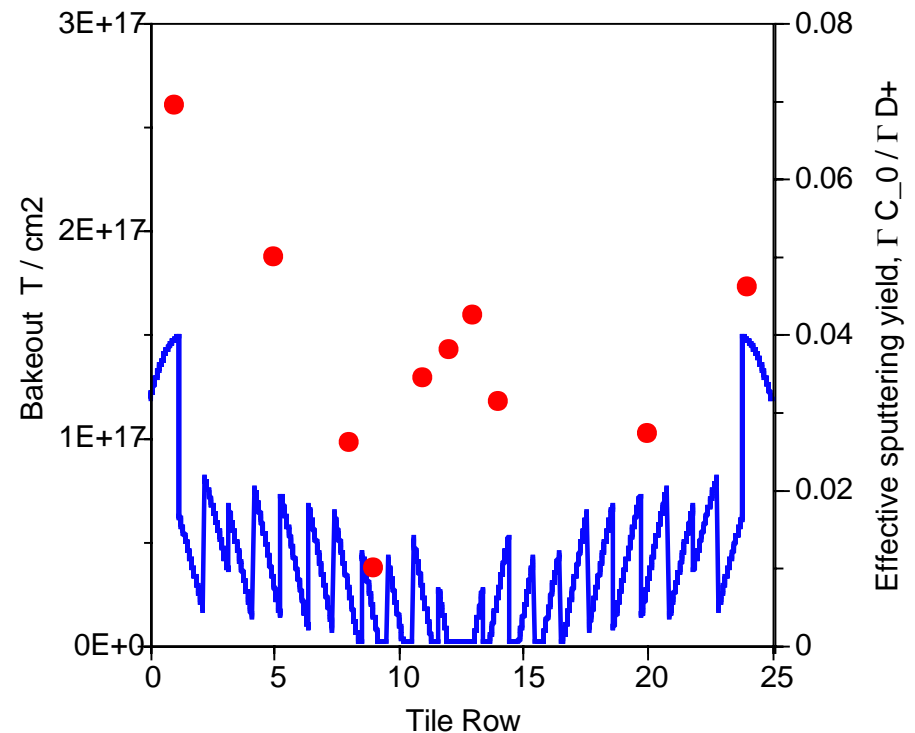


# Modelling appears to be on right track

- Higher effective sputtering yield at high latitudes and prompt local redeposition leads to high codeposition in these areas
- Data consistent with considerable number of TFTR discharges with large ( $\approx 10\text{cm}$ ) radial decay length of the  $\text{D}^+$  flux due to inner wall recycling and flux amplification.
- Li deposition at same locations may enhance retention (Li used for wall conditioning).
- Observed tritium concentrations (measured after modeling predictions) suggest model is on right track.



Measure tritium in selected tiles

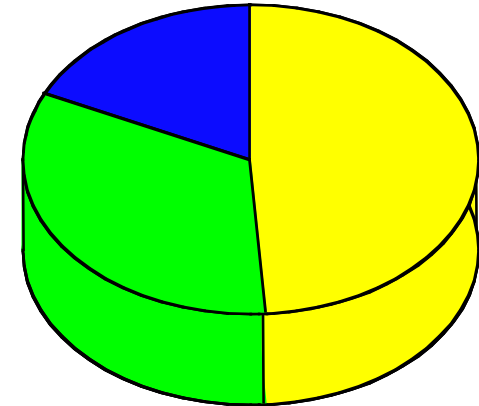


Row averaged tritium release / plasma facing area ( $\text{cm}^2$ ) compared to effective sputtering yield for # 76528

## Overview: Tritium retention in TFTR

Location:	Area (m <sup>2</sup> )	T conc. mg/m <sup>2</sup>	(g)
Bumper limiter	22	9	0.2
Outboard	110	3.3	0.36
Total			<b>0.56</b>
<i>cf. fueling - exhaust</i>			<b>0.64</b>

T on limiter 18%



T outboard 33%

T not retained 49%

- *Retention fraction  $\approx$  51% much much too high for ITER*
- *1/3 tritium on bumper limiter, 2/3 on outboard wall*
- *Remarkably good agreement between extrapolation from tile measurements and fueling less exhaust.*
- *Retained tritium consistent with modeling.*
- *0.5 - 1% of tritium in bulk of tile*
- *Some tritium trapped at fiber boundaries*