

# Plasma facing surface composition during Li evaporation on NSTX and LTX



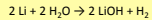
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## Astract:

• Evaporated lithium coatings react with residual water in the tokamak vacuum to produce lithium hydroxide and hydrogen:



• Since tokamaks typically do not have ultra-high vacuum (UHV) conditions, this process can occur in the time interval between lithium evaporation and the next discharge.

• The resulting PFC surface should be considered as a mixed material rather than a pure 'lithium coating'.

• We present calculations of the flux of water from the residual vacuum to PFCs in NSTX and LTX under various conditions.

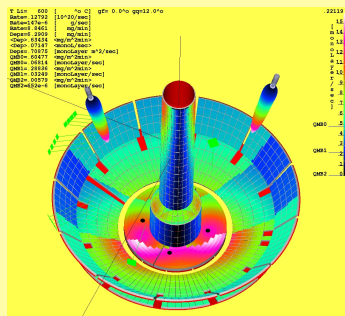
• To avoid reactions with residual vacuum gasses an ultra-high vacuum ( $\leq 1\text{e-}8$  Torr) is required and may be achievable by a large-scale lithium getter pump.

## Li deposition rate:

Q. How does the time scale to deliver a monolayer of lithium via evaporation compare to the time scale for the residual vacuum to deliver a monolayer of water to PFCs in LTX and NSTX?

One mole of lithium = 6.94 g and  $6.02\text{e}23$  atoms.  
Using the Li density =  $0.534 \text{ g/cm}^3$  a Li monolayer with cubic atom spacing has  $1.29\text{e}15$  atoms/cm<sup>2</sup> or  $1.3\text{e}20$  atoms or 1.5 mg over the NSTX deposition area of  $\sim 10 \text{ m}^2$ .

At a typical LITER evaporation rate of 20 mg/min depositing a Li monolayer would take 4.5 s.  
(note this is for a flat mirror-like surface - coating a more realistic rough surface with a monolayer of Li would take longer.)



Calculation by Cbebm code of lithium deposition from the LITER-1 lithium evaporator.  
[L. Zakharov et al., PPPL report PPPL-4187 Nov. 2006]

## H<sub>2</sub>O flux from base vacuum.

The flux of impinging gas molecules on a surface is given by kinetic theory:

$$\text{flux} = 3.513\text{e}22 [P/V(M^*T)] \text{ molec/cm}^2/\text{s}$$

where P is in Torr, M is in amu, and T is in K.

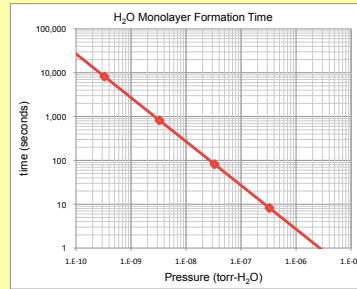
For water M=18, and using T=300K and P=1e-6 Torr

$$\text{Flux (H}_2\text{O)} = 4.78\text{e}14 \text{ molec/cm}^2/\text{s}.$$

The reaction probability for incident H<sub>2</sub>O molecules and surface Li atoms is near unity.

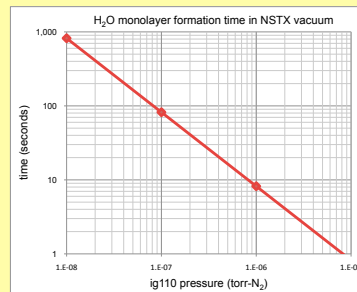
[1] A. Roth 'Vacuum Technology' 3rd ed. P. 3 table 1.2 and P. 36

Time to supply one molecule of water to every Li atom on a flat surface



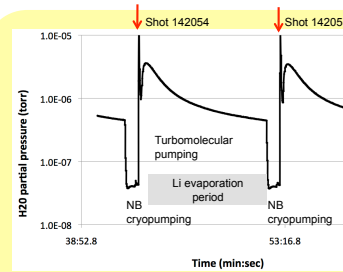
From NSTX ionization gauge and RGA calibrations:

$$\text{Water pressure (Torr)} = \text{IG110 reading (N}_2 \text{ calibration)} \times 0.33$$



Time to supply one molecule of water to every Li atom on a flat surface as a function of ig110 reading on NSTX

## NSTX water base pressure



Between shots the water monolayer formation time (6s) is similar to the Li monolayer impingement time (4.5 – 11.3 s) at 20 mg/min. 1 minute before shot, LITER was withdrawn and NB cryopump opened giving  $\sim 25$  sec with water flux lower than the Li flux on flat surface.

## Oxygen segregation to Li surface

Q. The majority of sputtered atoms originate at the surface. Can fresh lithium evaporation restore a pure Li surface over an oxidized lithium surface?

- Materials with a low surface tension such as LiOH tend to segregate to the top of liquid lithium surfaces leading to a large difference between surface and bulk compositions.
- Buried or dissolved LiOH (or Li<sub>2</sub>O) may resurface when Li melts.

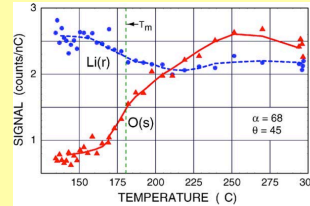


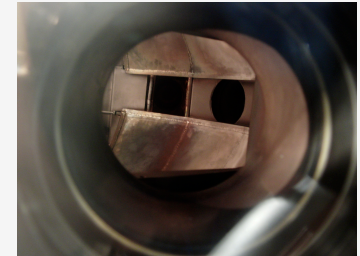
Fig. 3. Variation in oxygen signal intensity on a lithium surface as a function of temperature. The intensity of ion scattering from O and recoil emission from Li were monitored while the sample was bombarded with a 500 eV He<sup>+</sup> beam. Oxygen atoms appear more abundant on the liquid surface.

[Bastasz and Whaley, Fus. Eng. Des., (2004) 111]

- In a tokamak non-thermal mechanisms such as ion-induced diffusion and the formation of other lithium compounds may complicate this process.

## LTX operated with a Li-coated 300 °C shell :

- First full high temperature, high Z wall operation of a tokamak – Lithium evaporated into 5 mTorr helium fill to disperse coating



- Deposition rate  $\sim 0.75 \text{ g/hour/evaporator}$ 
  - 3 hour duration, plasma operations follow 1 ¼ h later
  - Estimated 1.6 micron average Li thickness.
  - Partial pressure of water estimated to be in 1e-8 Torr range
- Lithium coating darkens rapidly.
- No visual evidence of shiny metallic surface
  - Indicative of LiOH formation at the surface.
- Li evaporation onto cold walls produced large effect on peak Ip and discharge duration.
- With hot walls - no significant improvement so far over uncoated walls or walls with passivated Li coatings.

## Conclusions:

- At typical tokamak residual H<sub>2</sub>O pressures of 1e-6 – 1e-8 Torr after a discharge, evaporated lithium coatings will react with residual water to form LiOH at the surface on a time scale of 3 - 300 seconds.
- Buried oxygen-containing lithium compounds can segregate to surface of molten Li
- The resulting PFC surface should be considered as a mixed material rather than a pure 'lithium coating'.
- To avoid reactions with residual vacuum gasses an ultra-high vacuum ( $\leq 1\text{e-}8$  Torr) is required.

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