

Deuterium Ion Retention as Lithium Deuteride in Thin Li Films on Mo(110)

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Motivation and Overview

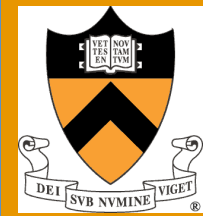


Motivation

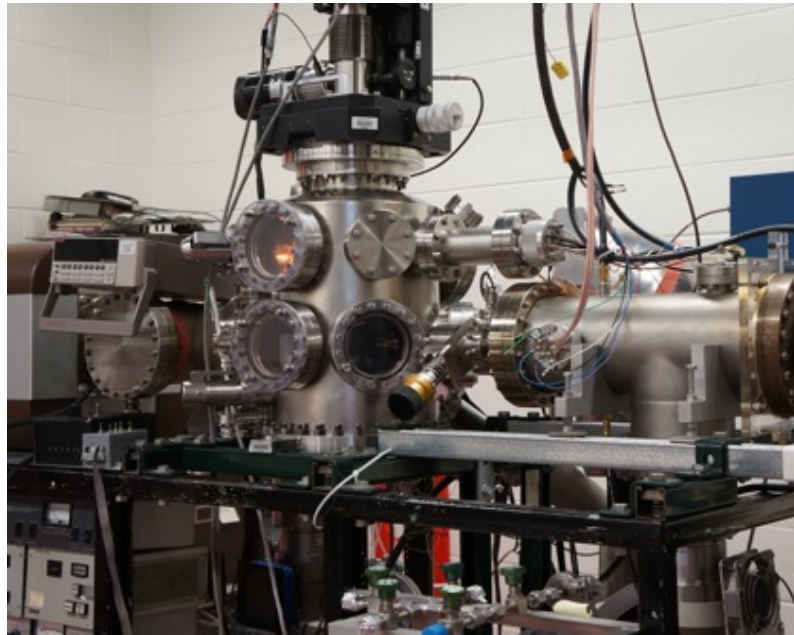
- Lithium coatings on plasma facing components have been shown to improve plasma performance
- Need to understand fundamental chemical interactions between plasma and plasma facing lithium films to help inform observations in more complex systems

Overview

- I. A surface science approach to understand D retention in Li
- II. Identifying and characterizing LiD formation in Li films
- III. D retention in monolayer and multilayer Li films
- IV. Quantifying D retention in Li films



Surface Science Approach to Understanding D Retention



Experimental Chamber in Surface Science & Technology Lab at PPPL

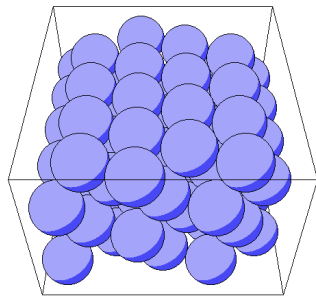
- Surface science experiments enable independent control of all variables --- This is something we cannot achieve in a tokamak or linear plasma device.
- Key variables affecting chemistry at surface are controlled:
 - Pressure (UHV, background 10^{-10} Torr)
 - Temperature (100 – 2000 K)
 - Composition (Mo, Li, D^+ , residual gases, etc.)
- Isolate effects of incident particles and their energies
- Start simple, i.e. single effects, and add complexity to bridge gap between model systems and tokamak environment



Studying Simple Model Systems



Simple Model Experiments



Mo(110) Crystal

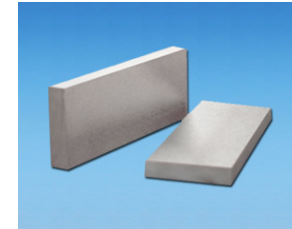
Grain boundaries

Alloying elements: Ti, Zr, C

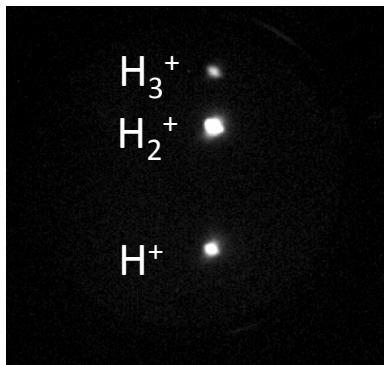
Surface roughness and different crystal facets

Adsorbed layers: O, C, etc.

More Complex Systems



TZM Alloy



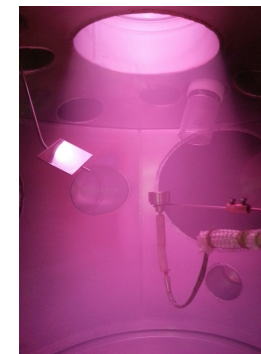
Monoenergetic Ion Beam
(image of ions on phosphor)

Multiple species: H^+ , H_2^+ , H_3^+

Contaminant molecules: H_2O

Increased flux: $10^{12} \rightarrow 10^{16}$
atoms/cm²/s

Atoms, ions, or atoms + ions



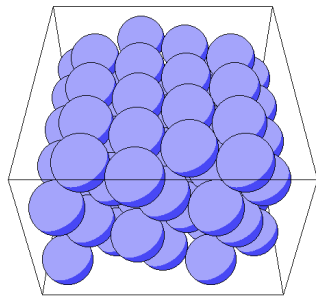
ECR Plasma Source



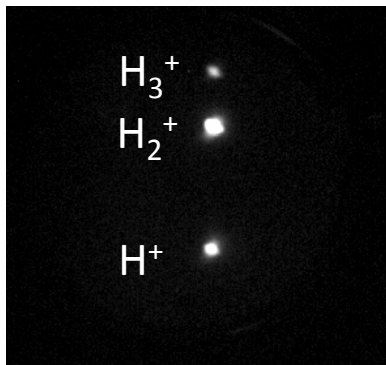
Studying Simple Model Systems



Simple Model Experiments



Mo(110) Crystal



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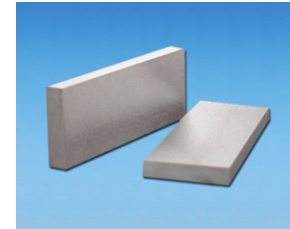
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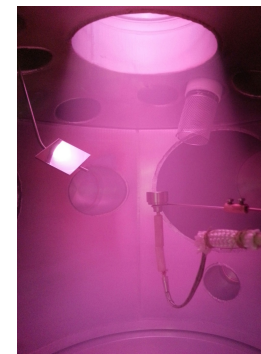
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More Complex Systems



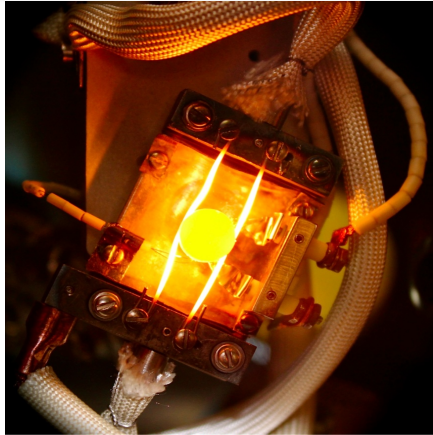
TZM Alloy



ECR Plasma Source

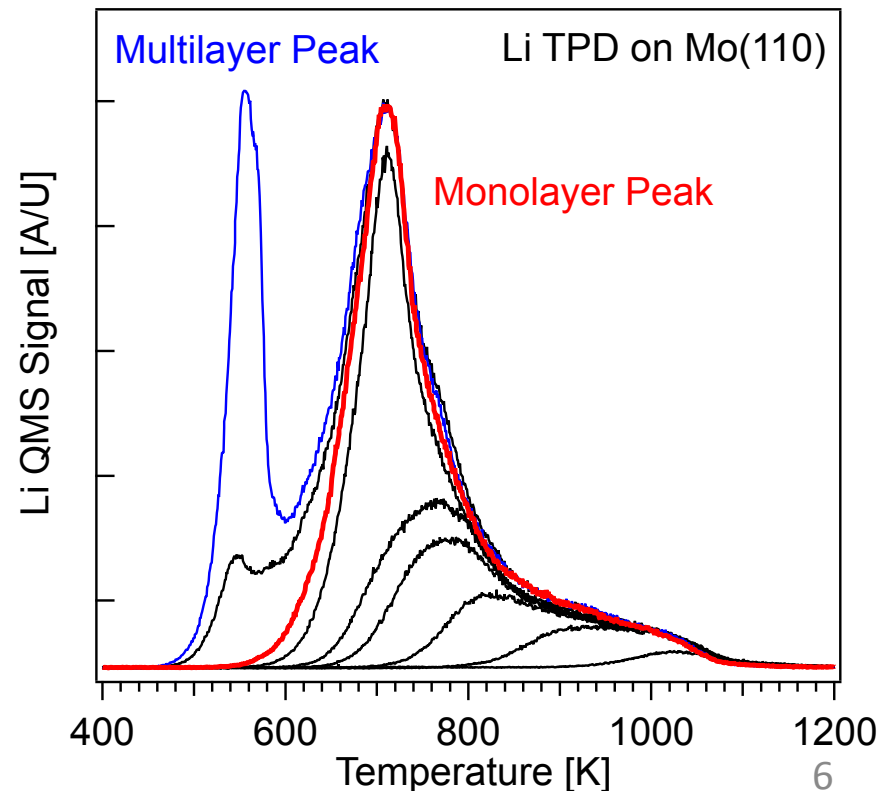
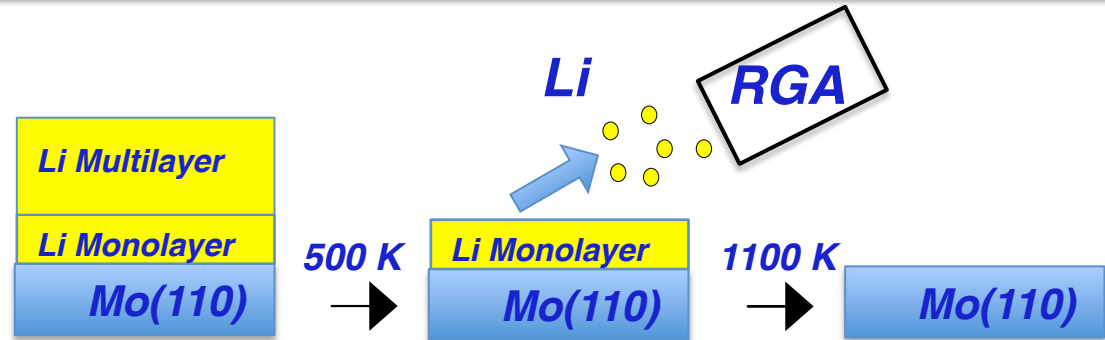


Thermal stability of Li on Mo(110)



Mo(110) resistively heated in UHV

- Temperature Programmed Desorption (TPD) is used to study Li films
- Area under TPD curve is proportional to atoms desorbed
- Sub-monolayer control of film thickness
- Desorption peaks correlated with binding (adsorption) energies

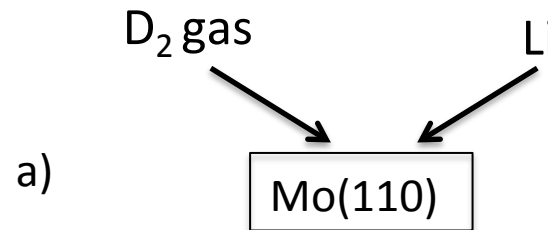




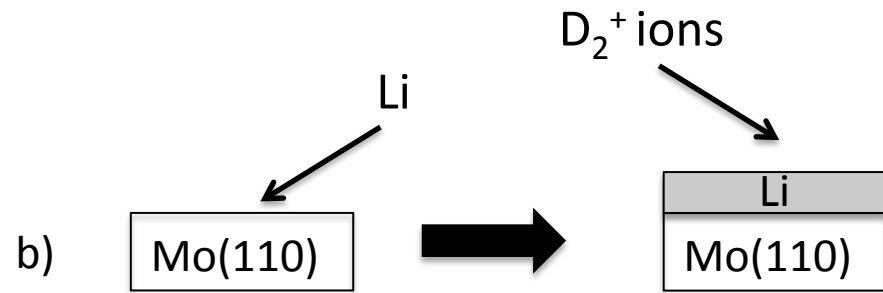
D⁺ Implantation Forms Lithium Deuteride



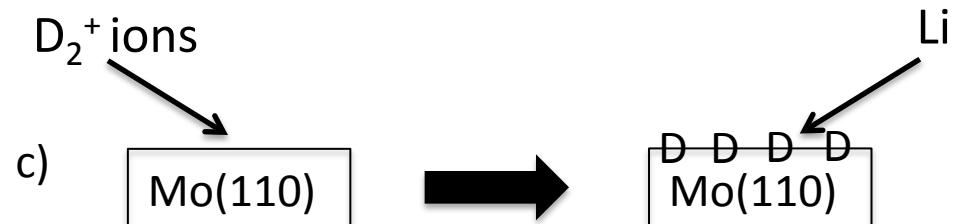
a) Lithium deposited with D₂ gas (P=10⁻⁶ Torr),
→ minimal LiD created



b) Deuterium ions implanted through Li film on Mo(110)
→ LiD created

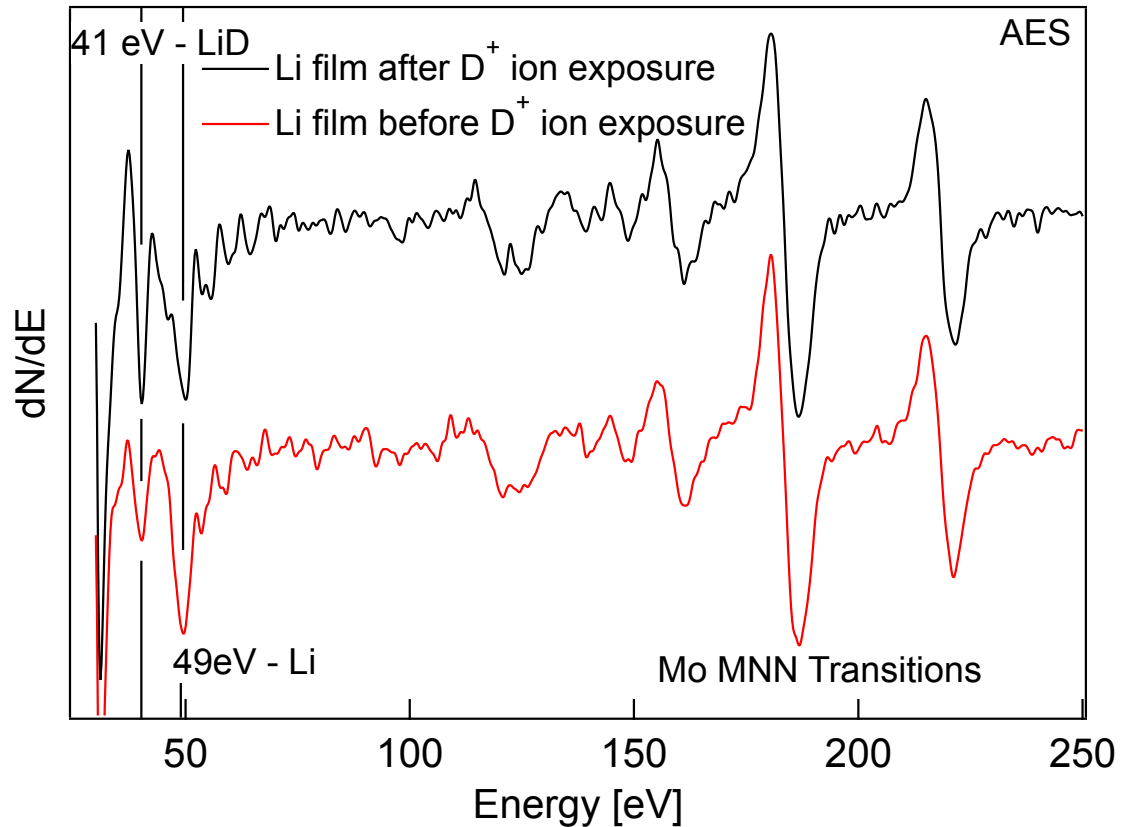
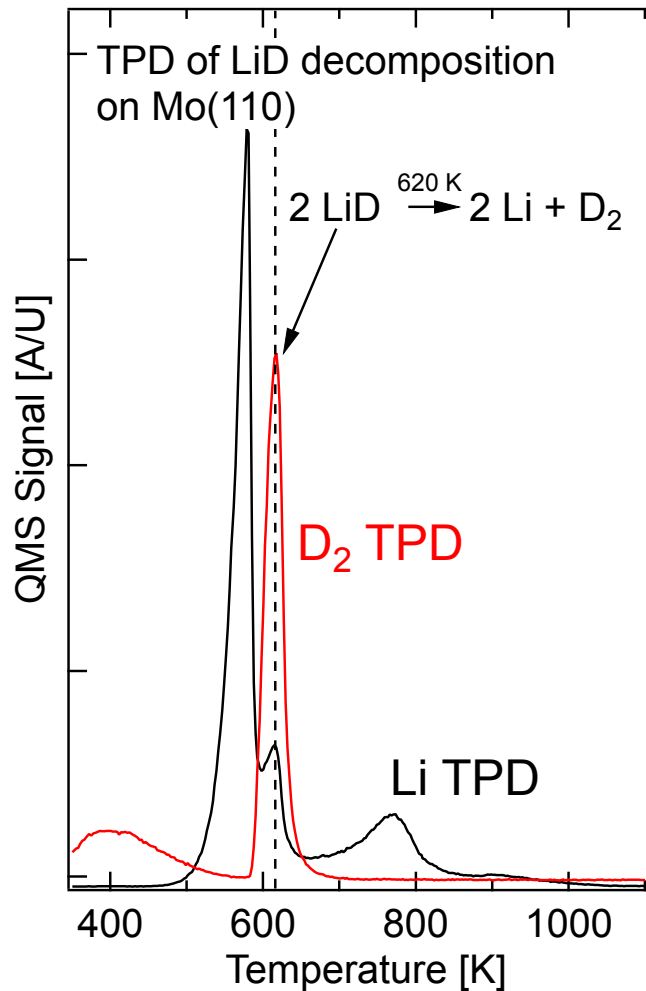


c) Deuterium ions implanted into Mo(110) crystal before Li film is deposited
→ LiD created





TPD and AES Indicate LiD Formation



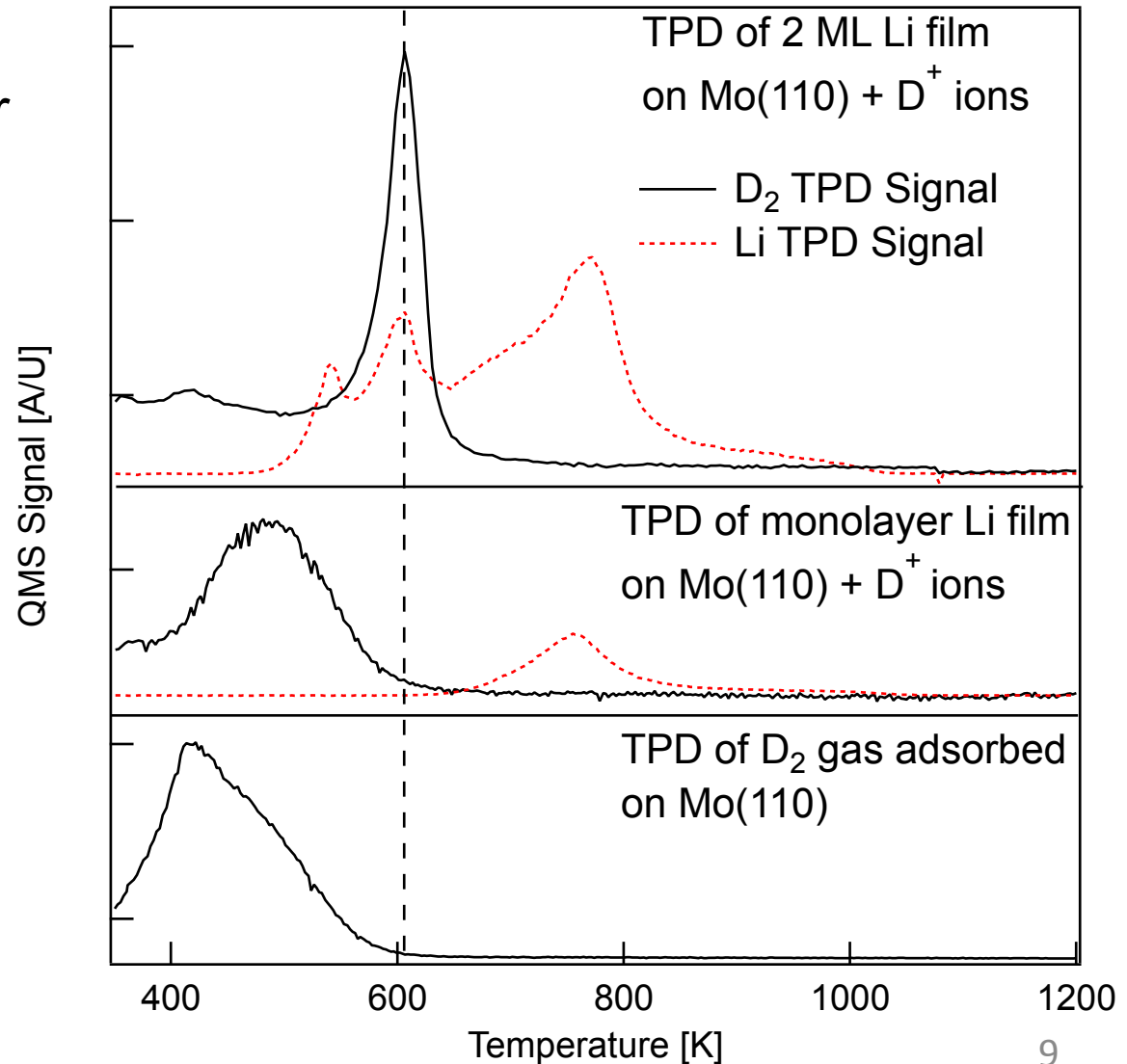
- Simultaneous desorption of D_2 and Li in TPD
- Shift in Li Auger peak from 49 eV to 41 eV



Li Monolayer Does Not Form LiD

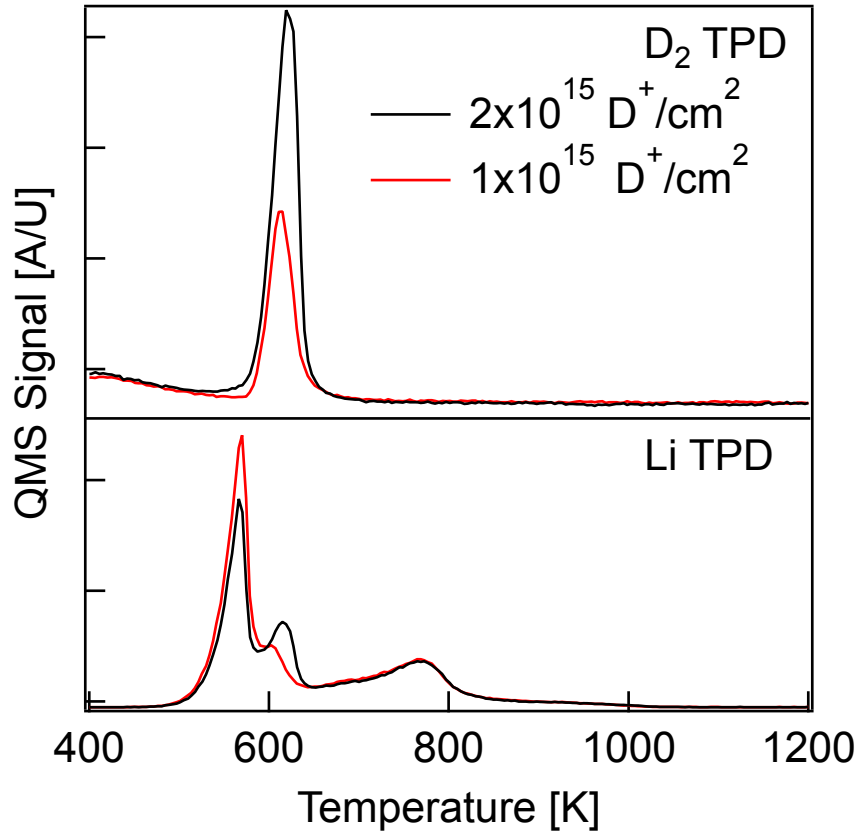


- D^+ implanted into monolayer and multilayer Li films on Mo(110)
- LiD forms when ions implanted into multilayer Li films
- LiD decomposition peak not observed in TPD of monolayer Li films
- D_2 desorption from monolayer Li film similar to D_2 desorbing from clean Mo(110) crystal



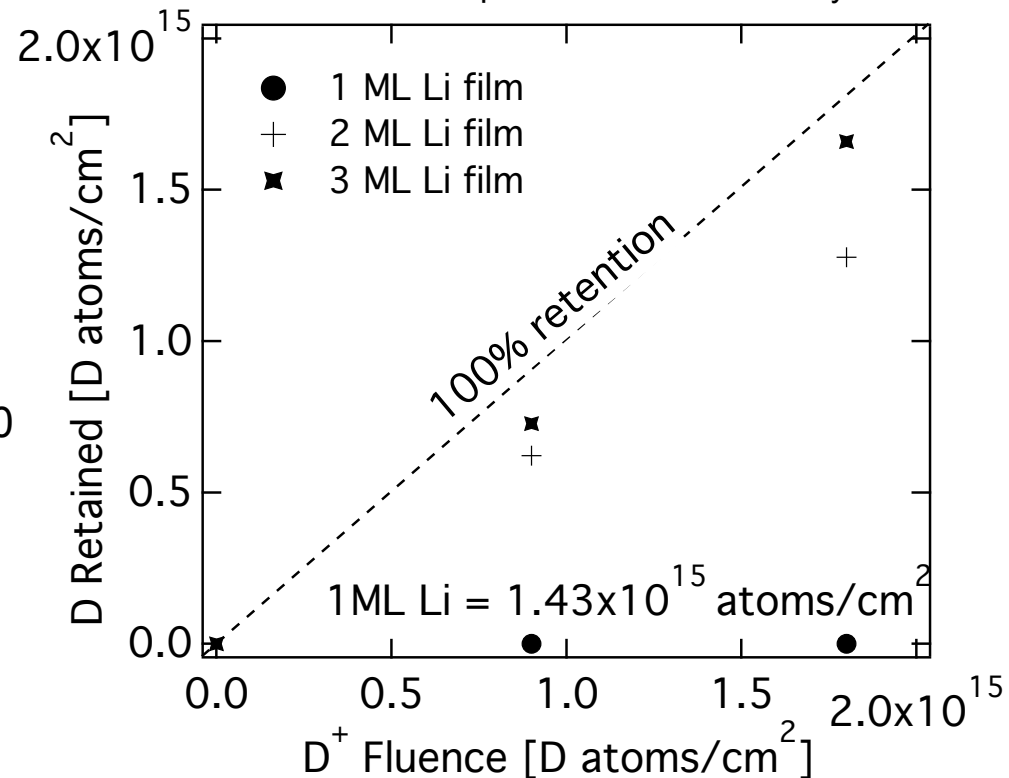


Using TPD to Quantify D Retention



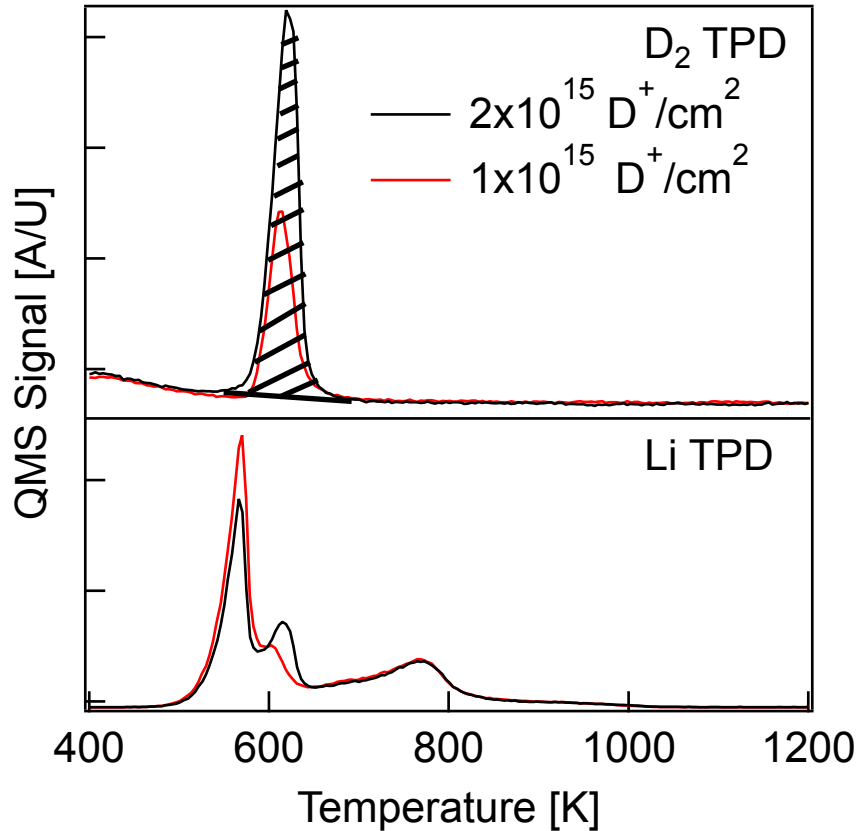
- D retention proportional to area under D_2 TPD curve
- Increased D^+ dose results in larger D_2 TPD area

- No LiD formed in 1 ML Li films
- Nearly all implanted D forms LiD in multilayer Li films
- Fraction of D forming LiD drops as $n_{D \text{ implanted}} \rightarrow n_{Li \text{ multilayer}}$



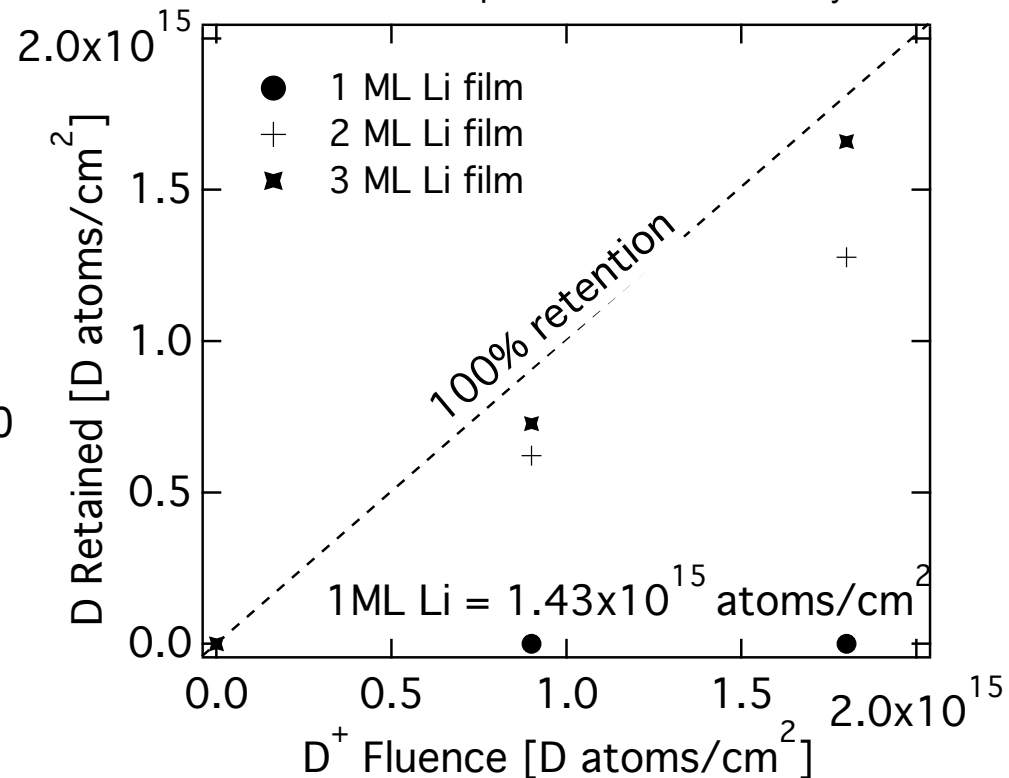


Using TPD to Quantify D Retention



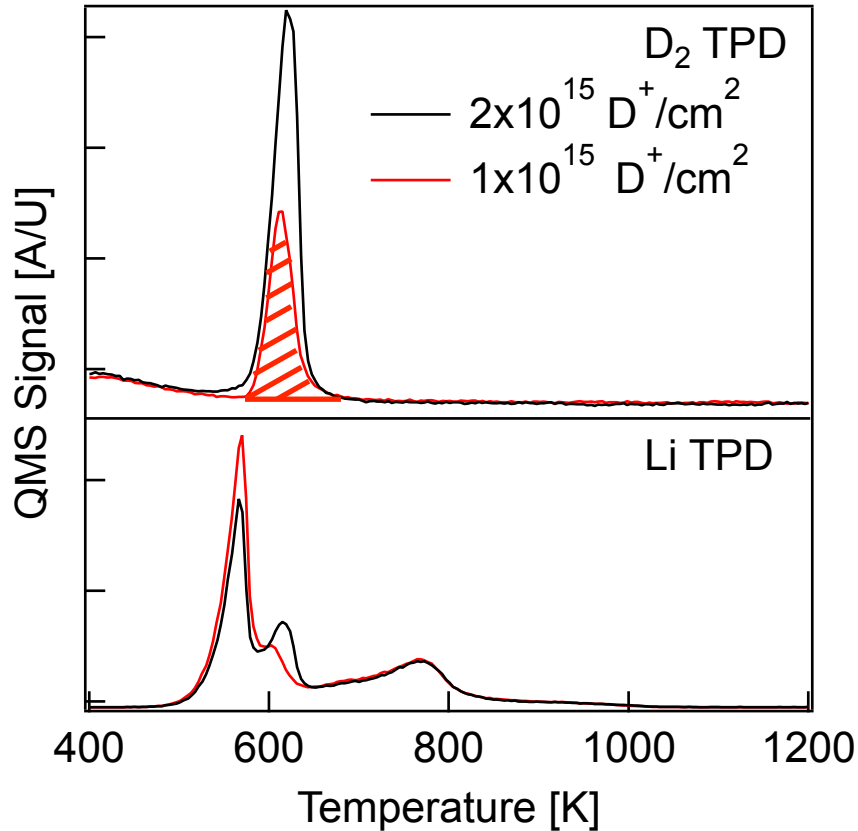
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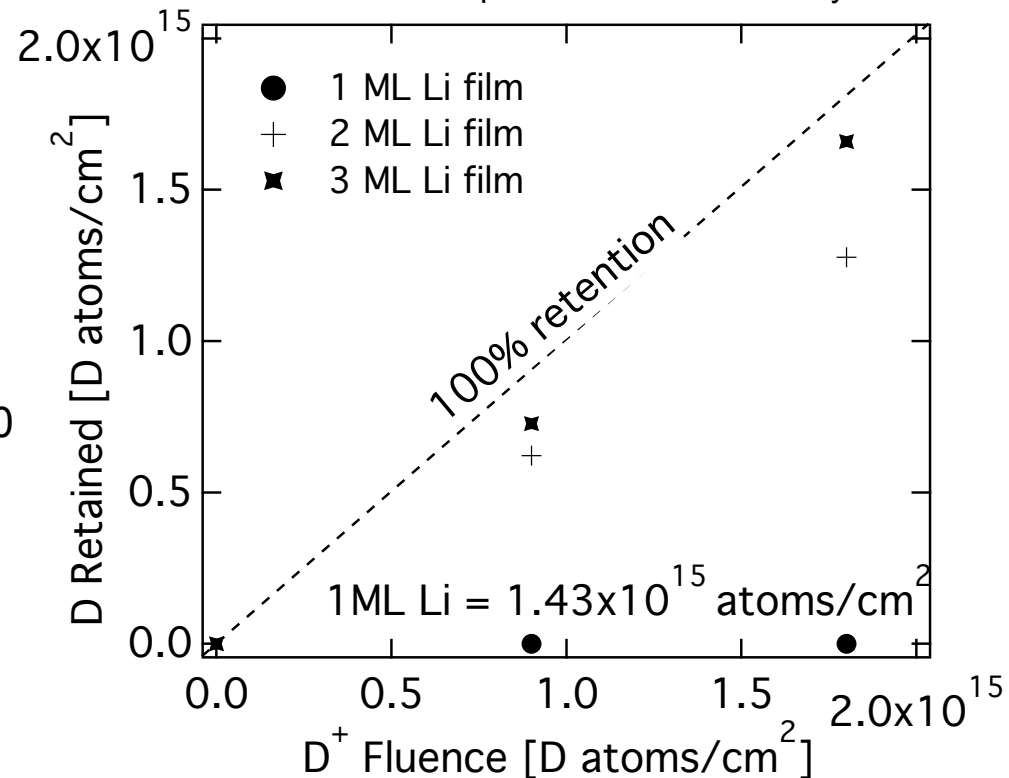


Using TPD to Quantify D Retention



- D retention proportional to area under D₂ TPD curve
- Increased D⁺ dose results in larger D₂ TPD area

- No LiD formed in 1 ML Li films
- Nearly all implanted D forms LiD in multilayer Li films
- Fraction of D forming LiD drops as $n_{D \text{ implanted}} \rightarrow n_{Li \text{ multilayer}}$





Conclusions



- D⁺ implanted into Li covered Mo(110) retained as LiD
- LiD is identified by shift in Li AES peak + simultaneous D₂ and Li TPD peaks
- LiD not formed in monolayer Li films
- Nearly 100% of D implanted forms LiD while $n_{\text{D implanted}} < n_{\text{Li-multilayer}}$
→ agrees with results of Baldwin et al.
Baldwin et al., Fusion Eng Des, 61-62 (2002) 231-236

See also

PP8.00081 : Suppressed gross erosion of high-temperature lithium films under high-flux deuterium bombardment: Wed afternoon posters

YI2.00003 : High Performance Discharges in the Lithium Tokamak eXperiment (LTX) with Liquid Lithium Walls: Friday 10:30

YI2.00005 : The Effects of Temperature and Oxidation on Deuterium Retention in Solid and Liquid Lithium Films on Molybdenum Plasma-Facing Components: Friday 11:30

JPR and BEK acknowledge support by DOE (FES) DE-SC0008598, Liquid Metals as Plasma-Facing Materials for Fusion Energy Systems: From Atoms to Tokamaks

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