

# MEASUREMENTS TO UNDERSTAND THE ORIGIN & EVOLUTION OF $\text{H}_2\text{O}$ AND OH ON THE ILLUMINATED MOON

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**How much, if any, of the ‘water’ observed on  
the illuminated Moon is  $\text{H}_2\text{O}$**

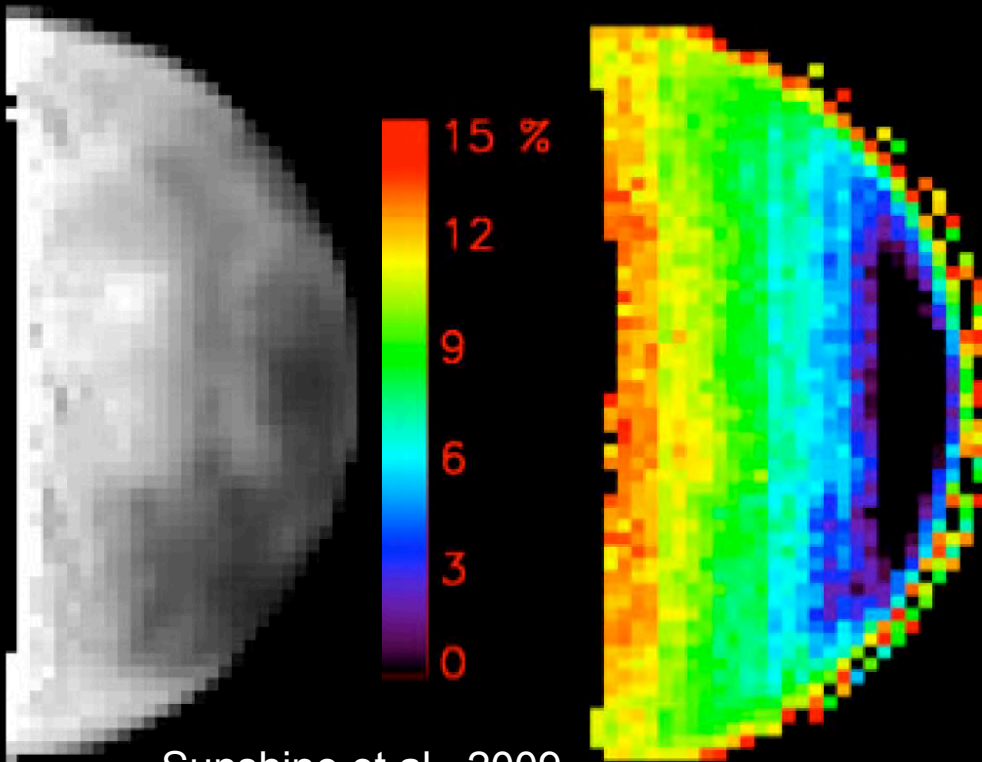
**APL**

*The Johns Hopkins University*  
**APPLIED PHYSICS LABORATORY**

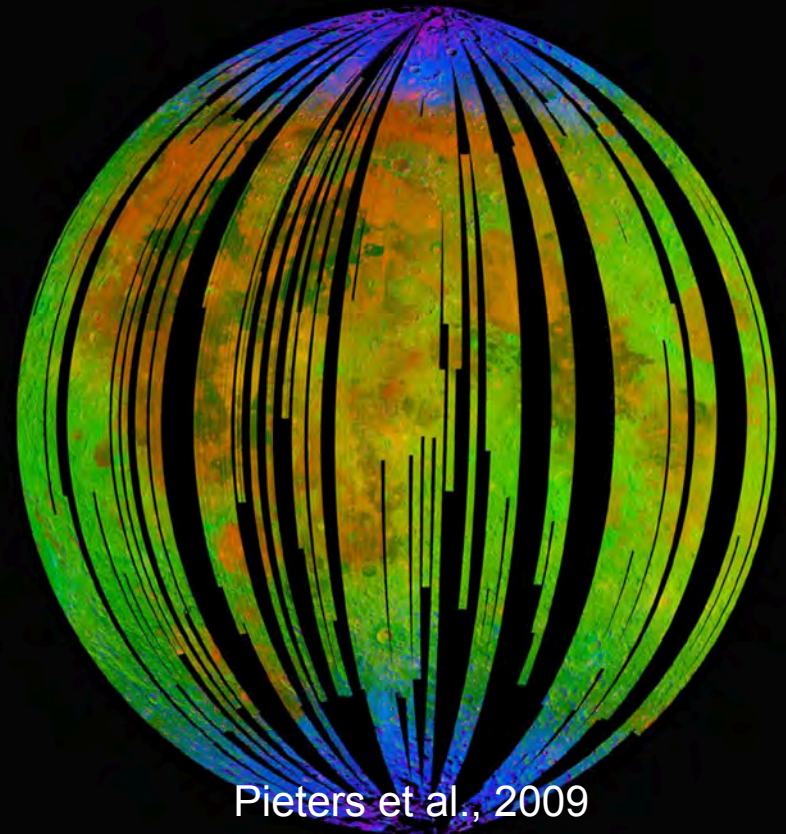
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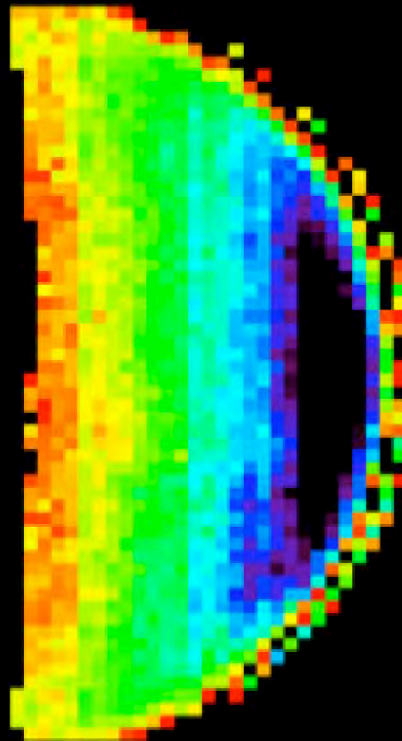
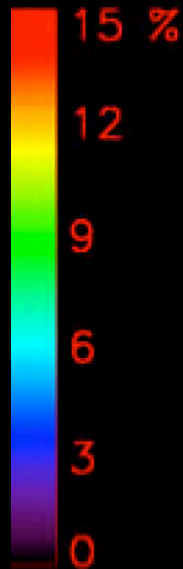
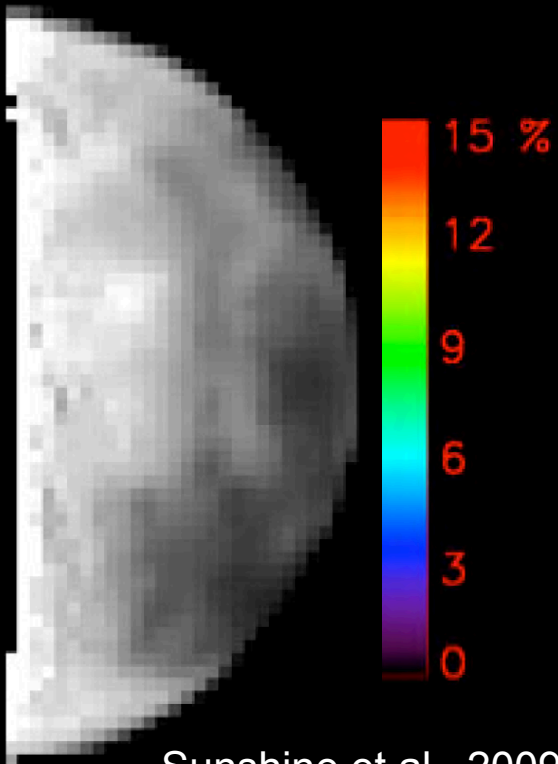
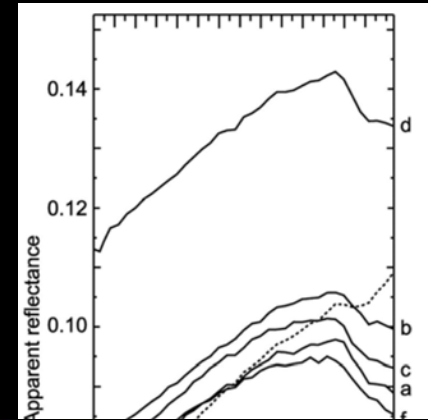
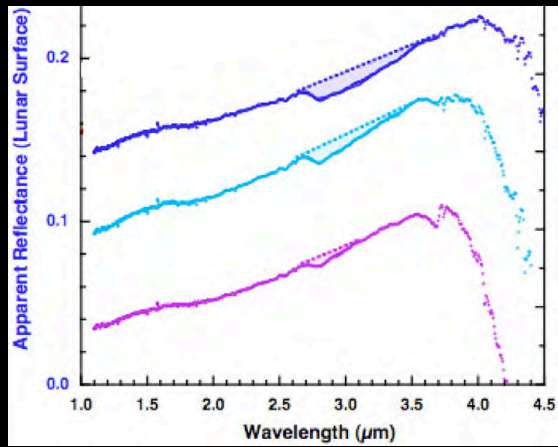
Sunshine et al., 2009



Pieters et al., 2009



# “Water” on the Illuminated Moon

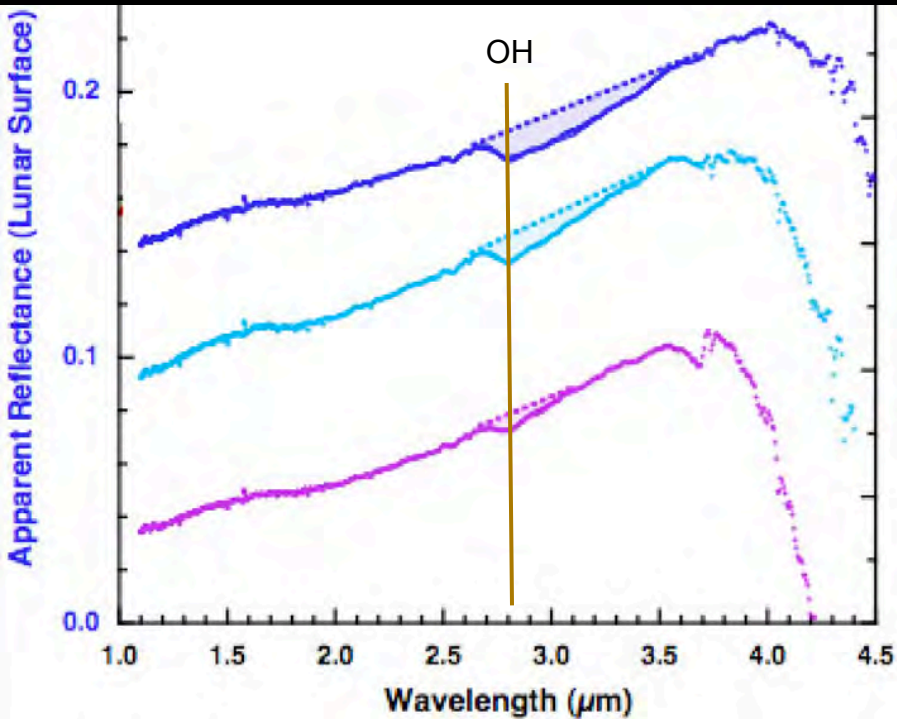


Sunshine et al., 2009

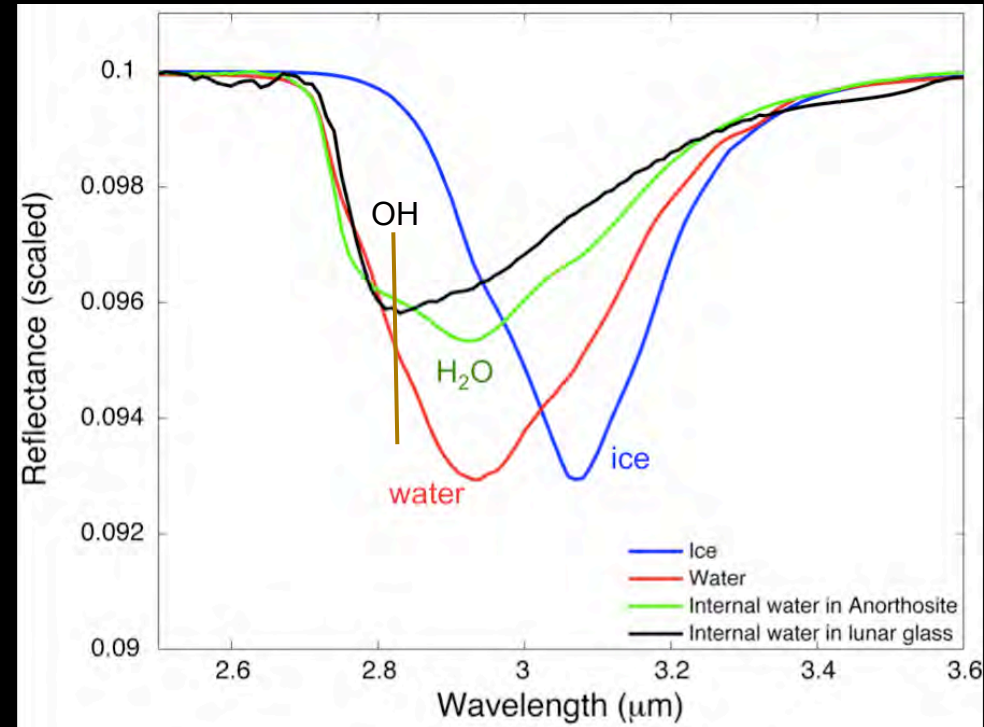
Pieters et al., 2009

# Hydroxyl (OH) does exist on the illuminated Moon

## EPOXI

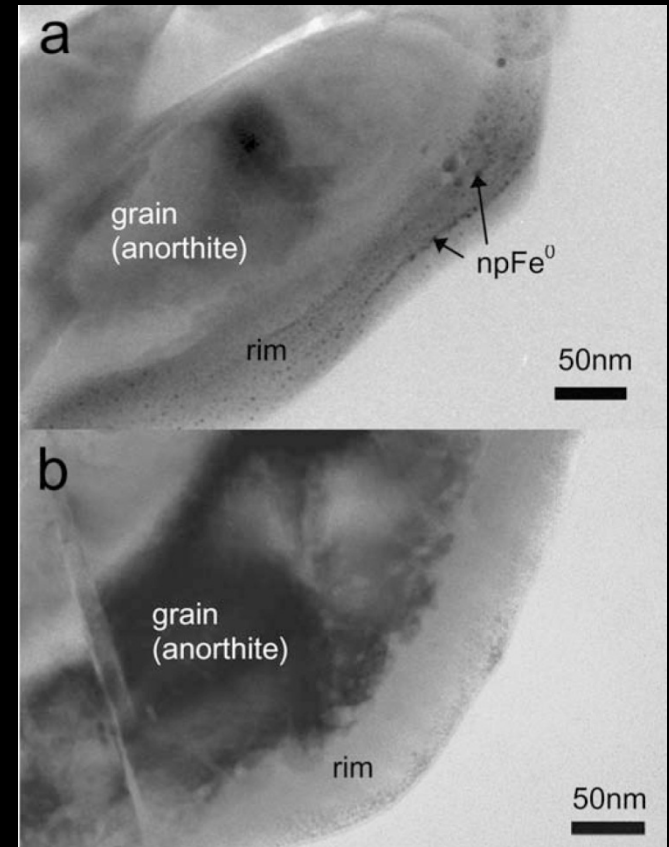
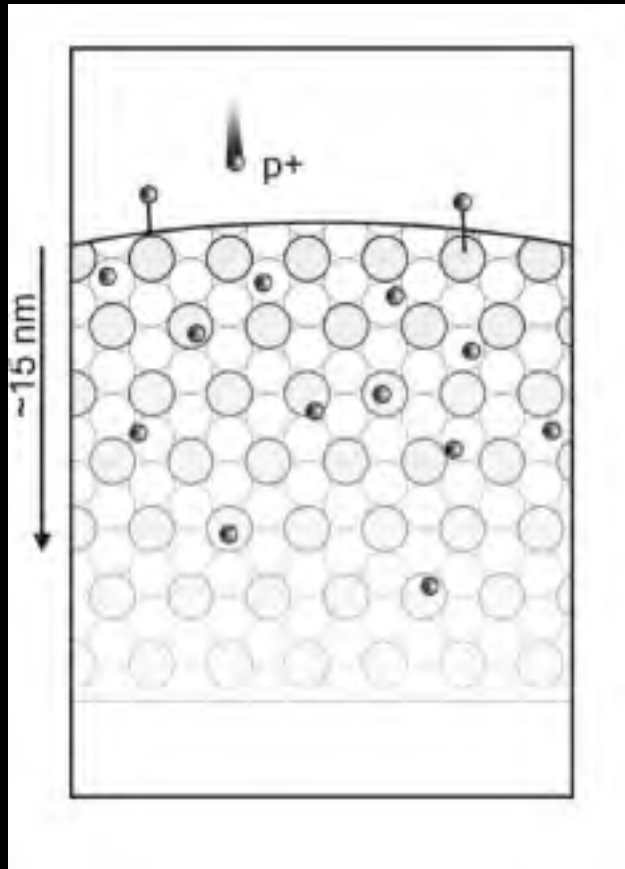


## Lab



# Possible formation mechanism of 'water' on the Illuminated Moon

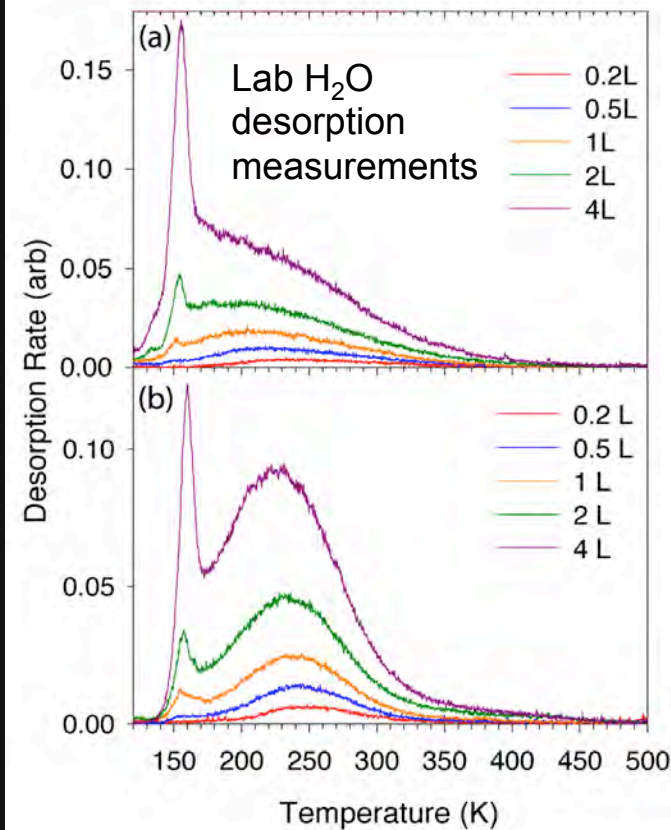
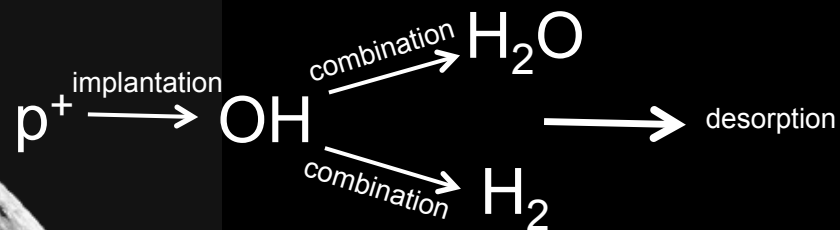
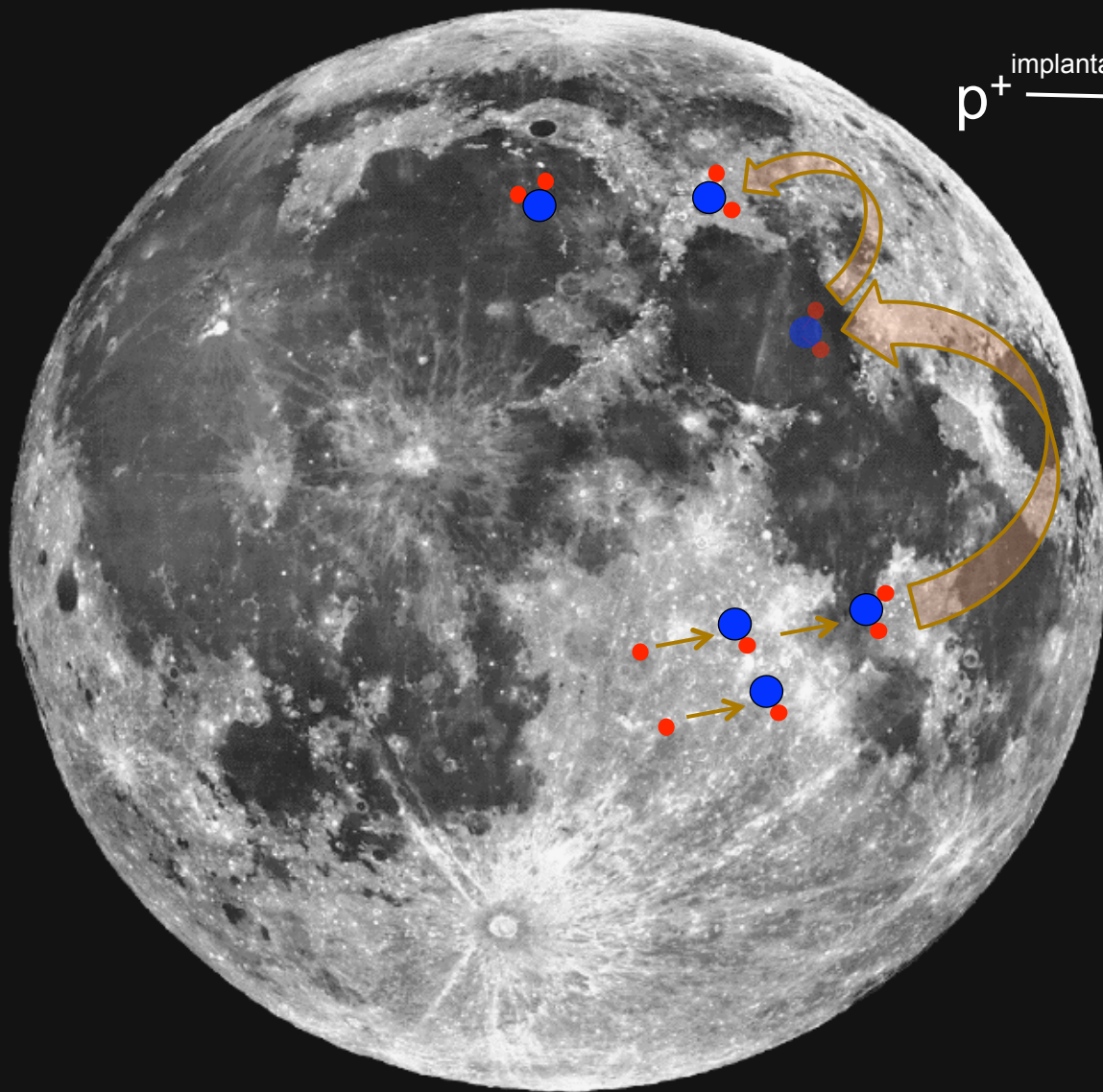
## Hydroxyl formation



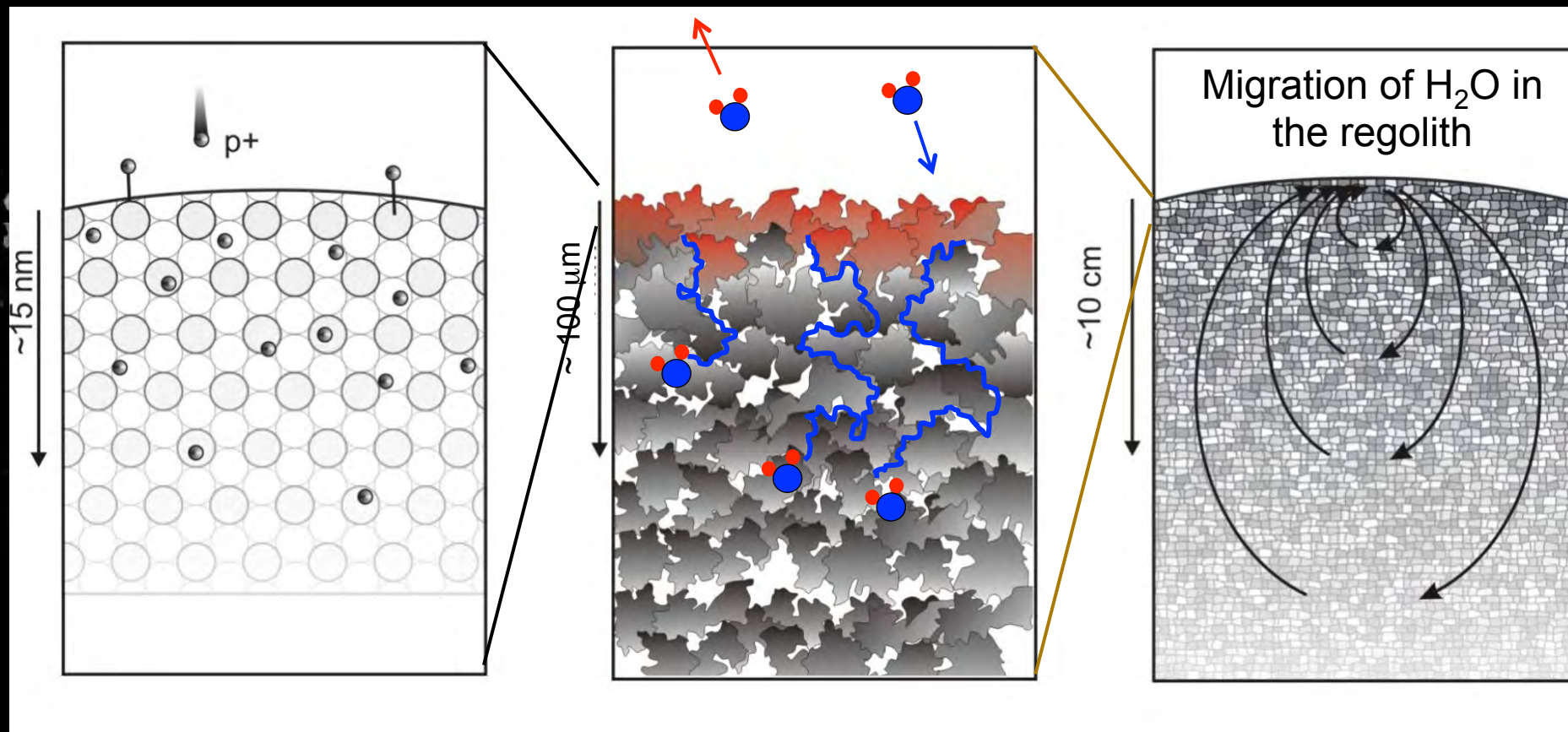
From Noble et al. (2005)



# Possible evolution of 'water' on the illuminated Moon



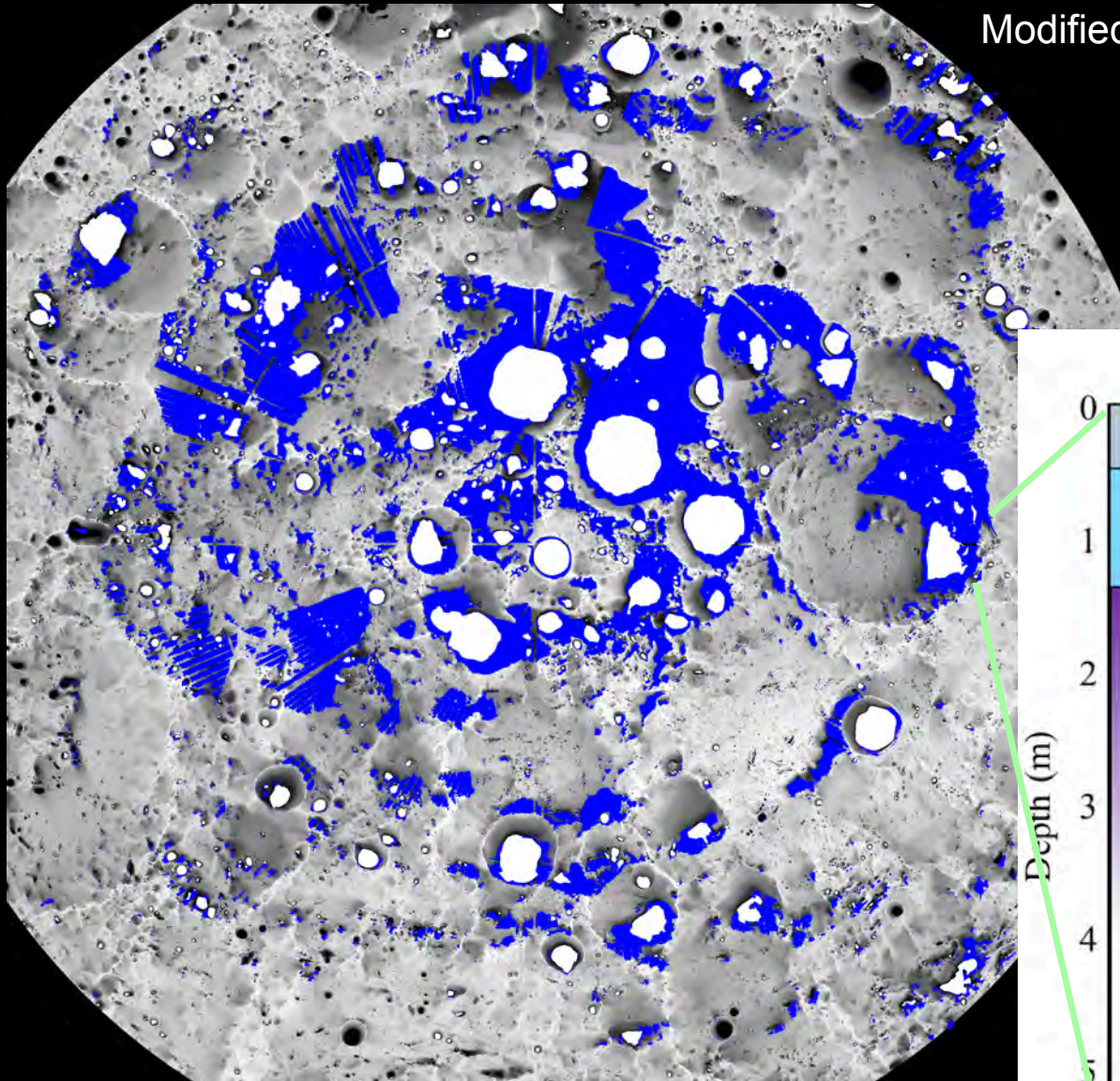
# Possible mechanisms for concentrating OH & H<sub>2</sub>O in the near surface



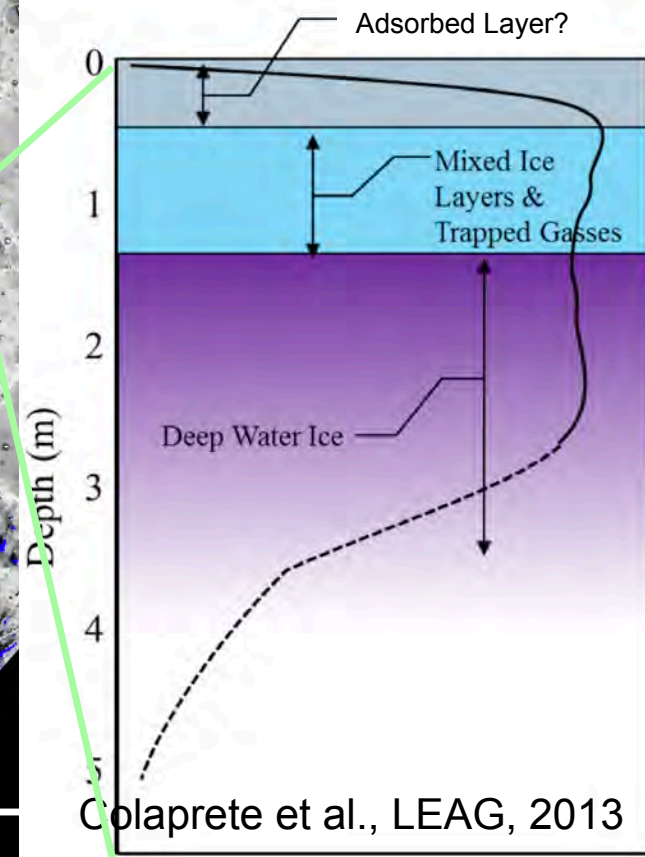


# Surface adsorbed water may exist...

Modified from McGovern, 2013



Adjacent to PSR

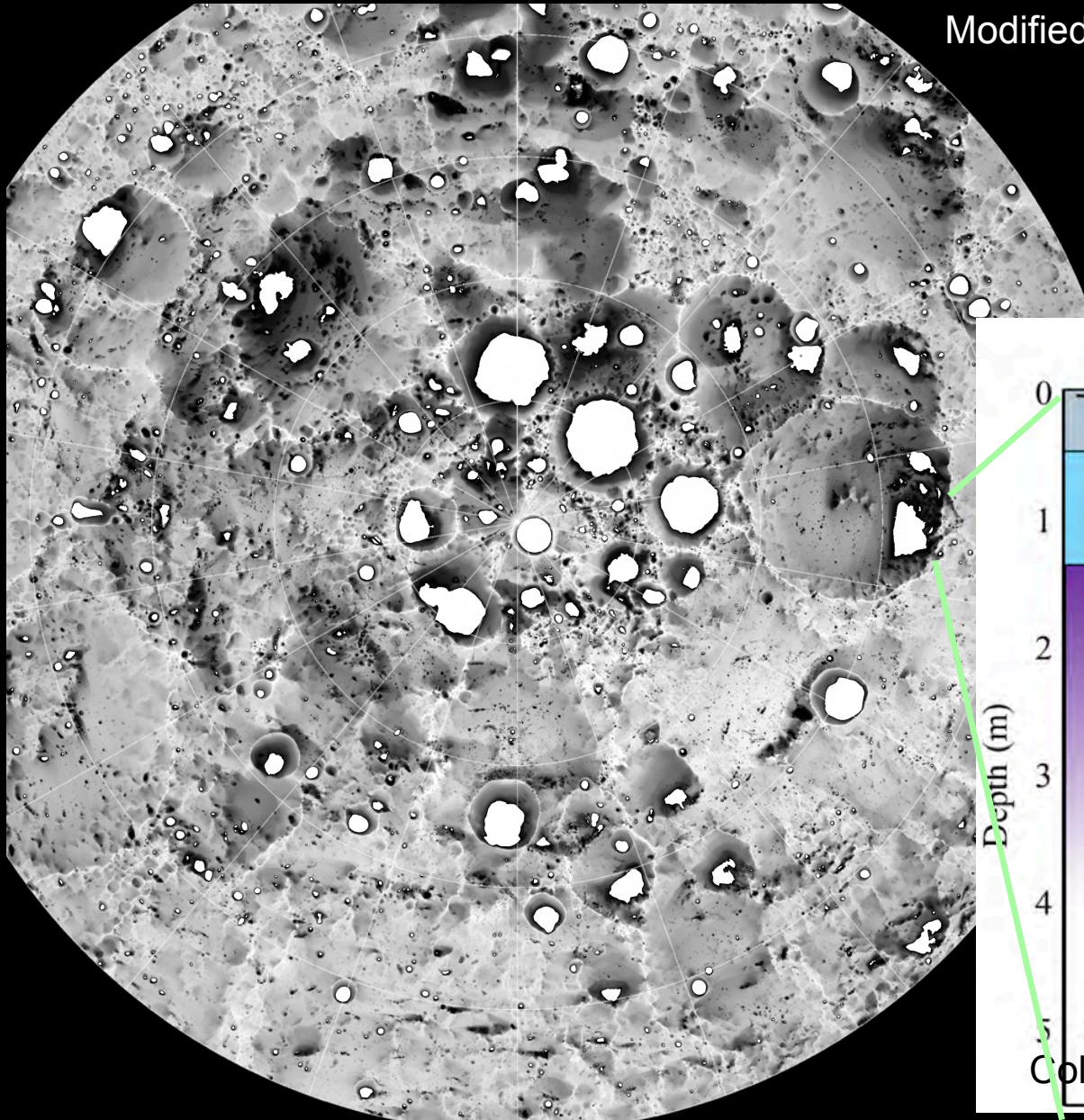


Colaprete et al., LEAG, 2013

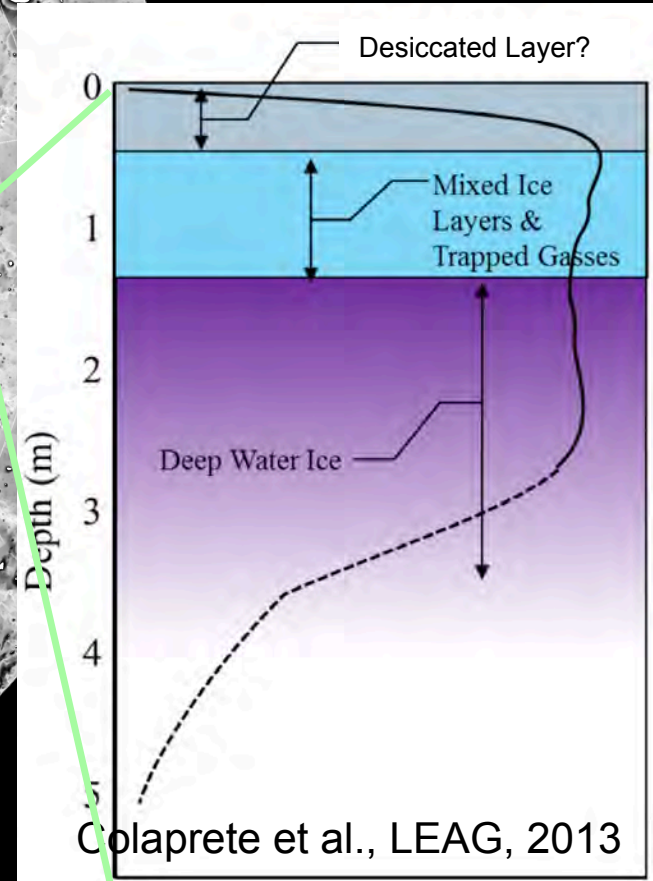


# Or the upper surface may be bone dry...

Modified from McGovern, 2013



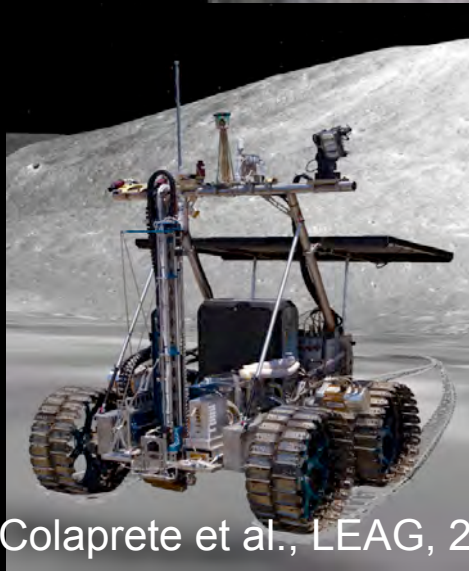
Adjacent to PSR





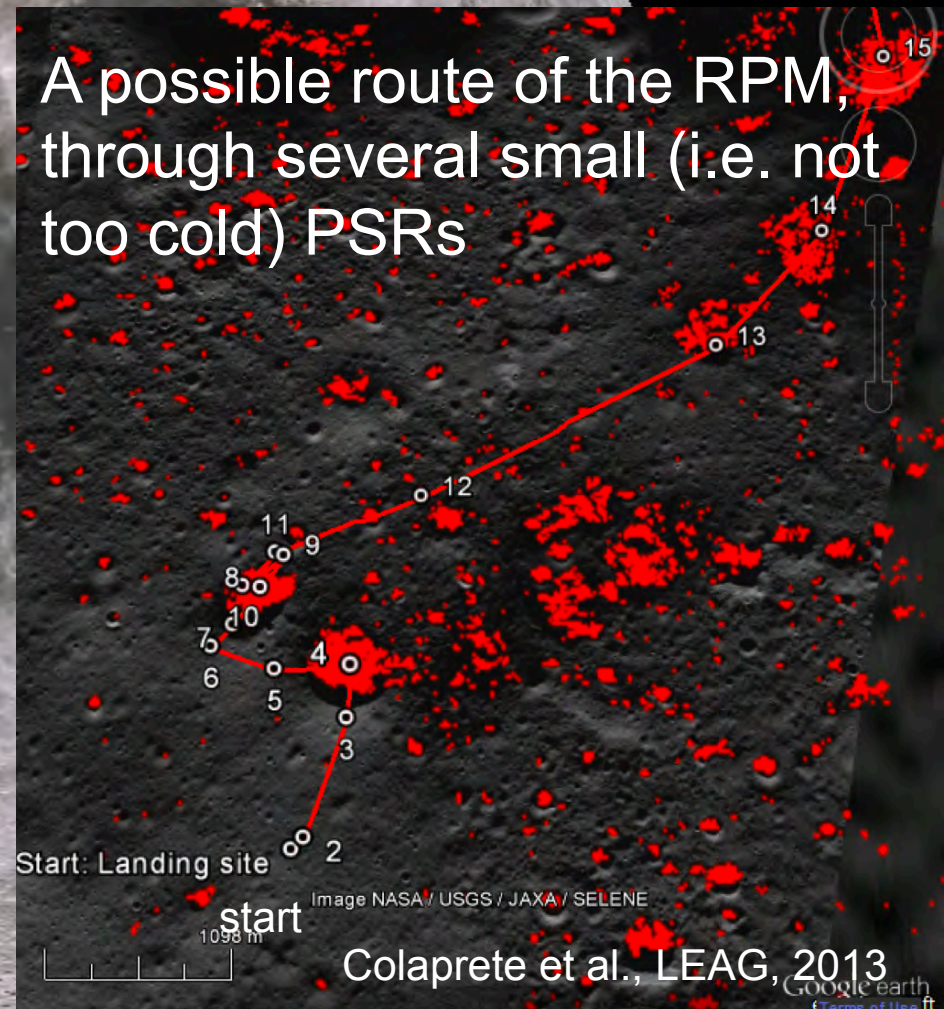
# A ground truth mission - RPM

...might not find water ice, but it may find significant adsorbed water



Colaprete et al., LEAG, 2013

A possible route of the RPM, through several small (i.e. not too cold) PSRs





## Goals for understanding water on the illuminated Moon

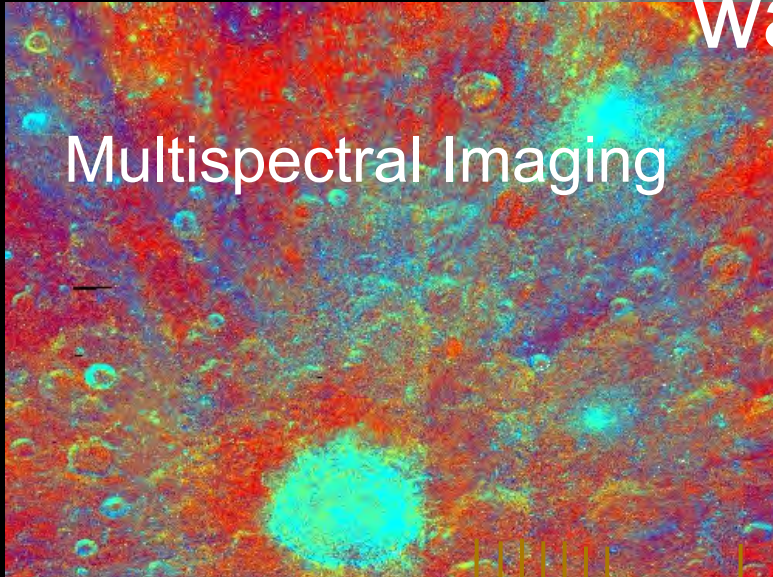
*Characterize the amount of, the chemical and physical state of, the distribution of, and evolution of (or life cycle of) water (OH and H<sub>2</sub>O) on the Moon.*

Scientifically interesting – answering these questions will have large implications for ‘water’ on ALL airless silicate bodies from Mercury to asteroids

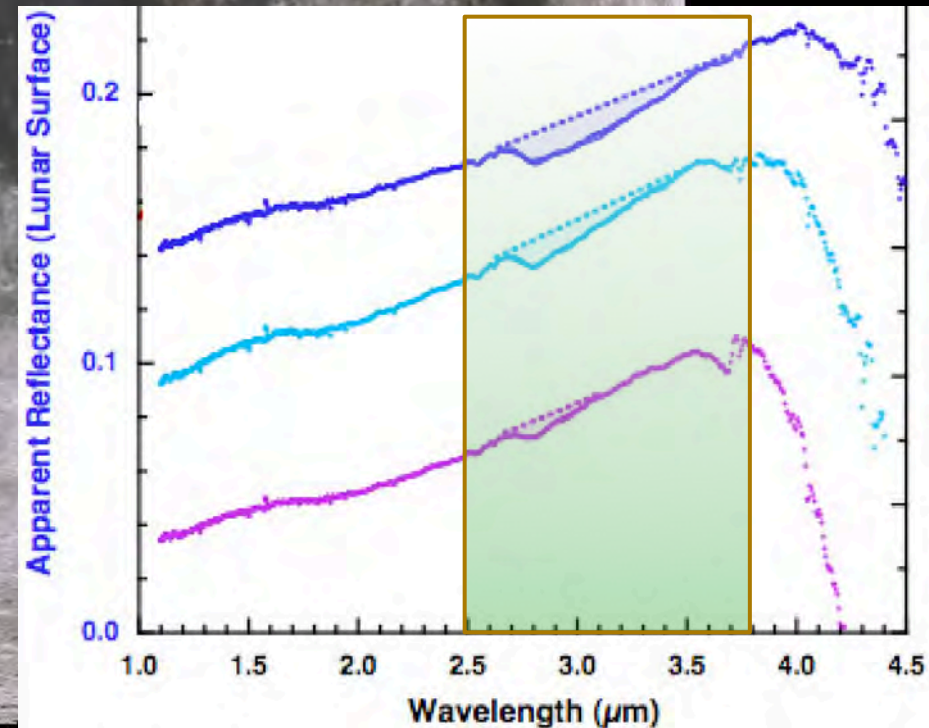
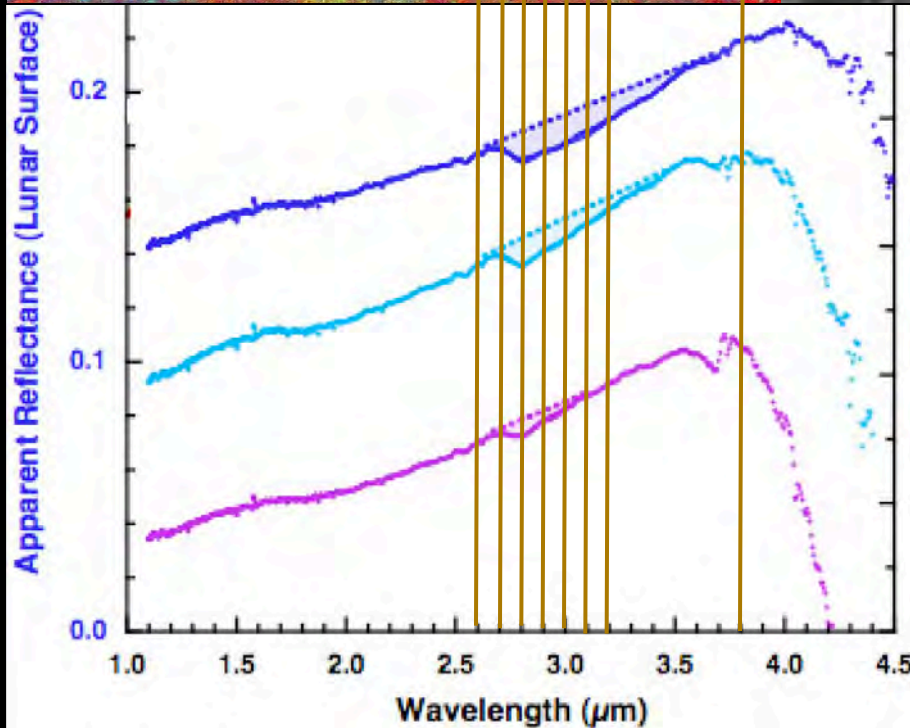
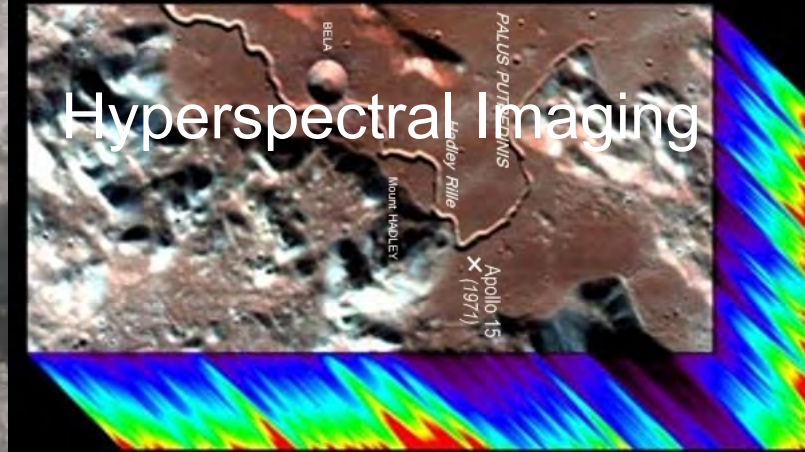
Exploration significance – two endmembers: extractable H<sub>2</sub>O or ephemeral OH. Knowing which exists (and relative abundance of each) has significant implication on our understanding of in-situ resources.

# Spectral measurements to characterize water cycle

Multispectral Imaging



Hyperspectral Imaging







## Long-duration stratospheric balloon payloads

5000 lb total payload  
instrumented for these measurements  
100K - 120K feet altitude (no telluric)  
~1" IFOV  
better than 1" stability  
2wk → 2mo duration  
no data volume limit  
potential 27/7 operations  
balmy summer Antarctic conditions



# Extra Slides



# Our understanding of the 'water'

## Observation

There is hydroxyl on the illuminated surface of the Moon

The band changes with time of day

Hydroxyl is found in pyroclastic deposits

Laboratory simulations produce hydroxyl in lunar samples and analogs

The flux of solar wind is  $3 \times 10^8$  /sec

## Inference

It is of solar wind origin

Either the 'hydroxyl' is being lost or it's a photometric effect

There is some primordial water in the Moon (few hundred ppm)

Solar wind can result in OH.

# Scientific and exploration relevant questions

## Unknown

Is the 'water' on the illuminated Moon due to solar wind implantation?

Is there molecular  $\text{H}_2\text{O}$ , in addition to OH, on the illuminated Moon (some tantalizing evidence from M3 and EPOXI)?

What is the evolution of the 'water'? Does its abundance cycle over a lunation? If the abundance decreases, by what mechanism does it devolve?

Is there a concentration of either OH or  $\text{H}_2\text{O}$  at higher latitudes, or are the observations a photometric effect?

## Implication

A potentially renewable resource. May also be ephemeral

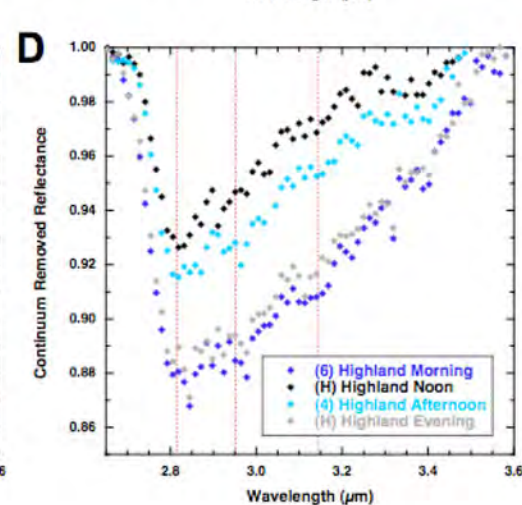
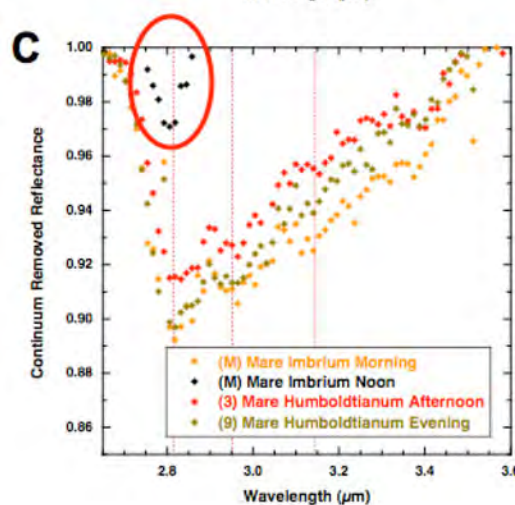
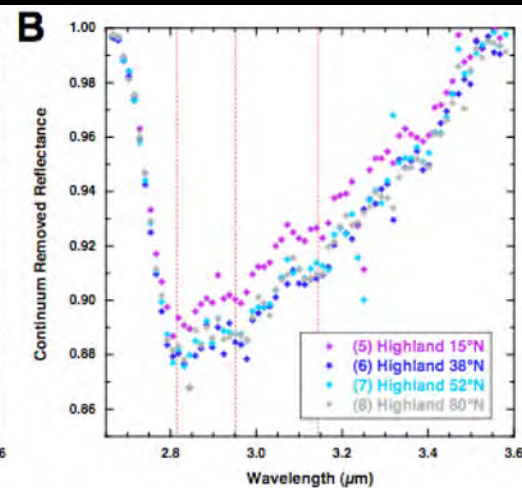
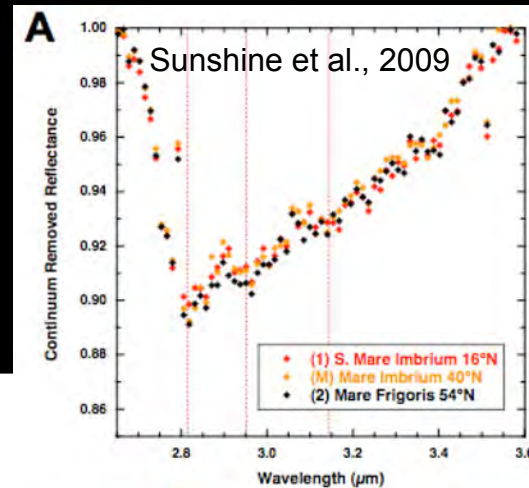
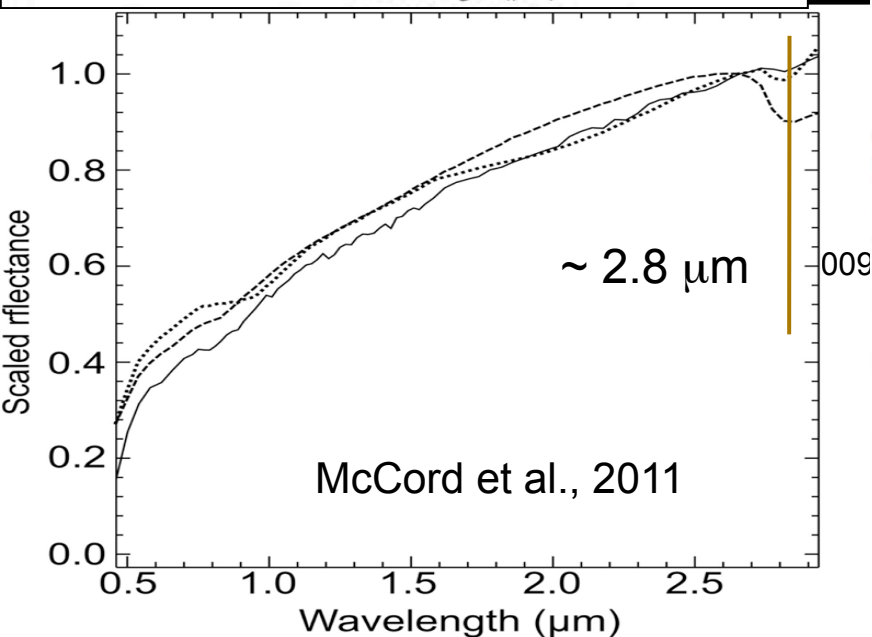
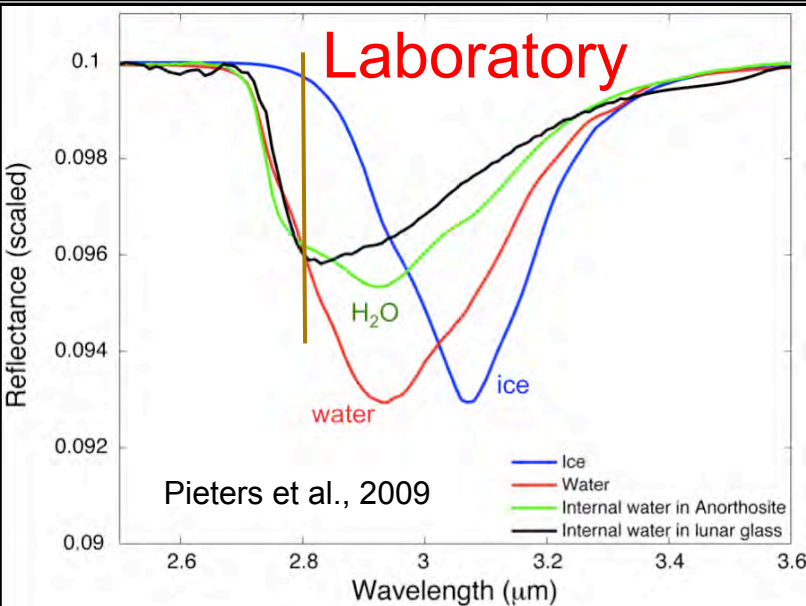
Potential easily extracted resource.

Evolution by  $\text{H}_2$  implies a poor resource. Evolution by  $\text{H}_2\text{O}$  implies a possible accumulation in cold traps.

If  $\text{H}_2\text{O}$  is present in illuminated high lats., may represent convenient resource, and possibly suggest continual formation of  $\text{H}_2\text{O}$



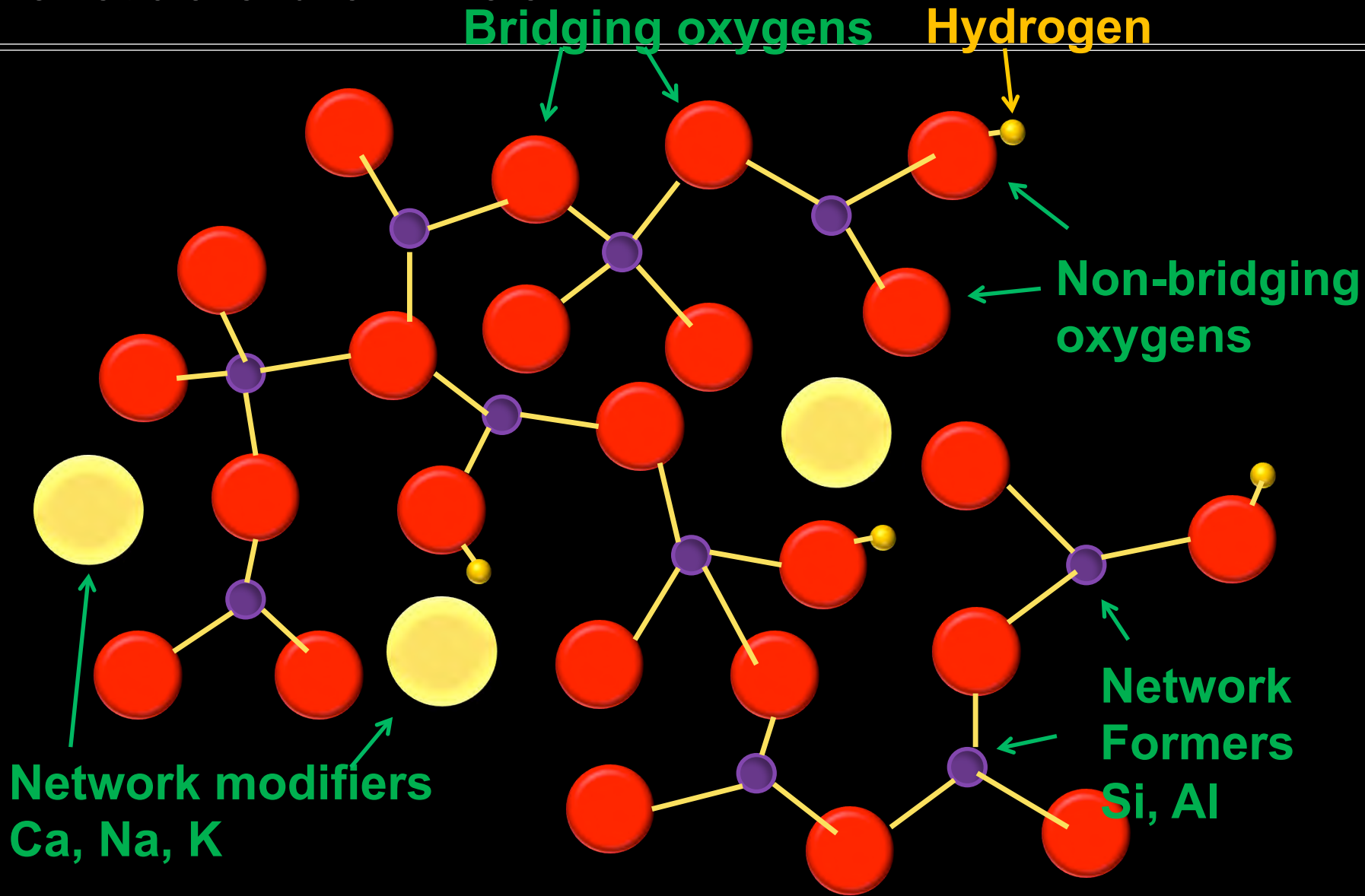
# Infrared Observations of Water (or at least OH)



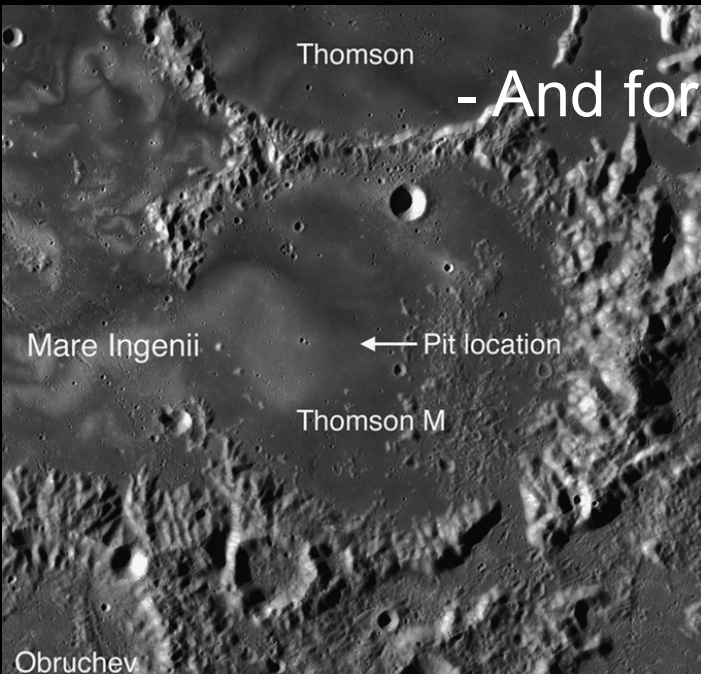




# Schematic of a lunar mineral



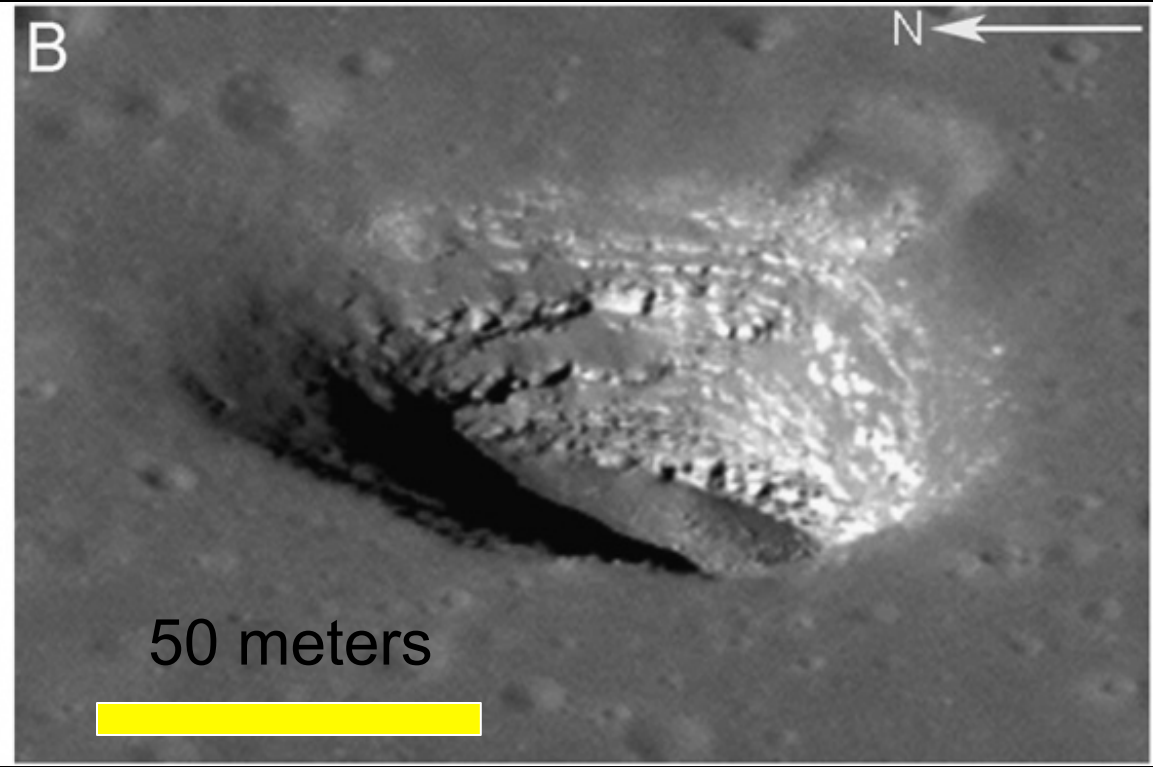
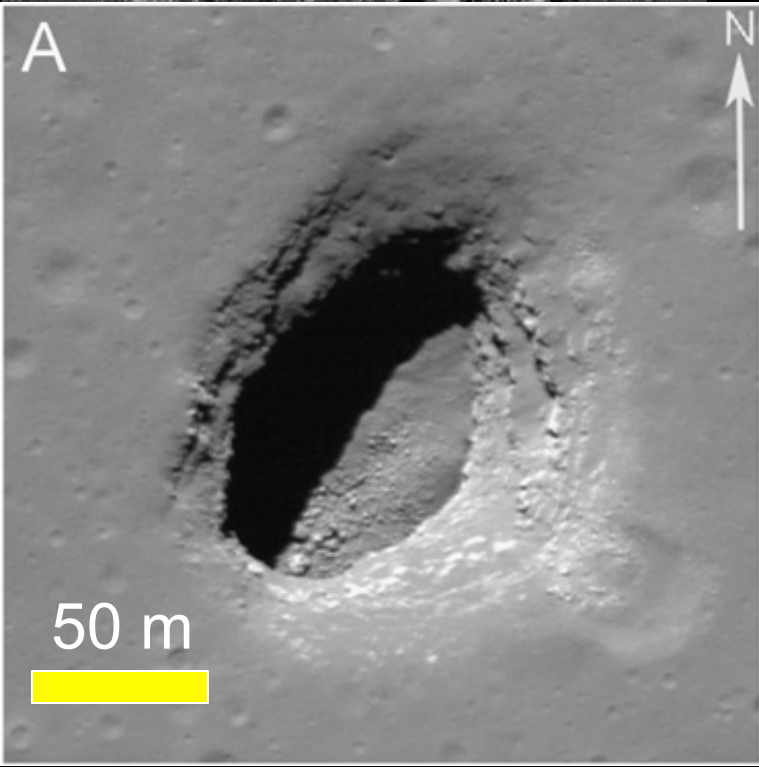
High alkali contents create non-bonded O where H can bond...more H.  
High alkali contents should imply high H adsorptivity!



- And for the not too adventurous water molecules

Pit craters may offer a shaded recesses cool enough to allow small amounts of water to accumulate

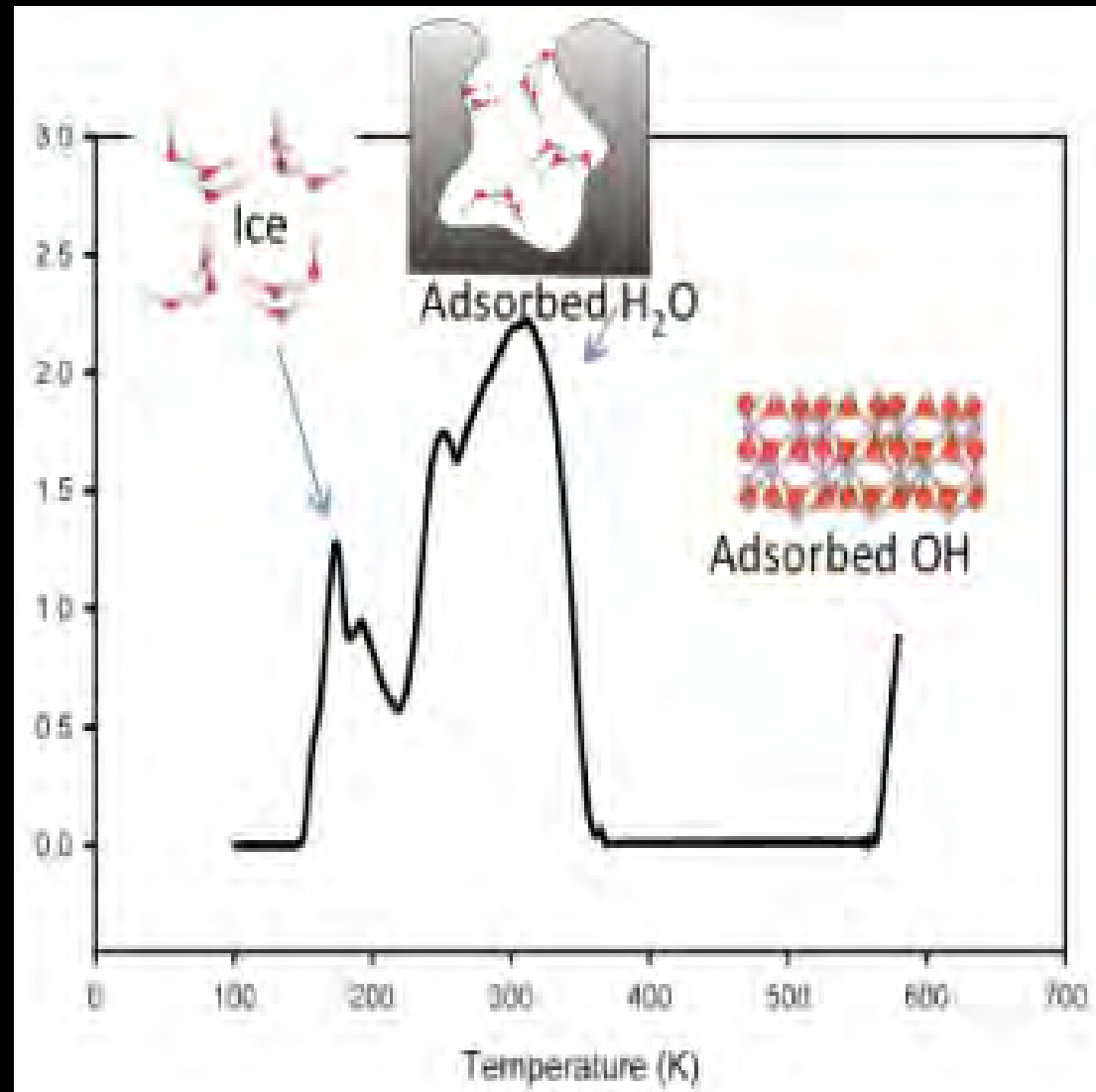
Robinson et al. (2012) *Planetary & Space Science* **69**, 18-27.





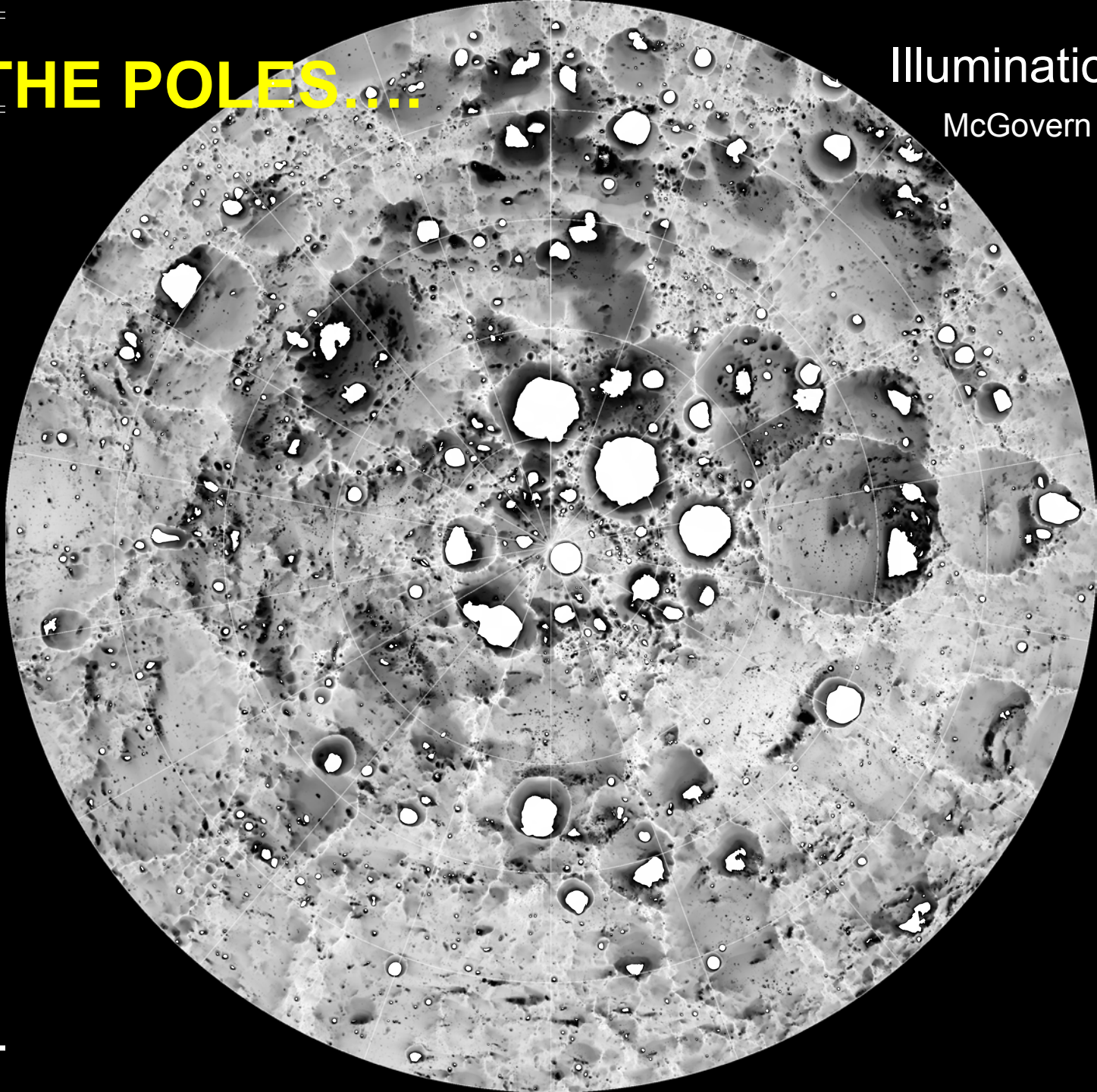
# In what ways does water stick to soil?

- Bulk physisorption of various phases ( $\sim 150\text{K}$ ).
- Low energy molecular chemisorption ( $\sim 150$  to  $250\text{K}$ ).
- High energy molecular chemisorption ( $\sim 250\text{K}$  to  $350\text{K}$ ).
- Dissociative chemisorption of  $-\text{OH}$  groups (usually  $>500\text{K}$ ).



# AT THE POLES....

Illumination map  
McGovern et al., 2012

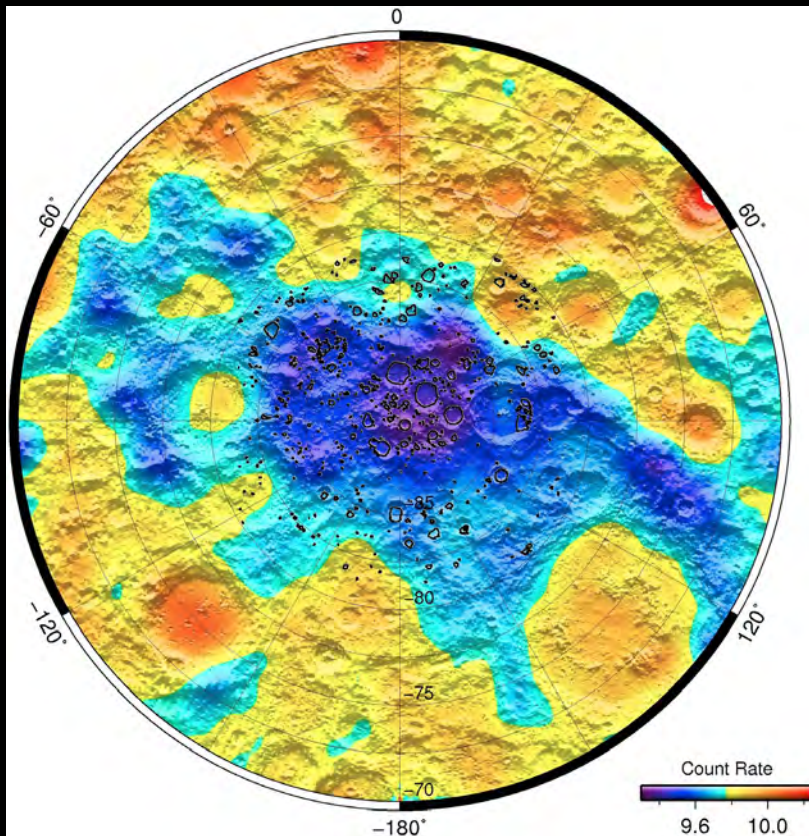




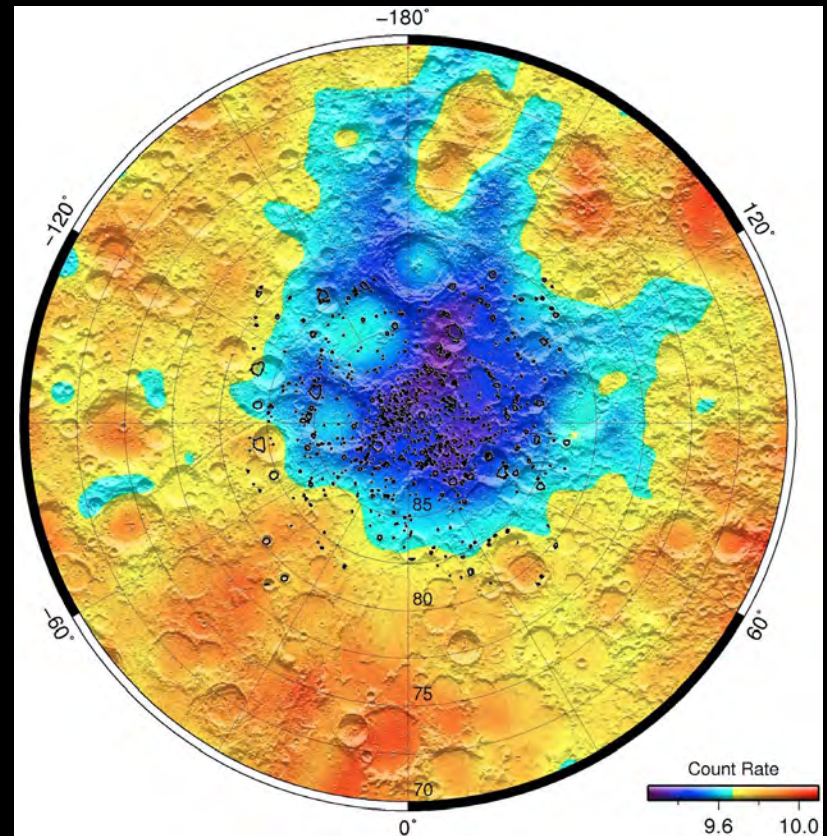
# Large Hydrogen concentrations exist even deeper

H-rich areas down to 70° latitude! (up to 1m deep)

South Pole



North Pole





# Diffusion and Desorption

**Diffusion:** 'slow', but happens more easily (at lower temperature). Requires "bridging" contact surface. This limiting area is the reason for the poor thermal conductivity.

**Desorption:** 'fast', but requires the  $\sim 2\times$  the energy.

## Diffusion:

1. at 250 K diffusion "lifetime"  $\sim 10^{-7}$  s  
path goes as particle radius<sup>2</sup>
2. Site to site hop distance  $\sim 3$  Angstroms
3. Assuming  $100\text{ }\mu\text{m}$  grain ( $10^{14}$  A) and assuming non-infinitesimal contact area between grains,
4. Result: can take  $\sim 3 \times 10^{18}$  hops or  $10^4$  seconds

**Desorption:** at 250 K  
desorption lifetime  $\sim 0.3$  s

