Understanding the Atmospheres of Hot Earths...

Progress in the lab!

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Recap of what we've been doing:

- Matrix isolation of samples using argon ice
- Fourier transform
 infrared spectroscopy
- Samples (komatiite):
 - SiO₂
 - MgO
 - CaO
 - Al_2O_3
 - FeO



Our (old) system

Focus GmbH EFM 4 UHV Evaporator (e-beam)





Adding H₂O

- Any water added to the crucible will evaporate preferentially long before the oxide rock sample even melts.
- Waters of hydration: evaporating water from a hydrate rock (efflorescence) with increasing temperature
- $(SiO_2)_{10} \cdot 14H_2O \rightarrow SiO_2 + 14H_2O$ (Water evaporates at only ~200 C)

Sealed Crucible

Idea: waters of hydration are confined to the crucible until the appropriate temperature.



 Pressure inside crucible causes foil seal to burst at a specific temperature.

 Seal between foil, crucible, and lid is made with a press.

Sealed Crucible

- Foil must be same material as crucible so it doesn't pop from thermal expansion
- Tested using a crucible with gas pressure added through the bottom.
- Problem: unable to make an airtight seal using refractory metal foils (ideas?)



Komatiite Diffusion Barrier

- Idea: creating a barrier of fused sample that will not permit diffusion of waters of hydration until the sample fully melts.
- This can be achieved by flashing the filled crucible at a temperature high enough to melt the surface grains in an inert atmosphere (argon).



[type of laser] laser. Not yet tested.

SiO₂ Atmosphere IR Spectrum

SiO₂ was evaporated and deposited into an Ar ice.



SiO₂ Atmosphere IR Spectrum

- Each peak appears as a triplet due to the presence of Si isotopes:
 - Si28 (1218 cm⁻¹)
 - Si29 (1222 cm⁻¹)
 - Si30 (1226 cm⁻¹)



SiO₂ Atmosphere IR Spectrum

- Measured constituents of pure silica atmosphere by peak intensity:
 - SiO dimer, SiO, SiO trimer, SiO₂, Si₂O₄, SiO₃
 - Possible SiH_4 and O_3 .

 Oxygen gas cannot be detected through IR spectroscopy but thin-film deposition and EDS/EDX measurements demonstrate a depletion of oxygen from the material in the crucible (preferential evaporation of oxygen).

Is this representative of an equilibrium atmosphere?

- Matrix isolation spectroscopy is claimed to work at ratios of 1000:1 argon to sample and at low temperatures
- We need to verify that no diffusion occurs in order to present matrix isolated results as equivalent to material in the crucible that is still at an equilibrium.
- Experiment: time evolution of spectra at various temperatures and pressures of argon

Problems!

Contamination feature began to appear:



- Identified as tungsten oxide, most likely residue from thoriated tungsten filament.
- Need to develop a cleaning technique to remove contamination from evaporator.

More Problems!

- Power supply failure: both power supplies (EVC100 and NGEFM) are unable to provide the desired current and voltage outputs to obtain high temperatures
- Solution: design our own evaporator that doesn't constantly fail and contaminate itself!
 - Thoriated tungsten electron source for e-beam evaporator doubles as a contamination source.
 - E-beam evaporator power supplies are expensive to repair and would not be able to melt some of our materials even when fixed.

New Designs

New vacuum chamber geometry will allow for simultaneous growth of samples, measurement of sample thickness, and infrared spectroscopy.



Optical System



Optical System

- Multi-pass optics to increase signal for inbeam spectroscopy.
- "Traditional" matrix isolation spectroscopy can still be done, as well.
- Still in the design phase.



Evaporator

- Resistive heating:
 - Higher temperature range
 - More reliable
- Crucible:
 - Larger sizes possible for better volume:aperture ratio or higher flux
 - On a bottom-loading flange like previous design
 - Water-cooling shroud



We could use some help with...

- The problem of adding water to a rock atmosphere remains.
- Water must be same temperature as komatiite sample.
- Nate's design: inner/outer crucible... how does it work?

Diffusion studies in Ar matrix isolation at low temperatures

- We have demonstrated that at temperatures below 18 K, no diffusion of CH_4 atoms takes place within an argon matrix over a period of 20+ hours, even at ratios of less than 1000:1 Ar:CH₄.
- Beginning at 19 K under ultra-high vacuum pressures (approx. 10⁻⁹ torr), argon atoms begin to leave the ice matrix allowing diffusion to take place.

Temperature-correlated diffusion in argon ice matrix

8.38 -8.36



Other Work

 MATLAB code for modeling planetary atmospheres as seen by telescope CCD





Other Work

 MATLAB code for modeling planetary atmospheres as seen by telescope CCD





Other Work



• Filters/bins modeled