

Thermochemical Approaches for the Characterization of Materials

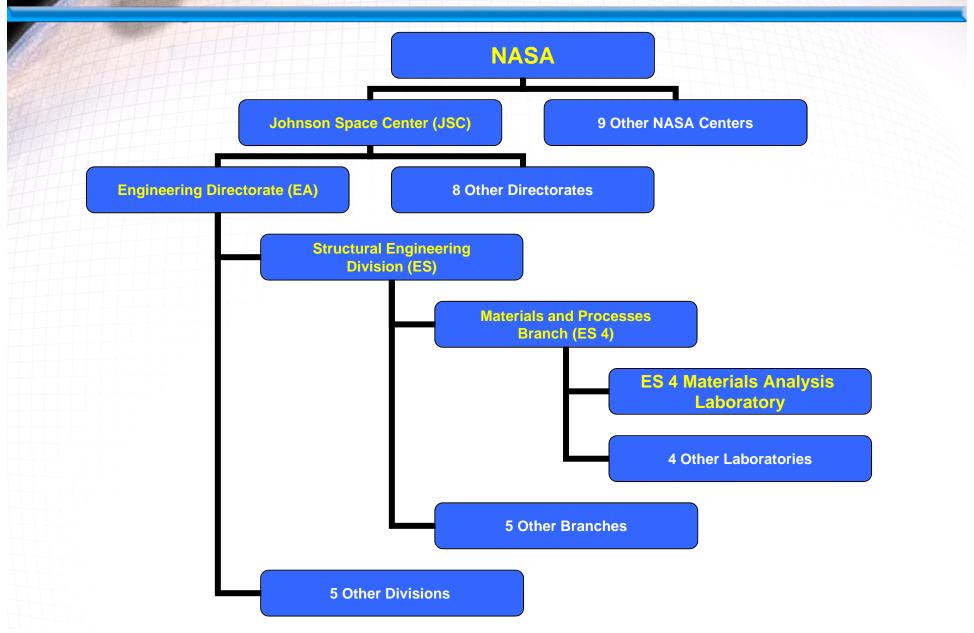
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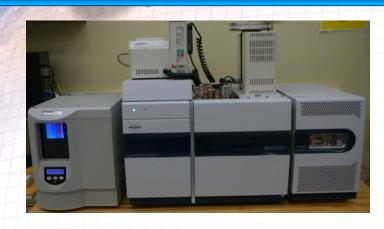


National Aeronautics and Space Administration





Analytical Chemistry Laboratory Equipment







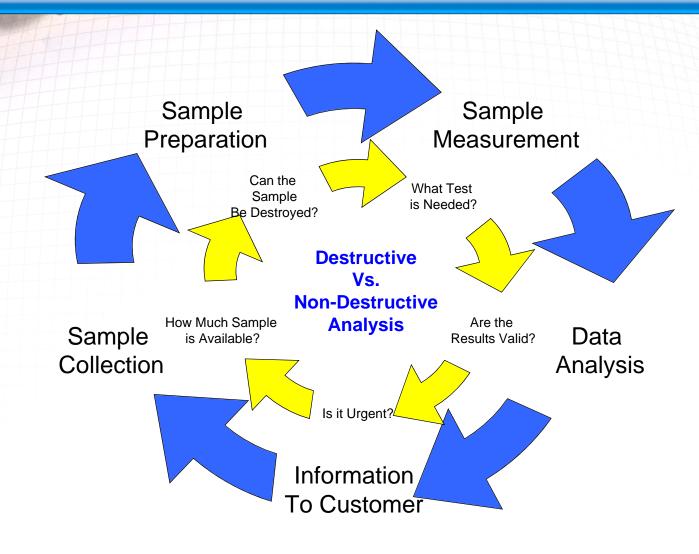
Key Laboratory Equipment

- Optical Instrumentation
 - UV-Vis, Fluorimeter, Solar Reflectance, Infrared Emittance, Raman
- Thermal Analysis Instrumentation
 - DSC, DMA, TGA, TMA, LFA, Rheometer
- Chemical Analysis Instrumentation
 - FT-IR, Ion trap GC-MS, Py-GC-MS, TGA-MS, TGA-IR



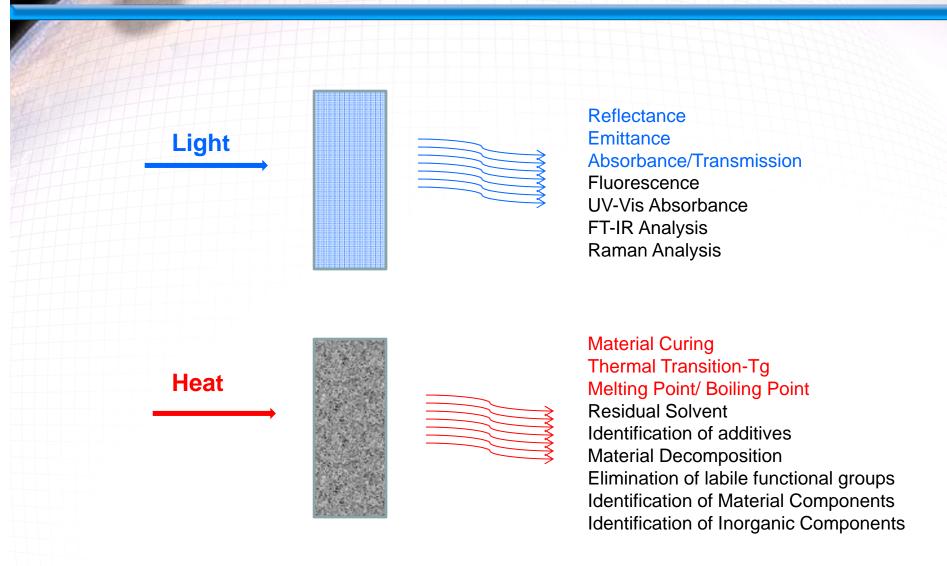


The Analytical Chemistry Cycle





Optical Vs. Thermal Techniques



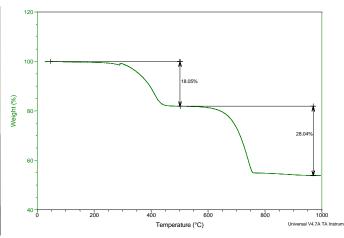


Thermogravimetric Analysis (TGA)

- A TGA instrument consists of an analytical balance and a furnace.
- A small sample of material is heated and its change in mass is measured as a function of temperature.
- Experiments can be conducted under inert or oxidizing atmospheres.
- Information gained from TGA includes:
 - Thermal stability for conducting additional thermal analysis
 - Identification of the number of components in the sample if the decomposition temperatures are different
 - Residual mass for assessing the extent of inorganic additives

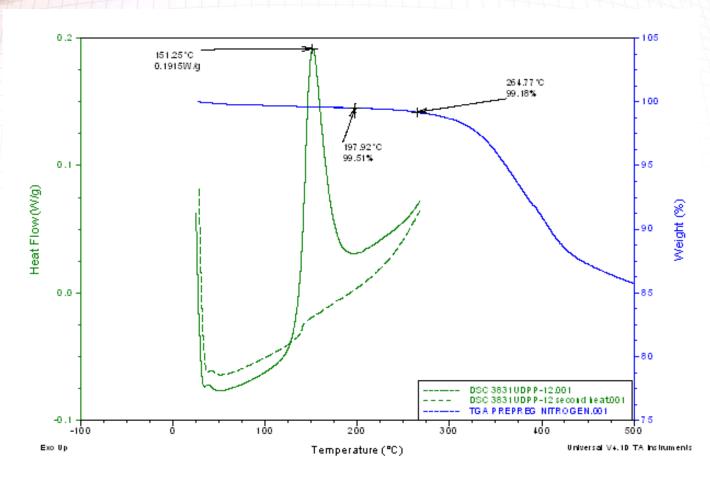








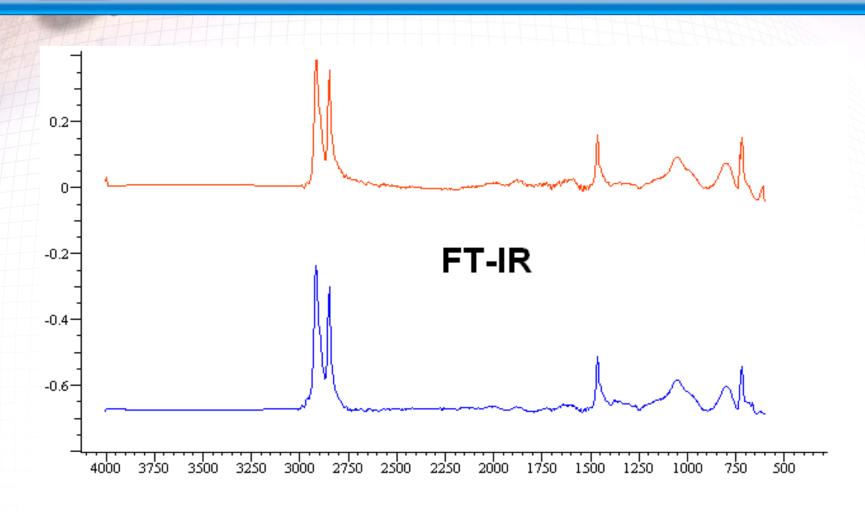
Thermal Analysis of Composite



Traditional thermal analysis of materials is performed by DSC (Differential Scanning Calorimetry) and by TGA (Thermogravimetric Analysis) instrumentation. Most Thermal Analysis is performed at temperatures below the onset of decomposition. Thermochemical Techniques for Material Characterization will utilize elevated temperatures until the material is fully degradated.



Spectral Analysis of HDPE and LDPE



Many industrial laboratories have only one technique available for characterization of the manufactured product. In many situations, one type of analytical technique is not adequate for assessing the product.

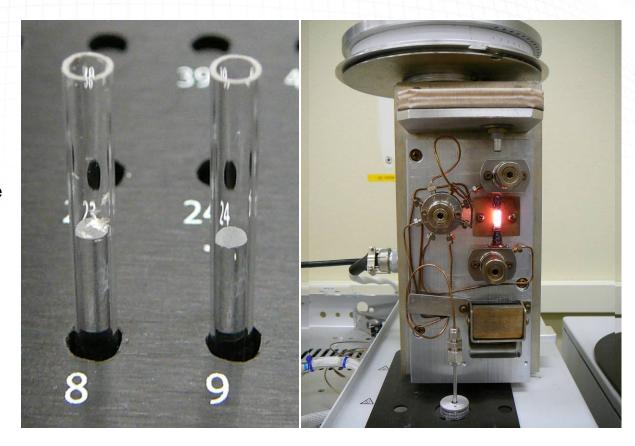


Pyrolysis for GC-MS of Solids

Sample size is relatively small:

50 to 200 μg is sufficient for solids 50 to 200 nL is sufficient for liquids

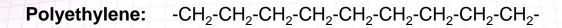
- Sample preparation is easy:
 Place sample inside 1.5 inch quartz tube containing filler tube and plug with glass wool.
- Samples can be solids, gels, viscous liquids, greases, crystalline, emulsions, foams, fabrics
- Pyrolysis temperatures are almost instantaneous
- Sample components can be quantified with the use of software

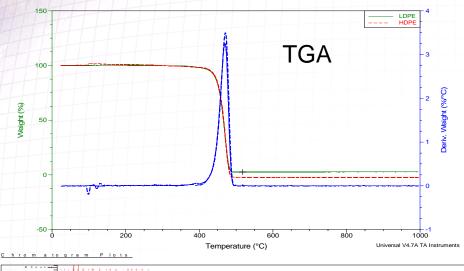


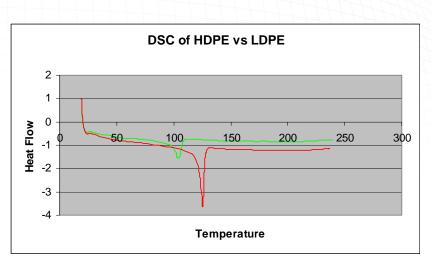
Pyrolysis is the thermal degradation of any substance through the fast application of heat.

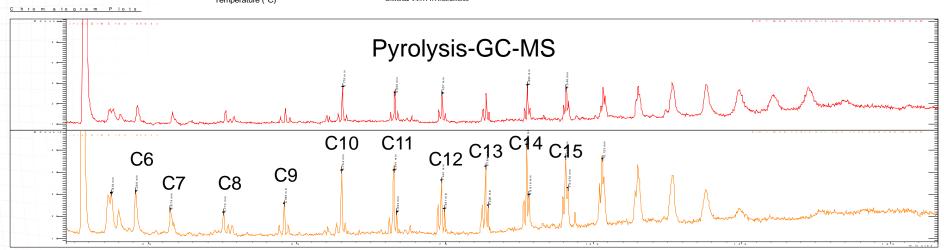


Thermal Analysis of HDPE and LDPE



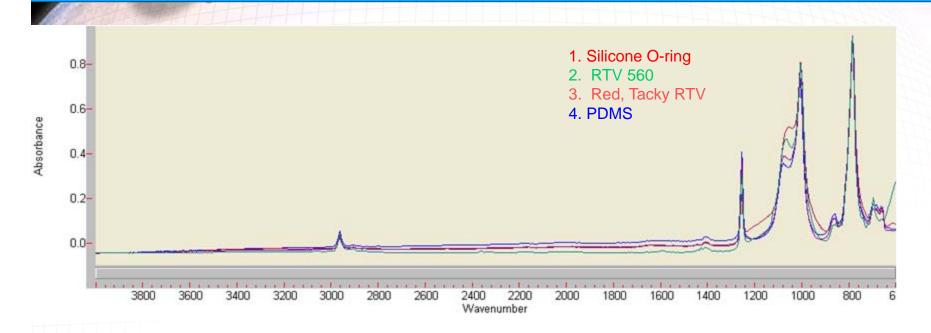




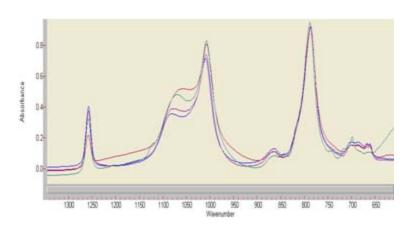




FT-IR Analysis of Silicone Materials

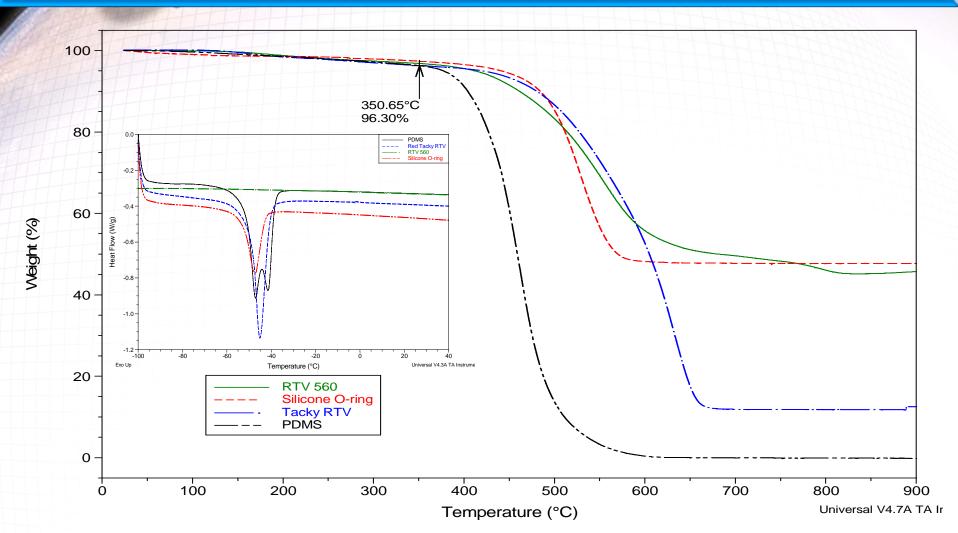


FT-IR is a non-destructive technique that is very diagnostic. However, if infrared light cannot penetrate the sample, any signal obtained through reflectance is only valid for the external surface of a sample.





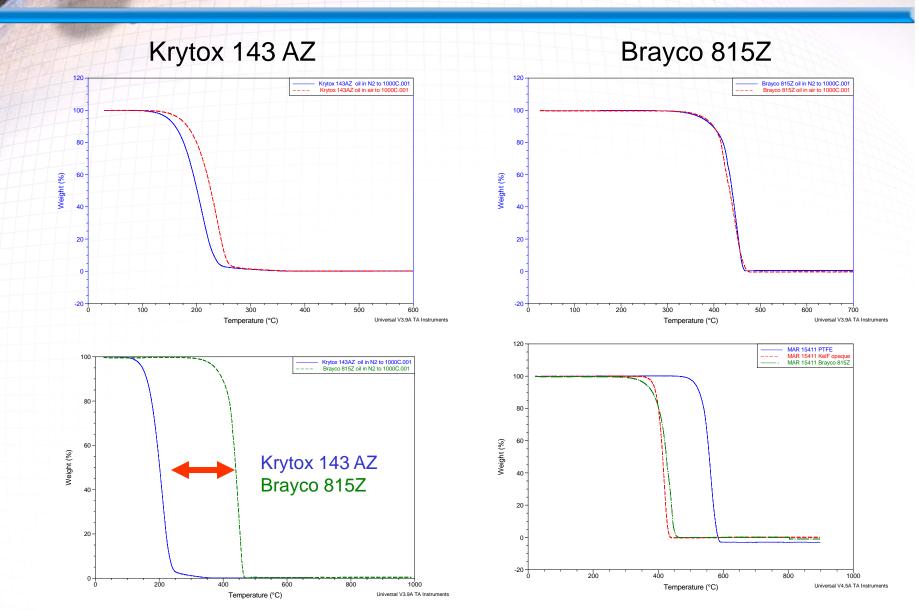
Thermal Analysis of Silicone Materials



The Silicone samples that were nearly identical by FT-IR displayed very different properties by thermal analysis.

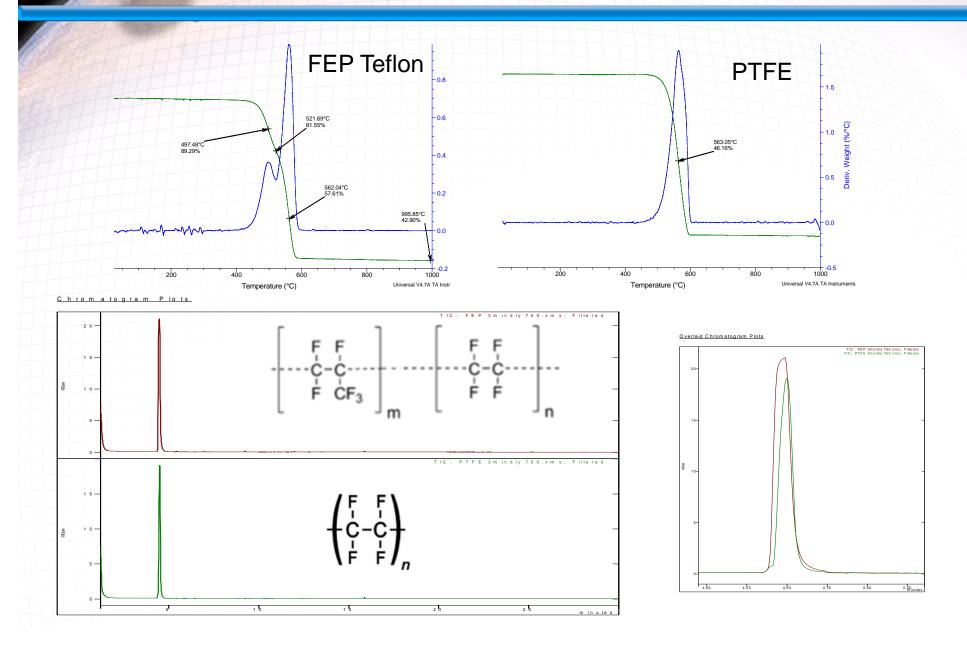


TGA Analysis of Fluorinated Materials



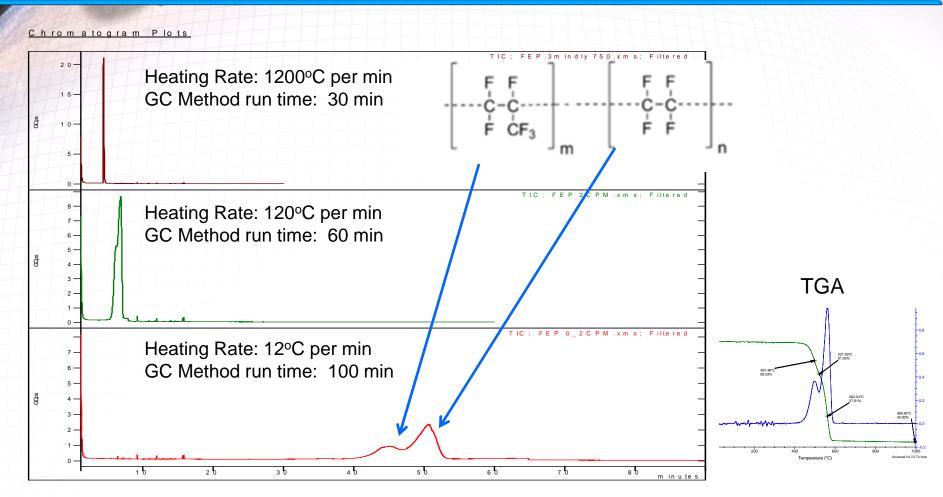


FEP Vs. PTFE Teflon



NASA

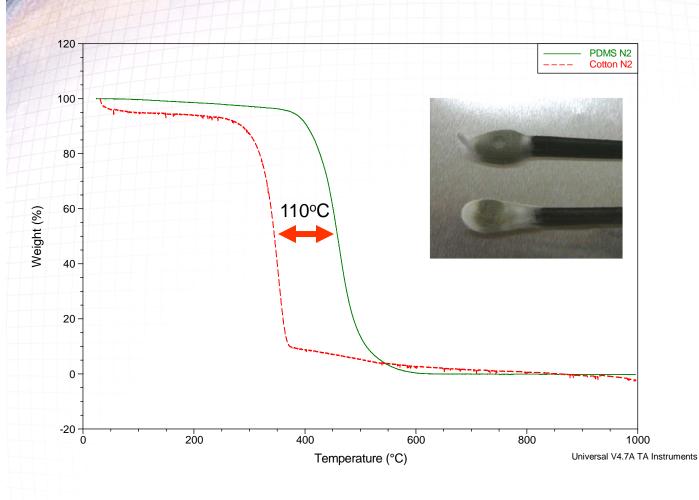
FEP Teflon Heated at Different Rates



During pyrolysis, materials undergo thermal degradation via chemical pathways dictated by the thermal stability of the components. When pyrolysis is slowed to simulate TGA conditions, a thermal response pattern similar to what was observed with TGA first derivative plot.



Cotton Vs. Silicone

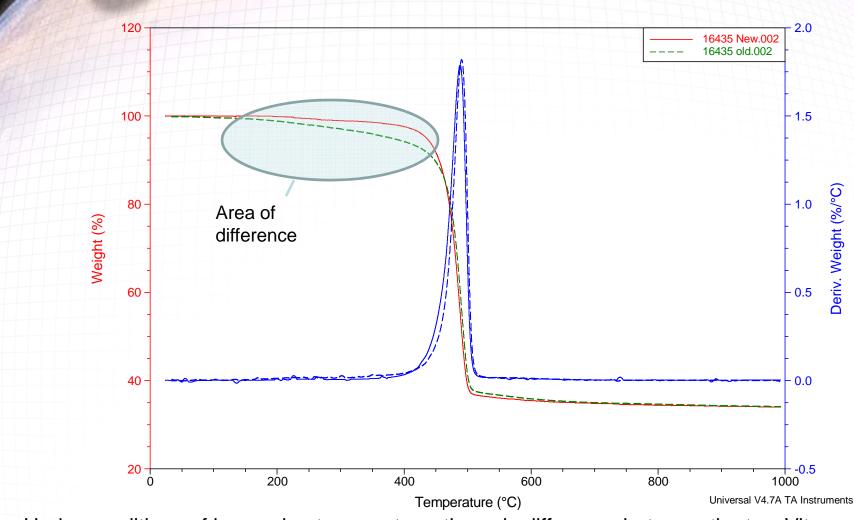


The large difference in thermal stability between cotton and silicones can be used to easily characterize the silicone sample collected on a cotton swab.

The cotton may be completely decomposed by application of heat without adversely affecting the silicone.



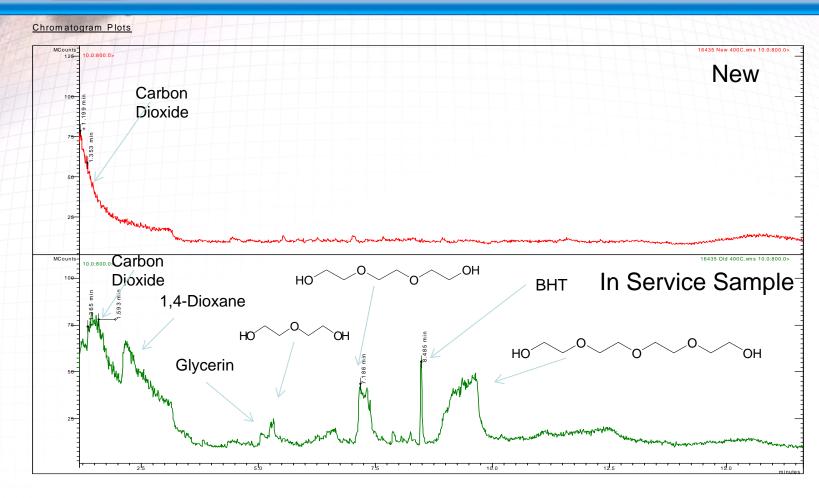
TGA Comparison of Gaskets



Under conditions of increasing temperature, the only difference between the two Viton Gaskets was found below 400°C, where the old sample lost a larger percentage of its mass compared to the new sample.



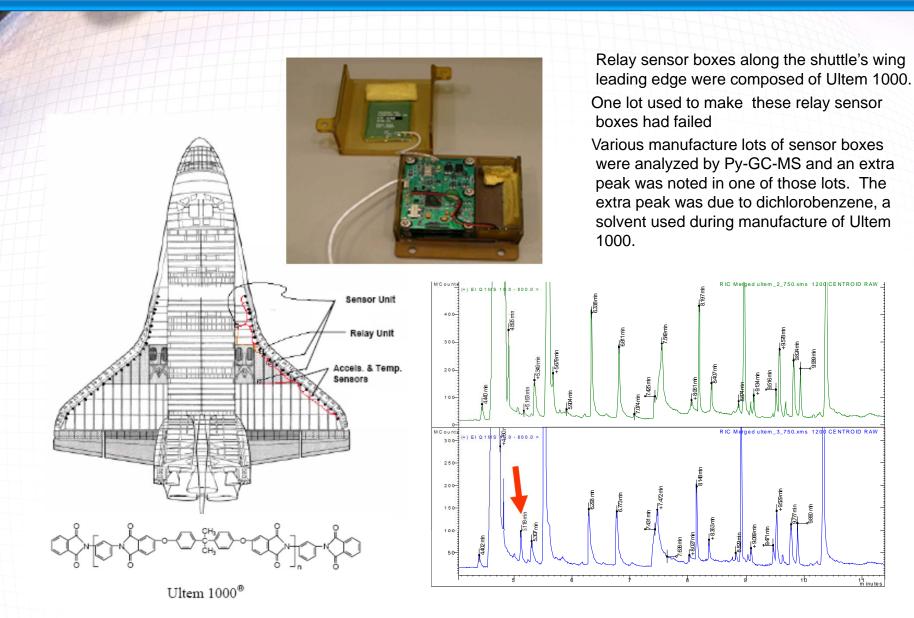
Thermal Extraction of Samples



Thermal extraction of the two samples was performed to account for the difference observed in the TGA experiments at temperatures below 400°C. Such an experiment indicated the Old sample contained various fragments that are attributed to polyethylene oxide. Other substances found included Glycerin and Butylated hydroxy toluene (BHT).

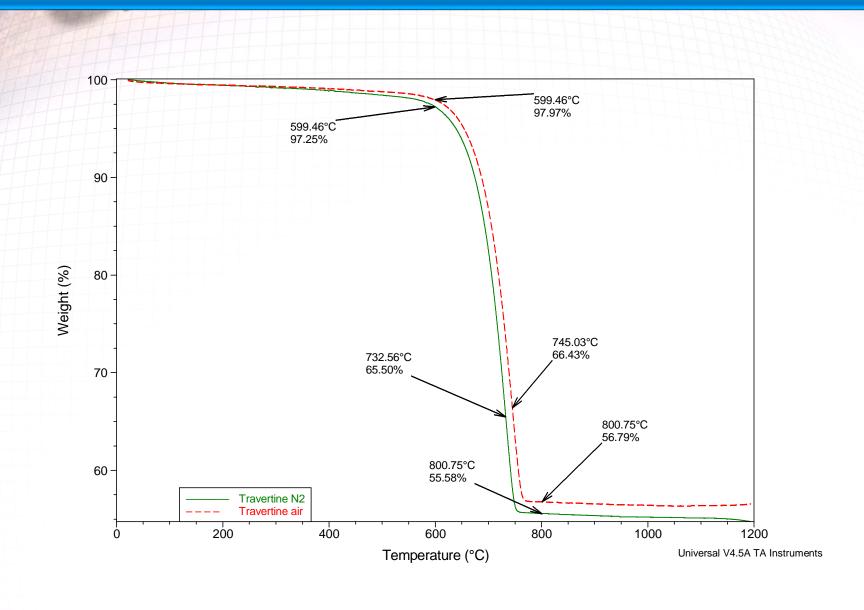


Pyrolysis-GC-MS of Ultem 1000



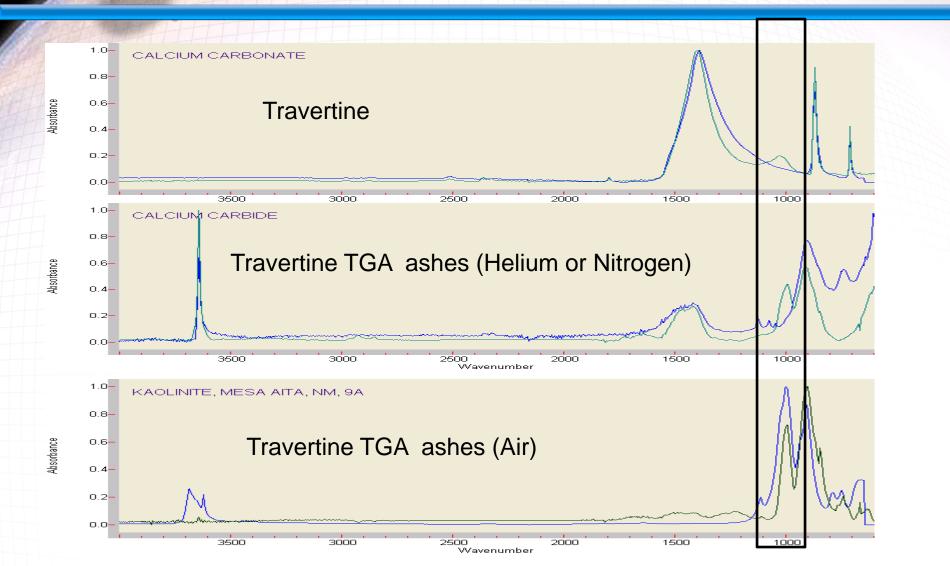


Thermal Response of Travertine in Different Atmospheres



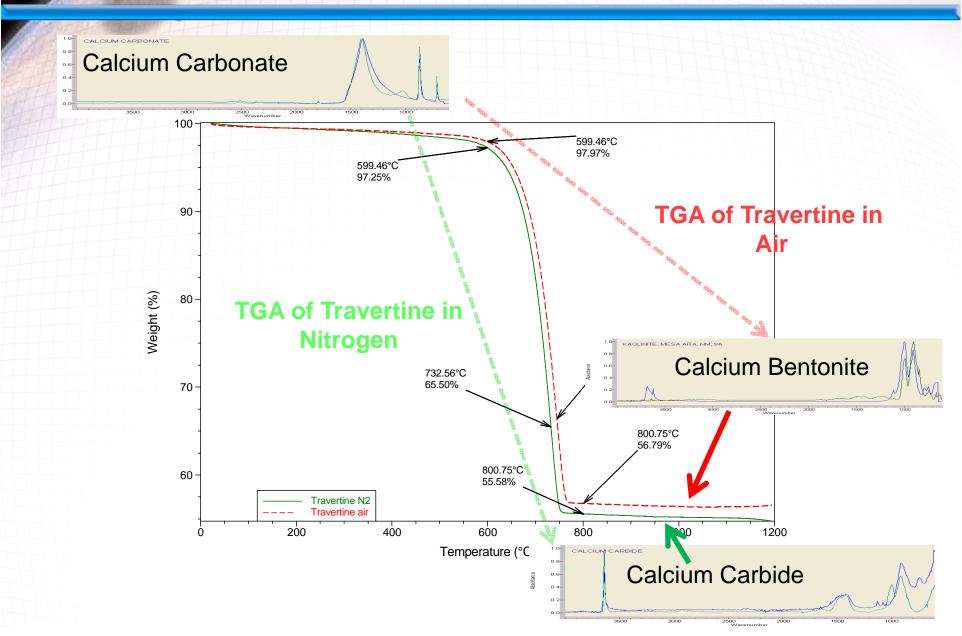


CaCO₃ → CaC₂ or Calcium Bentonite



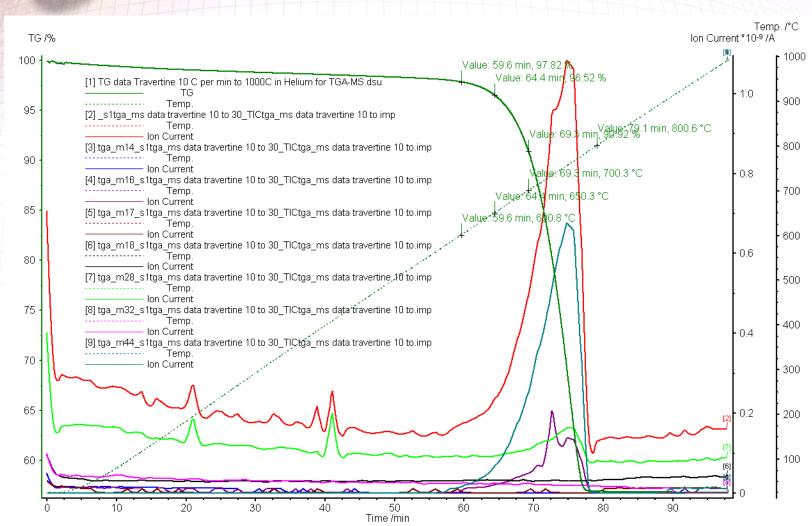


The Role of Gaseous Atmosphere During Thermal Decomposition of Travertine





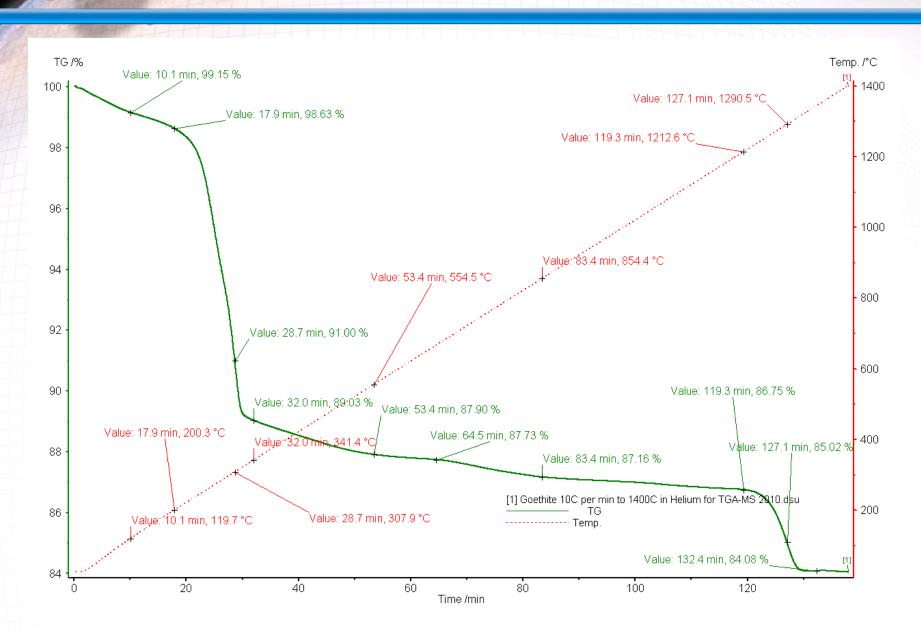
TGA-MS Analysis of Travertine



Substances being measured during mass loss near 700°C include CO₂, CO, and O'

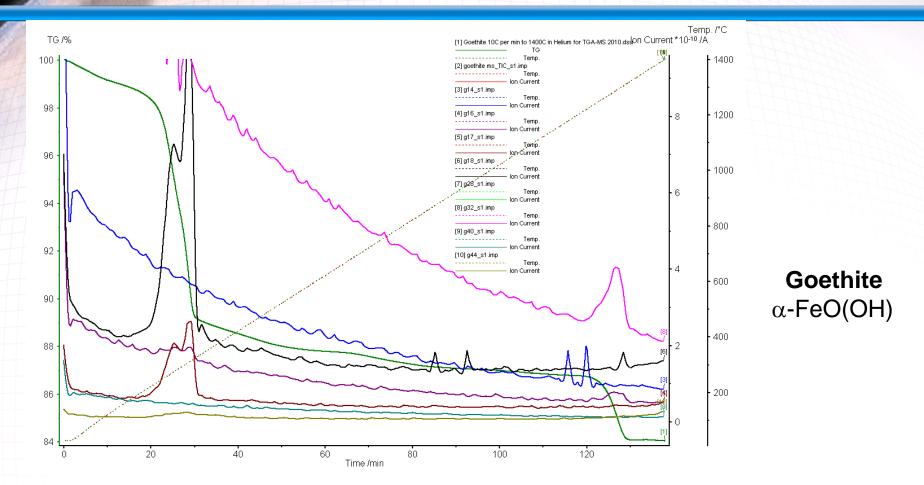


TGA Analysis of Geothite in Helium





Detected Mass Losses of Goethite



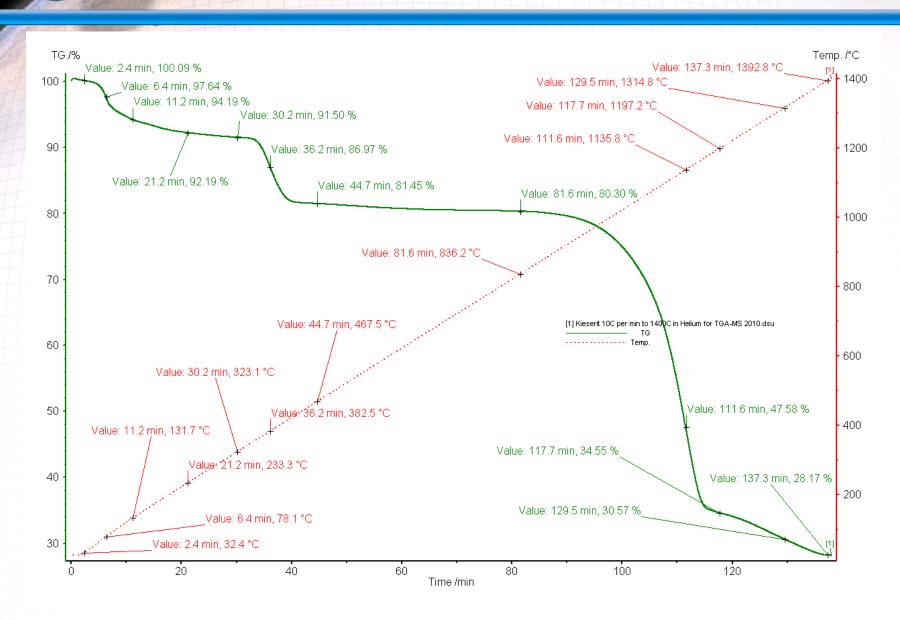
At 120°C, Mass losses include: *m/z* 14 (CH₂), 16 (O), 32 (O₂)

At 308°C, Mass losses include: *m/z* 17 (OH), 18 (H₂O), 32 (O₂)

At 1290°C, Mass losses include: *m/z* 16 (O), 18 (H₂O), 32 (O₂)

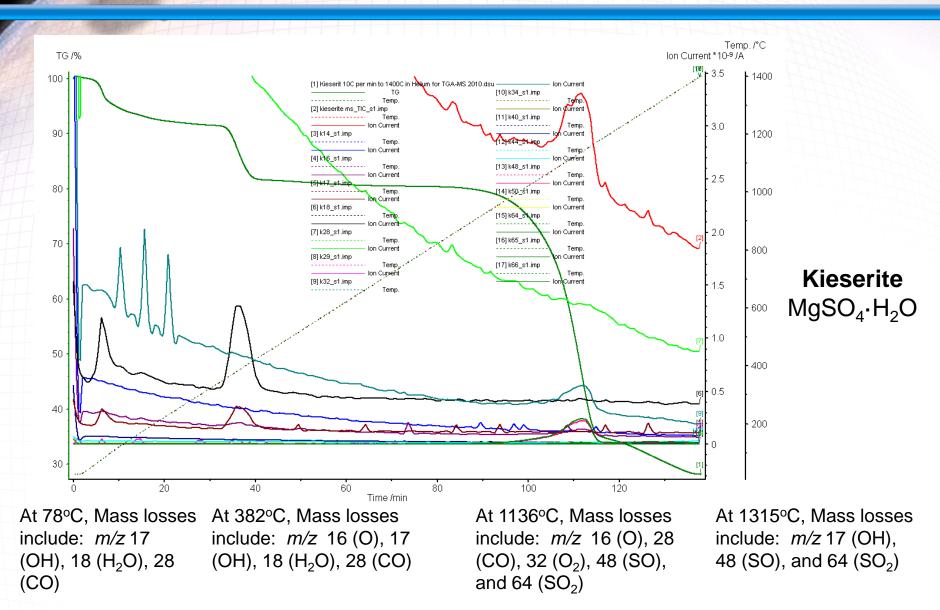


TGA Analysis of Kieserite





Detected Mass Losses of Kieserite

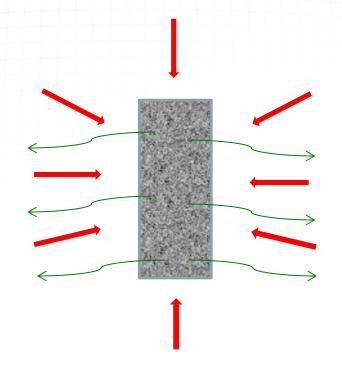




Applying Thermal Energy to Extract Chemical Information

Using Thermal Energy:

- •How much Thermal Energy do we add
- How fast do we add the Thermal Energy
- •What atmosphere do we use
- •How much sample do we use



Chemical Information

- Trapped solvent
- Organic additives
- •Labile Functional Groups
- Monomer identification
- Off-gassing information
- Inorganic additives

TGA

Pyrolysis-GC-MS

TGA-MS-IR