



Challenges for epoxy cure characterization and long term high temperature performance

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Epoxy Selection and Performance

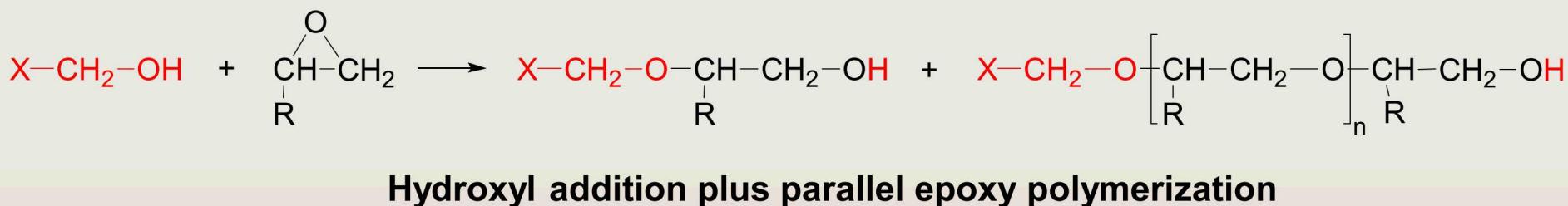
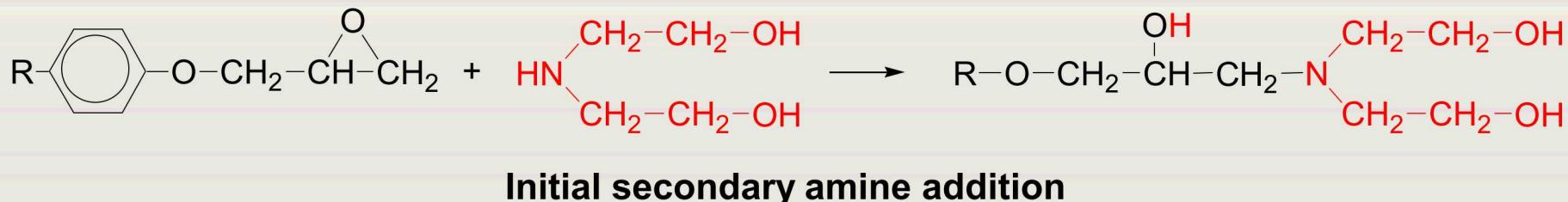
- Depends on cure behavior, cure state and understanding of degradation reactions

Research goals and approach:

- Better understand cure behavior in epoxies
- Identify reasons for unusual cure behavior in Epon 828/DEA
- Better approaches for quantification of cure conversion states
- High temperature epoxy degradation phenomena
- Develop diagnostic tools and lifetime prediction models

Key words: Materials Characterization, Spectroscopy, Mechanistic Evaluation

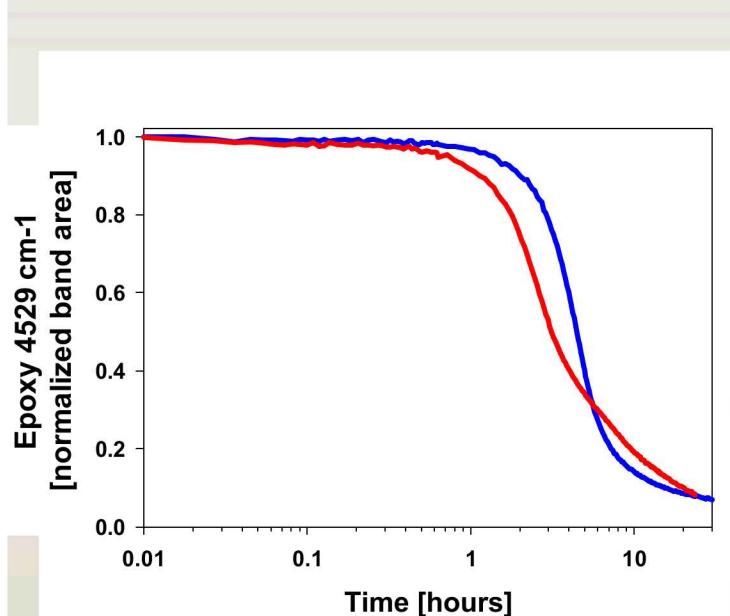
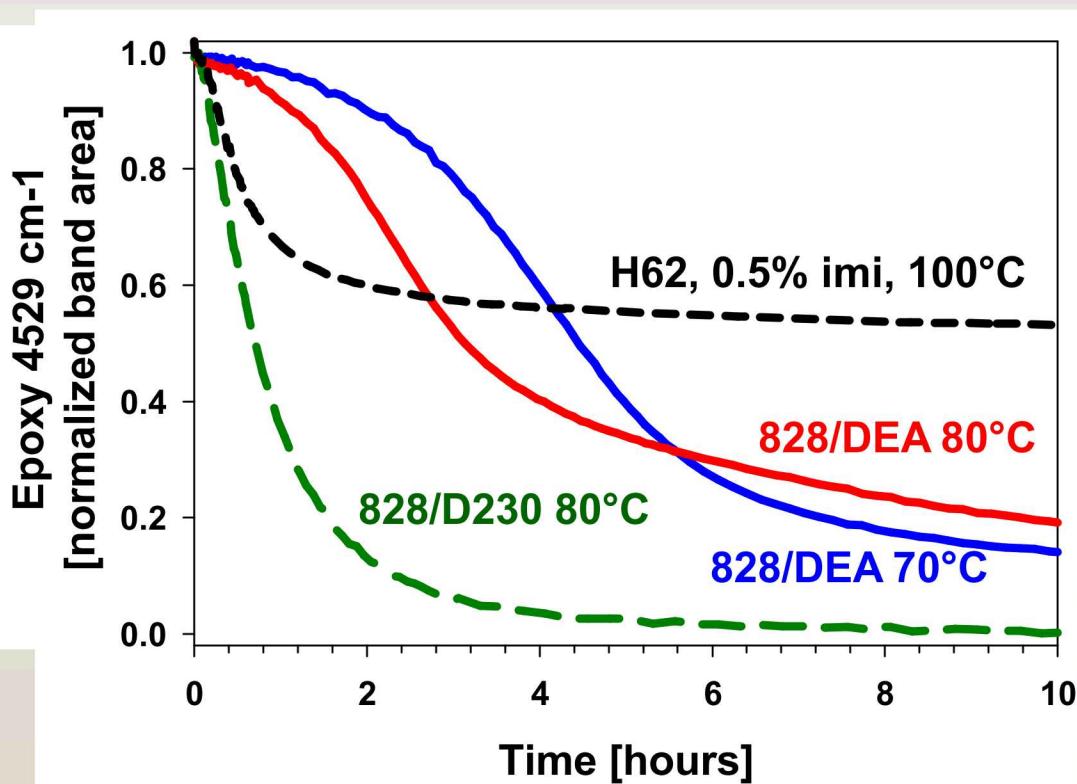
Epon 828 cured with DEA



- Excess epoxy group reactions:
- A) Epoxy groups can homo-polymerize (exact mechanism unclear)
- B) Epoxy reactions with existing hydroxyl (ethanol and epoxy addition OH)
- Competitive chemistry, is either process dominant? Combination?

Unexpected Reaction Kinetics in 828/DEA

- 828/DEA cure does not follow consistent Arrhenius behavior
- Expectation: Higher temperature should accelerate reaction and push cure conversion

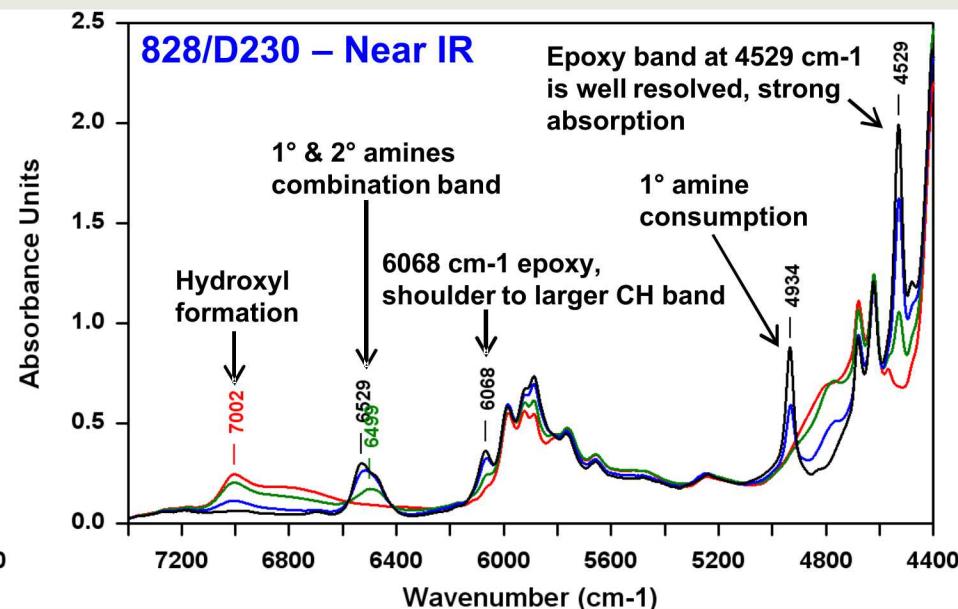
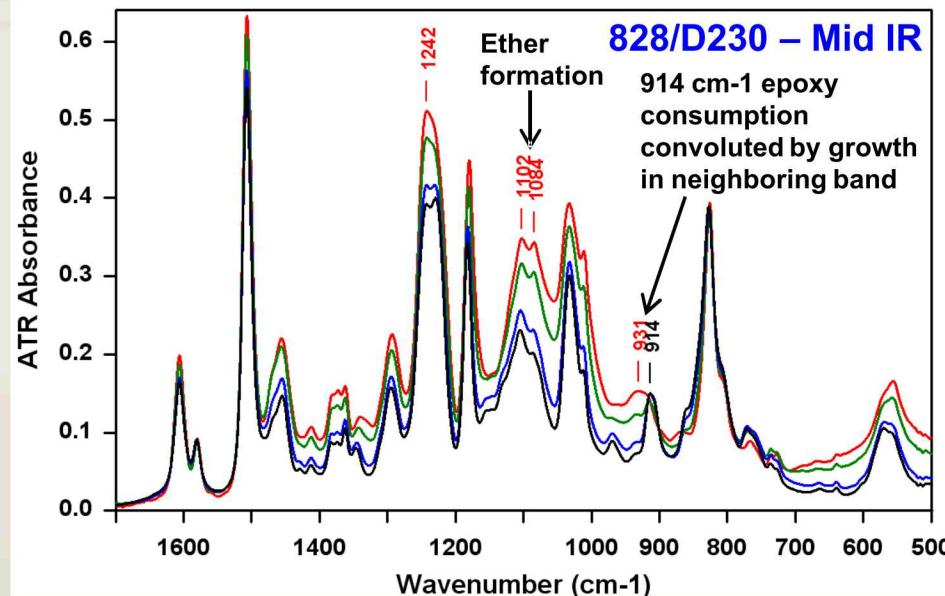


The cure reaction starts faster, but is then slower at higher temperature

Mid IR – NIR 828/D230

- Epoxy band in some systems (ie. 828/DEA) is convoluted in mid IR (914)
- NIR offers better resolved epoxy absorbance at 6068 and 4529 cm^{-1}

MIR and NIR spectra were acquired at ~8 seconds, 10 mins, 45 mins, and 15 hours at 80°C

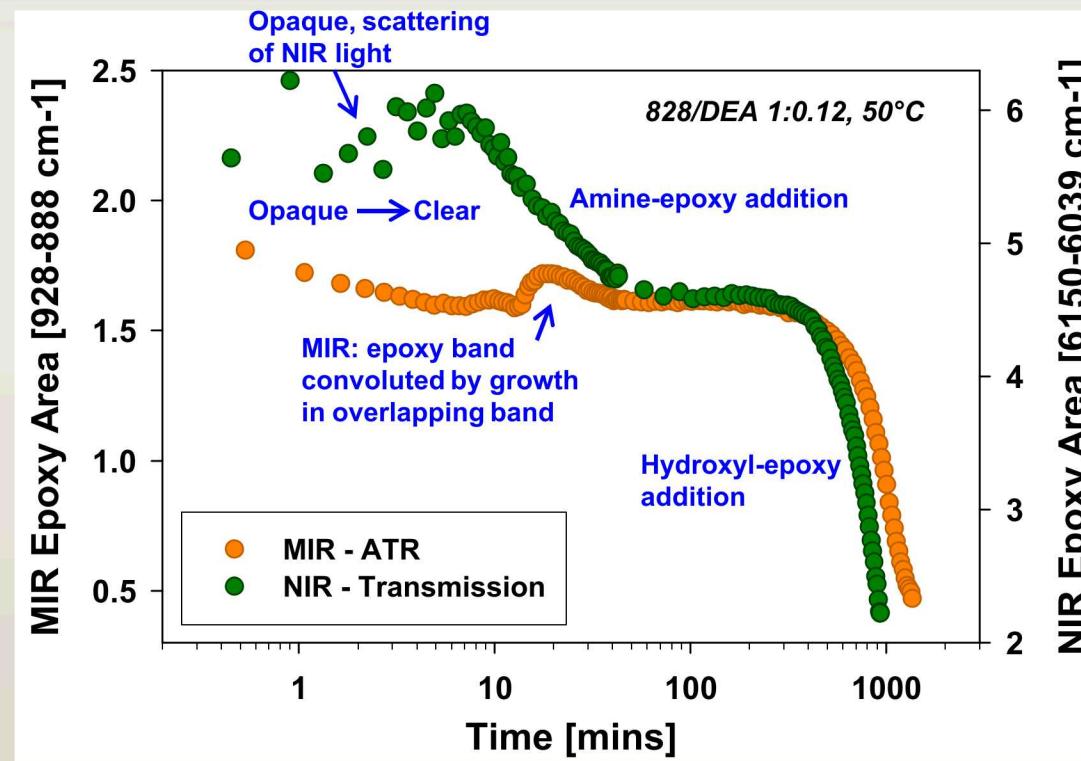


Epoxy as well as amine bands are resolved in NIR

E Duemichen, U Braun et al, Thermochimica Acta, 616 (2015) 49
 'NIR monitoring of epoxy cure with variable heating rate'

Mid IR - NIR

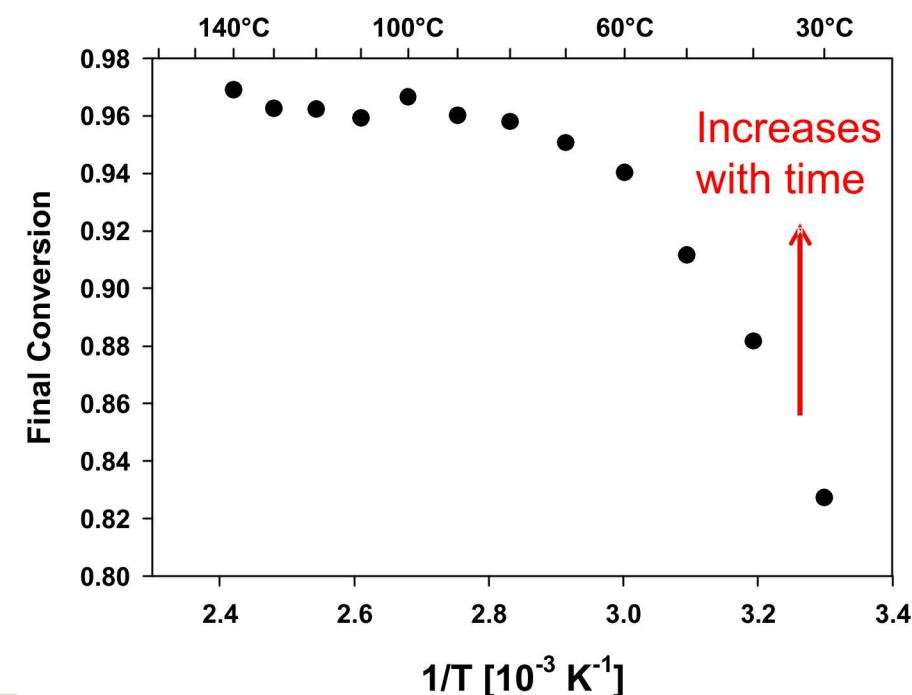
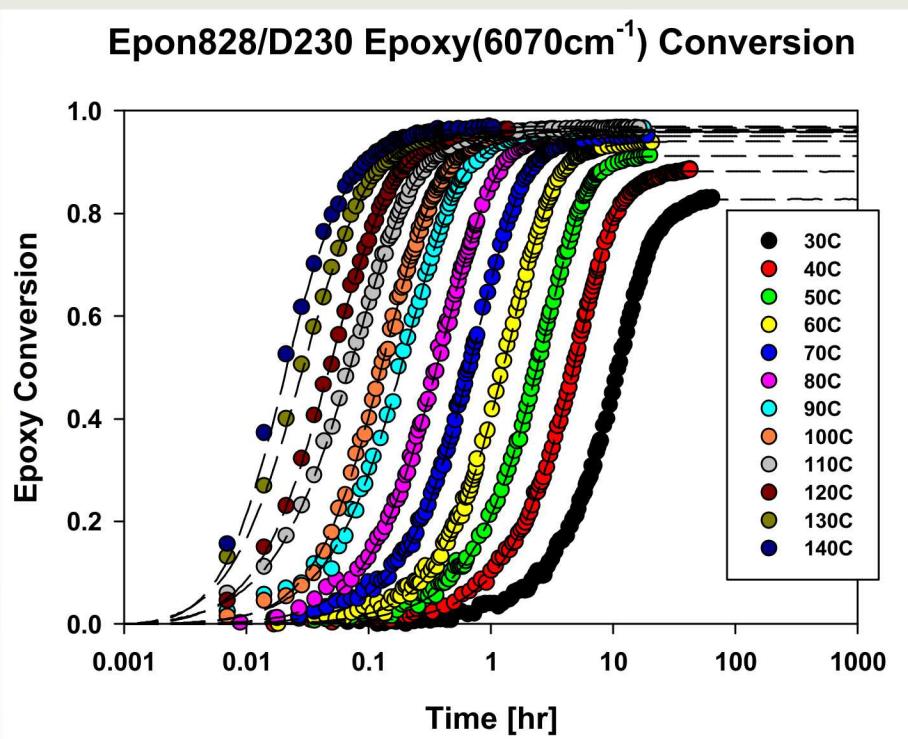
- Epoxy band consumption for 914 (mid-IR) and 6068 cm-1 (NIR) for 828/DEA cure at 50°C
- 914 is integrated between 928 and 888 (not deconvoluted)



NIR also more easily picks up initial sec-amine epoxy addition

Epon 828/D230 *t-T* Cure Behavior

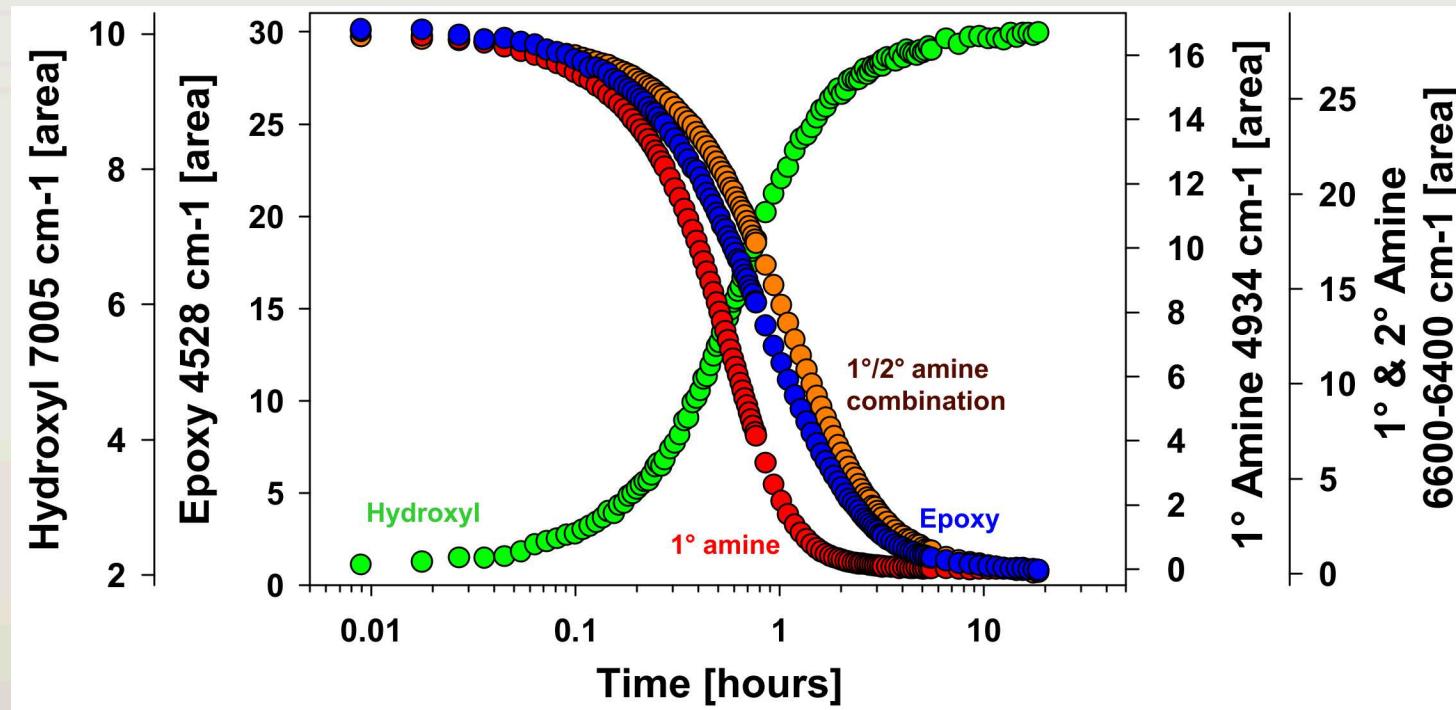
- Traditional epoxy cure
- Final conversion levels drop at lower T, but an ongoing power law driven cure reaction is evident



Cure speed increases and higher final conversion is obtained with T

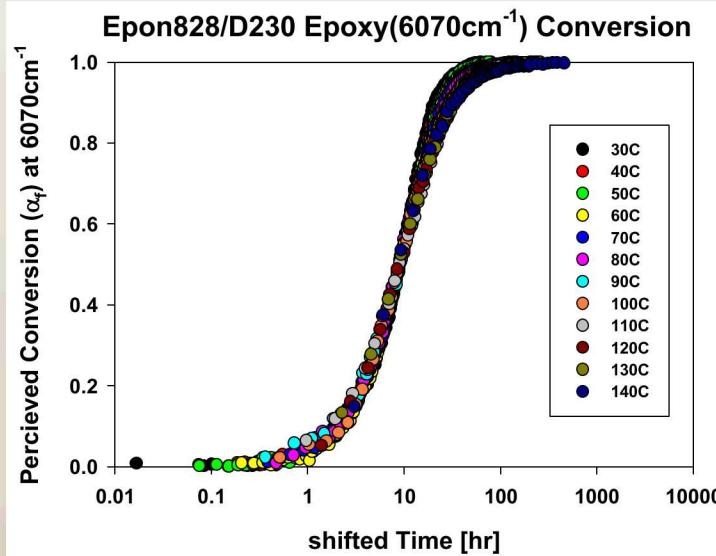
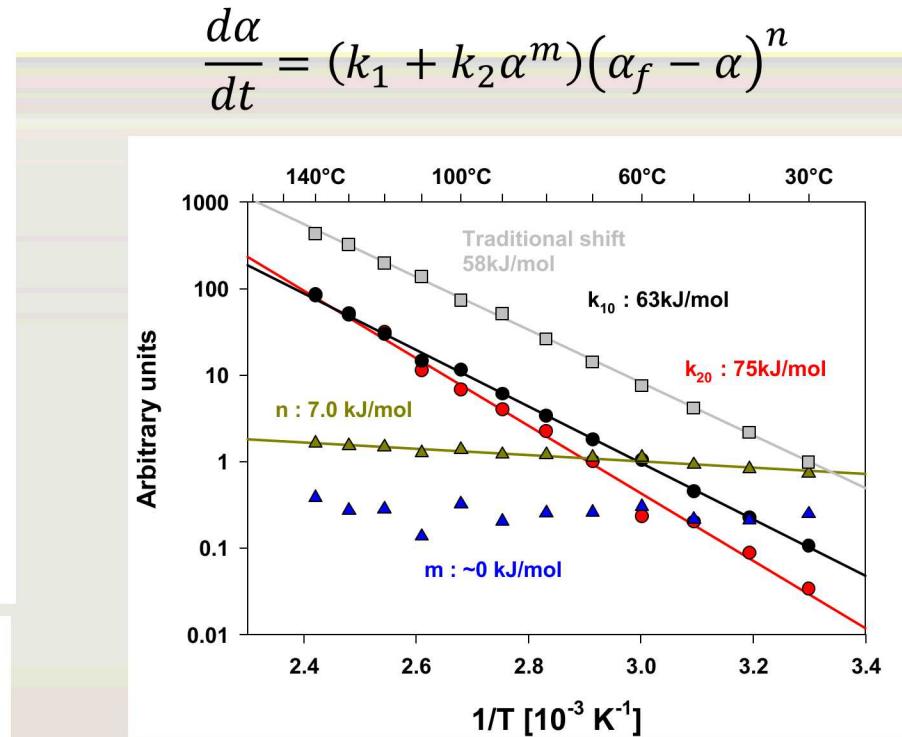
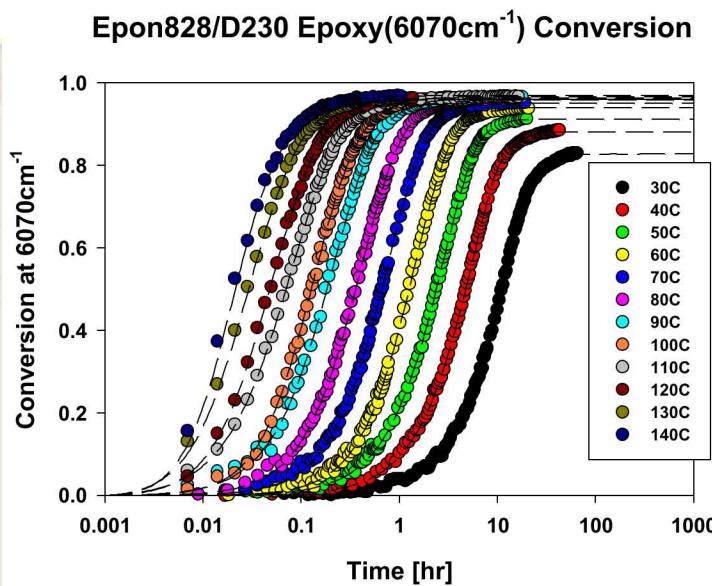
Epon 828 / D230 – 70°C Cure

- Epoxy consumption can be compared with amine loss and hydroxyl formation



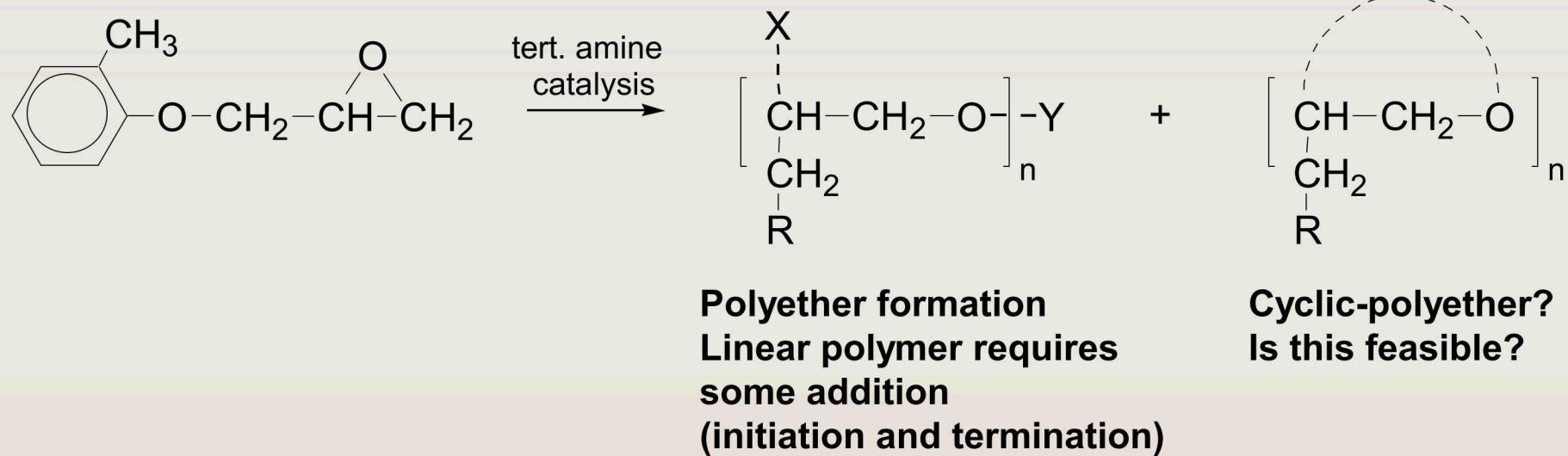
This system can be examined with multiple spectroscopy approaches

Epon 828 - D230



- Traditional shifting does not account for variance of kinetic regime at high conversion and is only available after data is corrected for final cure state (α_f)
- Kamal equation accounts for this with two modified rate constants and temperature dependent power.

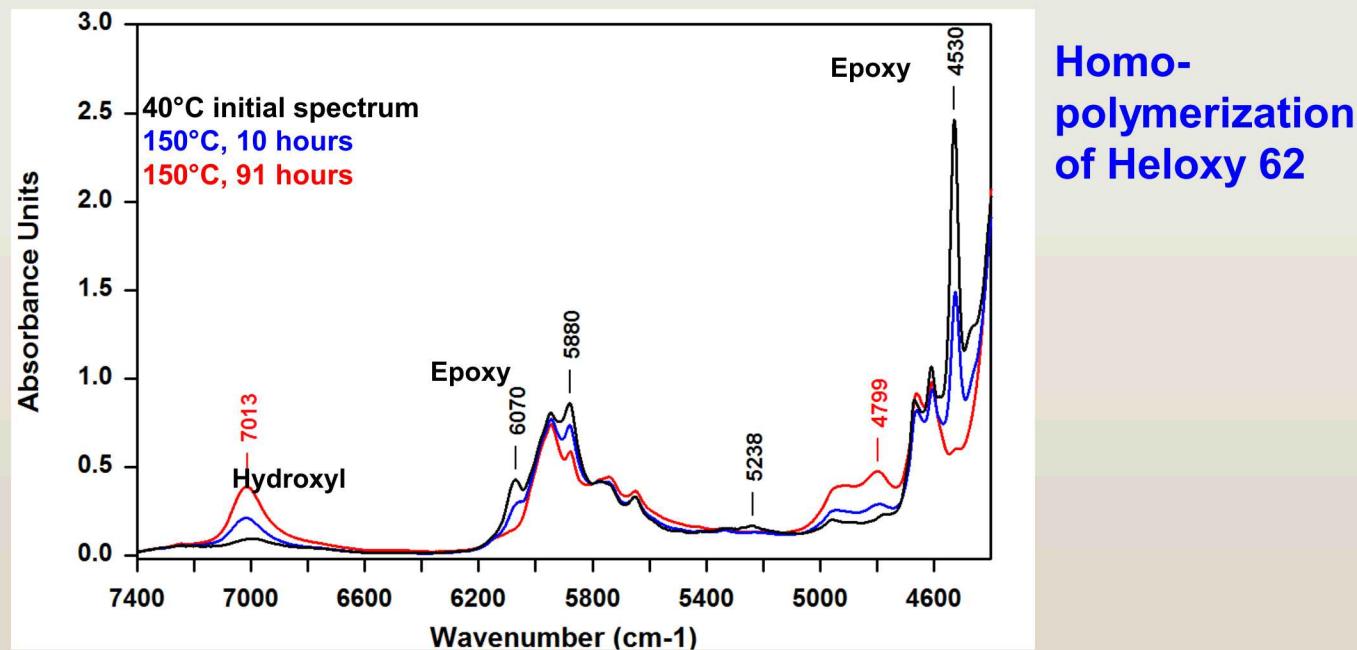
Mono-functional Epoxy Polymerization



How does this reaction proceed?
What is conversion with time?
Can water interfere/contribute?

NIR Epoxy Quantification Efforts

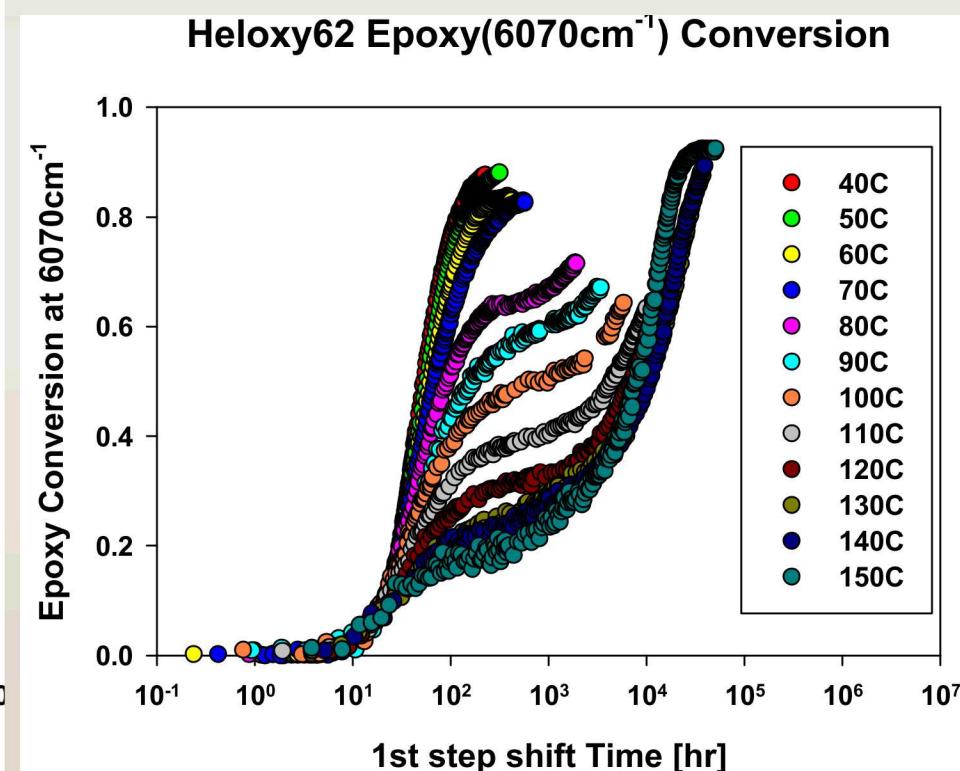
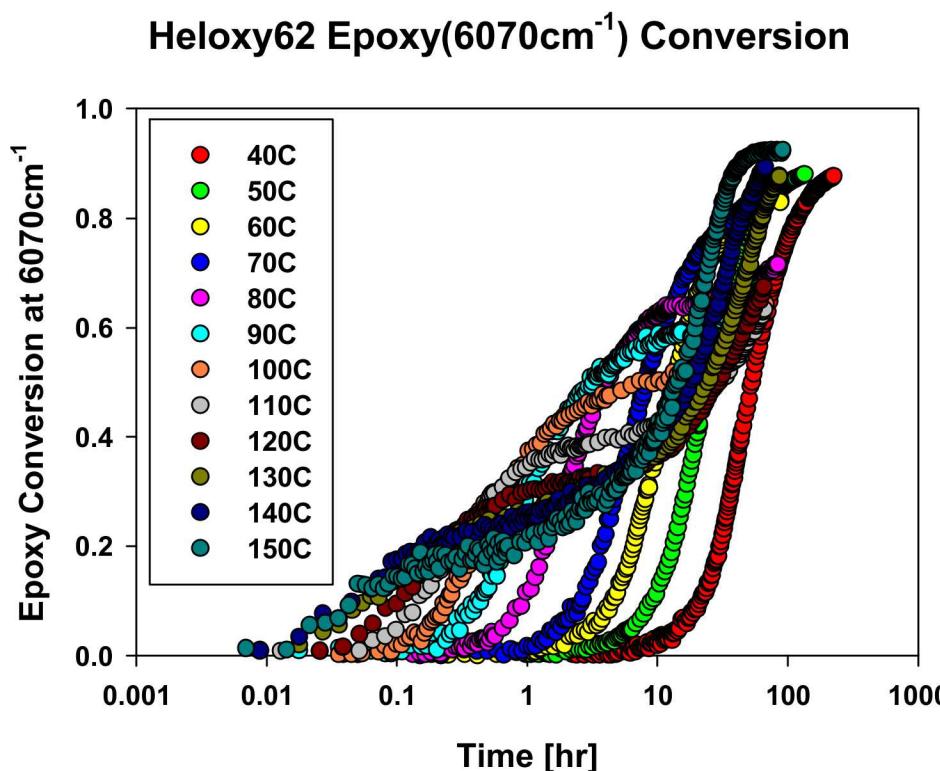
- Baseline and spectral correction for 'physics' T effects
- Band deconvolution with local baseline optimization (Matlab)
- Gaussian peaks with fixed peak position help the analysis
- Individual band integration and normalization for relative conversion
- Boundaries for conversion, initial reagent concentration ($\alpha = 0$), complete reaction from high T annealing above Tg ($\alpha = 1$)



Challenge is peak separation, integration, and conversion boundaries

Heloxy 62 – Complicated Cure Behavior

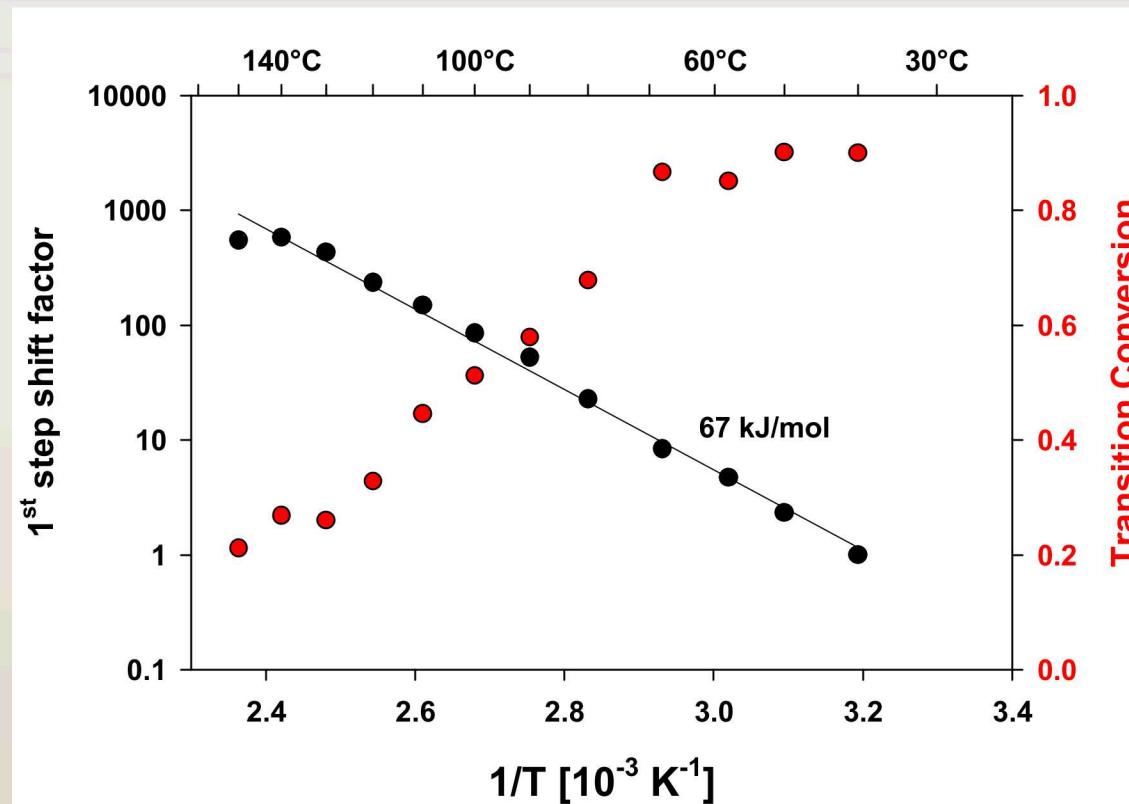
- No easy t-T superposition, highly complex cure behavior dependent on T
- Two cure stages result in complicated kinetic cure behavior and model
- Conversion state for transition between cure regimes depends on T



Delayed cure conversion at high temperatures
What do limited conversion states indicate?

Early Reaction Kinetics

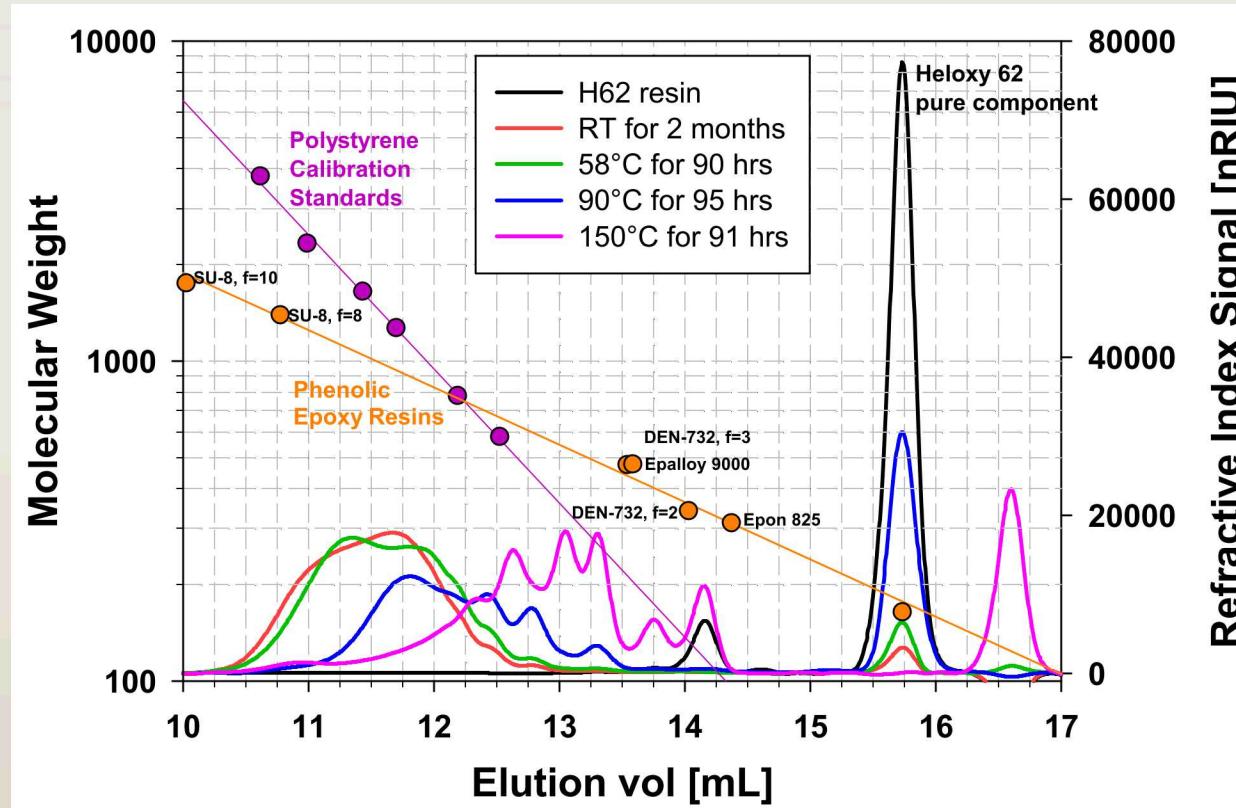
- Early stage t-T superposition yields 67 kJ/mol over wide T range
- Is limited to specific conversion level, that decreases with T



What do limited early conversion states indicate?

Homopolymerization

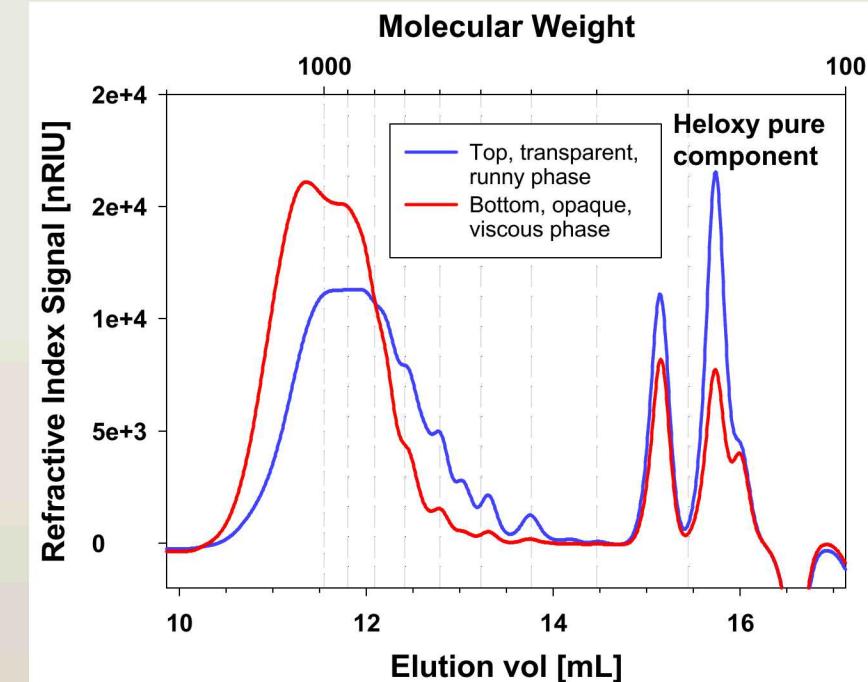
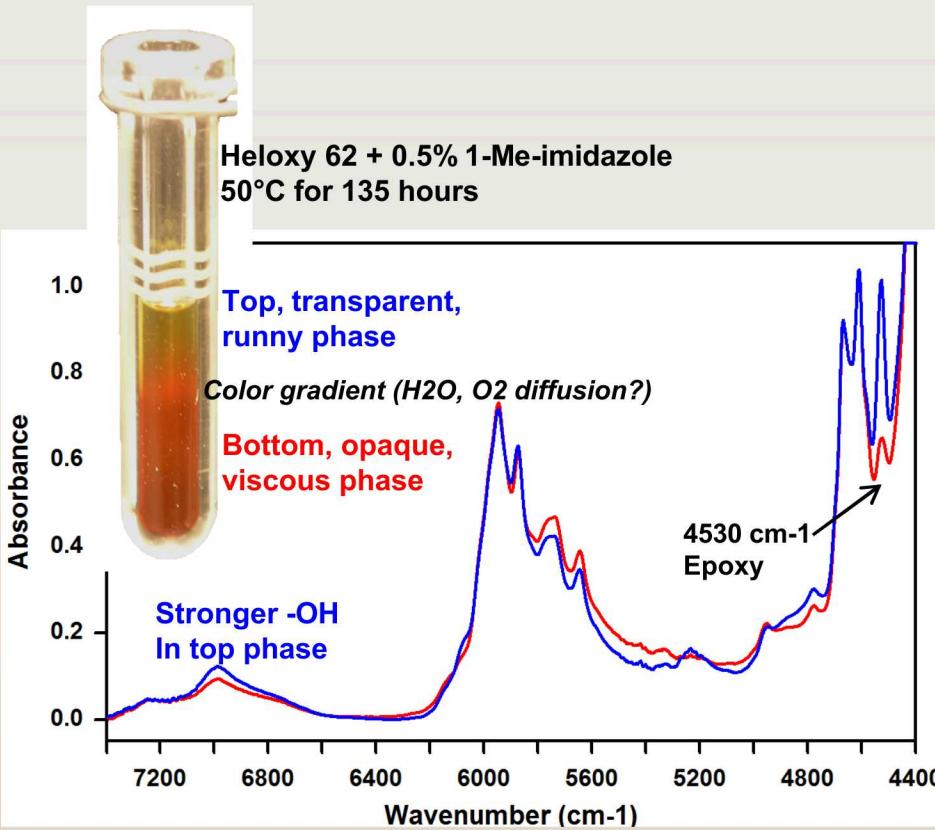
- Evidence for ‘ceiling’ temperature behavior
- What is M_w (polymerization degree) for linear polymerization?



- GPC proves limited chain propagation behavior
- Higher Mw at lower temperatures

Cure Inhibition - Water

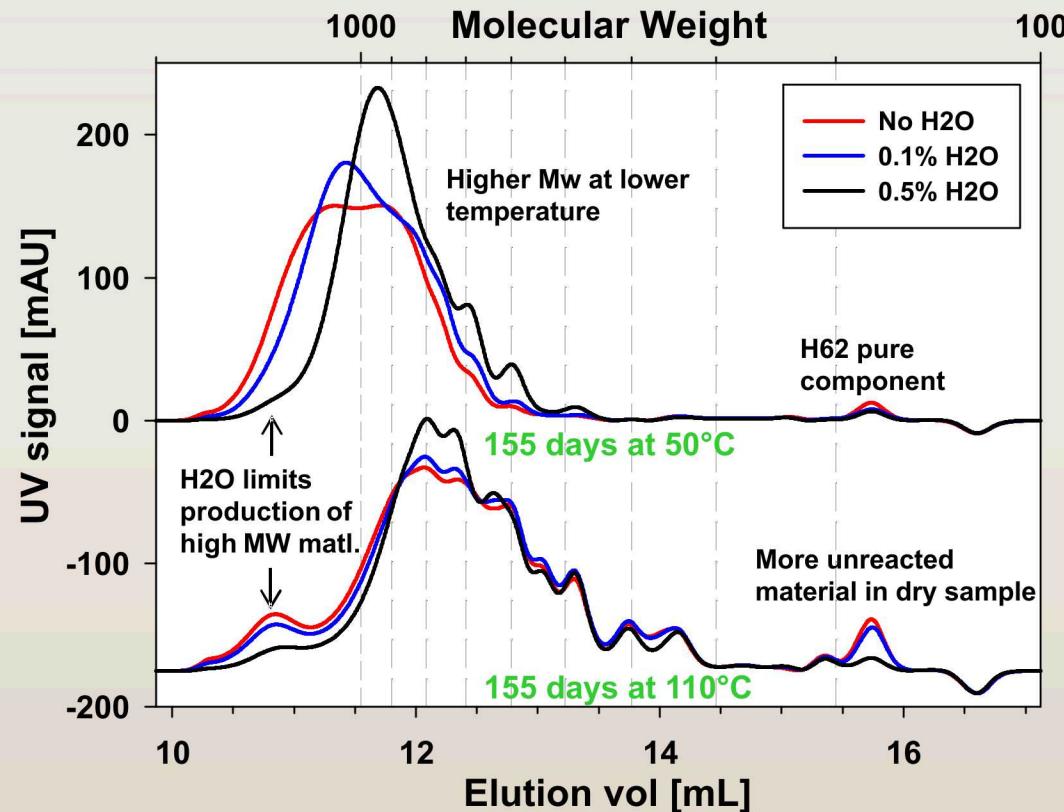
- Does water contribute?
- Can water favor linear polymers (recall addition needs)?



Clear evidence that atmospheric exposure affects cure behavior

Cure Inhibition - Water

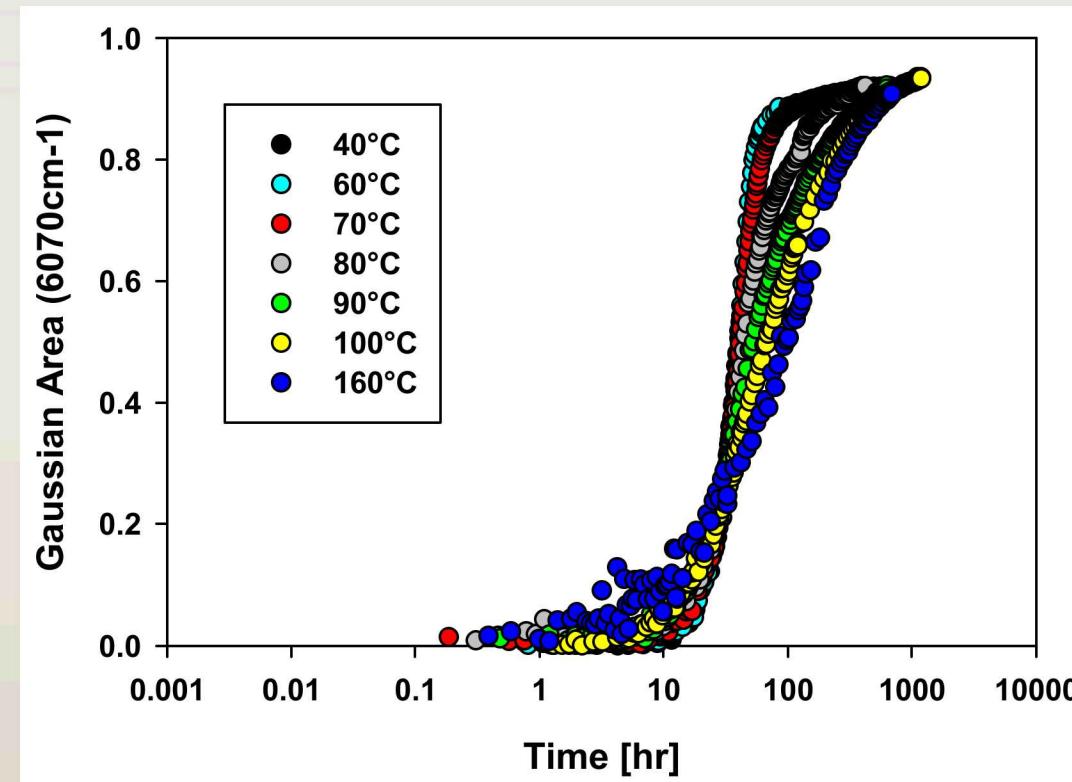
- 6 cure conditions: 0.1% & 0.5% H₂O addition vs. control at 50°C & 110°C
- H₂O is miscible with epoxy. Rxn carried out in sealed vials over 155 days.



- H₂O lowers chain propagation or increases termination
- Much higher Mw formation at lower temperature

Epon 828/DEA Cure Behavior

- Preliminary NIR data for t-T epoxy conversion (initial guidance)
- Peak quantification can be further optimized



Influence of temperature (anomalous cure) is clearly apparent
Pure mathematical data superposition may not be the best approach

Epon 828/DEA Cure Mechanisms

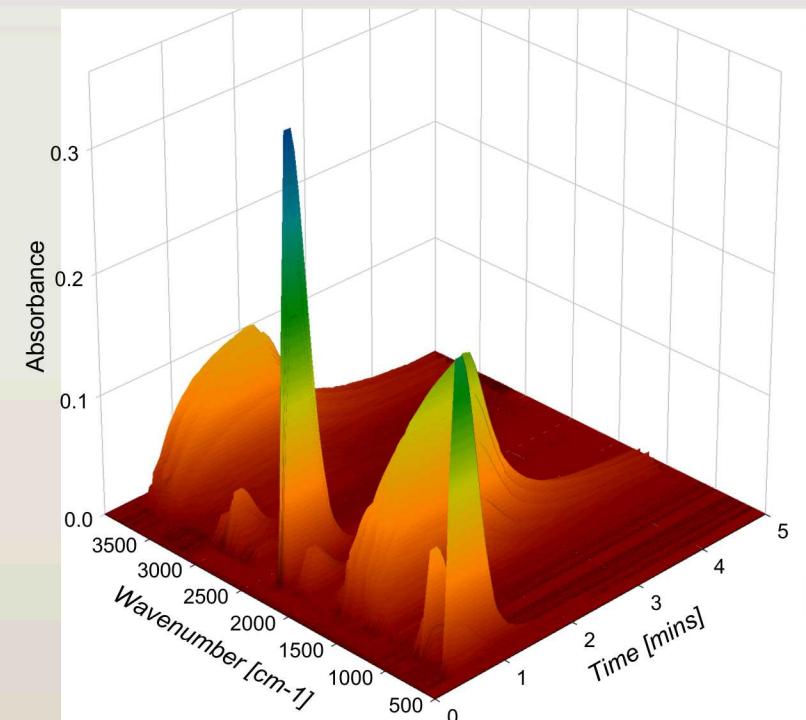
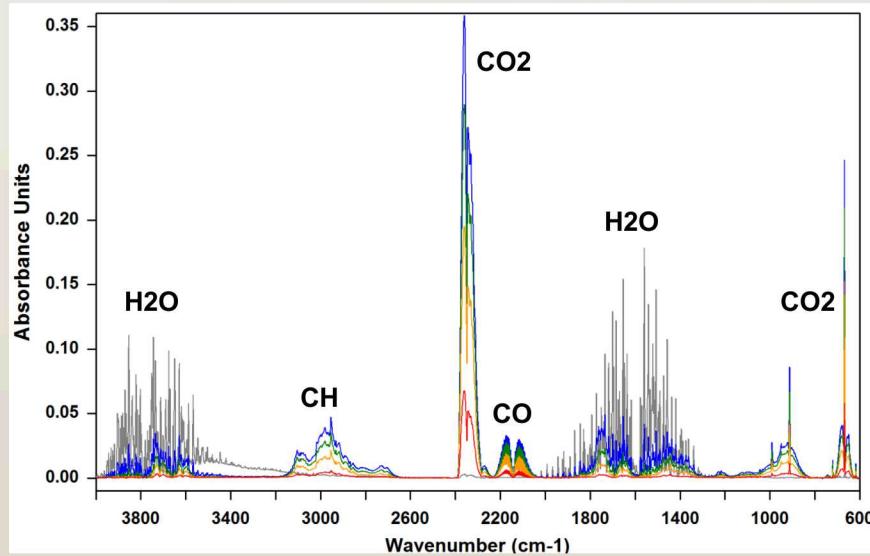
- Extensive literature review with collaborator John McCoy – NM Tech

“Cure mechanisms of Diglycidyl Ether of Bisphenol A (DGEBA) Epoxy with Diethanolamine” John D. McCoy*, Windy B. Ancipink, Caitlyn M. Clarkson, Jamie M. Kropka**, Mathias C. Celina**, Nicholas H. Giron**, Lebelo Hailesilassie, Narjes Fredj, under review “Polymer”

- At low temperature the DGEBA/DEA gelation reaction is “activated” (shows a pronounced induction time, similar to autocatalytic behavior) by the tertiary amine in the adduct.
- At high temperature, the activated nature of the reaction disappears.
- Upper stability temperature of the zwitterion initiator of the activated gelation reaction
- Reaction rate of epoxide consumption cannot be generically represented as a function only of temperature (T) and degree of epoxy conversion (α)
- Requires specific consideration of the dilute intermediates in the reaction sequence

High Temperature Performance of Epoxy

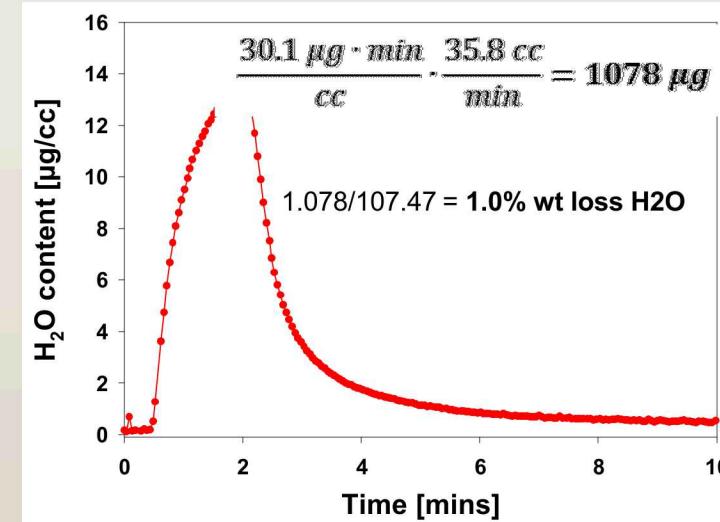
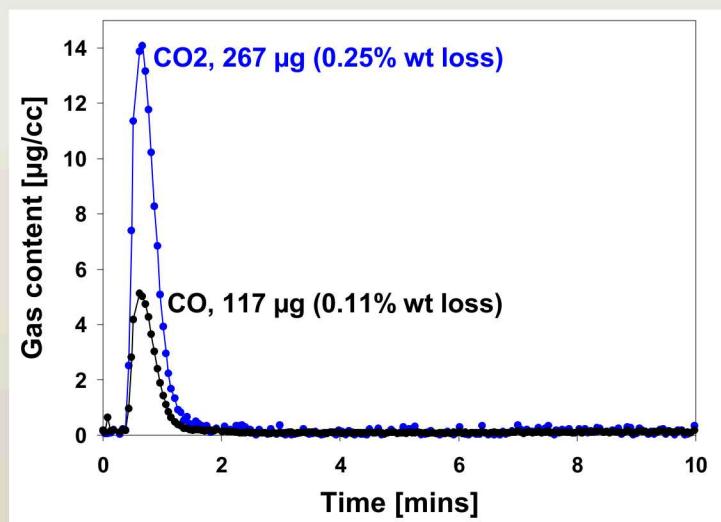
- Thermally induced decomposition (pyrolysis chemistry)
- IR based analysis of gaseous decomposition products
- Flow through approach, sealed ampoules, rapid scans



Quantification of H₂O and CO₂ yields via spectral and time integration

IR Analysis of Degradation Products

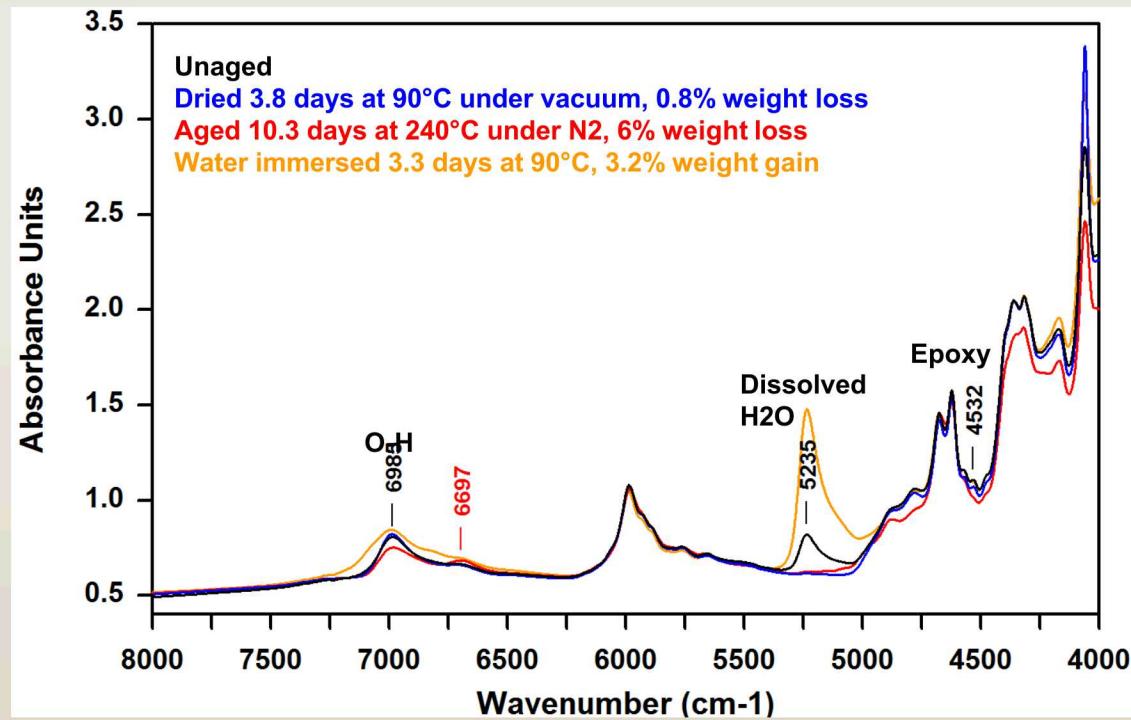
- Proof of principle epoxy aging test in sealed ampoule
- ~ 100 mg epoxy sample aged for 3.75 d at 240C, ~ 1% expected weight loss
- Rapid IR spectral acquisition of flushed out gas with decomposition products
 - Evidence for significant water, plus some CO₂ and CO as volatiles, some CH signatures
 - Integrate water peak between 1588 and 1518 cm⁻¹
 - Use corrected calibration for limited water spectral range; content in µg/cc
 - Determined 1.078 mg H₂O in sealed ampoule; actual weight loss was 2.4 mg (2.2%)



- At 240C and early on, the epoxy yields ~ 45% weight loss as water₂₀

Physically Absorbed Water

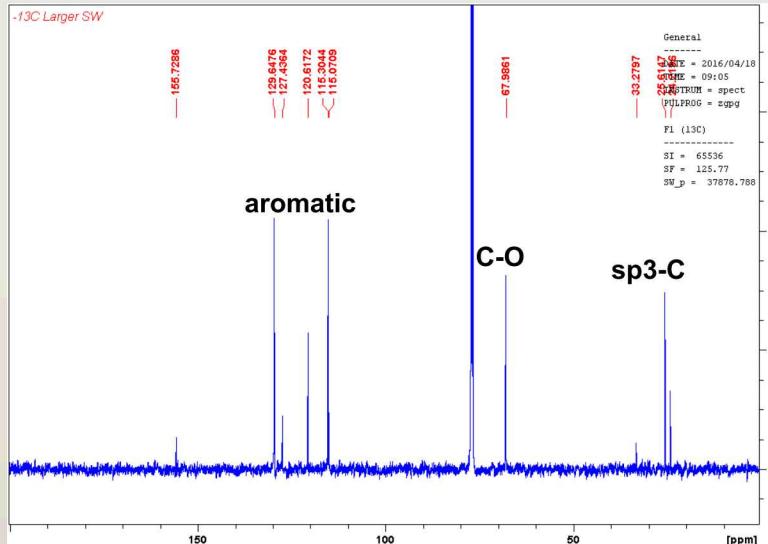
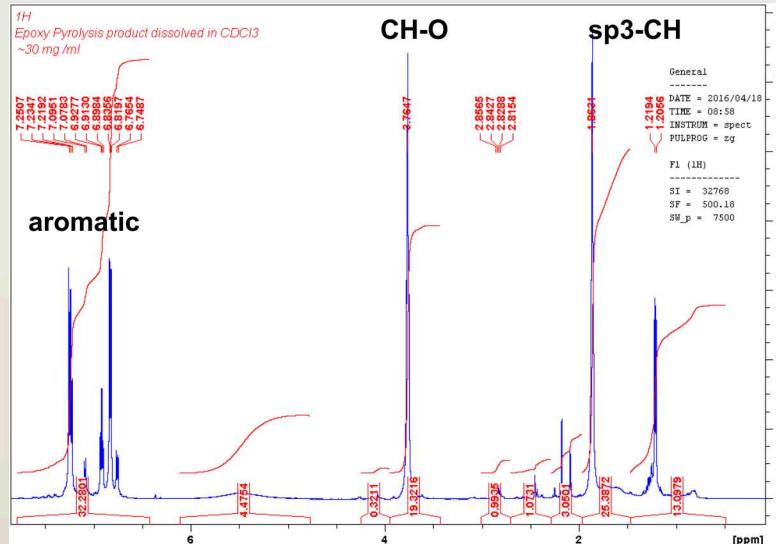
- Amine cured epoxy has a tendency to absorb water (somewhat hydroscopic)
- NIR can be used to quantify dissolved water in an epoxy
- Water uptake will depend on geometry and permeability (thin lab samples are expected to more easily pick up water)



Presence of dissolved water and its chemical formation

High T Decomposition Products

- High T exposure induces the beginning of pyrolysis (degradation chemistry)
- Question: Could limited pyrolysis still apply at 150°C/160°C (over many years)?
- Pyrolysis leads to weight loss, contraction, volatile formation (H₂O, CO₂, CO, CH)
- Epoxy fragments, liquid and solid at RT, can also be detected.

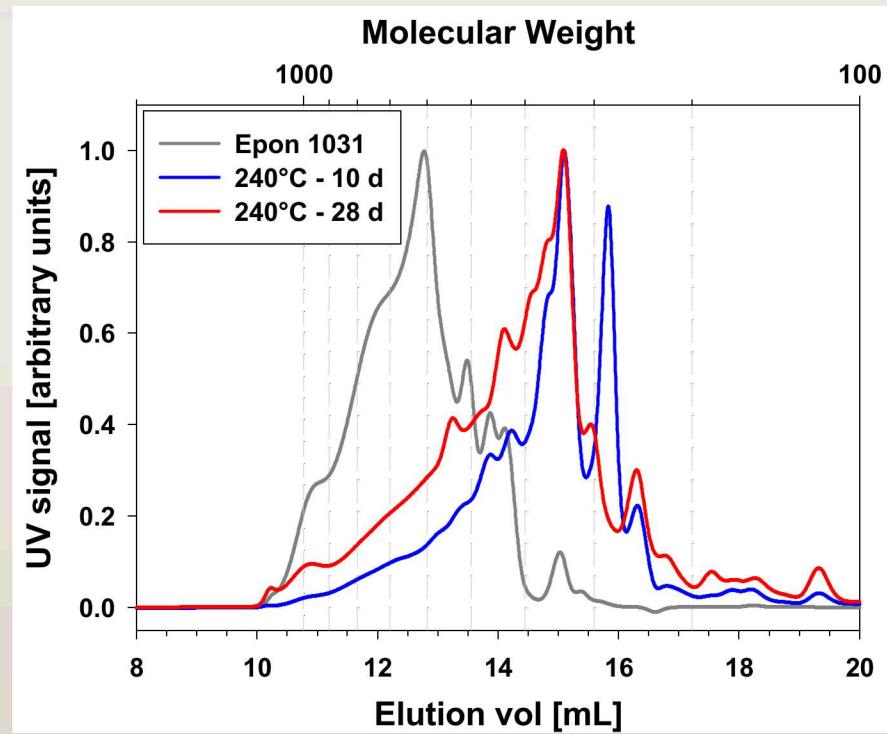


1H and 13C NMR by Todd Alam

- High T pyrolysis products include epoxy fragments

Liquid High T Products - Mw

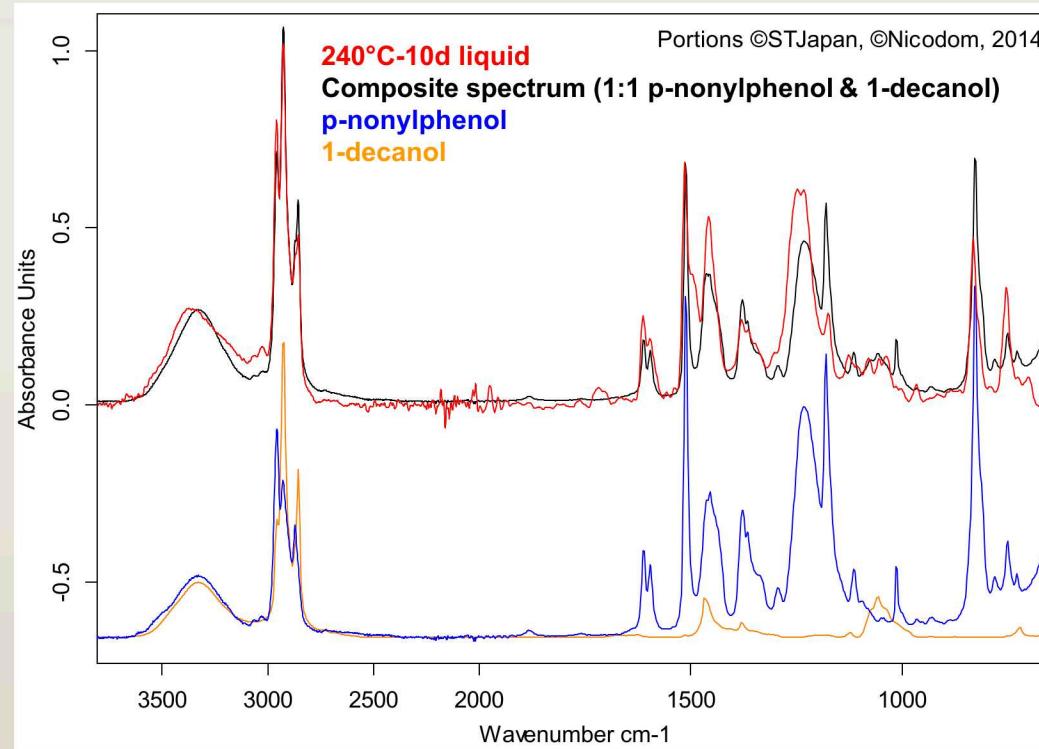
- Pyrolysis leads to some volatile material that can be liquid and solid upon cooling
- GPC was used for guidance on molecular weight of CDCl₃ soluble residue
- Two samples: 240°C-10d and 240°C-28d



- High T pyrolysis products yield 'organic material' in 10²-10³ Mw range

Liquid High T Products - FTIR

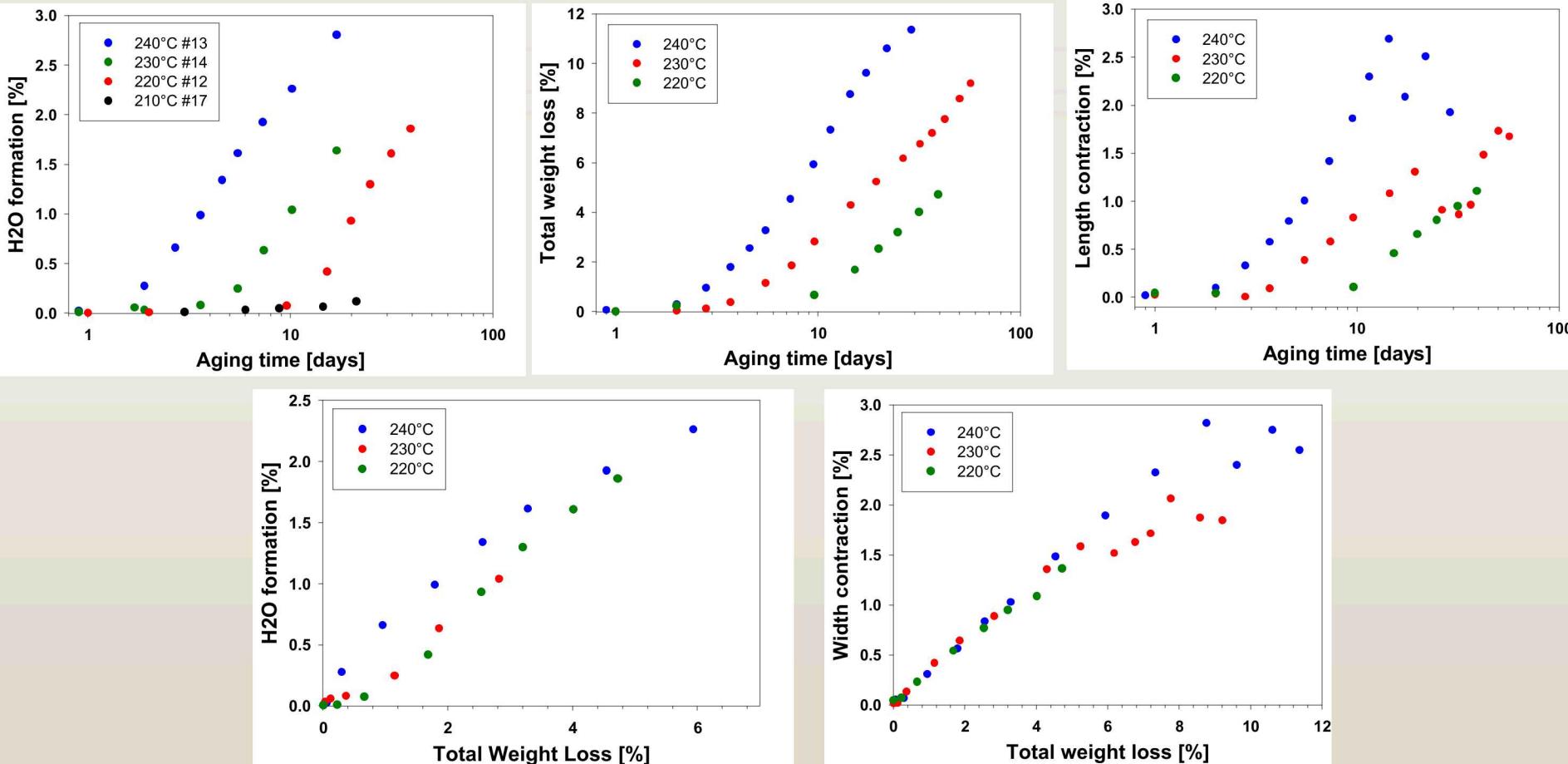
- FTIR spectrum of 240°C-10d liquid has strong OH, aliphatic CH, and aromatic bands
- A Bruker OPUS IR library search found several hits for aliphatic and aromatic alcohols
- IR is consistent with pyrolytic fragmentation of epoxy network



- FTIR of liquid product shows strong OH, CH, and aromatic bands

Volatile Yields - Weight Loss - Contraction

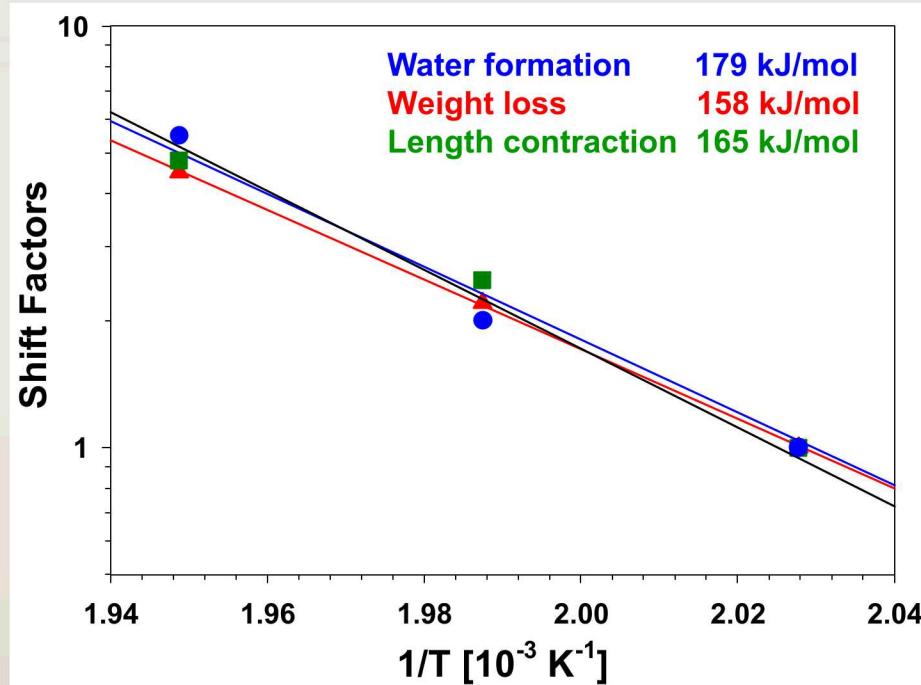
- Weight loss – Water formation – Material Contraction
- Stress and changes in local chemical environment



Aging studies to be continued at lower temperature – Arrhenius ?

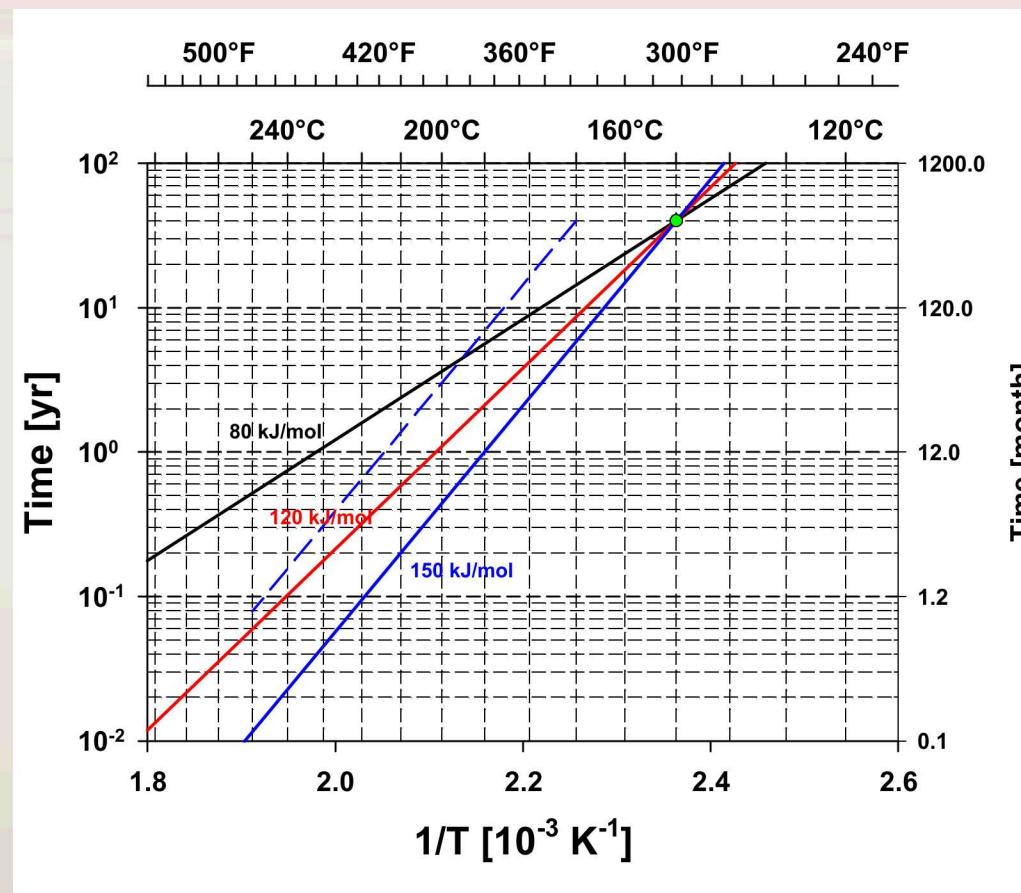
Kinetic Models – Basis for Extrapolations

- Shift factors were computed by time-temperature superposition referenced to 220°C



Initial trends: activation energies range from 158-179 kJ/mol.

Lifetime Prediction Models



Need to establish E_a for high temperature epoxy degradation processes through extensive t-T data sets

Summary

- NIR coupled with mid-IR spectroscopy are excellent cure monitoring tools
- Epoxy polymerization can be unexpectedly complex
- 828/DEA contains excess epoxy and shows anomalous cure behavior
- Elevated temperature may not favor rapid increased cure-conversion
- Small amounts of water can mechanistically interfere
- High temperature applications ($>150^{\circ}\text{C}$) will induce epoxy degradation
- Issues are volatile formation, weight loss and material contraction (stress)
- **Ongoing work: Extensive aging studies for lifetime prediction studies**
- **Impact: Improved methods for cure characterization and aging characterization**