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Interactions Between Structure and Processing that Control Moisture Uptake in High-Performance Polycyanurates

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Outline: Basic Studies of Moisture Uptake in Cyanate Ester Networks



- Background / Motivation
- SOTA Theories of Moisture Uptake in Thermosetting Networks
- New Tools and New Discoveries
- Unresolved Issues and Ways to Address
 Them

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Importance of Moisture Uptake in Composite Component Performance





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- Water can add significantly to launch or take-off weight (3% water in composite resins = about 50 lbs of extra weight on an large SRM)
- Items with high water content can fail catastrophically when suddenly heated
- Long-term exposure to water can facilitate many mechanisms of chemical degradation, necessitating substantial "knock down" factors in design allowables
- Though more stable than epoxy resins, cyanate esters can degrade on long-term exposure to hot water

Model High-Temperature Thermosetting Polymers: Cyanate Esters



- Glass transition temperatures at full cure of 200 400°C
- Uncured resins exist as low-melting solids, or low to moderate viscosity liquids, making them ideal for processes such as filament winding
- Broad compatibility with co-monomers, thermoplastic tougheners, or nanoparticles for control of physical and mechanical characteristics
- Single species reaction chemistry is "cleaner" than epoxy resin and well-understood; enables development of superior predictive models for failure; readily catalyzed to cure at reasonable temperatures

Cyanate Esters: Next-Generation High-Performance Composite Resin





• Many opportunities for technical transition beyond SRM cases ...

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• In addition to glass transition temperature, many properties, such as density and water uptake, are mainly functions of conversion and monomer type

Basic Understanding of Moisture Uptake in Highly Cross-linked Networks





- Cure results in:
 - Net Shrinkage
 - Less permeability
 - Higher modulus
 - Brittleness



Increasing conversion joins together more "loose ends" in the network, eliminating free space where water can be absorbed, therefore water uptake is expected to decrease with increasing conversion

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Proposed Role of Vitrification in Controlling Moisture Uptake





- Cure results in:
 - Net Expansion
 - Higher permeability
 - Lower modulus
 - **Toughness**



Increasing conversions join "loose ends", but in the process they "freeze" the individual molecular centers-of-mass in place, resulting in exactly the opposite of the effects predicted by the traditional picture.

New Tools for Understanding Network



Typical computed DSC baseline with estimated zones of uncertainty

- BADCy catalyzed with 2 phr nonylphenol + 160 ppm Cu in Cu(II)Acac: $dV_m/dn_{tr} = 46 \pm 4$ cc/mol vs. previously determined value (2012) of 37 ± 15 cc/mol
- New DSC method allows for more precise and objective assessment of conversion, combined with T_G assessment for determination of conversion to within 0.01.
- New density technique adapted for use by Mr. Michael Ford allows for 100x faster assessment with high precision.

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- In accordance with properties of water in cyanurate networks (i.e. higher miscibility at lower temperatures, no clustering)
- Recent NASA study of cryogenic toughness of cyanate ester / carbon fiber composites showed that residual stress minimization (by minimization of cure temperature) was the most important factor in achieving good toughness at cryogenic temperatures



Moisture Uptake Is Likely a Minor Influence on Cryogenic Toughness



							-			
Material	B - Thickness		W - Height		a - Notch + Crack Length		P _Q - Applied Load		K _q (K _{1c})	
Specimen ID	(in)	(mm)	(in)	(mm)	(in)	(mm)	(lbf)	(N)	(psi-in ^{1/2})	(MPa-m ^{1/2})
Catalyzed LECy Panel Cured 1-6-14 Water Soaked-Tested 4-16-14										
STL300-1	0.1429	3.630	0.3920	9.957	0.1895	4.813	24.24	107.8	2,739	3.01
STL300-2	0.1456	3.698	0.4270	10.85	0.1985	5.042	27.70	123.2	2,782	3.06
STL300-3	0.1452	3.688	0.4280	10.87	0.2150	5.461	19.89	88.47	2,247	2.47
STL300-4	0.1508	3.830	0.4640	11.79	0.2195	5.575	27.76	123.5	2,647	2.91
STL300-5	0.1423	3.614	0.4310	10.95	0.2115	5.372	27.67	123.1	3,064	3.37
								Average	2,696	2.96
Catalyzed LECy (1-10-14) Tested 4-17-14										
STL300-6	0.1413	3.589	0.3575	9.081	0.1815	4.610	22.30	99.19	2,881	3.17
STL300-7	0.1472	3.739	0.3335	8.471	0.1810	4.597	19.89	88.47	2,868	3.15
STL300-8	0.1424	3.617	0.3545	9.004	0.1625	4.128	21.75	96.74	2,404	2.64
STL300-9	0.1408	3.576	0.3575	9.081	0.1710	4.343	27.20	120.99	3,216	3.53
STL300-10	0.1425	3.620	0.3520	8.941	0.1605	4.077	19.82	88.16	2,181	2.40
								Average	2,710	2.98

- LECy samples were soaked for 28 days at room temperature so as to maximize water uptake (1.5 wt%) while minimizing carbamate formation ($\Delta V/V \sim 0.2\%$).
- No significant difference between exposed and control samples in fracture toughness at RT; same glassy state dynamics should apply at cryogenic temperatures
- The water in cyanate ester networks appears to be neither clustered nor strongly bound



Nanoscale Reinforcement and Interphases





- A slight amount of water does accumulate at hydrophilic interphases in cyanate esters, but other effects such as poor bonding and damage tend to overwhelm these effects
- Nanoscale reinforcements such as graphene oxide with extreme water transport characteristics can alter the water uptake significantly.
- Generally speaking, blends and co-networks tend to follow linear rules of mixtures



Effect of Cyanurate Density on Moisture Uptake





3



- Each monomer has a different water uptake as a function of cyanurate density, although all show an increasing trend
- The free volume / cyanurate density relationships are similar for all monomers

4



Effect of Methylation on Moisture Uptake





- Addition of a methyl group near the cyanurate oxygen causes a significant reduction in the tendency of moisture uptake to increase at high conversions
- Addition of a methyl group far from the cyanurate oxygen has no effect on water uptake as a function of conversion
- Methyl groups near the cyanurate oxygen block the favored sites for water uptake
- Vitrification appears not to influence the results
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Effect of Silicon Substitution on Moisture Uptake





- Substitution of a silicon atom for a carbon atom in di(cyanate ester) network segments reduces moisture uptake by up to 50%
- In the corresponding tri(cyanate ester) networks, the same substitution increases moisture uptake by ~100% (i.e. effect is not intrinsic property of Si atom)
- Differences in vitrification also do not explain these effects, as they cause little change in moisture uptake

Summary: Basic Studies of Moisture Uptake in Cyanate Ester Networks



- Many aspects of moisture uptake (and its minimization) in thermosetting polymer networks have been clarified through recent research efforts at AFRL/RQRP
- New tools and techniques for quantifying structure and properties of thermosetting networks have allowed for significant new insights into structureproperty relationships in thermosetting networks
- Several unresolved issues relating to our understanding of moisture uptake in cyanate ester networks remain; approaches based on molecular modeling may provide insights leading to significant application payoffs

QUESTIONS?

