Thermal/Mechanical Properties of Wood-PVC Composites – Effect of Maleation

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# Introduction

Maleation in wood-polymer composites helps create chemical bridges at the interface.

- Improving compatibility between polar wood and non-polar polymer
- Helping transfer stresses at the interface
- Improving interfacial adhesion strength

 Maleation influences mechanical and thermal properties of resultant composites.

Heat flow, heat capacity, and enthalpy
Glass transition
Moduli and bonding strength

# Objectives

To investigate thermal/mechanical characteristics of maleated wood-PVC composites.

To study the relationship between measured properties and coupling agent performance in resultant composites.

# Background

Thermal/Mechanical Analysis Techniques

### **Temperature-molecular Mass Diagram**

#### Semi-polymers (e.g., PVC and Lignin)



#### **Transition Temperatures**



### **Glass Transition Temperature T**g





### Stress-strain Relationship Under Dynamic (sinusoidal) Loading



## <u>Dynamic Stress-strain</u> <u>Relationship</u>

Stress  $\sigma(t)$  under a sinusoidal load:

Strain  $\gamma(t)$  by a phase angle  $\delta$  corresponding to the stress  $\sigma(t)$ :

Dynamic modulus E\*:

Relationship among complex, storage, and loss moduli:

Phase angle  $\delta$ :

 $\sigma(t) = \sigma_0 \sin(\omega t + \delta)$ 

 $\gamma(t) = \gamma_0 \sin(\omega t)$ 

$$E^*(\omega) = \frac{\sigma(t)}{\gamma(t)}$$

 $E^{*}(\omega) = E'(\omega) + iE''(\omega)$ 

 $\tan \delta = \frac{E''(\omega)}{E'(\omega)}$ 

#### **Thermal/Mechanical Properties**

Glass transition temperature T<sub>g</sub>
 DSC and DMA
 Melting temperature T<sub>m</sub>

- DSC

Heat flow (dQ/dt) and enthalpy (AH)

- DSC

Bonding moduli (E', E'', and  $E^*$ ) and the phase angle ( $\delta$ )

– DMA

Thermal stability (weight loss under heat)
 - TGA

# Experimental

#### Materials

Wood Veneer - Yellow poplar (0.91 mm Thick)
PVC film - Clear (0.0762 mm Thick)
Maleated polypropylene (MAPP)

Epolene E-43 (Mw =9,100)
Epolene G-3015 (Mw =47,000)

Initiator - Benzyol peroxide
Solvent - Toluene

Sohxlet Extraction ASTM standard D1105-96. Wood specimens were extracted for 4 hours with two sets of solvents. Coupling Treatment Wood specimens were dipped in the coupling solutions of 0, 12.5, 25, and 50 g/L MAPP at 100°C for 5 min under a continuous stirring with a magnetic stirrer.

#### Wood Veneer under Sohxlet extraction



Manufacture of wood-PVC composites Pressure: 0.276 MPa Pressing procedure: Heating 3 min at 178°C and then cooling at 70°C for 1 min Shear strength measurement Shear tests followed the ASTM standards D3163 and D3165

### Wood-PVC Laminates under Shear Testing



# DMA (Seiko Instruments, Model DMS 110)



### DMA Procedure - Using three cycles

			Temperature [°C]		Rate
Specimen	Test mode	Test cycle	Start	Stop	[°C/min]
		First heating	20	220	0.50
Wood	Bending	First cooling	220	30	0.25
		Second heating	30	220	0.50
		First heating	20	100	0.50
PVC	Bending	First cooling	100	30	0.25
		Second heating	30	100	0.50
		First heating	20	150	0.50
Woo-PVC	Bending	First cooling	150	30	0.25
composites		Second heating	30	150	0.50

## TGA system (TA Instruments, Model TGA2950)



Procedure: Heating from 25°C to 600°C under a N<sub>2</sub> flux at a pressure of 8 KPa

### **DSC** (TA Instruments, Model DSC2920)





#### Procedure

For interphase samples, heating from  $25^{\circ}$ C to  $200^{\circ}$ C under a  $N_2$  flux at a pressure of 8 KPa

For modified wood veneer and wood-PVC composite samples, cooling at -10°C for a while and then heating up to 200°C in a  $N_2$ flux

#### Summary Results on Thermal/Mechanical Properties

Material	E' (GPa) <sup>a</sup>	E" (GPa) <sup>a</sup>	Glass transition (°C) <sup>a</sup>	tanð <sup>a</sup>	Shear strength (MPa)	Enthalpy (J/g) <sup>b</sup>	TG at 600°C (%)	DTG <sub>max</sub> (%/°C) <sup>c</sup>
PVC	5.73	0.44	76.1	0.39	-	0.81 @79.6°C	10.3	2.37 @257°C
Wood	10.43	0.41	67.2	0.05	-	21.69 @50.7°C	18.8	1.47 @356°C
Wood-PVC composites:								
0% MAPP	7.85	1.04	85.1	0.22	3.14	-	16.8	0.80 @266°C,
2.95% E-43	7.96	0.97	85.9	0.22	2.90	-	17.9	0.65 @329°C 0.75 @270°C, 0.62 @340°C
4.12% E-43	9.45	1.23	83.0	0.24	3.03	15.95 @88.0°C	18.2	$0.69 @266^{\circ}C,$
6.83% E-43	9.16	1.15	82.6	0.23	3.32	-	16.5	0.67 @275°C, 0.63 @344°C
2.17% G-3015	7.08	0.8	85.5	0.23	2.90	-	16.5	0.72 @266°C,
3.64% G-3015	8.98	1.16	83.9	0.24	2.94	15.99 @81.8°C	15.4	0.75 @344°C 0.72 @258°C,
6.35% G-3015	8.56	1.09	84.3	0.24	3.61	-	17.0	0.71 @276°C, 0.58 @335°C

<sup>a</sup> The value was measured in first heating at 1 Hz;

<sup>b</sup> The value was measured at the glass transition;

<sup>c</sup> Two maximum peaks were selected for wood-PVC composites.

# DMA Results

#### Glass Transitions of Wood-PVC Composites



# Influence of Frequency on E' and tan $\delta$ of Wood-PVC Composites with 6.83% E-43



#### Influence of MAPP Retention on E' and tan $\delta$ of Wood-PVC Composites (Freq = 1 Hz)



# TGA Results

#### Influence of Maleation on Decomposition of Wood-PVC Composites by TG



#### Influence of Maleation on Decomposition of Wood-PVC Composites by DTG



#### Comparisons on DTG Decomposition of Wood-PVC Composites with and without Maleation



# DSC Results

#### Heat Flow (dQ/dt) vs. Temperature For Wood-PVC Composites



# Derivative DSC spectra for PVC, modified wood veneer, and wood-PVC composites



#### **DSC Spectra of PVC-MAPP Interphases**



### Conclusions

- Maleation significantly influenced the thermal behavior of wood-PVC composites.
- E'and E\* increased with MAPP retention and graft rate. However, tanδwas independent of retention and graft rate.
- Wood-PVC composites with MAPP had significant shifts in DMA, DSC, and TG/DTG spectra compared with those without MAPP.