



TECHNOLOGY DRIVEN. WARFIGHTER FOCUSED.

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5th Triennial International Aircraft Fire and Cabin Safety Conference, Atlantic City, NJ, November 1, 2007

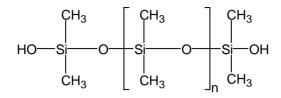




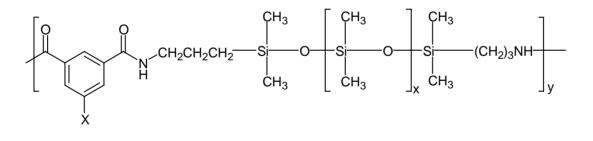
- Success of military operations severely impacted by burn injuries (loss of personnel and high medical treatment costs \$10's of millions/year)
- Polymers such as Kevlar and Nomex offer a high degree of flame retardancy, but at a high cost
- Manufacture and processing requires the use of hazardous ingredients and solvents
- ✤ Highly insoluble and used as fibers, not easily coated onto objects
- Although current FR materials (Halogenated, Phosphorous, nitrogen and Inorganic compounds) additives such as halogenated, aluminium trihydroxide, antimony oxides may be a cost effective solution in many FR applications, but they have long term ecological problems due to the release of toxic gases upon combustion











1: X = OH; **2**: X = NH₂

To Save Lives while making or using the FR materials — We do not want to kill people or cause damage to the environment. Do We?

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- Mild reaction conditions
- Efficient and economical
- Servironmentally benign
 Chemistry
- Easy work up
- Regioselective





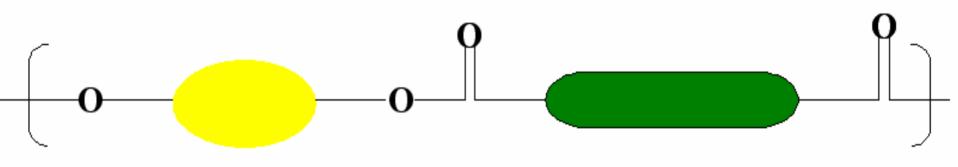
- Regio-, Chemo-, Enantio-, Diastereo-selectivity
- Proven track record of catalyzing transesterification, (trans)amidation and imidation reactions
- Flexible reaction conditions
 - Temperatures (40 90°C)
 - No solvent required
 - If solvent necessary, its compatible



Lipases in Polymer Synthesis

SARMY NATICA

Condensation Polymerization



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RMYNA Lipase Catalyzed Synthesis of RDECOM Siloxane Based Aromatic Co-Polymers CH₃ X-(H₂C)₃--(CH₂)₃-X MeO OMe ĊH₂ ĊH₃ ĊH₃ OH (OCH₂CH₂)OH X = X = NH_2 Novozyme-435, Bulk (No solvent) 48 hrs 90°C, vacuum

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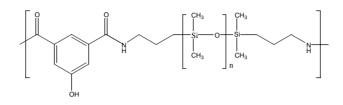
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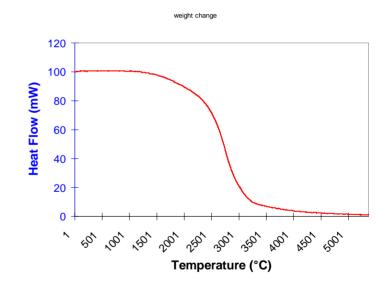


Polysiloxane co-Polyamide





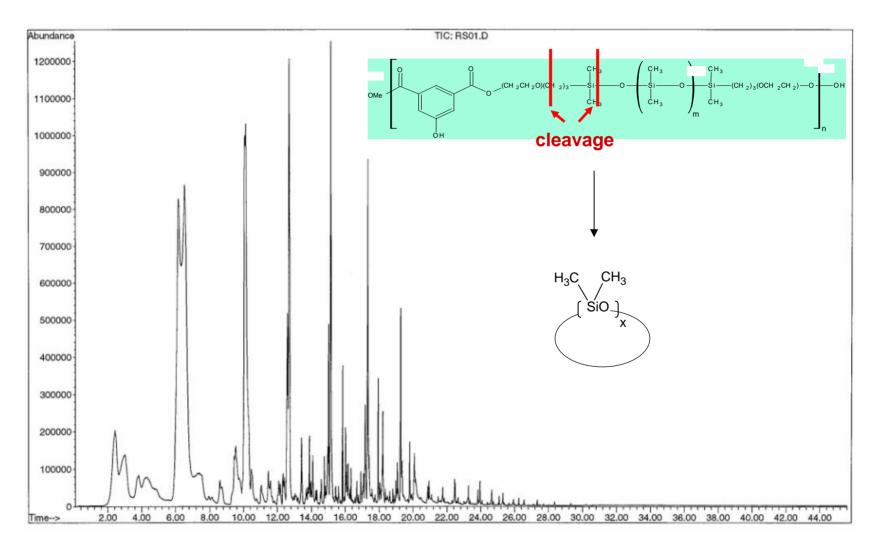
TGA	
T _{onset} (°C)	403
T _{max} (°C)	462
Char yield (%)	10.2
PCFC	
Heat release capacity (J/g.K)	260
Total heat release (KJ/g)	21.2



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RDECOM Py GC/MS of Siloxane copolyester

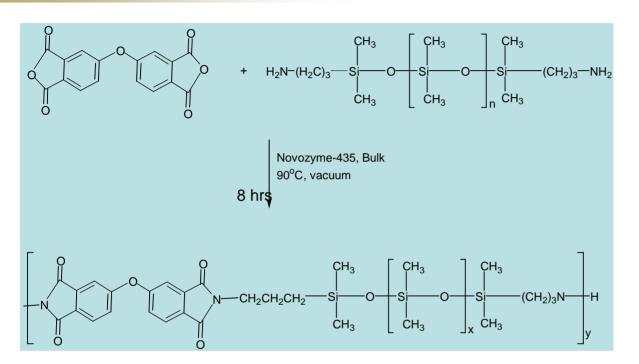


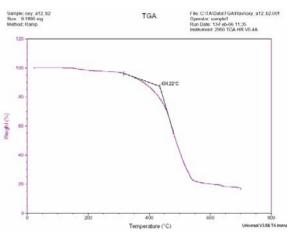


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RDECOM Enzymatic Synthesis of Polyimides







Direct Polyimide formation without polyamic acid intermediate

$$Yield = ~80\%$$



Only 8 hrs (48 hours for Polyesters and Polyamides)

Ave. GPC MW = 75K, PD = 1.7 (prior 20K was highest)

 $T_{dec} = 460 \text{ °C}$ (10wt% wt loss) HRC = 313 J/g K

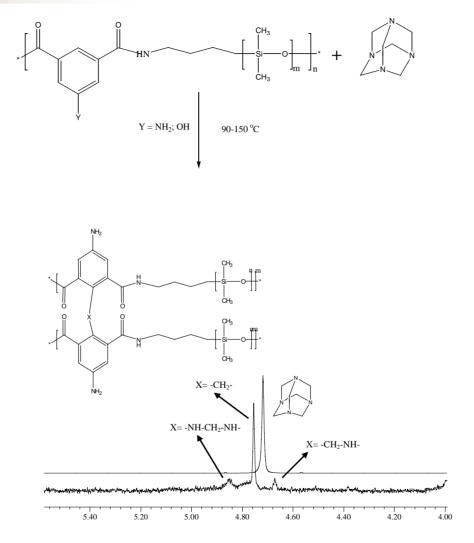
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Mosurkal et al, *Macromolecules* **2007**, *40*, 7742. **10**



Crosslinking of Polyamide





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FR Properties of HMTA cross-linked Polyamide



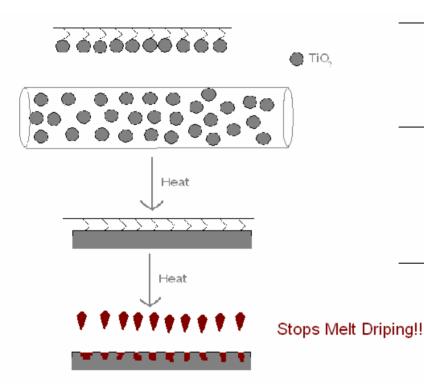
Crosslinking %HMTA (by weight)	HR capacity (J/g*K)	char yield %
0%	194	11
1%	173	12.3
5%	156	14.4
10%	108	14.9
15%	125	12.5
20%	90	13.2

Mosurkal et al, Polym. Prepr. (Am. Chem. Soc., Div. Polym. Chem.) 2006, 47, 1110.

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Polymer-TiO₂ Nanocomposites for Improved FR materials





 Covalent attachment of metal oxide nanoparticles (TiO₂) to FR polymer

→ Nanoparticle incorporated into FR film or fiber

Upon heat, nanoparticles sinter to form porous metal oxide layer which serves to absorb polymer melt, suffocate heat and increase heat of decomposition.

Advantages

- Controllable covalent attachment of nanoparticle to the polymer
- Increase the heat of decomposition
- Suffocates heat
- Melt absorber minimize melt drip
- Environmentally benign
- Inexpensive (paints, toothpaste, fillers)

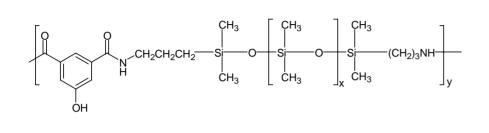
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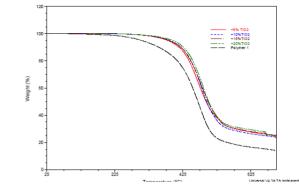
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RDECOM Poly-TiO₂ nanocomposites

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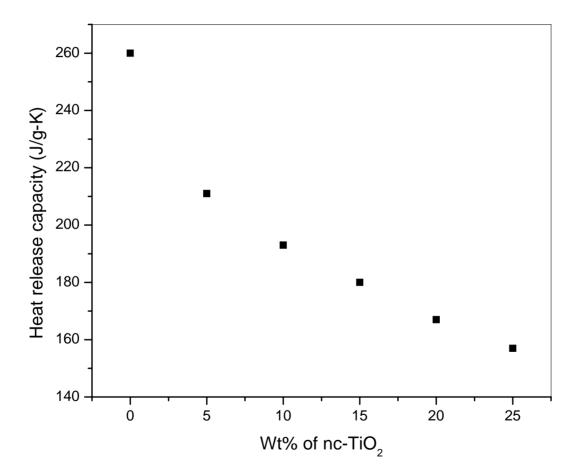


wt%	PC	FC	TGA	N
of TiO ₂	HRC (Jg¹ _κ ¹)	Total HR (KJg ⁻¹)	T _{dec} (10% wt loss) (° _{C)}	Char Yield⁵
0	260.2	21.2	408	14.0
5	211.1	11.0	413	23.9
10	192.7	10.7	417	24.1
15	179.7	10.0	418	24.9
20	167.5	9.70	425	28.0

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Heat release vs. Wt% of TiO₂



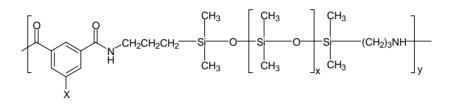
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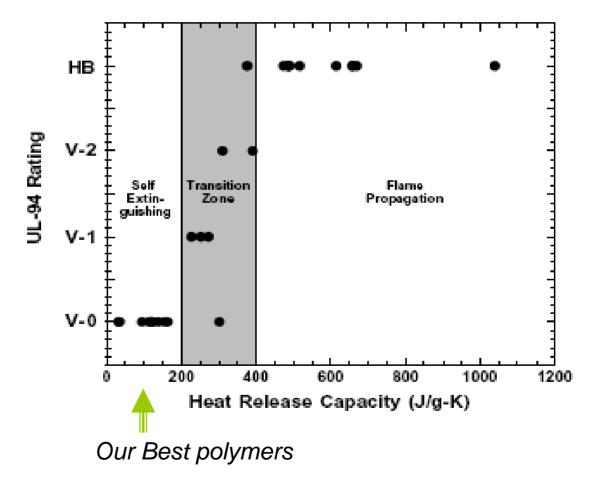
1: X = OH; **2**: X = NH₂

Polymer	P	CFC		TGA
	HRC (Jg ⁻¹ κ ¹)	Total HR (KJg ⁻¹)	T _{dec} ^a (° _{C)}	Char Yield⁵
1	260.2	21.1	408	14.0
+TiO ₂ ^c	167.5	9.7	425	31.2
2	194.0	15.8	407	15.0
+TiO ₂ ^c	128.9	14.7	426	33.7

a at 10% weight loss; b o c 20wt% of TiO2

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HR Capacity vs. UL-94 rating



Ref: Lyon et al, FAA

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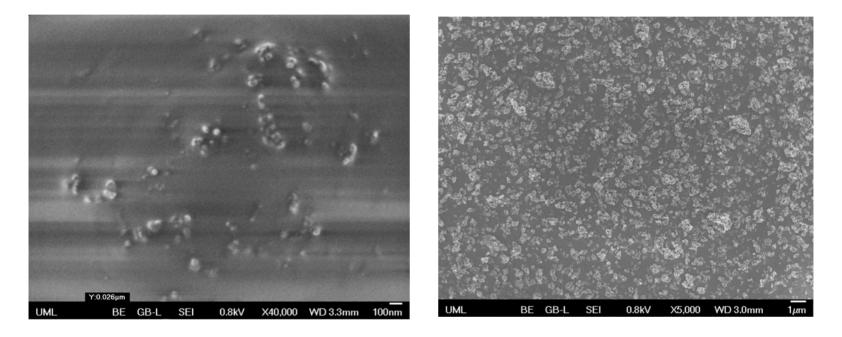
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Thin film spincoated on ITO glass plate



Before Burn

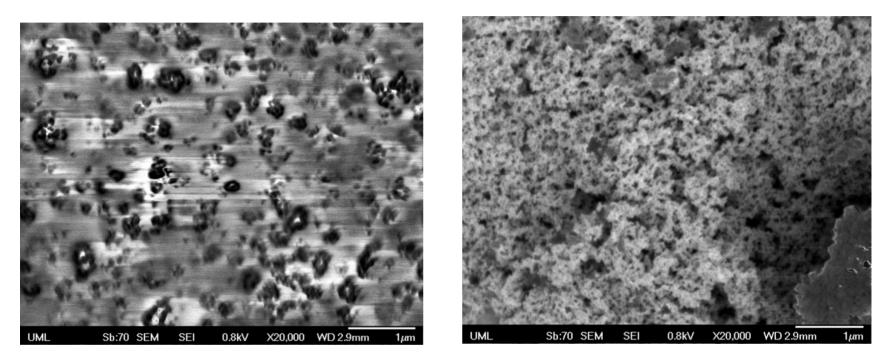
After Burn (500 ^OC /1hr)

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STORE R POCE CENTS

Thick film spincoated on ITO glass plate and burnt at 500 °C for 1 hr



Char underneath surface (white)

Char-W

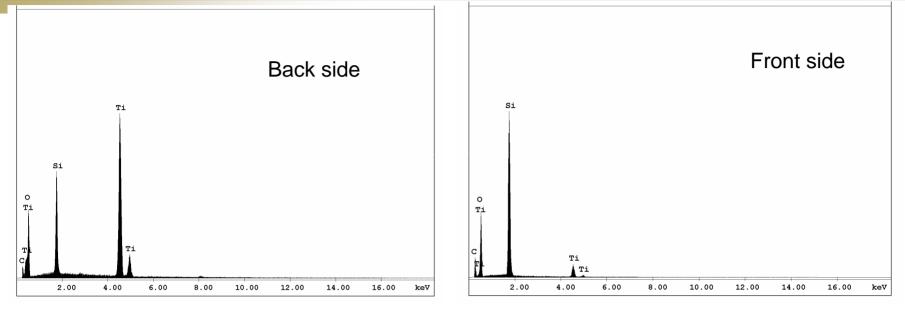
Char upside Surface (black)

Char-B

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Char-W

Char-B

Atomic %

	С	0	Si	Ti
Char-W	13.8	57.7	9.2	19.3
Char-B	39.4	43.2	15.7	1.7

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Polymer/TiO₂ Composite Coated Camouflage Fabric (cotton/nylon blend)



A12 - 20wt%TiO₂ on 3 x 1 in. fabric



Burning of A12-TiO2 coated camouflage fabric Self extinguish within 25sec. No melt drip Not ASTM standards

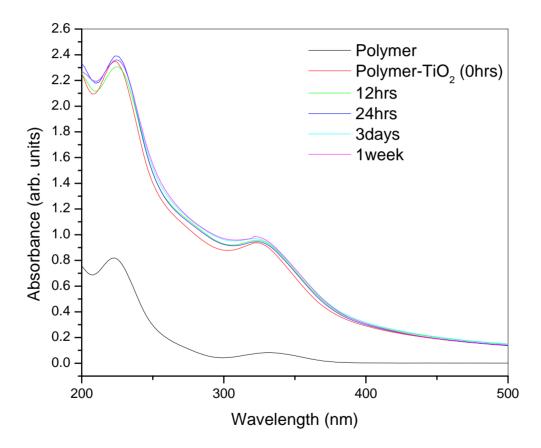
Protective swelled char formation around the fabric after burnt

When broken the protective Layer, fabric piece intact

Char is swelled and heat/mass transfer to the underlying material is hindered

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RDECOM Conclusions on TiO2-Polymer nanocomposites



- Increased thermal stability as TiO₂ limit the thermal conduction to the polymer inside and thus kinetics of degradation.
- Restricted mobility of polymer chains results from steric hindrance due to the presence of additive solid particles.
- Due to the increased viscosity of the melt with amount of TiO₂, the gas emission is hindered.
- Promotes char formation containing Si atoms (Silicate) on the surface which also hinders the heat transfer to the combustible gases.
- Photostability of the polymer in the presence of TiO_2 is good.



FR Material	Degradation Temp. (°C)	Heat release Capacity (J/g.K)	Environ- mental effects	Process- ability	Synthesis	Cost
KEVLAR	500-550	302	No toxic byproducts?	Highly Insoluble difficult to coat	Hazard ingredients Multi-step	Too expensive to provide every Soldier
NOMEX	500-550	52	No toxic byproducts?	Highly Insoluble difficult to coat	Hazard ingredients Multi-step	Too expensive to provide every Soldier
Our Best Performing Polymers	(Polyamide) 400 428** (Polyimide) 450	(Polyamide) 194 <u>132**</u> <u>90*</u> (Polyimide) 313	No toxic byproducts No toxic byproducts	Highly soluble and easy to coat Highly soluble to partially soluble	Enzymatic Synthesis No solvents, 1-step Enzymatic Synthesis No solvents, 1-step	Potential to be very cost effective

*cross-linked with 20% hexamethylenetetramine; **with 20% TiO₂ nanoparticles

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Polysiloxane-copolymers are a promising alternative to halogenated and other expensive fire safe polymers

- Flame retardancy comparable to well known FR polymers (high degradation temperatures, low heat release capacity)
- Environmentally benign synthesis
 (single step, enzymes as catalysts and no solvents)
- No toxic byproducts (cyclic siloxanes)
- Potentially processable for coatings, fibers, polymer blends
- Polymer-TiO₂ nanocomposites show improved FR and photostable
- SEM images showed that TiO₂ is helping to form Silicate char on the surface
- Potential to be low cost

RNFFA



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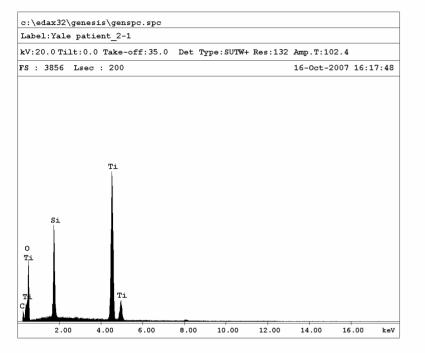






"Everything in this Universe burn at some temperatures, But if we can understand 'why' and 'how' a material burn, it may be possible to create an efficient, economical and more importantly environmentally safe Fire Proof material which can save lives" **RDECOM** EDAX composition

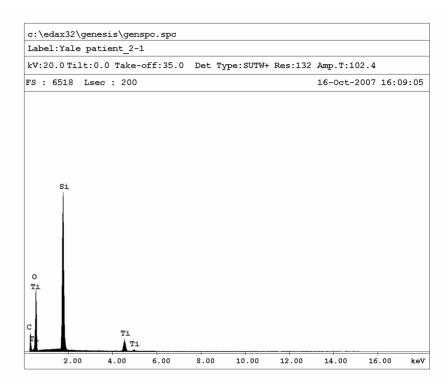




Filename	СК	ок	SiK	TiK
genspc.spc	28.03	40.98	26.22	4.77
genspc.spc	7.30	40.65	11.38	40.67

Atomic % by Element

2				
Filename	СК	ок	SiK	TiK
genspc.spc	39.37	43.20	15.75	1.68
genspc.spc	13.80	57.71	9.20	19.29



Weight % by Element

Filename	ск	ок	SiK	TiK
genspc.spc	28.03	40.98	26.22	4.77

Atomic % by Element Filename C K O K SiK TiK genspc.spc 39.37 43.20 15.75 1.68

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