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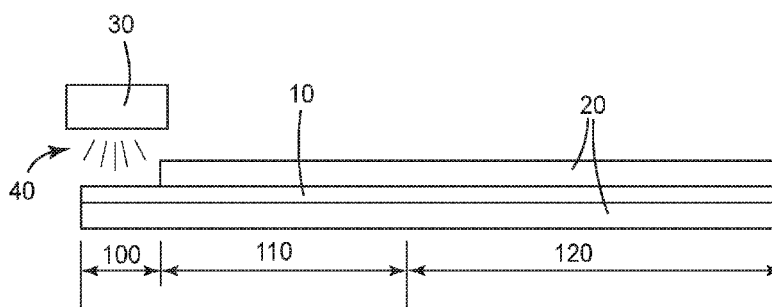


FIG. 1

(57) Abstract: Compositions that are curable to polythioether polymers are provided, comprising: a) a dithiol monomer; b) a diene monomer; c) a radical cleaved photoinitiator; d) a peroxide; and e) an amine; where the peroxide and amine together are a peroxide-amine redox initiator. In some embodiments, the amine is a tertiary amine. In some embodiments, the amine is selected from the group consisting of dihydroxyethyl-p-toluidine, N,N-diisopropylethylamine, and N, N, N', N', N''-pentamethyl-diethylenetriamine. In some embodiments, the peroxide is selected from the group consisting of di-tert-butyl peroxide, methyl ethyl ketone peroxide, and benzoyl peroxide. In some embodiments, the composition may additionally comprise a polythiol monomer having three or more thiol groups.

DUAL CURE POLYTHIOETHER

Field of the Disclosure

This disclosure relates to compositions that cure to form polythioether polymers
5 and that include a dual cure curing mechanism, as well as sealants comprising the same.

Summary of the Disclosure

Briefly, the present disclosure provides compositions that are curable to
10 polythioether polymers, comprising: a) a dithiol monomer; b) a diene monomer; c) a radical cleaved photoinitiator; d) a peroxide; and e) an amine; where the peroxide and amine together are a peroxide-amine redox initiator. In some embodiments, the amine is a tertiary amine. In some embodiments, the amine is selected from the group consisting of dihydroxyethyl-p-toluidine, N,N-diisopropylethylamine, and N, N, N',
15 N'', N''-pentamethyl-diethylenetriamine. In some embodiments, the peroxide is selected from the group consisting of di-tert-butyl peroxide, methyl ethyl ketone peroxide, and benzoyl peroxide. In some embodiments, the composition may additionally comprise a polythiol monomer having three or more thiol groups. In some embodiments, the composition may additionally comprise one or more fillers. In some
20 embodiments, the composition may additionally comprise one or more nanoparticle fillers. In some embodiments, the composition may additionally comprise calcium carbonate. In some embodiments, the composition may additionally comprise nanoparticle calcium carbonate. In some embodiments, the composition may be cured by application of light from an actinic light source. In some embodiments, the
25 composition may be cured by application of light from a blue light source. In some embodiments, the composition may be cured by application of light from a UV light source.

In another aspect, the present disclosure provides sealants comprising curable compositions according to the present disclosure. In another aspect, the present
30 disclosure provides seals obtained by cure of such sealants.

In another aspect, the present disclosure provides polythioether polymers obtained by cure of any the compositions according to the present disclosure. In some

embodiments, the polythioether polymer has a Tg less than -50°C. In some embodiments, the polythioether polymer exhibits high jet fuel resistance characterized by a volume swell of less than 30% and a weight gain of less than 20% when measured according to Society of Automotive Engineers (SAE) International Standard AS5127/1.

5 All scientific and technical terms used herein have meanings commonly used in the art unless otherwise specified.

As used in this specification and the appended claims, the singular forms “a”, “an”, and “the” encompass embodiments having plural referents, unless the content clearly dictates otherwise.

10 As used in this specification and the appended claims, the term “or” is generally employed in its sense including “and/or” unless the content clearly dictates otherwise.

As used herein, “have”, “having”, “include”, “including”, “comprise”, “comprising” or the like are used in their open ended sense, and generally mean “including, but not limited to.” It will be understood that the terms “consisting of” and
15 “consisting essentially of” are subsumed in the term “comprising,” and the like.

Brief Description of the Drawing

FIG. 1 is a diagram representing the use of a dual cure polythioether according to the present invention as a sealant between two substrates.

20

Detailed Description

The present disclosure provides a dual cure polythioether material such as a sealant. In some embodiments the sealant has both cure on demand properties and dark cure properties. In some embodiments the sealant may be cured by exposure to actinic
25 radiation in 120 seconds or less, in some embodiments 90 seconds or less, in some embodiments 60 seconds or less, in some embodiments 30 seconds or less, and in some embodiments 20 seconds or less. In some embodiments cure on demand can be initiated with UV light, in some with visible light, in some with LED sourced UV light, and in some with LED sourced visible light. Furthermore, in some embodiments the
30 initially photoinitiated cure propagates by a dark cure mechanism into adjacent portions of the polythioether material that are without light access. In some embodiments cure propagates by at least 2.5 cm into the dark zone with 90% or greater conversion in 120

seconds or less, in some embodiments 90 seconds or less, in some embodiments 60 seconds or less, in some embodiments 30 seconds or less, and in some embodiments 20 seconds or less. In some embodiments cure propagates by at least 5.0 cm into the dark zone with 90% or greater conversion in 120 seconds or less, in some embodiments 90
5 seconds or less, in some embodiments 60 seconds or less, in some embodiments 30 seconds or less, and in some embodiments 20 seconds or less.

With reference to FIG. 1, in one embodiment of a sealant according to the present disclosure, sealant **10** is applied between opaque substrates **20**. Actinic light **40** from actinic light source **30** is used to initiate cure of a portion of sealant **10** exposed to
10 said light, in exposed zone **100**. However, sealant **10** in dark zones **110** and **120** is not exposed to actinic light **40**. Use of the dual cure initiation system of the present disclosure allows the initially photoinitiated cure to propagate by a dark cure mechanism into adjacent dark zones **110** and **120**. In some embodiments, sealant in dark zone **110**, representing the 2 inches (5 cm) immediately adjacent to exposed zone
15 **100**, is fully cured within 20 seconds, and cure continues further into dark zone **120**.

In some embodiments, the initiation system includes a) a radical cleaved photoinitiator and b) a peroxide-amine redox initiator.

In some embodiments, the cured material has low glass transition temperature, in some embodiments less than -20 °C, in some embodiments less than -30 °C, in some
20 embodiments less than -40 °C, and in some embodiments less than -50 °C. In some embodiments, the cured material has excellent fuel resistance properties. In some embodiments, the cured material combines low glass transition temperature of less than -50 °C with excellent fuel resistance properties. Thus, in certain embodiments this dual cure technology can be applied to aircraft or automobile sealant applications and may
25 result in greater ease and speed of vehicle manufacture.

Selected Embodiments

The following numbered embodiments are intended to further illustrate the present disclosure but should not be construed to unduly limit this disclosure.

30

1. A composition that is curable to a polythioether polymer, comprising:
a) a dithiol monomer;

- b) a diene monomer;
- c) a radical cleaved photoinitiator;
- d) a peroxide; and
- e) an amine;

5 where the peroxide and amine together are a peroxide-amine redox initiator.

2. The composition according to embodiment 1 where the amine is a tertiary amine.

10 3. The composition according to any of the preceding embodiments where the amine is selected from the group consisting of dihydroxyethyl-p-toluidine, N,N-diisopropylethylamine, and N, N, N', N'', N'''-pentamethyl-diethylenetriamine.

15 4. The composition according to any of the preceding embodiments where the peroxide is selected from the group consisting of di-tert-butyl peroxide, methyl ethyl ketone peroxide, and benzoyl peroxide.

5. The composition according to any of the preceding embodiments additionally comprising:

20 f) a polythiol monomer having three or more thiol groups.

6. The composition according to any of the preceding embodiments additionally comprising:

g) at least one filler.

25

7. The composition according to any of the preceding embodiments additionally comprising:

h) at least one nanoparticle filler.

30 8. The composition according to any of the preceding embodiments additionally comprising:

j) calcium carbonate.

9. The composition according to any of the preceding embodiments additionally comprising:

k) nanoparticle calcium carbonate.

5

10. The composition according to any of the preceding embodiments which is curable by actinic light source.

11. The composition according to any of the preceding embodiments which is curable by blue light source.

10

12. The composition according to any of the preceding embodiments which is curable by UV light source.

13. A sealant comprising the composition according to any of the preceding embodiments.

15

14. A polythioether polymer obtained by cure of any the composition according to any of embodiments 1-12.

20

15. The polythioether polymer according to embodiment 14 having a Tg less than -50°C.

16. The polythioether polymer according to embodiment 14 or 15 which exhibits high jet fuel resistance characterized by a volume swell of less than 30% and a weight gain of less than 20% when measured according to Society of Automotive Engineers (SAE) International Standard AS5127/1.

25

Objects and advantages of this disclosure are further illustrated by the following examples, but the particular materials and amounts thereof recited in these examples, as well as other conditions and details, should not be construed to unduly limit this disclosure.

30

Examples

Unless otherwise noted, all reagents were obtained or are available from Sigma-Aldrich Company, St. Louis, Missouri, or may be synthesized by known methods.

5 Unless otherwise reported, all ratios are by weight percent.

The following abbreviations are used to describe the examples:

	°C:	degrees Centigrade
	cm:	centimeter
10	LED:	light emitting diode
	mL:	milliliter
	Mn:	Molecular weight
	mW:	milliWatt
	nm:	nanometer
15	T _g :	glass transition temperature
	UV:	ultraviolet

Definitions

Maximum Processing Time: The time taken for the thiol-ene curable composition to
20 begin gelling.

Photo Cure Time: The cure time of the portion of thiol-ene curable composition exposed to light.

Dark Cure Time: The cure time of the portion of thiol-ene curable composition not exposed to light.

25 Redox Cure Time: The cure time for the portion of thiol-ene curable composition initiated by redox mechanism.

Dark Cure Distance: The length of thiol-ene composition dark cured, as measured from the leading edge of the opaque silicone rubber sheet.

30

Materials.

Abbreviations for the materials used in the examples are as follows:

- A-200: A hydrophilic fumed silica, obtained under the trade designation “AEROSIL 200” from Evonik Industries AG, Essen, Germany.
- BPO: Benzoyl peroxide.
- 5 CMP: 3-chloro-2-methyl-1-propene.
- DABCO: 1,4-Diazabicyclo[2.2.2]octane, obtained under the trade designation “DABCO” from Air Products & Chemicals, Inc., Allentown, Pennsylvania.
- DHEPT: dihydroxyethyl-p-toluidine.
- 10 DIPEA: N,N-diisopropylethylamine, obtained from Alfa Aesar, Ward Hill, Massachusetts.
- DMDO: 1,8-Dimercapto-3,6-dioxaoctane, obtained from Arkena, Inc., King of Prussia, Pennsylvania.
- DSW: An aluminosilicate clay, obtained under the trade designation “DRAGONITE SELECT WHITE” from Applied Minerals, Inc., New York, New York.
- 15 DVE-3: Triethyleneglycol divinylether, obtained under the trade designation “RAPI-CURE DVE-3” from Ashland Specialty Ingredients, Wilmington, Delaware.
- 20 E-8220: A diglycidylether of bisphenol F, obtained under the trade designation “EPALLOY 8220” from Emerald Performance Materials, LLC, Cuyahoga Falls, Ohio.
- I-819: Phenylbis(2,4,6-trimethylbenzoyl)phosphine oxide, obtained under the trade designation “IRGACURE 819” from BASF Corp., Florham Park, New Jersey.
- 25 MEHQ: Hydroquinone monomethyl ether.
- MEKP: Methyl ethyl ketone peroxide, obtained from 3M Company, St. Paul, Minnesota.
- PMETA: N, N, N', N'', N'''-pentamethyl-diethylenetriamine, obtained from TCI America, Portland, Oregon..
- 30 S-31: Nanoparticle (70 – 100 nm) calcium carbonate, obtained under the trade designation “SOCAL 31” from Solvay Chemicals, Inc., Houston, Texas.

- TAC: Triallylcyanurate, obtained from Sartomer, Inc., Exton, Pennsylvania.
- TBPO: di-tert-butyl peroxide, obtained from TCI America.
- TEMPO: 2,2,6,6-tetramethyl-1-piperidinyloxy, obtained from Oakwood Products, Inc., West Columbia, South Carolina.
- 5 TPTMP: Trimethylolpropane tris(3-mercaptopropionate).
- TVCH: 1,2,4-Trivinylcyclohexane, obtained from BASF Corp., Florham Park, New Jersey.
- TVBT: tris[4-(vinylxy)butyl] trimellitate, obtained from Allied Signal, Inc., Morristown, New Jersey.
- 10 VAZO 52: 2,2'-azobis(2,4-dimethyl-pentanenitrile), obtained under the trade designation "VAZO 52" from E.I. du Dupont de Nemours and Company, Wilmington, Delaware.
- VAZO 67: 2,2'-azobis-(2-methylbutyronitrile), obtained under the trade designation "VAZO 67" from E.I. du Dupont de Nemours and Company.

15

Polythioethers were synthesized as follows:

- PTE-1. Into a 1000-mL round bottom flask equipped with an air-driven stirrer, thermometer, and a dropping funnel, was added 392.14 grams (2.15 moles) DMDO and
- 20 82.23 grams (0.25 moles) E-8220. To this mixture was added 0.15 grams DABCO. The system was flushed with nitrogen, then mixed and heated for four hours at 60 - 70°C. 12.5 grams (0.05 moles) TAC was added, followed by approximately 0.15 grams VAZO 67. With continuous stirring, the mixture was heated to 60°C, and held at this temperature for approximately 30 - 45 minutes. 313.13 grams (1.55 moles)
- 25 DVE-3 were slowly added drop-wise to the flask over a period of 45 - 60 minutes, keeping the temperature at approximately between 68 - 80°C. Additional VAZO 67 was added in approximately 0.15 gram increments over approximately 6 hours, for a total of approximately 0.6 grams. The temperature was raised to 100°C and the material degassed for approximately 10 minutes. The resultant polythioether was
- 30 approximately 3200 Mn with a 2.2 functionality.

PTE-2. Into a 500 mL four-neck, round bottom flask fitted with a stirrer, thermometer, chilled water condenser and a pressure equalizing addition funnel was added 206.54 grams of a 20% aqueous solution of sodium hydroxide (1.033 moles). To this was added, drop wise with stirring, 94.08 grams (0.51 moles) DMDO, and the mixture then allowed to cool to approximately 21°C. 96.4 grams (1.065 moles) CMP was added drop wise with vigorous stirring, and stirring continued for another 2 hours. The mixture was then held at 21°C for approximately 16 hours, after which 150 grams of a clear layer was decanted. NMR analysis confirmed the decanted layer to be CMP diene.

Into a 100-mL round bottom flask equipped with an air-driven stirrer, thermometer, and a dropping funnel, was added 39.64 grams (0.22 moles) DMDO and 4.10 grams (0.0125 moles) E-8220. To this mixture was added 0.02 grams DABCO. The system was flushed with nitrogen, then mixed and heated for 1.5 hours at 60 - 70°C. 3.66 grams (0.0125 moles) CMP diene was added followed by approximately 0.01 grams VAZO 52. With continuous stirring, the mixture was heated to 60°C, and held at this temperature for approximately 1.5 hrs. 0.83 grams (0.005 mole) TVCH were added and the temperature maintained for another 1.5 hrs. 31.80 grams (0.157 moles) DVE-3 were slowly added drop-wise to the flask over a period of 45 - 60 minutes, keeping the temperature at approximately 70°C. Additional VAZO 52 was added in approximately 0.01 gram increments over approximately 16 hours, for a total amount of about 0.4 grams. The temperature is raised to 100°C and the material degassed for approximately 10 minutes. The resultant polythioether was approximately 3200 Mn with a 2.2 functionality.

Example 1

Part A was prepared by dissolving 0.0139 grams BPO and 0.0300 grams I-819 in 3.0000 grams DVE-3 in a 20 mL amber vial, on a roll mill for 40 minutes at 21°C. Part B was prepared by dissolving 0.0139 grams DHEPT and 0.0394 grams I-819 in 3.9407 grams TPTMP in a 20 mL amber vial, also on a roll mill for 8 hours at 21°C. Part A was then added to Part B and manually stirred for one minute until homogeneously dispersed.

Examples 2-8

The procedure generally described in Example 1 for preparing homogeneous mixtures of peroxide, photo initiator and vinyl monomer in Part A, and amine, photo initiator
5 and thiol monomer in Part B, was repeated according to the formulations listed in Tables 1A and 1B.

Example 9

Part A was prepared by dissolving 0.0347 grams BPO and 0.1549 grams I-819 in
10 15.0000 grams DVE-3 in a 20 mL amber vial, on a roll mill for 40 minutes at 21°C. The solution was transferred to a speed mixer jar. 0.1000 grams A-200 and 5.3111 grams clay were added to the solution and homogeneously dispersed by means of a high speed mixer at 2,000 rpm for one minute. Part B was prepared by dissolving 0.0347 grams DIPEA and 0.1974 grams I-819 in 19.7034 grams TPTMP in a 40 mL
15 amber vial, also on a roll mill for 40 minutes at 21°C. The solution was transferred to a speed mixer jar. 0.1000 grams A-200 and 1.8916 grams clay were added to the solution and homogeneously dispersed by means of the high speed mixer at 2,000 rpm for one minute. Part A and Part B were homogeneously dispersed through a static mixer.

20

Example 10

The procedure generally described in Example 9 for preparing homogeneous mixtures of Part A and Part B was repeated according to the formulations listed in Tables 1A and 1B.

25

Example 11

The procedure generally described in Example 1 for preparing homogeneous mixtures of Part A and Part B was repeated according to the formulations listed in Tables 1A and 1B.

30

Example 12

0.0694 grams I-819 was dissolved in a mixture of in 3.0000 grams DVE-3 and 3.9407 grams TPTMP in a 20 mL amber vial, on a roll mill for 40 minutes at 21°C

Example 13

- 5 The procedure generally described in Example 1 for preparing homogeneous mixtures of Part A and Part B was repeated according to the formulations listed in Tables 1A and 1B.

Example 14

- 10 0.0765 grams I-819 was dissolved in a mixture of in 4.0000 grams DMDO and 3.3467 grams TAC in a 20 mL amber vial, on a roll mill for 40 minutes at 21°C

Example 15

- The procedure generally described in Example 1 for preparing homogeneous mixtures
15 of Part A and Part B was repeated according to the formulations listed in Tables 1A and 1B, wherein Part B was mixed for 24 hours rather than 8 hours.

Example 16

- The procedure generally described in Example 1 for preparing homogeneous mixtures
20 of Part A and Part B was repeated according to the formulations listed in Tables 1A and 1B.

TABLE 1A

Example	Part A Composition (grams)														
	BPO	MEKP	I-819	DVE-3	MEHQ	TEMPO	TBPO	TVBT	A-200	DSW	S-31	DIPEA	DMDO	TAC	TMPTP
1	0.0139	0	0.0300	3.0000	0	0	0	0	0	0	0	0	0	0	0
2	0.0139	0	0.0300	3.0000	0.0018	0	0	0	0	0	0	0	0	0	0
3	0.0278	0	0.0300	3.0000	0	0.0009	0	0	0	0	0	0	0	0	0
4	0.0139	0	0.0300	3.0000	0	0	0	0	0	0	0	0	0	0	0
5	0.0070	0	0.0300	3.0000	0	0	0	0	0	0	0	0	0	0	0
6	0	0	0.0300	3.0000	0	0	0.0139	0	0	0	0	0	0	0	0
7	0.0171	0	0.0554	0	0	0	0	5.5359	0	0	0	0	0	0	0
8	0.0171	0	0.0554	0	0	0	0	5.5359	0	0	0	0	0	0	0
9	0.0347	0	0.1549	15.0000	0	0	0	0	0.1000	5.3111	0	0	0	0	0
10	0.0532	0	0.1116	11.5000	0	0	0	0	0.3000	0	2.9669	0	0	0	0
11	0.0139	0	0.0600	6.0000	0	0	0	0	0	0	0	0	0	0	0
12	0	0	0.0694	3.0000	0	0	0	0	0	0	0	0	0	0	0
13	0	0	0.0400	0	0	0	0	0	0	0	0	0	0	0	3.9407
14	0	0	0.0765	0	0	0	0	0	0	0	0	0.0153	4.0000	0	0
15	0.0134	0	0.0035	0	0	0	0	0	0	0	0	0	4.0000	3.6467	0
16	0	0.0230	0.0300	0.0624	0	0	0	0.3468	0	0	0	0	0	0	0
														0.1197	0

TABLE 1B

Example	Part B Composition (grams)												
	DHEPT	PMETA	I-819	TPTMP	DIPEA	DMDO	A-200	DSW	S-31	BPO	TAC	PTE-1	PTE-2
1	0.0139	0	0.0394	3.9407	0	0	0	0	0	0	0	0	0
2	0.0139	0	0.0394	3.9407	0	0	0	0	0	0	0	0	0
3	0.0278	0	0.0394	3.9407	0	0	0	0	0	0	0	0	0
4	0	0	0.0394	3.9407	0/0139	0	0	0	0	0	0	0	0
5	0	0	0.0394	3.9407	0.0070	0	0	0	0	0	0	0	0
6	0	0	0.0394	3.9407	0.0139	0	0	0	0	0	0	0	0
7	0.0171	0	0.0300	0	0	3.0000	0	0	0	0	0	0	0
8	0	0	0.0300	0	0.0171	3.0000	0	0	0	0	0	0	0
9	0	0	0.1974	19.7034	0.0347	0	0.1000	1.8916	0	0	0	0	0
10	0	0	0.1516	15.1059	0.0532	0	0.3000	0	1.0631	0	0	0	0
11	0	0	0.0788	7.8814	0	0	0	0	0	0	0	0	0
12	0	0	0	0	0	0	0	0	0	0	0	0	0
13	0	0	0.0365	0	0	0	0	0	0	0.0153	3.6467	0	0
14	0	0	0	0	0	0	0	0	0	0	0	0	0
15	0.0134	0	0.0300	0	0	0	0	0	0	0	0	3.0000	0
16	0	0.0160	0.0300	0	0	0	0	0	0	0	0	0	3.0000

Evaluations

The following molds were used for curing evaluations:

5 Glass mold. An elongate 25 cm by 1.27 cm by 0.1 cm deep silicone rubber mold over a glass base, with an opaque silicone rubber sheet covering all but 1.27 cm of one end of the mold.

10 Aluminum, frit glass and black coated wood molds. An elongate 10 cm by 1.27 cm by 0.1 cm deep silicone rubber mold over an aluminum, frit glass or black coated wood base, with an opaque silicone rubber sheet covering all but 1.27 cm of one end of the mold.

15 Teflon™ mold. A 8.4 cm by 3.2 cm by 0.2 cm deep silicone rubber mold over a Teflon™ base, with an opaque silicone rubber sheet covering all but 2 cm of one end of the mold.

20 The curable composition was applied to the mold, an opaque silicone rubber sheet was then laid over the curable composition according to the dimensions described above. The remaining exposed area of the composition was then exposed to a 88 mW 455 nm LED light source, at a distance of 1.27 cm, for between 10 – 60 seconds. The following thiol-ene curing evaluations are listed in Tables 2 and 3.

TABLE 2

Example	Mold	Cure Length (cm) After 60 second Exposure	
		Fully Cured	Partially Cured
5	Glass	4 cm	14 cm
11	Glass	3 cm	8 cm
12	Teflon	7.6 cm	N/A
	Aluminum	2.2 cm	N/A
	Black frit primed glass	1.3 cm	N/A
	Glass	8 cm	13 cm
	Black coated wood surface	4 cm	4 cm
13	Glass	4 cm	4.5 cm
	Black coated wood surface	1 cm	3 cm
14	Black coated wood surface	< 1 cm	1 cm

TABLE 3

Example	Approximate Maximum Processing Time (minutes)	Approximate Photo Cure Time (seconds)	Approximate Redox Cure Time (minutes)
1	0	N/A*	0.25
2	0	N/A*	0.25
3	3	10	10
4	30	10	180 - 240
5	60	10	180 - 300
6	120 - 240	10	720 - 960
7	0	N/A*	0.25
8	15	10	180 - 240
9	60	60	180 - 300
10	30	60	180 - 240
15	0	N/A*	5
16	30	10	720 - 840

* N/A: *The thiol-ene composition cured during the mixing step and could not be*
5 *applied to the mold.*

The Tg of photo-initiated and redox-initiated Examples 1, 7, and 15 were measured using a model “DSC Q2000” differential scanning calorimeter, obtained from TA Instruments, New Castle, Delaware. Results are listed in Table 4.

TABLE 4

Example	T _g (°C)	
	Photo initiation	Redox initiation
1	-44°C	-46°C to -41°C
7	-44°C	-51°C to -50°C
15	-55°C	-55°C

Various modifications and alterations of this disclosure will become apparent to
5 those skilled in the art without departing from the scope and principles of this
disclosure, and it should be understood that this disclosure is not to be unduly limited
to the illustrative embodiments set forth hereinabove.

We claim:

1. A composition that is curable to a polythioether polymer, comprising:

- a) a dithiol monomer;
- b) a diene monomer;
- 5 c) a radical cleaved photoinitiator;
- d) a peroxide; and
- e) an amine;

where the peroxide and amine together are a peroxide-amine redox initiator.

10 2. The composition according to claim 1 where the amine is a tertiary amine.

3. The composition according to any of the preceding claims where the amine is selected from the group consisting of dihydroxyethyl-p-toluidine, N,N-diisopropylethylamine, and N, N, N', N'', N'''-pentamethyl-diethylenetriamine.

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4. The composition according to any of the preceding claims where the peroxide is selected from the group consisting of di-tert-butyl peroxide, methyl ethyl ketone peroxide, and benzoyl peroxide.

20 5. The composition according to any of the preceding claims additionally comprising:

f) a polythiol monomer having three or more thiol groups.

6. The composition according to any of the preceding claims additionally comprising:

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g) at least one filler.

7. The composition according to any of the preceding claims additionally comprising:

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h) at least one nanoparticle filler.

8. The composition according to any of the preceding claims additionally comprising:

j) calcium carbonate.

5 9. The composition according to any of the preceding claims additionally comprising:

k) nanoparticle calcium carbonate.

10 10. The composition according to any of the preceding claims which is curable by actinic light source.

11. The composition according to any of the preceding claims which is curable by blue light source.

15 12. The composition according to any of the preceding claims which is curable by UV light source.

13. A sealant comprising the composition according to any of the preceding claims.

20 14. A polythioether polymer obtained by cure of any the composition according to any of claims 1-12.

15. The polythioether polymer according to claim 14 having a Tg less than -50°C.

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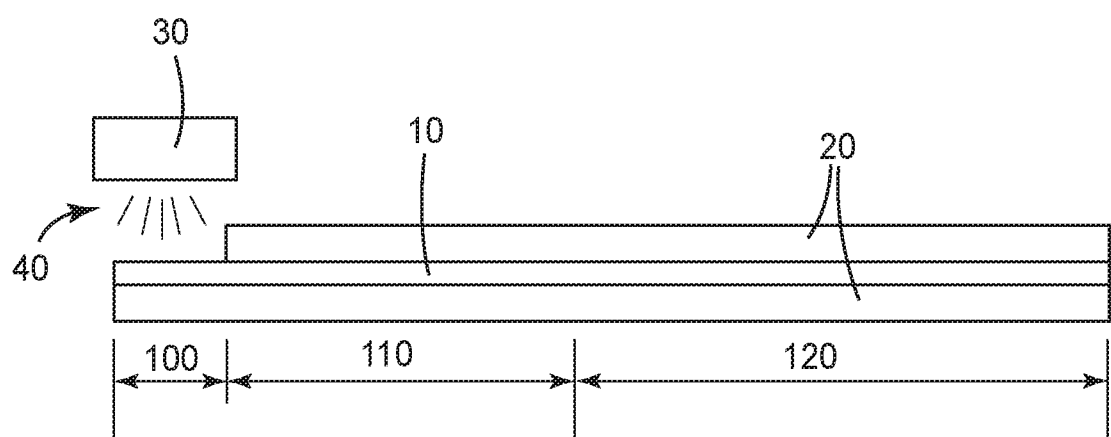


FIG. 1

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2015/067450

A. CLASSIFICATION OF SUBJECT MATTER

INV. C08G75/00 B05D3/02 B32B37/00 B64D45/02 C08F8/34
C08G18/04 C08G75/02 C09J181/02 C08L81/00 C08L81/02
F16B37/14

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

C08G B05D B32B B64D C08F C09J C08L F16B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-Internal, BEILSTEIN Data, CHEM ABS Data, PAJ, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 2014/164244 A1 (3M INNOVATIVE PROPERTIES CO [US]) 9 October 2014 (2014-10-09) page 1, line 5 - page 3, line 12; claims 1-26; examples -----	1-15
Y	WO 2014/172305 A1 (3M INNOVATIVE PROPERTIES CO [US]) 23 October 2014 (2014-10-23) page 1, line 10 - page 5, line 19; claims 1-44; examples -----	1-15
Y	US 2004/247792 A1 (SAWANT SURESH [US] ET AL) 9 December 2004 (2004-12-09) paragraph [0001] - paragraph [0106]; claims 1-81; examples ----- -/-	1-15



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

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"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

10 March 2016

Date of mailing of the international search report

17/03/2016

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INTERNATIONAL SEARCH REPORT

International application No

PCT/US2015/067450

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
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Y	US 3 923 748 A (HUTT JACK W ET AL) 2 December 1975 (1975-12-02) column 1, line 13 - column 7, line 37; claims 1-12; examples -----	1-15

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