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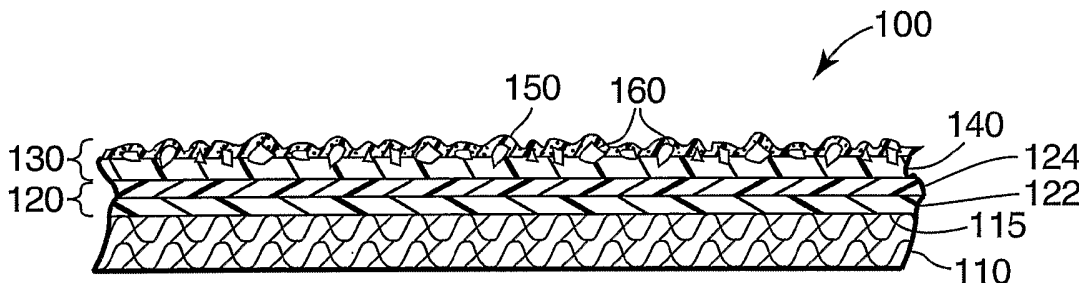
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(54) Title: COATED ABRASIVE ARTICLE WITH COMPOSITE TIE LAYER, AND METHOD OF MAKING AND USING THE SAME



(57) Abstract: Coated abrasive articles have a composite tie layer. The composite tie layer is preparable by disposing a first polymerizable composition comprising a polyfunctional aziridine on a backing, and disposing a second polymerizable composition comprising at least one acidic free-radically polymerizable monomer and at least one oligomer having at least two pendant free-radically polymerizable groups on the first polymerizable composition, at least partially polymerizing the first and second polymerizable compositions, wherein homopolymerization of the oligomer results in a polymer having a glass transition temperature of less than 50 degrees Celsius.

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**COATED ABRASIVE ARTICLE WITH COMPOSITE TIE LAYER, AND  
METHOD OF MAKING AND USING THE SAME**

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**BACKGROUND**

In general, coated abrasive articles have abrasive particles secured to a backing. More typically, coated abrasive articles comprise a backing having two major opposed surfaces and an abrasive layer secured to one of the major surfaces. The abrasive layer is typically comprised of abrasive particles and a binder, wherein the binder serves to secure the abrasive particles to the backing.

One common type of coated abrasive article has an abrasive layer which comprises a make layer, a size layer, and abrasive particles. In making such a coated abrasive article, a make layer comprising a first binder precursor is applied to a major surface of the backing. Abrasive particles are then at least partially embedded into the make layer (for example, by electrostatic coating), and the first binder precursor is cured (that is, crosslinked) to secure the particles to the make layer. A size layer comprising a second binder precursor is then applied over the make layer and abrasive particles, followed by curing of the binder precursors.

Another common type of coated abrasive article comprises an abrasive layer secured to a major surface of a backing, wherein the abrasive layer is provided by applying a slurry comprised of binder precursor and abrasive particles onto a major surface of a backing, and then curing the binder precursor.

In another aspect, coated abrasive articles may further comprise a supersize layer covering the abrasive layer. The supersize layer typically includes grinding aids and/or anti-loading materials.

Optionally, backings used in coated abrasive articles may be treated with one or more applied coatings. Examples of typical backing treatments are a backsize layer (that is, a coating on the major surface of the backing opposite the abrasive layer), a presize layer or a tie layer (that is, a coating on the backing disposed between the abrasive layer and the backing), and/or a saturant that saturates the backing. A subsize is similar to a saturant, except that it is applied to a previously treated backing.

However, depending on the particular choice of abrasive layer and backing (treated or untreated), the abrasive layer may partially separate from the backing during abrading resulting in the release of abrasive particles. This phenomenon is known in the abrasive art as "shelling". In most cases, shelling is undesirable because it results in a loss of performance.

In one approach, a tie layer disposed between the backing and the abrasive layer has been used to address the problem of shelling in some coated abrasive articles.

Yet, despite such advances, there remains a continuing need for new materials and methods that can reduce the problem of shelling in coated abrasive articles.

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### SUMMARY

In one aspect, the present invention provides a method of making a coated abrasive article comprising:

disposing a first polymerizable composition on at least a portion a backing, the first polymerizable composition comprising an isotropic composition comprising at least one polyfunctional aziridine;

disposing a second polymerizable composition comprising at least one acidic free-radically polymerizable monomer and at least one oligomer having at least two pendant free-radically polymerizable groups on at least a portion of the first polymerizable composition, wherein homopolymerization of the oligomer results in a polymer having a glass transition temperature of less than 50 degrees Celsius;

at least partially polymerizing the first and second polymerizable compositions to form a composite tie layer;

disposing a polymerizable make resin precursor on the composite tie layer;

embedding abrasive particles in the make resin precursor;

at least partially polymerizing the make resin precursor;

disposing a polymerizable size resin precursor on the at least partially polymerized make resin precursor; and

at least partially polymerizing the size resin precursor.

In another aspect, the present invention provides a method of making a coated abrasive article comprising:

disposing a first polymerizable composition on at least a portion of a backing, the first polymerizable composition comprising an isotropic composition comprising at least one polyfunctional aziridine and at least one acidic free-radically polymerizable monomer;

5 disposing a second polymerizable composition comprising at least one oligomer having at least two pendant free-radically polymerizable groups on at least a portion of the first polymerizable composition, wherein homopolymerization of the oligomer results in a polymer having a glass transition temperature of less than 50 degrees Celsius;

at least partially polymerizing the first and second polymerizable compositions to form a composite tie layer;

10 providing a tool having a surface with plurality of precisely-shaped cavities therein, and urging a slurry into at least a portion of the cavities, the slurry comprising at least one binder precursor and abrasive particles;

contacting the slurry with the composite tie layer; and

at least partially polymerizing the slurry.

15 In yet another aspect, the present invention provides a coated abrasive article comprising:

a backing having a major surface;

an inhomogeneous composite tie layer secured to at least a portion of the major surface, wherein the composite tie layer is preparable by

20 disposing a layer of first polymerizable composition on at least a portion of the major surface, the first polymerizable composition comprising an isotropic composition comprising at least one polyfunctional aziridine, and

25 disposing a second polymerizable composition comprising at least one acidic free-radically polymerizable monomer and at least one oligomer having at least two pendant free-radically polymerizable groups on at least a portion of the first polymerizable composition, wherein homopolymerization of the oligomer results in a polymer having a glass transition temperature of less than 50 degrees Celsius, and

30 at least partially polymerizing the first and second polymerizable compositions; and

an abrasive layer secured to at least a portion of the composite tie layer.

Coated abrasive articles according to the present invention are typically useful for abrading a workpiece, and may exhibit low levels of controlling shelling during abrading processes.

As used herein, the term "(meth)acryl" includes both "acryl" and "methacryl".

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#### BRIEF DESCRIPTION OF THE DRAWING

FIG. 1 is a cross-sectional view of an exemplary coated abrasive article;

FIG. 2 is a cross-sectional view of another exemplary coated abrasive article; and

FIG. 3 is a cross-sectional view of another exemplary coated abrasive article.

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#### DETAILED DESCRIPTION

Coated abrasive articles according to present invention comprise a backing having a major surface, a composite tie layer secured to at least a portion of the major surface, and an abrasive layer secured to at least a portion of the composite tie layer.

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Suitable backings include those known in the art for making coated abrasive articles. Typically, the backing has two opposed major surfaces. The thickness of the backing generally ranges from 0.02 to 5 millimeters, desirably from 0.05 to 2.5 millimeters, and more desirably from 0.1 to 0.4 millimeter, although thicknesses outside of these ranges may also be useful.

20

The backing may be flexible or rigid, and may be made of any number of various materials including those conventionally used as backings in the manufacture of coated abrasives. Examples include paper, cloth, film, polymeric foam, vulcanized fiber, woven and nonwoven materials, combinations of two or more of these materials. The backing may also be a laminate of two materials (for example, paper/film, cloth/paper, film/cloth).

25

Exemplary flexible backings include polymeric film (including primed films) such as polyolefin film (for example, polypropylene including biaxially oriented polypropylene, polyester film, polyamide film, cellulose ester film), metal foil, mesh, scrim, foam (for example, natural sponge material or polyurethane foam), cloth (for example, cloth made from fibers or yarns comprising polyester, nylon, silk, cotton, and/or rayon), paper, vulcanized paper, vulcanized fiber, nonwoven materials, and combinations thereof. Cloth backings may be woven or stitch bonded.

30

The backing may be a fibrous reinforced thermoplastic such as described, for example, as described, for example, in U.S. Pat. No. 5,417,726 (Stout et al.), or an endless spliceless belt, for example, as described, for example, in U.S. Pat. No. 5,573,619 (Benedict et al.). Likewise, the backing may be a polymeric substrate having hooking stems projecting therefrom such as that described, for example, in U.S. Pat. No. 5,505,747 (Chesley et al.). Similarly, the backing may be a loop fabric such as that described, for example, in U.S. Pat. No. 5,565,011 (Follett et al.).

Exemplary rigid backings include metal plates, and ceramic plates. Another example of a suitable rigid backing is described, for example, in U.S. Pat. No. 5,417,726 (Stout et al.).

The backing may be a treated backing having one or more treatments applied thereto such as, for example, a presize, a backsize, a subsize, and/or a saturant. Additional details regarding backing treatments can be found in, for example, U.S. Pat. Nos. 5,108,463 (Buchanan et al.); 5,137,542 (Buchanan et al.); 5,328,716 (Buchanan); and 5,560,753 (Buchanan et al.).

The composite tie layer is typically prepared by at least partially polymerizing a composite tie layer precursor. The composite tie layer precursor is typically prepared according to a two-step process.

In a first step, a first polymerizable composition is applied to at least a portion a backing. The first polymerizable composition is isotropic and comprises at least one polyfunctional aziridine. The first polymerizable composition may further comprise surfactant (for example, cationic, anionic and/or nonionic surfactant) to aid in wetting the backing. Typically, the first polymerizable composition includes water and/or organic solvent (for example, methyl ethyl ketone, glyme, propanol) to reduce the viscosity and/or solids content of the first polymerizable composition to a level that is suitable for the chosen method of application (for example, knife coating, roll coating, gravure coating, or spray coating), although this is not a requirement. If present, the water or other solvent is then typically at least partially removed (for example, by evaporation) prior to the second step, although this is not a requirement. Optionally, a period of at least 10, 20, or 30 seconds or even longer, may elapse prior to commencing the second step.

The first polymerizable composition is typically coated on the backing so as to achieve a dried add on weight in a range of from 0.1 grams/meter<sup>2</sup> (gsm) up to 10 gsm, although higher and lower dry add on weights may also be used.

5 In a second step, a second polymerizable composition is applied to at least a portion of the coated (and optionally dried) first polymerizable composition. The second polymerizable composition comprises at least one acidic free-radically polymerizable monomer and at least one oligomer having at least two pendant free-radically polymerizable groups, wherein homopolymerization of the oligomer results in a polymer having a glass transition temperature of less than 50 degrees Celsius. The second  
10 polymerizable composition may include water or other solvent and/or at least one reactive diluent to reduced the viscosity and/or solids content of the first polymerizable composition to a level that is suitable for the chosen method of application (for example, knife coating, roll coating, gravure coating, or spray coating), although this is not a requirement. The second polymerizable composition may, optionally, further comprise a  
15 curative that is capable of inducing free-radical polymerization. If present, the water or other solvent is then typically at least partially removed (for example, by evaporation) prior to the second step to form a composite tie layer precursor, although this is not a requirement. After an optional period of at least 30 seconds, the composite tie layer precursor is at least partially polymerized.

20 The second polymerizable composition is typically coated on the at least partially dried coated first polymerizable composition so as to achieve a dried add on weight in a range of from 0.1 grams/meter<sup>2</sup> (gsm) up to 400 gsm, more typically 110 gsm, although higher and lower dry add on weights may also be used.

25 Some intermixing of the polyfunctional aziridine, acidic free-radically polymerizable monomer, and oligomer having at least two pendant free-radically polymerizable groups may occur during the two-step process leading to a two layer composite tie layer, or a one-layer composite tie layer having a concentration gradient (for example of polyfunctional aziridine) across its thickness, but the process is typically not be carried out such that the polyfunctional aziridine, acidic free-radically polymerizable  
30 monomer and oligomer having at least two pendant free-radically polymerizable groups intermix to form an isotropic tie layer precursor. Thus, the term "composite tie-layer" is

intended to draw attention to the two-step nature of the composite tie layer manufacture rather than to imply that the composite tie layer includes two discreet layers.

Typically, composite tie layer weight is in a range of from 0.1 gsm up to 400 gsm, more typically, typically 110 gsm, although higher and lower weights may also be used.

5 As used herein, the term "polyfunctional aziridine" refers to a species having a plurality of aziridinyl groups. Suitable polyfunctional aziridines include, for example, those disclosed in U.S. Pat. Nos. 3,225,013 (Fram); 4,769, 617 (Canty); and 5,534,391 (Wang). Specific examples include trimethylolpropane tris[3-aziridinyl propionate]; trimethylolpropane tris[3-(2-methylaziridinyl)propionate]; trimethylolpropane tris[2-aziridinylbutyrate]; tris(1-aziridinyl)phosphine oxide; tris(2-methyl-1-aziridinyl)phosphine  
10 oxide; pentaerythritol tris[3-(1-aziridinyl)propionate]; and pentaerythritol tetrakis[3-(1-aziridinyl)propionate]. Combinations of more than one polyfunctional aziridine may also be used.

Commercially available polyfunctional aziridines include those available under the  
15 trade designations "XAMA-2" (believed to be trimethylolpropane tris[3-(2-methylaziridinyl)propanoate]) and "XAMA-7" (believed to be pentaerythritol tris(beta-(N-aziridinyl)propionate)) from EIT, Inc. Corporation, Lake Wylie, South Carolina; "HYDROFLEX XR2990" (believed to be trimethylolpropane tris[3-(2-methylaziridinyl)propanoate]) from H.B. Fuller Co., Vadnais Heights, Minnesota; and  
20 "NEOCRYL CX-100" (believed to be trimethylolpropane tris[3-(2-methylaziridinyl)propanoate]) from Zeneca Resins, Wilmington, Massachusetts.

The amount of polyfunctional aziridine incorporated into the composite tie layer precursor is generally in a range of from at least 0.1, 0.5, 1, or 2 percent by weight up to and including 4, 6, 8, or even 10 percent by weight, or more, based on the total weight of  
25 polyfunctional aziridine, acidic free-radically polymerizable monomer, and oligomer having at least two pendant free-radically polymerizable groups.

The acidic free-radically polymerizable monomer has both an acidic group and a group (for example, a (meth)acryl group) that is free-radically polymerizable. The acidic group may be, for example, carbon-, sulfur-, or phosphorus-based, and may be the free  
30 acid or in a partially or fully neutralized state. The acidic free-radically polymerizable monomer may have more than one acidic groups and/or free-radically polymerizable groups.



Useful carbon-based acidic free-radically polymerizable monomers include, for example, (meth)acrylic acid, maleic acid, monoalkyl esters of maleic acid, fumaric acid, monoalkyl esters of fumaric acid, itaconic acid, isocrotonic acid, crotonic acid, citraconic acid, and beta-carboxyethyl acrylate.

5 Useful sulfur-based acidic free-radically polymerizable monomers include, for example, 2-sulfoethyl methacrylate, styrene sulfonic acid, and 2-acrylamido-2-methylpropanesulfonic acid.

Useful phosphorus-based acidic free-radically polymerizable monomers include, for example, vinyl phosphonic acid.

10 Acidic, free-radically polymerizable monomers are commercially available, for example, under the trade designations "PHOTOMER 4173" from Cognis Corp., Cincinnati, Ohio, and "CN118", "CD9050", "CD9051" and "CD9052" all from Sartomer Co., Exton Pennsylvania.

15 The amount of acidic free-radically polymerizable monomer incorporated into the composite tie layer precursor is generally in a range of from at least 1, or 2 percent by weight up to and including 5, 10, 20, 30, or even 45 percent by weight, or more, based on the total weight of polyfunctional aziridine, acidic free-radically polymerizable monomer, and oligomer having at least two pendant free-radically polymerizable groups.

20 The oligomer having at least two pendant free-radically polymerizable groups is selected such that free-radical homopolymerization of the oligomer (for example, by photo- or thermal initiation) results in a polymer having a glass transition temperature at or below 50 degrees Celsius ( $^{\circ}\text{C}$ ). As used herein, the term "oligomer" refers to molecule composed of a small number of linked monomer units. Oligomers generally have less than one hundred monomer units and more typically less than thirty.

25 Useful oligomers having at least two pendant free-radically polymerizable groups include, for example, aliphatic and aromatic urethane (meth)acrylate oligomers, polybutadiene (meth)acrylate oligomer, acrylic (meth)acrylate oligomers, polyether (meth)acrylate oligomers, aliphatic and aromatic polyester (meth)acrylate oligomers, epoxy (meth)acrylate oligomers, and combinations thereof.

30 Methods for making such oligomers are well known in the art, and many useful free-radically polymerizable oligomers are commercially available. Examples include aliphatic and aromatic urethane (meth)acrylate oligomers such as those available from

UCB Chemicals Corp., Smyrna, Georgia, under the trade designations "EBECRYL 270", "EBECRYL 8804", "EBECRYL 8807", "EBECRYL 4827", "EBECRYL 6700", "EBECRYL 5129", or "EBECRYL 8402" and those available from Sartomer Co., Exton, Pennsylvania, under the trade designations "CN 1963", "CN 934", "CN 953B70", "CN 984", "CN 962", "CN 964", "CN 965", "CN 972", "CN 978"; polyester (meth)acrylate oligomers such as those available from UCB Chemicals Corp. under the trade designations "EBECRYL 80", "EBECRYL 81", "EBECRYL 657", "EBECRYL 810", "EBECRYL 450", "EBECRYL 870", or "EBECRYL 2870" and that available from Sartomer Co. under the trade designation "CN 292"; polyether (meth)acrylate oligomers such as those available from Sartomer Co. under the trade designations "CN 501", "CN 502", "CN 550", "CN 551"; acrylic oligomers such as those available from Sartomer Co. under the trade designations "CN 816", "CN 817", "CN 818"; epoxy (meth)acrylate oligomers such as that available from Sartomer Co. under the trade designation "CN119", and "CN121"; and polybutadiene (meth)acrylate oligomers such as that available from Sartomer Co. under the trade designation "CN 301".

The amount of oligomer incorporated into the composite tie layer precursor is generally in a range of from at least 30, 35, 40, or 45 percent by weight up to and including 50, 60, 70, 80, 90, or even 95 percent by weight, or more, based on the total weight of polyfunctional aziridine, acidic free-radically polymerizable monomer, and oligomer having at least two pendant free-radically polymerizable groups. The composite tie layer precursor may, optionally, further comprise one or more curatives that are capable of at least partially polymerizing the composite tie layer precursor. Useful curatives include free-radical initiators such as, for example, photoinitiators and/or thermal initiators for free-radical polymerization. Blends of photo-and/or thermal initiators may be used.

Useful photoinitiators include those known as useful for photocuring free-radically polyfunctional acrylates. Exemplary photoinitiators include benzoin and its derivatives such as alpha-methylbenzoin; alpha-phenylbenzoin; alpha-allylbenzoin; alpha-benzylbenzoin; benzoin ethers such as benzil dimethyl ketal (for example, as commercially available under the trade designation "IRGACURE 651" from Ciba Specialty Chemicals, Tarrytown, New York), benzoin methyl ether, benzoin ethyl ether, benzoin n-butyl ether; acetophenone and its derivatives such as 2-hydroxy-2-methyl-1-phenyl-1-propanone (for example, as commercially available under the trade designation

"DAROCUR 1173" from Ciba Specialty Chemicals) and 1-hydroxycyclohexyl phenyl ketone (for example, as commercially available under the trade designation "IRGACURE 184" from Ciba Specialty Chemicals); 2-methyl-1-[4-(methylthio)phenyl]-2-(4-morpholinyl)-1-propanone (for example, as commercially available under the trade designation "IRGACURE 907" from Ciba Specialty Chemicals); 2-benzyl-2-(dimethylamino)-1-[4-(4-morpholinyl)phenyl]-1-butanone (for example, as commercially available under the trade designation "IRGACURE 369" from Ciba Specialty Chemicals).

Other useful photoinitiators include, for example, pivaloin ethyl ether, anisoin ethyl ether, anthraquinones (for example, anthraquinone, 2-ethylanthraquinone, 1-chloroanthraquinone, 1,4-dimethylanthraquinone, 1-methoxyanthraquinone, or benzanthraquinone), halomethyltriazines, benzophenone and its derivatives, iodonium salts and sulfonium salts, titanium complexes such as bis(eta<sup>5</sup>-2,4-cyclopentadien-1-yl)-bis[2,6-difluoro-3-(1H-pyrrol-1-yl)phenyl]titanium (for example, as commercially available under the trade designation "CGI 784DC" from Ciba Specialty Chemicals); halomethylnitrobenzenes (for example, 4-bromomethylnitrobenzene), mono- and bis-acylphosphines (for example, as commercially available from Ciba Specialty Chemicals under the trade designations "IRGACURE 1700", "IRGACURE 1800", "IRGACURE 1850", and "DAROCUR 4265").

One or more spectral sensitizers (for example, dyes) may be added to the composite tie layer precursor in combination with the optional photoinitiator, for example, in order to increase sensitivity of the photoinitiator to a specific source of actinic radiation.

Examples of suitable thermal free-radical polymerization initiators include peroxides such as benzoyl peroxide, dibenzoyl peroxide, dilauryl peroxide, cyclohexane peroxide, methyl ethyl ketone peroxide; hydroperoxides such as tert-butyl hydroperoxide and cumene hydroperoxide; dicyclohexyl peroxydicarbonate; 2,2'-azobis(isobutyronitrile); and t-butyl perbenzoate. Examples of commercially available thermal free-radical polymerization initiators include initiators available from E. I. du Pont de Nemours and Co., Wilmington, Delaware, under the trade designation "VAZO" (for example, "VAZO 64" and "VAZO 52") and from Elf Atochem North America, Philadelphia, Pennsylvania, under the trade designation "LUCIDOL 70".

If present, the curative is typically used in an amount effective to facilitate polymerization, for example, in an amount in a range of from 0.01 percent by weight up to

10 percent by weight, based on the total amount of tie layer precursor, although amounts outside of these ranges may also be useful.

In addition to other components, the composite tie layer precursor of the present invention may contain optional additives, for example, to modify performance and/or appearance. Exemplary additives include, fillers, solvents, plasticizers, wetting agents, 5 surfactants, pigments, coupling agents, fragrances, fibers, lubricants, thixotropic materials, antistatic agents, suspending agents, pigments, and dyes.

Reactive diluents may also be added to the composite tie layer precursor, for example, to adjust viscosity and/or physical properties of the cured composition. 10 Examples of suitable reactive diluents include diluents mono and polyfunctional (meth)acrylate monomers (for example, ethylene glycol di(meth)acrylate, hexanediol di(meth)acrylate, triethylene glycol di(meth)acrylate, trimethylolpropane tri(meth)acrylate, tripropylene glycol di(meth)acrylate), vinyl ethers (for example, butyl vinyl ether), vinyl esters (for example, vinyl acetate), and styrenic monomers (for example, styrene).

15 The application of the tie layer precursor to the backing can be performed in a variety of ways including, for example, such techniques as brushing, spraying, roll coating, curtain coating, gravure coating, and knife coating. Organic solvent may be added to the isotropic polymerizable composition to facilitate the specific coating technique used. The coated backing may then be processed for a time at a temperature sufficient to 20 dry (if organic solvent is present) and at least partially polymerize the coating thereby securing it to the backing.

After an optional period of at least 30 seconds, the tie layer precursor is typically at least partially polymerized, for example, by any of a number of well-known techniques such as, for example, by exposure electron beam radiation, actinic radiation (that is, 25 ultraviolet and/or visible electromagnetic radiation), and thermal energy. If actinic radiation is used, at least one photoinitiator is typically present in the tie layer precursor. If thermal energy is used, at least one thermal initiator is typically present in the tie layer precursor. The polymerization may be carried out in air or in an inert atmosphere such as, for example, nitrogen or argon.

30 In one exemplary embodiment, abrasive layer comprises a make layer comprising a first binder resin, abrasive particles embedded in the make layer, and a size layer comprising a second binder resin secured to the make layer and abrasive particles.

Referring to FIG. 1, exemplary coated abrasive article 100 according to the present invention has backing 110, composite tie layer 120 according to the present invention secured to major surface 115 of backing 110, and abrasive layer 130 secured to composite tie layer 120. Composite tie layer 120 comprises first and second, optionally interdiffused, layers 122 and 123, respectively. First layer 122 comprises polyfunctional aziridine, and second layer 124 comprises an acidic free-radically polymerizable monomer and at least one oligomer having at least two pendant free-radically polymerizable groups on at least a portion of the first polymerizable composition, wherein homopolymerization of the oligomer results in a polymer having a glass transition temperature of less than 50 degrees Celsius. Abrasive layer 130, includes abrasive particles 160 secured to composite tie layer 120 by make layer 140 and size layer 150.

The make and size layers may comprise any binder resin that is suitable for use in abrading applications. Typically, the make layer is prepared by coating at least a portion of the backing (treated or untreated) with a make layer precursor. Abrasive particles are then at least partially embedded (for example, by electrostatic coating) in the make layer precursor comprising a first binder precursor, and the make layer precursor is at least partially polymerized. Next, the size layer is prepared by coating at least a portion of the make layer and abrasive particles with a size layer precursor comprising a second binder precursor (which may be the same as, or different from, the first binder precursor), and at least partially curing the size layer precursor. In one embodiment, the make layer precursor may be partially polymerized prior to coating with abrasive particles and further polymerized at a later point in the manufacturing process.

In one embodiment, a supersize may be applied to at least a portion of the size layer. Useful first and second binder precursors are well known in the abrasive art and include, for example, free-radically polymerizable monomer and/or oligomer, epoxy resins, phenolic resins, melamine-formaldehyde resins, aminoplast resins, cyanate resins, or combinations thereof.

Useful abrasive particles are well known in the abrasive art and include for example, fused aluminum oxide, heat treated aluminum oxide, white fused aluminum oxide, black silicon carbide, green silicon carbide, titanium diboride, boron carbide, tungsten carbide, titanium carbide, diamond, cubic boron nitride, garnet, fused alumina zirconia, sol gel abrasive particles, silica, iron oxide, chromia, ceria, zirconia, titania,

silicates, metal carbonates (such as calcium carbonate (for example, chalk, calcite, marl, travertine, marble and limestone), calcium magnesium carbonate, sodium carbonate, magnesium carbonate), silica (for example, quartz, glass beads, glass bubbles and glass fibers) silicates (for example, talc, clays, (montmorillonite) feldspar, mica, calcium  
5 silicate, calcium metasilicate, sodium aluminosilicate, sodium silicate) metal sulfates (for example, calcium sulfate, barium sulfate, sodium sulfate, aluminum sodium sulfate, aluminum sulfate), gypsum, aluminum trihydrate, graphite, metal oxides (for example, tin oxide, calcium oxide), aluminum oxide, titanium dioxide) and metal sulfites (for example, calcium sulfite), metal particles (for example, tin, lead, copper), plastic abrasive particles  
10 formed from a thermoplastic material (for example, polycarbonate, polyetherimide, polyester, polyethylene, polysulfone, polystyrene, acrylonitrile-butadiene-styrene block copolymer, polypropylene, acetal polymers, polyvinyl chloride, polyurethanes, nylon), plastic abrasive particles formed from crosslinked polymers (for example, phenolic resins, aminoplast resins, urethane resins, epoxy resins, melamine-formaldehyde, acrylate resins,  
15 acrylated isocyanurate resins, urea-formaldehyde resins, isocyanurate resins, acrylated urethane resins, acrylated epoxy resins), and combinations thereof.

In another exemplary embodiment of a coated abrasive article according to the present invention, the abrasive layer may comprise abrasive particles dispersed in a binder. Referring now to FIG. 2, exemplary coated abrasive article 200 has backing 210,  
20 composite tie layer 220 according to the present invention secured to major surface 215 of backing 210, and abrasive layer 230 secured to composite tie layer 220. Composite tie layer comprises first and second, optionally interdiffused, layers 222 and 223, respectively. First layer 222 comprises polyfunctional aziridine, and second layer 224  
25 comprises an acidic free-radically polymerizable monomer and at least one oligomer having at least two pendant free-radically polymerizable groups on at least a portion of the first polymerizable composition, wherein homopolymerization of the oligomer results in a polymer having a glass transition temperature of less than 50 degrees Celsius. Abrasive layer 230 includes abrasive particles 260 dispersed in binder 240.

In making such a coated abrasive article, a slurry comprising a binder precursor  
30 and abrasive particles is typically applied to a major surface of the backing, and the binder precursor is then at least partially cured. Suitable binder precursors and abrasive particles include, for example, those listed hereinabove.

In another exemplary embodiment, a coated abrasive article according to the present invention may comprise a structured abrasive article. Referring now to FIG. 3, exemplary structured abrasive article 300 has backing 310, composite tie layer 320 according to the present invention secured to major surface 315 of backing 310, and  
5 abrasive layer 330 secured to composite tie layer 315. Composite tie layer 320 comprises first and second, optionally interdiffused, layers 322 and 323, respectively. First layer 322 comprises polyfunctional aziridine, and second layer 324 comprises an acidic free-radically polymerizable monomer and at least one oligomer having at least two pendant free-radically polymerizable groups on at least a portion of the first polymerizable  
10 composition, wherein homopolymerization of the oligomer results in a polymer having a glass transition temperature of less than 50 degrees Celsius. Abrasive layer 330 includes a plurality of precisely-shaped abrasive composites 355. The abrasive composites comprise abrasive particles 360 dispersed in binder 350.

In making such a coated abrasive article, a slurry comprising a binder precursor  
15 and abrasive particles may be applied to a tool having a plurality of precisely-shaped cavities therein. The slurry is then at least partially polymerized and adhered to the composite tie layer, for example, by adhesive or addition polymerization of the slurry. Suitable binder precursors and abrasive particles include, for example, those listed hereinabove.

20 The abrasive composites may have a variety of shapes including, for example, those shapes selected from the group consisting of cubic, block-like, cylindrical, prismatic, pyramidal, truncated pyramidal, conical, truncated conical, cross-shaped, and hemispherical.

Optionally, coated abrasive articles may further comprise, for example, a backsize  
25 (that is, a coating on the major surface of the backing opposite the major surface having the abrasive coat), a presize and/or subsize (that is, a coating between the composite tie layer and the major surface to which the composite tie layer is secured), and/or a saturant which coats both major surfaces of the backing. Coated abrasive articles may further  
30 comprise a supersize covering at least a portion of the abrasive coat. If present, the supersize typically includes grinding aids and/or anti-loading materials.

Coated abrasive articles according to the present invention may be converted, for example, into belts, rolls, discs (including perforated discs), and/or sheets. For belt

applications, two free ends of the abrasive sheet may be joined together using known methods to form a spliced belt.

Further description of techniques and materials for making coated abrasive articles may be found in, for example, U.S. Pat. Nos. 4,314,827 (Leitheiser et al.); 4,518,397  
5 (Leitheiser et al.); 4,588,419 (Caul et al.); 4,623,364 (Cottringer et al.); 4,652,275 (Bloecher et al.); 4,734,104 (Broberg); 4,737,163 (Larkey); 4,744,802 (Schwabel); 4,751,138 (Tumey et al.); 4,770,671 (Monroe et al.); 4,799,939 (Bloecher et al.); 4,881,951 (Wood et al.); 4,927,431 (Buchanan et al.); 5,498,269 (Larmie); 5,011,508 (Wald et al.); 5,078,753 (Broberg et al.); 5,090,968 (Pellow); 5,108,463 (Buchanan et al.);  
10 5,137,542 (Buchanan et al.); 5,139,978 (Wood); 5,152,917 (Pieper et al.); 5,201,916 (Berg et al.); 5,203,884 (Buchanan et al.); 5,227,104 (Bauer); 5,304,223 (Pieper et al.); 5,328,716 (Buchanan); 5,366,523 (Rowenhorst et al.); 5,378,251 (Culler et al.); 5,417,726 (Stout et al.); 5,429,647 (Larmie); 5,436,063 (Follett et al.); 5,490,878 (Peterson et al.); 5,492,550 (Krishnan et al.); 5,496,386 (Broberg et al.); 5,520,711 (Helmin); 5,549,962  
15 (Holmes et al.); 5,551,963 (Larmie); 5,556,437 (Lee et al.); 5,560,753 (Buchanan et al.); 5,573,619 (Benedict et al.); 5,609,706 (Benedict et al.); 5,672,186 (Chesley et al.); 5,700,302 (Stoetzel et al.); 5,851,247 (Stoetzel et al.); 5,913,716 (Mucci et al.); 5,942,015 (Culler et al.); 5,954,844 (Law et al.); 5,961,674 (Gagliardi et al.); 5,975,988 (Christianson); 6,059,850 (Lise et al.); and 6,261,682 (Law).

20 Abrasive articles according to the present invention are useful for abrading a workpiece in a process wherein at least a portion of the abrasive layer of a coated abrasive article is frictionally contacted with the abrasive layer with at least a portion of a surface of the workpiece, and then at least one of the coated abrasive article or the workpiece is moved relative to the other to abrade at least a portion of the surface. The abrading process  
25 may be carried out, for example, by hand or by machine. Optionally, liquid (for example, water, oil) and/or surfactant (for example, soap, nonionic surfactant) may be applied to the workpiece, for example, to facilitate the abrading process.



Objects and advantages of this invention are further illustrated by the following non-limiting 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 invention.

5

### EXAMPLES

Unless otherwise noted, all parts, percentages, ratios, etc. in the examples and the rest of the specification are by weight, and all reagents used in the examples were obtained, or are available, from general chemical suppliers such as, for example, Sigma-Aldrich Co., Saint Louis, Missouri, or may be synthesized by conventional methods.

10

The following abbreviations are used throughout the Examples.

TABLE OF ABBREVIATIONS

A1	silane methacrylate commercially available from GE Silicones, Friendly, West Virginia under the trade designation "SILANE A-174NT"
A2	silicon dioxide commercially available from Degussa Corp., Parsippany, New Jersey under the trade designation "SILICONE DIOXIDE OX-50 AEROSIL"
ACR1	trimethylolpropane triacrylate, commercially available under the trade designation "TMPTA-N" from UCB Group, Springfield, Massachusetts
AFR1	modified epoxy acrylate, commercially available under the trade designation "CN118" from Sartomer Co., Exton, Pennsylvania
AFR2	monofunctional acid ester acrylate, commercially available under the trade designation "CD9050" from Sartomer Co.
AFR3	trifunctional acid ester acrylate, commercially available under the trade designation "CD9052" from Sartomer Co.
AFR4	acidic aromatic acrylate oligomer, commercially available under the trade designation "PHOTOMER 4173" from Cognis Corp., Cincinnati, Ohio
AZ1	polyfunctional aziridine commercially available under the trade designation from "HYDROFLEX XR-2990" from H.B. Fuller Co.
BK1	a treated fabric backing, prepared according to the following procedure: follows: EPR1 (11,306, grams (g)) was mixed with 1507 g of ACR1 and 151 g of PI2 at 20 °C until homogeneous using a mechanical stirrer. The mixture was

	<p>then heated at 50 °C in an oven for 2 hours. After removing the mixture from the oven, 1206 grams DICY was added and with stirring for 10 minutes. Next, 754 g of NOV1 was added and stirring continued for 10 minutes. 114 g of CUR1 was added and stirring continued until dissolved. A 30.5 cm wide coating knife obtained from the Paul N. Gardner Co., Pompano Beach, Florida, and a 30 cm x 30 cm x 2.5 cm machined stainless steel coating platform were heated to 66 °C. The knife was set to a minimum gap of 225 micrometers. A 100% polyester 4/1 sateen fabric made from open-end spun yarns weighing 326 grams/meter<sup>2</sup>, commercially available under the trade designation "POWERSTRAIGHT" from Milliken and Co., Spartanburg, South Carolina, was placed under the coating knife. The resin composition was poured onto the polyester fabric and then the fabric was pulled by hand under the knife to form a presize coat on the fabric. The pre-sized fabric was then irradiated by passing once through a UV processor obtained under the trade designation "UV PROCESSOR", obtained from Fusion UV Systems, Gaithersburg, Maryland, using a "FUSION D" bulb at 761 Watts/inch<sup>2</sup> (118 W/cm<sup>2</sup>) and 16.4 feet/minute (5 m/min), then thermally cured at 160 °C for 5 minutes. The resultant pre-size coating weight was 106 g of/meter<sup>2</sup>. A resin blend was prepared, by mixing until homogeneous at 20 °C, 55 percent by weight FL1; 43 percent by weight RPR1 and a small amount of red Fe<sub>2</sub>O<sub>3</sub> (2 percent by weight) for color. The backside of the fabric was then coated with this resin blend and cured at 90 °C for 10 minutes, then at 105 °C for 15 minutes. The resultant backsize coating weight was 111.5 grams/meter<sup>2</sup>.</p>
BR1	acrylated aliphatic urethane, commercially available under the trade designation "EBECRYL 8402" from UCB Group
BR2	acrylated polyester, obtained under the trade designation "EBECRYL 810" from UCB Group
BR3	aliphatic polyurethane, obtained under the trade designation "EBECRYL 270" from UCB Group
BR4	polyether dimethacrylate obtained under the trade designation "SR 210" from Sartomer Co.

CUR1	2-propylimidazole, commercially available under the trade designation "ACTIRON NXJ-60 LIQUID" from Synthron, Morganton, North Carolina
DICY	dicyandiamide (having an average particle size of less than 10 micrometers), commercially available under the trade designation "AMICURE CG-1400" from Air Products and Chemicals
EPR1	epoxy resin commercially available under the trade designation "EPON 828" from Resolution Performance Products, Houston, Texas
FL1	calcium carbonate filler commercially available from J.W. Huber Corp., Atlanta, Georgia, under the trade designation "HUBERCARB Q325"
LA1	hot melt adhesive, commercially available under the trade designation "JET-MELT HOT MELT ADHESIVE PG3779" from 3M Company
MN2	sol-gel abrasive grain, commercially available under the trade designation "GRADE JIS 400 3M CUBITRON 321" from 3M Company
NOV1	novolac resin, commercially available under the trade designation "RUTAPHEN 8656F" from Bakelite AG, Frielendorf, German
pbw	parts by weight
PI1	2-benzyl-2-(dimethylamino)-1-[4-(4-morpholinyl)phenyl]-1-butanone, commercially available under the trade designation "IRGACURE 369" from Ciba Specialty Chemicals, Hawthorne, New York
PI2	2,2-dimethoxy-2-phenylacetophenone, commercially available under the trade designation "IRGACURE 651" from Ciba Specialty Chemicals
RPR1	resole phenolic (a phenol-formaldehyde resin, having phenol to formaldehyde ratio of 1.5-2.1/1, catalyzed with 2.5 percent potassium hydroxide)

### 90° PEEL ADHESION TEST

5 A coated abrasive article to be tested is converted into an 8 cm wide by 25 cm long piece. One-half the length of a wooden board (17.8 cm by 7.6 cm by 0.6 cm) is coated with Laminating Adhesive 1 (LA1) applied with a hot melt glue gun (commercially available under the trade designation "POLYGUN II HOT MELT APPLICATOR" from 3M Company). The entire width of, but only the first 15 cm of the length of, the coated abrasive article is coated with laminating adhesive on the side bearing the abrasive particles. The side of the coated abrasive article bearing the abrasive particles is attached

to the side of the board containing the laminating adhesive coating in such a manner that the 10 cm of the coated abrasive article not bearing the laminating adhesive overhangs from the board. Pressure is applied such that the board and the coated abrasive article become intimately bonded. Operating at 25 °C, the abrasive article to be tested is cut  
5 along a straight line on both sides of the article such that the width of the coated abrasive article is reduced to 5.1 cm. The resulting abrasive article/board composite is mounted horizontally in a fixture attached to the upper jaw of a tensile testing machine, commercially available under the trade designation "SINTECH 6W" from MTS Systems Corp., Eden Prairie, Minnesota. Approximately 1 cm of the overhanging portion of the  
10 coated abrasive article was mounted into the lower jaw of the machine such that the distance between the jaws was 12.7 cm. The machine separated the jaws at a rate of 0.05 centimeter/second (cm/sec), with the coated abrasive article being pulled at an angle of 90° away from the wooden board so that a portion of the coated abrasive article separated from the board. The force required for such separation (that is, stripback force) is reported in  
15 kilograms/centimeter (kg/cm).

#### General Method for Preparation of Backing with Composite Tie Layer

The backing is coated with a solution of 98 g of water, 2 g of AZ1, 1 drop of nonionic surfactant (commercially available under the trade designation "Triton X-100" commercially available from Dow Chemical Co., Midland, Michigan). The solution was  
20 coated on the backing at using a handheld knife coater set at zero gap, and drawn across the backing at a rate of about 1 foot per second (0.3 m/sec). The coated backing is allowed to air dry.

Next, a second coating of a 100 percent solids mixture of free-radically  
25 polymerizable acidic monomer and oligomer is applied onto the AZ1-coated surface of the backing using a 4-inch (1.6-cm) wide hand-held coating knife, available from the Paul N. Gardner Company, Pompano Beach, Florida. The knife gap is set at 225 micrometers. The resultant tie layer precursor-coated backing is then passed once through a UV processor having the trade designation "UV PROCESSOR", obtained from Fusion UV  
30 Systems, Gaithersburg, Maryland, using a "FUSION D" bulb at 761 Watts/inch<sup>2</sup> (118 W/cm<sup>2</sup>) and 16.4 feet/minute (5 m/min), then heated at 120 °C for 10 to 20 minutes to

give a backing having a tie layer secured thereto. The nominal coating weight of the resultant tie layer is 110 grams/m<sup>2</sup>.

#### Preparation of Slurry Resin 1 (SR1)

5           A one-gallon (4-L) plastic container was charged with 1917 g of ACR1, 19 g of PI1, 1738 g of F2, 2235 of MN2, 74 g of A1 and 17 g of A2. The resin was mechanically stirred at 25 °C for 1 hour.

#### General Method for Bonding an Abrasive layer to a Tie Layer

10           Slurry 1 is coated onto the tie layer using a handheld coating knife at a coating thickness of 2-3 mils (101 micrometers) onto a tool having precisely-shaped cavities therein as described in Example 1 of U.S. Pat. Appl. No. 10/668,736 (Collins et al.), and then transferred to tie layer. The slurry is passed once through two UV processors obtained under the trade designation "UV PROCESSOR", obtained from Fusion UV Systems,  
15           Gaithersburg, Maryland, using a "FUSION D" bulb at 761 Watts/inch<sup>2</sup> (118 W/cm<sup>2</sup>) and 50 feet/minute (15 m/min), and then heated at 120 °C for 24 hours.

#### EXAMPLES 1-12

20           As indicated in Table 1, backings having composite tie layers were prepared according to the General Method for Preparation of Backing with Composite Tie Layer. An Abrasive Layer was then applied to the composite tie layer. The resultant coated abrasive articles were subjected to the 90° Peel Adhesion Test. In Table 1, the coated abrasives failed within the coated abrasive.

TABLE 1

Example	Tie Layer Precursor Components				Backing	Abrasive Binder Precursor	Laminating Adhesive	Stripback Force, kg/cm
	Oligomer / amount, pbw	Acidic monomer / amount, pbw	Curative / amount, pbw	AZ1, pbw				
1	BR1 / 90	AFR3 / 10	PI1 / 1	2	BK1	SL1	LA1	4.53
2	BR1 / 80	AFR1 / 20	PI1 / 1	2	BK1	SL1	LA1	3.08
3	BR1 / 90 + BR4 / 5	AFR4 / 5	PI2 / 1	2	BK1	SL1	LA1	4.44, 4.83
4	BR1 / 85	AFR3 / 10, AFR4 / 5	PI2 / 1	2	BK1	SL1	LA1	3.82, 4.89
5	BR1 / 85	AFR3 / 10, AFR4 / 5	PI2 / 1	5	BK1	SL1	LA1	3.78
6	BR1 / 85	AFR3 / 10, AFR4 / 5	PI2 / 1	8	BK1	SL1	LA1	3.10
7	BR2 / 90	AFR3 / 10	PI2 / 1	2	BK1	SL1	LA1	4.83
8	BR2/90	AFR3/10	PI2/1	2	BK1	SL1	LA1	5.01

9	BR2/90	AFR2/10	PI2/1	2	BK1	SL1	LA1	4.19
10	BR2/90	AFR2/10	PI2/1	2	BK1	SL1	LA1	4.24
11	BR2/95	AFR4/5	PI2/1	2	BK1	SL1	LA1	3.01
12	BR2/95	AFR4/5	PI2/1	2	BK1	SL1	LA1	3.69

Various modifications and alterations of this invention may be made by those skilled in the art without departing from the scope and spirit of this invention, and it should be understood that this invention is not to be unduly limited to the illustrative embodiments set forth herein.



What is claimed is:

1. A method of making a coated abrasive article comprising:
  - disposing a first polymerizable composition on at least a portion a backing, the first  
5 polymerizable composition comprising an isotropic composition comprising at least one  
polyfunctional aziridine;
  - disposing a second polymerizable composition comprising at least one acidic free-  
radically polymerizable monomer and at least one oligomer having at least two pendant  
free-radically polymerizable groups on at least a portion of the first polymerizable  
10 composition, wherein homopolymerization of the oligomer results in a polymer having a  
glass transition temperature of less than 50 degrees Celsius;
  - at least partially polymerizing the first and second polymerizable compositions to  
form a composite tie layer;
  - disposing a polymerizable make resin precursor on the composite tie layer;
  - 15 embedding abrasive particles in the make resin precursor;
  - at least partially polymerizing the make resin precursor;
  - disposing a polymerizable size resin precursor on the at least partially polymerized  
make resin precursor; and
  - at least partially polymerizing the size resin precursor.
- 20 2. A method according to claim 1, wherein the first and second polymerizable  
compositions are simultaneously at least partially polymerized.
3. A method according to claim 1, wherein the first and second polymerizable  
25 compositions are sequentially at least partially polymerized.
4. A method according to claim 1, wherein at least one of the first or second  
polymerizable compositions further comprises a curative.
- 30 5. A method according to claim 4, wherein the curative comprises at least one free-  
radical photoinitiator.

6. A method according to claim 4, wherein the curative comprises at least one free-radical thermal initiator.

7. A method according to claim 1, wherein based on the total weight of polyfunctional aziridine, acidic free-radically polymerizable monomer, and oligomer having at least two free-radically polymerizable groups, the amount of polyfunctional aziridine is in a range of from 0.5 to 10 percent, and wherein the amount of acidic free-radically polymerizable monomer is in a range of from 1 to 45 percent.

8. A method according to claim 1, wherein based on the total weight of polyfunctional aziridine, acidic free-radically polymerizable monomer, and oligomer having at least two free-radically polymerizable groups, the amount of polyfunctional aziridine is in a range of from 2 to 4 percent, and wherein the amount of acidic free-radically polymerizable monomer is in a range of from 2 to 20 percent.

9. A method according to claim 1, wherein the polyfunctional aziridine is selected from the group consisting of trimethylolpropane tris[3-aziridinyl propionate], trimethylolpropane tris[3(2-methyl-aziridinyl)-propionate], trimethylolpropane tris[2-aziridinyl butyrate], tris(1-aziridinyl)phosphine oxide, tris(2-methyl-1aziridinyl)phosphine oxide, pentaerythritol tris-3-(1-aziridinyl propionate), pentaerythritol tetrakis-3-(1-aziridinyl propionate), and combinations thereof.

10. A method according to claim 1, wherein the acidic free-radically polymerizable monomer is selected from the group consisting of (meth)acrylic acid, maleic acid, monoalkyl esters of maleic acid, fumaric acid, monoalkyl esters of fumaric acid, itaconic acid, isocrotonic acid, crotonic acid, citraconic acid, and beta-carboxyethyl acrylate, 2-sulfoethyl methacrylate, styrene sulfonic acid, and 2-acrylamido-2-methylpropanesulfonic acid, vinyl phosphonic acid, and combinations thereof.

11. A method according to claim 1, wherein the oligomer having at least two pendant free-radically polymerizable groups is selected from the group consisting of aliphatic and aromatic urethane (meth)acrylate oligomers, polybutadiene (meth)acrylate

oligomer, acrylic (meth)acrylate oligomers, polyether (meth)acrylate oligomers, aliphatic and aromatic polyester (meth)acrylate oligomers, epoxy (meth)acrylate oligomers, and combinations thereof.

- 5 12. A method according to claim 1, wherein at least one of the make or size resin precursor comprises at least one free-radically polymerizable monomer, free-radically polymerizable oligomer, epoxy resin, phenolic resin, melamine-formaldehyde resin, aminoplast resin, cyanate resin, or a combination thereof.
- 10 13. A method according to claim 1, wherein the backing comprises a treated backing comprising at least one treatment selected from the group consisting of a presize, a backsize, a subsize, and a saturant.
14. A method according to claim 1, further comprising disposing a supersize resin  
15 precursor on at least a portion of the at least partially polymerized size resin precursor, and at least partially polymerizing the supersize resin precursor.
15. A method of making a coated abrasive article comprising:  
disposing a first polymerizable composition on at least a portion of a backing, the  
20 first polymerizable composition comprising an isotropic composition comprising at least one polyfunctional aziridine and at least one acidic free-radically polymerizable monomer;  
disposing a second polymerizable composition comprising at least one oligomer  
having at least two pendant free-radically polymerizable groups on at least a portion of the  
first polymerizable composition, wherein homopolymerization of the oligomer results in a  
25 polymer having a glass transition temperature of less than 50 degrees Celsius;  
at least partially polymerizing the first and second polymerizable compositions to  
form a composite tie layer;  
providing a tool having a surface with plurality of precisely-shaped cavities  
therein, and urging a slurry into at least a portion of the cavities, the slurry comprising at  
30 least one binder precursor and abrasive particles;  
contacting the slurry with the composite tie layer; and  
at least partially polymerizing the slurry.

16. A method according to claim 15, wherein the first and second polymerizable compositions are simultaneously at least partially polymerized.
- 5 17. A method according to claim 15, wherein the first and second polymerizable compositions are sequentially at least partially polymerized.
18. A method according to claim 15, wherein at least one of the first or second polymerizable compositions further comprises a curative.
- 10 19. A method according to claim 18, wherein the curative comprises a free-radical photoinitiator.
20. A method according to claim 18, wherein the curative comprises at least one free-  
15 radical thermal initiator.
21. A method according to claim 15, wherein based on the total weight of polyfunctional aziridine, acidic free-radically polymerizable monomer, and oligomer having at least two free-radically polymerizable groups, the amount of acidic free-  
20 radically polymerizable monomer is in a range of from 1 to 45 percent.
22. A method according to claim 15, wherein based on the total weight of polyfunctional aziridine, acidic free-radically polymerizable monomer, and oligomer having at least two free-radically polymerizable groups, the amount of polyfunctional  
25 aziridine is in a range of from 2 to 4 percent, and wherein the amount of acidic free-radically polymerizable monomer is in a range of from 2 to 20 percent.
23. A method according to claim 15, wherein the polyfunctional aziridine is selected from the group consisting of trimethylolpropane tris[3-aziridiny] propionate],  
30 trimethylolpropane tris[3-(2-methylaziridiny]propionate], trimethylolpropane tris[2-aziridiny]butyrate], tris(1-aziridiny]phosphine oxide, tris(2-methyl-1-aziridiny]phosphine

oxide, pentaerythritol tris[3-(1-aziridinyl)propionate], and pentaerythritol tetrakis[3-(1-aziridinyl)propionate], and combinations thereof.

5 24. A method according to claim 15, wherein the acidic free-radically polymerizable monomer is selected from the group consisting of (meth)acrylic acid, maleic acid, monoalkyl esters of maleic acid, fumaric acid, monoalkyl esters of fumaric acid, itaconic acid, isocrotonic acid, crotonic acid, citraconic acid, and beta-carboxyethyl acrylate, 2-sulfoethyl methacrylate, styrene sulfonic acid, and 2-acrylamido-2-methylpropanesulfonic acid, vinyl phosphonic acid, and combinations thereof.

10

25. A method according to claim 15, wherein the oligomer having at least two pendant free-radically polymerizable groups is selected from the group consisting of aliphatic and aromatic urethane (meth)acrylate oligomers, polybutadiene (meth)acrylate oligomer, acrylic (meth)acrylate oligomers, polyether (meth)acrylate oligomers, aliphatic and aromatic polyester (meth)acrylate oligomers, epoxy (meth)acrylate oligomers, and combinations thereof.

15

26. A method according to claim 15, wherein at least one of the make or size resin precursor comprises at least one free-radically polymerizable monomer, free-radically polymerizable oligomer, epoxy resin, phenolic resin, melamine-formaldehyde resin, aminoplast resin, cyanate resin, or a combination thereof.

20

27. A method according to claim 15, wherein the backing comprises a treated backing comprising at least one treatment selected from the group consisting of a presize, a backsize, a subsize, and a saturant.

25

28. A coated abrasive article comprising:  
a backing having a major surface;  
an inhomogeneous composite tie layer secured to at least a portion of the major surface, wherein the composite tie layer is preparable by

30

disposing a layer of first polymerizable composition on at least a portion of the major surface, the first polymerizable composition comprising an isotropic composition comprising at least one polyfunctional aziridine, and

5 disposing a second polymerizable composition comprising at least one acidic free-radically polymerizable monomer and at least one oligomer having at least two pendant free-radically polymerizable groups on at least a portion of the first polymerizable composition, wherein homopolymerization of the oligomer results in a polymer having a glass transition temperature of less than 50 degrees Celsius, and

10 at least partially polymerizing the first and second polymerizable compositions; and

an abrasive layer secured to at least a portion of the composite tie layer.

29. A coated abrasive article according to claim 28, wherein the abrasive layer  
15 comprises make and size layers and abrasive particles.

30. A coated abrasive article according to claim 28, wherein the abrasive layer comprises abrasive particles dispersed in a binder.

20 31. A coated abrasive article according to claim 30, wherein the abrasive layer comprises a precisely-shaped abrasive composites.

32. A coated abrasive article according to claim 28, wherein at least one of the first or second polymerizable compositions further comprises a curative.

25 33. A coated abrasive article according to claim 32, wherein the curative comprises at least one free-radical photoinitiator.

30 34. A coated abrasive article according to claim 32, wherein the curative comprises at least one free-radical thermal initiator.

35. A coated abrasive article according to claim 28, wherein based on the total weight of polyfunctional aziridine, acidic free-radically polymerizable monomer, and oligomer having at least two free-radically polymerizable groups, the amount of polyfunctional aziridine is in a range of from 0.5 to 10 percent, and wherein the amount of acidic free-  
5 radically polymerizable monomer is in a range of from 1 to 45 percent.

36. A coated abrasive article according to claim 28, wherein based on the total weight of polyfunctional aziridine, acidic free-radically polymerizable monomer, and oligomer having at least two free-radically polymerizable groups, the amount of polyfunctional  
10 aziridine is in a range of from 2 to 4 percent, and wherein the amount of acidic free-radically polymerizable monomer is in a range of from 2 to 20 percent.

37. A coated abrasive article according to claim 28, wherein the polyfunctional aziridine is selected from the group consisting of trimethylolpropane tris[3-aziridinyl  
15 propionate], trimethylolpropane tris[3-(2-methylaziridinyl)propionate], trimethylolpropane tris[2-aziridinylbutyrate], tris(1-aziridinyl)phosphine oxide, tris(2-methyl-1-aziridinyl)phosphine oxide, pentaerythritol tris[3-(1-aziridinyl)propionate], and pentaerythritol tetrakis[3-(1-aziridinyl)propionate], and combinations thereof.

38. A coated abrasive article according to claim 28, wherein the acidic free-radically  
20 polymerizable monomer is selected from the group consisting of (meth)acrylic acid, maleic acid, monoalkyl esters of maleic acid, fumaric acid, monoalkyl esters of fumaric acid, itaconic acid, isocrotonic acid, crotonic acid, citraconic acid, and beta-carboxyethyl acrylate, 2-sulfoethyl methacrylate, styrene sulfonic acid, and 2-acrylamido-2-  
25 methylpropanesulfonic acid, vinyl phosphonic acid, and combinations thereof.

39. A coated abrasive article according to claim 28, wherein the oligomer having at least two pendant free-radically polymerizable groups is selected from the group consisting of aliphatic and aromatic urethane (meth)acrylate oligomers, polybutadiene  
30 (meth)acrylate oligomer, acrylic (meth)acrylate oligomers, polyether (meth)acrylate oligomers, aliphatic and aromatic polyester (meth)acrylate oligomers, epoxy (meth)acrylate oligomers, and combinations thereof.

40. A coated abrasive article according to claim 28, wherein at least one of the make or size resin precursor comprises at least one free-radically polymerizable monomer, free-radically polymerizable oligomer, epoxy resin, phenolic resin, melamine-formaldehyde resin, aminoplast resin, cyanate resin, or a combination thereof.

41. A coated abrasive article according to claim 28, wherein the backing comprises a treated backing comprising at least one treatment selected from the group consisting of a presize, a backsize, a subsize, and a saturant.

10

42. A method of abrading a workpiece comprising method of abrading a workpiece comprising:

frictionally contacting at least a portion of the abrasive layer of a coated abrasive article according to claim 28 with at least a portion of a surface of the workpiece; and

15 moving at least one of the coated abrasive article or the workpiece relative to the other to abrade at least a portion of the surface.

43. A method of abrading a workpiece comprising method of abrading a workpiece comprising:

20 frictionally contacting at least a portion of the abrasive layer of a coated abrasive article according to claim 29 with at least a portion of a surface of the workpiece; and

moving at least one of the coated abrasive article or the workpiece relative to the other to abrade at least a portion of the surface.

25

44. A method of abrading a workpiece comprising method of abrading a workpiece comprising:

frictionally contacting at least a portion of the abrasive layer of a coated abrasive article according to claim 30 with at least a portion of a surface of the workpiece; and

30 moving at least one of the coated abrasive article or the workpiece relative to the other to abrade at least a portion of the surface.



45. A method of abrading a workpiece comprising method of abrading a workpiece comprising:

frictionally contacting at least a portion of the abrasive layer of a coated abrasive article according to claim 31 with at least a portion of a surface of the workpiece; and

5 moving at least one of the coated abrasive article or the workpiece relative to the other to abrade at least a portion of the surface.

1/1

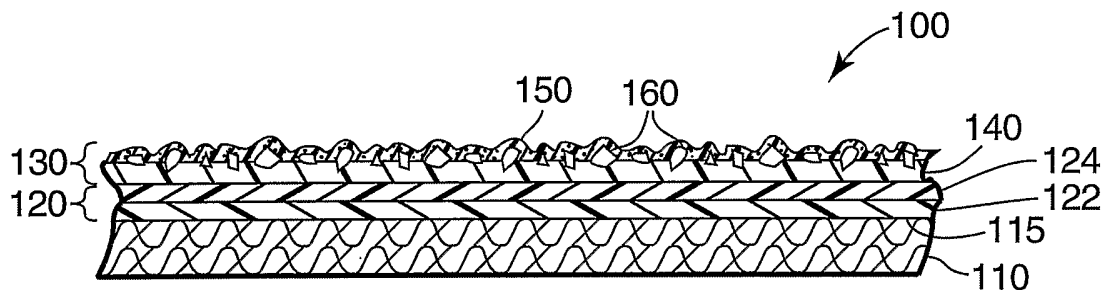


Fig. 1

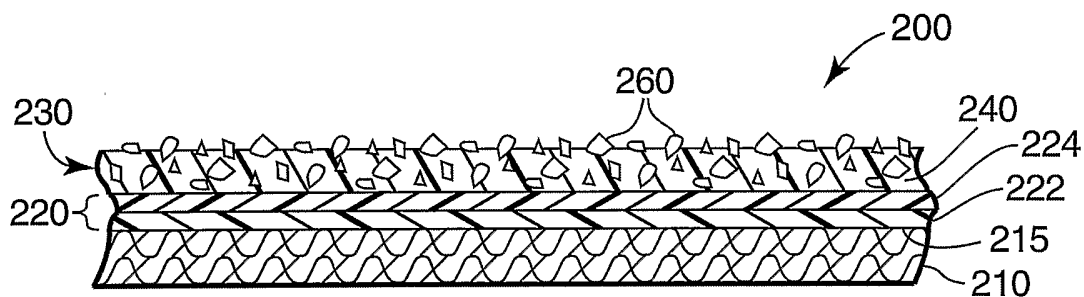


Fig. 2

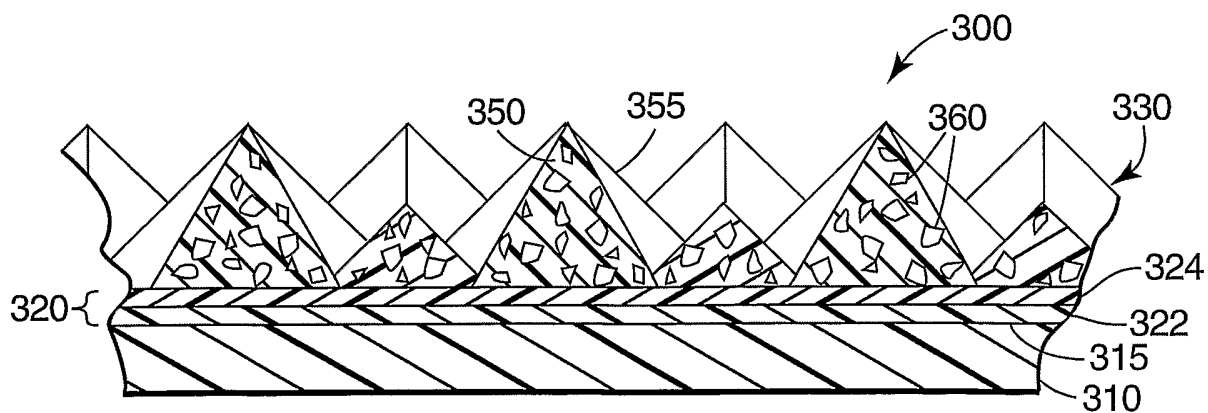


Fig. 3

**INTERNATIONAL SEARCH REPORT**

Internat	Application No
	PCT/US2005/015217

**A. CLASSIFICATION OF SUBJECT MATTER**  
 IPC 7 B24D3/00 B24D11/00

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)  
 IPC 7 B24D

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 practical, search terms used)  
 EPO-Internal

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 4 939 008 A (KEMSKI ET AL) 3 July 1990 (1990-07-03) column 4, lines 38-50; example 22	1-45
A	EP 0 590 665 A (MINNESOTA MINING AND MANUFACTURING COMPANY) 6 April 1994 (1994-04-06) example 3	1-45
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A	US 5 611 825 A (ENGEN ET AL) 18 March 1997 (1997-03-18) column 16, line 9 - column 17, line 17	
	-/--	

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

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Date of the actual completion of the international search

Date of mailing of the international search report

21 July 2005

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INTERNATIONAL SEARCH REPORT

Internat	Application No
PCT/US2005/015217	

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 2004/029511 A1 (KINCAID DON H ET AL) 12 February 2004 (2004-02-12) claims 27-29,33 -----	
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