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(54) Title: CYCLIC OLEFIN COMPOSITIONS FOR TEMPORARY WAFER BONDING

(57) Abstract: New compositions and methods of using those compositions as bonding compositions are provided. The compositions comprise a cycloolefin copolymer dispersed or dissolved in a solvent system, and can be used to bond an active wafer to a carrier wafer or substrate to assist in protecting the active wafer and its active sites during subsequent processing and handling. The compositions form bonding layers that are chemically and thermally resistant, but that can also be softened or dissolved to allow the wafers to slide or be pulled apart at the appropriate stage in the fabrication process.



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CYCLIC OLEFIN COMPOSITIONS FOR TEMPORARY WAFER BONDING

BACKGROUND OF THE INVENTION

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Government Funding

This invention was made with government support under contract number FA8650-05-D-5807 awarded by the Air Force Research Laboratory (AFRL). The United States Government has certain rights in the invention.

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Field of the Invention

The present invention is broadly concerned with novel compositions and methods of using those compositions to form bonding compositions that can support active wafers on a carrier wafer or substrate during wafer thinning and other processing.

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Description of the Prior Art

Wafer (substrate) thinning has been used to dissipate heat and aid in the electrical operation of integrated circuits (IC). Thick substrates cause an increase in capacitance, requiring thicker transmission lines, and, in turn, a larger IC footprint. Substrate thinning increases impedance while capacitance decreases impedance, causing a reduction in transmission line thickness, and, in turn, a reduction in IC size. Thus, substrate thinning facilitates IC miniaturization.

Geometrical limitations are an additional incentive for substrate thinning. Via holes are etched on the backside of a substrate to facilitate frontside contacts. In order to construct a via using common dry-etch techniques, geometric restrictions apply. For substrate thicknesses of less than 100 μm , a via having a diameter of 30-70 μm is constructed using dry-etch methods that produce minimal post-etch residue within an acceptable time. For thick substrates, vias with larger diameters are needed. This requires longer dry-etch times and produces larger quantities of post-etch residue, thus significantly reducing throughput. Larger vias also require larger quantities of metallization, which is more costly. Therefore, for backside processing, thin substrates can be processed more quickly and at lower cost.

Thin substrates are also more easily cut and scribed into ICs. Thinner substrates have a smaller amount of material to penetrate and cut and therefore require less effort. No matter what

method (sawing, scribe and break, or laser ablation) is used, ICs are easier to cut from thinner substrates. Most semiconductor wafers are thinned after frontside operations. For ease of handling, wafers are processed (i.e., frontside devices) at their normal full-size thicknesses, e.g., 600-700 μm . Once completed, they are thinned to thicknesses of 100-150 μm . In some cases (e.g., when hybrid substrates such as gallium arsenide (GaAs) are used for high-power devices) thicknesses may be taken down to 25 μm .

Mechanical substrate thinning is performed by bringing the wafer surface into contact with a hard and flat rotating horizontal platter that contains a liquid slurry. The slurry may contain abrasive media along with chemical etchants such as ammonia, fluoride, or combinations thereof. The abrasive provides "gross" substrate removal, i.e., thinning, while the etchant chemistry facilitates "polishing" at the submicron level. The wafer is maintained in contact with the media until an amount of substrate has been removed to achieve a targeted thickness.

For a wafer thickness of 300 μm or greater, the wafer is held in place with tooling that utilizes a vacuum chuck or some means of mechanical attachment. When wafer thickness is reduced to less than 300 μm , it becomes difficult or impossible to maintain control with regard to attachment and handling of the wafer during further thinning and processing. In some cases, mechanical devices may be made to attach and hold onto thinned wafers, however, they are subject to many problems, especially when processes may vary. For this reason, the wafers ("active" wafers) are mounted onto a separate rigid (carrier) substrate or wafer. This substrate becomes the holding platform for further thinning and post-thinning processing. Carrier substrates are composed of materials such as sapphire, quartz, certain glasses, and silicon, and usually exhibit a thickness of 1000 μm . Substrate choice will depend on how closely matched the coefficient of thermal expansion (CTE) is between each material. However, most of the currently available adhesion methods do not have adequate thermal or mechanical stability to withstand the high temperatures encountered in backside processing steps, such as metallization or dielectric deposition and annealing. Many current methods also have poor planarity (which contributes excessive total thickness variation across the wafer dimensions), and poor chemical resistance.

One method that has been used to mount an active wafer to a carrier substrate is via a thermal release adhesive tape. This process has two major shortcomings. First, the tapes have limited thickness uniformity across the active wafer/carrier substrate interface, and this limited

uniformity is often inadequate for ultra-thin wafer handling. Second, the thermal release adhesive softens at such low temperatures that the bonded wafer/carrier substrate stack cannot withstand many typical wafer processing steps that are carried out at higher temperatures.

Thermally stable adhesives, on the other hand, often require excessively high bonding pressures or bonding temperatures to achieve sufficient melt flow for good bond formation to occur. Likewise, too much mechanical force may be needed to separate the active wafer and carrier wafer because the adhesive viscosity remains too high at practical debonding temperatures. Thermally stable adhesives can also be difficult to remove without leaving residues.

There is a need for new compositions and methods of adhering an active wafer to a carrier substrate that can endure high processing temperatures and that allow for ready separation of the wafer and substrate at the appropriate stage of the process.

SUMMARY OF THE INVENTION

The present invention overcomes the problems of the prior art by broadly providing a wafer bonding method, which includes providing a stack comprising first and second substrates bonded together via a bonding layer, and separating the first and second substrates. The bonding layer is formed from a composition comprising a cycloolefin copolymer (COC) dissolved or dispersed in a solvent system.

The invention also provides an article comprising first and second substrates and a bonding layer. The first substrate comprises a back surface and an active surface, which comprises at least one active site and a plurality of topographical features. The second substrate has a bonding surface. The bonding layer is bonded to the active surface of the first substrate and to the bonding surface of the second substrate. The bonding layer is formed from a composition comprising a cycloolefin copolymer dissolved or dispersed in a solvent system.

In a further embodiment, the invention is concerned with a composition useful for bonding two substrates together. The inventive composition comprises a cycloolefin copolymer and an ingredient dissolved or dispersed in a solvent system. The ingredient is selected from the group consisting of tackifier resins, low molecular weight cycloolefin copolymers, and mixtures thereof.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure (Fig.) 1 illustrates the inventive method of thinning and debonding two wafers according to the present invention;

Fig. 2 is a flow diagram showing the typical process steps followed in the examples;

Fig. 3 is a graph depicting the rheological analysis results of bonding compositions according to the invention debonded at 150°C;

Fig. 4 is a graph depicting the rheological analysis results for bonding compositions according to the invention debonded at 200°C; and

Fig. 5 is a graph depicting the rheological analysis results for bonding compositions according to the invention debonded at 250°C.

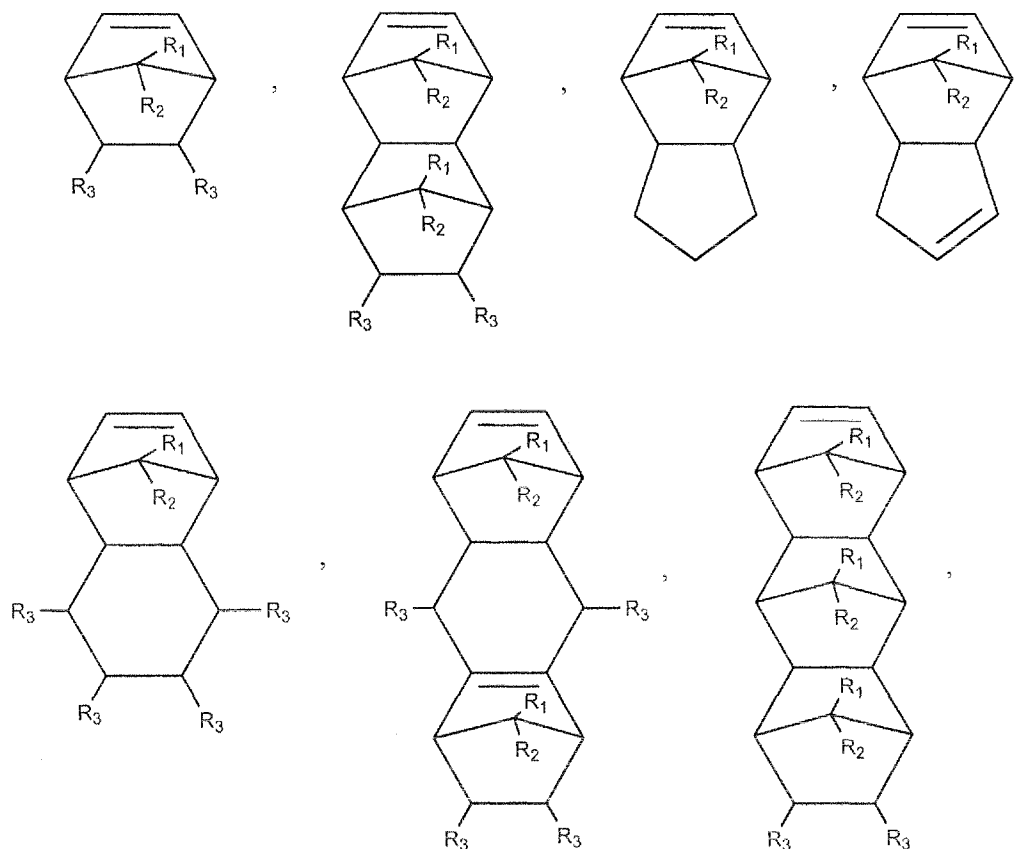
DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

In more detail, the inventive compositions comprise a cycloolefin copolymer (COC) dispersed or dissolved in a solvent system. The copolymer is preferably present in the composition at levels of from about 5% to about 85% by weight, more preferably from about 5% to about 60% by weight, and most preferably from about 10% to about 40% by weight, based upon the total weight of the composition taken as 100% by weight.

The preferred copolymers are thermoplastic and preferably have a weight average molecular weight (M_w) of from about 2,000 Daltons to about 200,000 Daltons, and more preferably from about 5,000 Daltons to about 100,000 Daltons. Preferred copolymers preferably have a softening temperature (melt viscosity at 3,000 Pa·S) of at least about 100°C, more preferably at least about 140°C, and even more preferably from about 160°C to about 220°C. Suitable copolymers also preferably have a glass transition temperature (T_g) of at least about 60°C, more preferably from about 60°C to about 200°C, and most preferably from about 75°C to about 160°C.

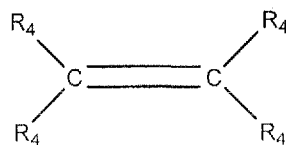
Preferred cycloolefin copolymers are comprised of recurring monomers of cyclic olefins and acyclic olefins, or ring-opening polymers based on cyclic olefins. Suitable cyclic olefins for use in the present invention are selected from the group consisting of norbornene-based olefins, tetracyclododecene-based olefins, dicyclopentadiene-based olefins, and derivatives thereof. Derivatives include alkyl (preferably C_1 - C_{20} alkyls, more preferably C_1 - C_{10} alkyls), alkylidene (preferably C_1 - C_{20} alkylidenes, more preferably C_1 - C_{10} alkylidenes), aralkyl (preferably C_6 - C_{30}

aralkyls, more preferably C₆-C₁₈ aralkyls), cycloalkyl (preferably C₃-C₃₀ cycloalkyls, more preferably C₃-C₁₈ cycloalkyls), ether, acetyl, aromatic, ester, hydroxy, alkoxy, cyano, amide, imide, and silyl-substituted derivatives. Particularly preferred cyclic olefins for use in the present invention include those selected from the group consisting of



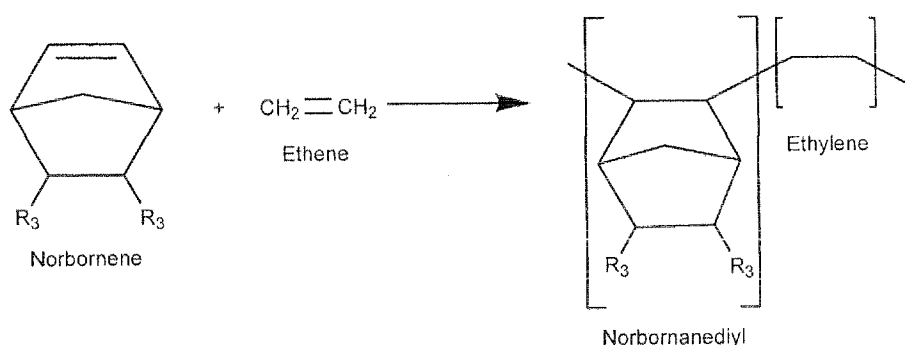
and combinations of the foregoing, where each R₁ and R₂ is individually selected from the group consisting of -H, and alkyl groups (preferably C₁-C₂₀ alkyls, more preferably C₁-C₁₀ alkyls), and each R₃ is individually selected from the group consisting of -H, substituted and unsubstituted aryl groups (preferably C₆-C₁₈ aryls), alkyl groups (preferably C₁-C₂₀ alkyls, more preferably C₁-C₁₀ alkyls), cycloalkyl groups (preferably C₃-C₃₀ cycloalkyl groups, more preferably C₃-C₁₈ cycloalkyl groups), aralkyl groups (preferably C₆-C₃₀ aralkyls, more preferably C₆-C₁₈ aralkyl groups such as benzyl, phenethyl, and phenylpropyl, etc.), ester groups, ether groups, acetyl groups, alcohols (preferably C₁-C₁₀ alcohols), aldehyde groups, ketones, nitriles, and combinations thereof.

Preferred acyclic olefins are selected from the group consisting of branched and unbranched C₂-C₂₀ alkenes (preferably C₂-C₁₀ alkenes). More preferably, suitable acyclic olefins for use in the present invention have the structure



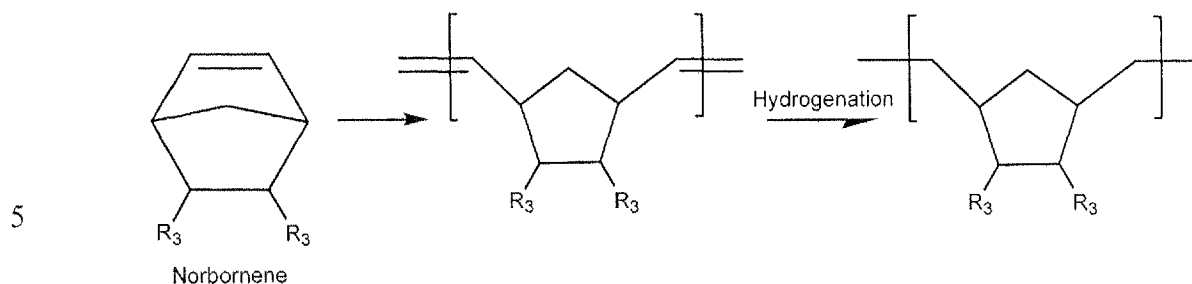
where each R₄ is individually selected from the group consisting of -H and alkyl groups (preferably C₁-C₂₀ alkyls, more preferably C₁-C₁₀ alkyls). Particularly preferred acyclic olefins for use in the present invention include those selected from the group consisting of ethene, propene, and butene, with ethene being the most preferred.

Methods of producing cycloolefin copolymers are known in the art. For example, cycloolefin copolymers can be produced by chain polymerization of a cyclic monomer with an acyclic monomer (such as norbornene with ethene as shown below).



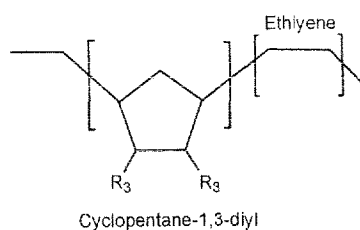
The reaction shown above results in an ethene-norbornene copolymer containing alternating norbornanediyl and ethylene units. Examples of copolymers produced by this method include TOPAS®, produced by Goodfellow Corporation and TOPAS Advanced Polymers, and APEL®, produced by Mitsui Chemicals. A suitable method for making these copolymers is disclosed in U.S. Patent No. 6,008,298, incorporated by reference herein.

Cycloolefin copolymers can also be produced by ring-opening metathesis polymerization of various cyclic monomers followed by hydrogenation as illustrated below.



The polymers resulting from this type of polymerization can be thought of conceptually as a copolymer of ethene and a cyclic olefin monomer (such as alternating units of ethylene and cyclopentane-1,3-diyl as shown below).

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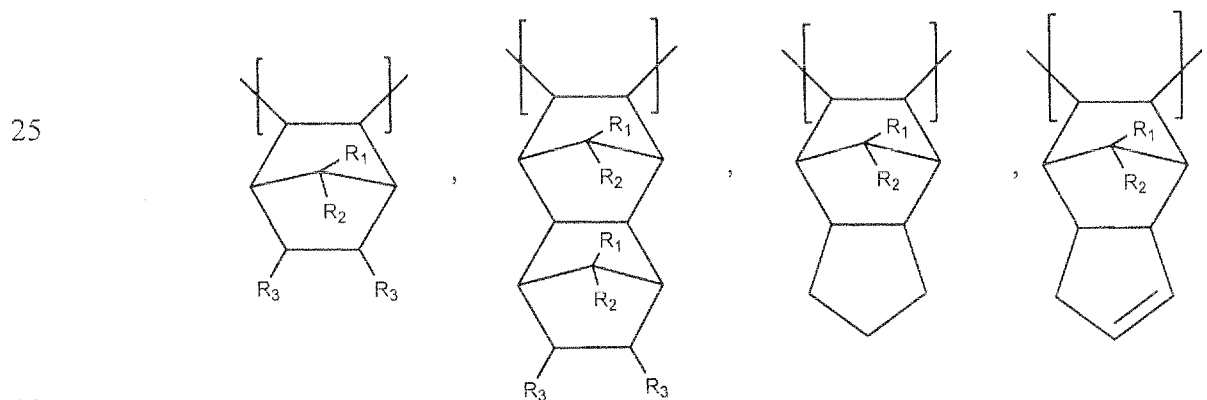


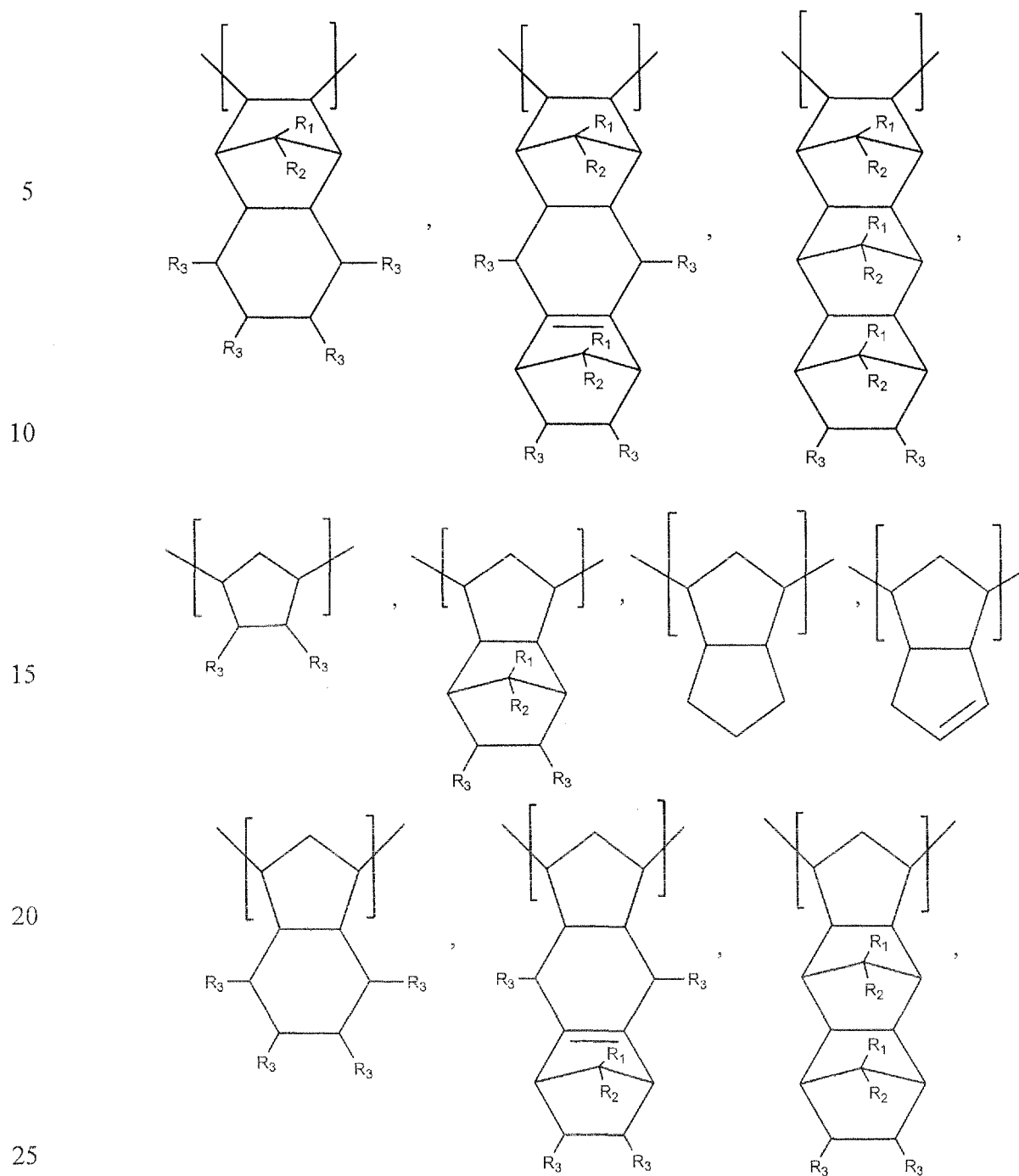
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Examples of copolymers produced by this method include ZEONOR® from Zeon Chemicals, and ARTON® from JSR Corporation. A suitable method of making these copolymers is disclosed in U.S. Patent No. 5,191,026, incorporated by reference herein.

20 Accordingly, copolymers of the present invention preferably comprise recurring monomers of:

(I):





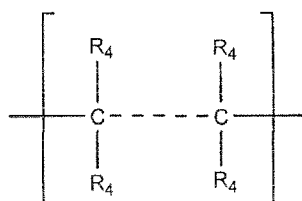
and combinations of the foregoing, where:

each R₁ and R₂ is individually selected from the group consisting of -H, and alkyl groups (preferably C₁-C₂₀ alkyls, more preferably C₁-C₁₀ alkyls), and

each R_3 is individually selected from the group consisting of -H, substituted and unsubstituted aryl groups (preferably C_6 - C_{18} aryls), alkyl groups (preferably C_1 - C_{20} alkyls, more preferably C_1 - C_{10} alkyls), cycloalkyl groups (preferably C_3 - C_{30} cycloalkyl groups, more preferably C_3 - C_{18} cycloalkyl groups), aralkyl groups (preferably C_6 - C_{30} aralkyls, more preferably C_6 - C_{18} aralkyl groups, such as benzyl, phenethyl, and phenylpropyl, etc.), ester groups, ether groups, acetyl groups, alcohols (preferably C_1 - C_{10} alcohols), aldehyde groups, ketones, nitriles, and combinations thereof;

and

(II):



where:

--- is a single or double-bond; and

each R_4 is individually selected from the group consisting of -H and alkyl groups (preferably C_1 - C_{20} alkyls, more preferably C_1 - C_{10} alkyls).

The ratio of monomer (I) to monomer (II) within the polymer is preferably from about 5:95 to about 95:5, and more preferably from about 30:70 to about 70:30.

The inventive compositions are formed by simply mixing the cycloolefin copolymer and any other ingredients with the solvent system, preferably at room temperature to about 150°C , for time periods of from about 1-72 hours.

The composition should comprise at least about 15% by weight solvent system, preferably from about 30% to about 95% by weight solvent system, more preferably from about 40% to about 90% by weight solvent system, and even more preferably from about 60% to about 90% by weight solvent system, based upon the total weight of the composition taken as 100% by weight. The solvent system should have a boiling point of from about 50 - 280°C , and preferably from about 120 - 250°C . Suitable solvents include, but are not limited to, methyl ethyl ketone (MEK) and cyclopentanone, as well as hydrocarbon solvents selected from the group consisting of limonene, mesitylene, dipentene, pinene, bicyclohexyl, cyclododecene,

1-tert-butyl-3,5-dimethylbenzene, butylcyclohexane, cyclooctane, cycloheptane, cyclohexane, methylcyclohexane, and mixtures thereof.

The total solids level in the composition should be at least about 5% by weight, preferably from about 5% to about 85% by weight, more preferably from about 5% to about 60% by weight, and even more preferably from about 10% to about 40% by weight, based upon the total weight of the composition taken as 100% by weight.

According to the invention, the composition can include additional ingredients, including low molecular weight cycloolefin copolymer (COC) resins and/or tackifier resins or rosins. The composition can also include a number of optional ingredients selected from the group consisting of plasticizers, antioxidants, and mixtures thereof.

When a low molecular weight COC resin is used in the composition, it is preferably present in the composition at a level of from about 2% to about 80% by weight, more preferably from about 5% to about 50% by weight, and even more preferably from about 15% to about 35% by weight, based upon the total weight of the composition taken as 100% by weight. The term "low molecular weight cycloolefin copolymer" is intended to refer to COCs having a weight average molecular weight (M_w) of less than about 50,000 Daltons, preferably less than about 20,000 Daltons, and more preferably from about 500 to about 10,000 Daltons. Such copolymers also preferably have a T_g of from about 50°C to about 120°C, more preferably from about 60°C to about 90°C, and most preferably from about 60°C to about 70°C. Exemplary low molecular weight COC resins for use in the present compositions are those sold under the name TOPAS® Toner TM (M_w 8,000), available from Topas Advanced Polymers.

When a tackifier or rosin is utilized, it is preferably present in the composition at a level of from about 2% to about 80% by weight, more preferably from about 5% to about 50% by weight, and even more preferably from about 15% to about 35% by weight, based upon the total weight of the composition taken as 100% by weight. The tackifiers are chosen from those having compatible chemistry with the cycloolefin copolymers so that no phase separation occurs in the compositions. Examples of suitable tackifiers include, but are not limited to, polyterpene resins (sold under the name SYLVAREST™ TR resin; Arizona Chemical), beta-polyterpene resins (sold under the name SYLVAREST™ TR-B resin; Arizona Chemical), styrenated terpene resins (sold under the name ZONATAC NG resin; Arizona Chemical), polymerized rosin resins (sold under the name SYLVAROS® PR resin; Arizona Chemical), rosin ester resins (sold under the name

EASTOTAC™ resin; Eastman Chemical), cyclo-aliphatic hydrocarbon resins (sold under the name PLASTOLYN™ resin; Eastman Chemical, or under the name ARKON™ resin; Arakawa Chemical), C5 aliphatic hydrocarbon resins (sold under the name PICCOTAC™ resin; Eastman Chemical), hydrogenated hydrocarbon resins (sold under the name REGALITE™ resin; Eastman Chemical), and mixtures thereof.

When an antioxidant is utilized, it is preferably present in the composition at a level of from about 0.1% to about 2% by weight, and more preferably from about 0.5% to about 1.5% by weight, based upon the total weight of the composition taken as 100% by weight. Examples of suitable antioxidants include those selected from the group consisting of phenolic antioxidants (such as pentaerythritol tetrakis(3-(3,5-di-tert-butyl-4-hydroxyphenyl)propionate sold under the name IRGANOX® 1010 by Ciba), phosphite antioxidants (such as tris(2,4-di-tert-butylphenyl)phosphite sold under the name IRGAFOS® 168 by Ciba), phosphonite antioxidants (such as tetrakis(2,4-di-tert-butylphenyl)[1,1-biphenyl]-4,4'-diylbisphosphonite sold under the name IRGAFOX® P-EPQ by Ciba), and mixtures thereof.

In alternative embodiments, it is preferred that the compositions are essentially free (less than about 0.1% and preferably about 0% by weight) of adhesion promoting agents, such as bis(trimethoxysilyl)ethylbenzene, aminopropyl tri(alkoxy silanes) (e.g., aminopropyl tri(methoxy silane), aminopropyl tri(ethoxy silanes), -phenyl aminopropyl tri(ethoxy silane)), and other silane coupling agents, or mixtures thereof. In some embodiments, the final composition is also thermoplastic (i.e., noncrosslinkable). Thus, in these alternative embodiments, the composition will be essentially free (less than about 0.1% by weight and preferably about 0% by weight) of crosslinking agents, such as POWDERLINK® by Cytec, and EPI-CURE™ 3200 by Hexion Specialty Chemicals.

According to one aspect, the melt viscosity (complex coefficient of viscosity) of the final composition will preferably be less than about 100 Pa·S, more preferably less than about 50 Pa·S, and even more preferably from about 1 Pa·S to about 35 Pa·S. For purposes of these measurements, the melt viscosity is determined via rheological dynamic analysis (TA Instruments, AR-2000, two parallel-plate configuration where the plates have a diameter of 25 mm). Furthermore, the melt viscosity is preferably determined at the preferred debonding temperature of the composition in question. As used herein, the term "preferred debonding temperature" of the composition is defined as the temperature at which the melt viscosity of the

composition is below 100 Pa·S, and is determined by dynamic measurement at 1 Hz oscillation frequency in temperature ramp. The compositions also preferably have a storage modulus (G') of less than about 100 Pa, preferably less than about 50 Pa, and even more preferably from about 1 Pa to about 26 Pa, when measured at the preferred debonding temperature of the composition.

- 5 The storage modulus is determined by dynamic measurement at 1 Hz oscillation frequency in temperature ramp.

The compositions are thermally stable up to about 350°C. There is also preferably less than about 5% by weight, and more preferably less than about 1.5% by weight, loss of the composition after one hour at the preferred debonding temperature plus 50°C (preferably at a
10 temperature of from about 200°C to about 300°C), depending upon the composition. In other words, very little to no thermal decomposition occurs in the composition at this temperature, as determined by thermogravimetric analysis (TGA), described herein.

Although the composition could be applied to either the carrier substrate or active wafer first, it is preferred that it be applied to the active wafer first. These compositions can be coated
15 to obtain void-free thick films required for bump wafer applications and to achieve the required uniformity across the wafer. A preferred application method involves spin-coating the composition at spin speeds of from about 500-5000 rpm (more preferably from about 1000-3500 rpm), at accelerations of from about 3000-10,000 rpm/second, and for spin times of from about 30-180 seconds. It will be appreciated that the application steps can be varied to achieve a
20 particular thickness.

After coating, the substrate can be baked (e.g., on a hot plate) to evaporate the solvents. Typical baking would be at temperatures of from about 70-250°C, and preferably from about 80-240°C for a time period of from about 1-60 minutes, and more preferably from about 2-10 minutes. The film thickness (on top of the topography) after bake will typically be at least
25 about 1 µm, and more preferably from about 10-200µm.

After baking, the desired carrier wafer is contacted with, and pressed against, the layer of inventive composition. The carrier wafer is bonded to this inventive composition by heating at a temperature of from about 100-300°C, and preferably from about 120-180°C. This heating is preferably carried out under vacuum and for a time period of from about 1-10 minutes, under
30 a bond force of from about 0.1 to about 25 kiloNewtons. The bonded wafer can be subjected to

backgrinding, metallization, patterning, passivation, via forming, and/or other processing steps involved in wafer thinning, as explained in more detail below.

Fig. 1(a) illustrates an exemplary stack 10 comprising active wafer 12 and carrier wafer or substrate 14. It will be appreciated that stack 10 is not shown to scale and has been exaggerated for the purposes of this illustration. Active wafer 12 has an active surface 18. As shown in Fig. 1(a), active surface 18 can comprise various topographical features 20a-20d. Typical active wafers 12 can include any microelectronic substrate. Examples of some possible active wafers 12 include those selected from the group consisting of microelectromechanical system (MEMS) devices, display devices, flexible substrates (e.g., cured epoxy substrates, roll-up substrates that can be used to form maps), compound semiconductors, low k dielectric layers, dielectric layers (e.g., silicon oxide, silicon nitride), ion implant layers, and substrates comprising silicon, aluminum, tungsten, tungsten silicide, gallium arsenide, germanium, tantalum, tantalum nitride, SiGe, and mixtures of the foregoing.

Carrier substrate 14 has a bonding surface 22. Typical carrier substrates 14 comprise a material selected from the group consisting of sapphire, ceramic, glass, quartz, aluminum, silver, silicon, glass-ceramic composites (such as products sold under the name Zerodur®, available from Schott AG), and combinations thereof.

Wafer 12 and carrier substrate 14 are bonded together via bonding composition layer 24. Bonding layer 24 is formed of the cycloolefin copolymer compositions described above, and has been applied and dried as also described above. As shown in the Fig. 1(a), bonding layer 24 is bonded to active surface 18 of wafer 12 as well as to bonding surface 22 of substrate 14. Unlike prior art tapes, bonding layer 24 is a uniform (chemically the same) material across its thickness. In other words, the entire bonding layer 24 is formed of the same composition.

It will be appreciated that, because bonding layer 24 can be applied to active surface 18 by spin-coating or spray-coating, the bonding composition flows into and over the various topographical features. Furthermore, the bonding layer 24 forms a uniform layer over the topography of active surface 18. To illustrate this point, Fig. 1 shows a plane designated by dashed line 26, at end portion 21 and substantially parallel to back surface 16. The distance from this plane to bonding surface 22 is represented by the thickness "T." The thickness "T" will vary by less than about 20%, preferably by less than about 10%, more preferably by less than about

5%, even more preferably by less than about 2%, and most preferably less than about 1% across the length of plane 26 and substrate 14.

The wafer package can then be subjected to subsequent thinning (or other processing) of the substrate as shown in Fig. 1(b), where 12' presents the wafer 12 after thinning. It will be appreciated that the substrates can be thinned to thicknesses of less than about 100 μm , preferably less than about 50 μm , and more preferably less than about 25 μm . After thinning, typical backside processing, including backgrinding, patterning (e.g., photolithography, via etching), passivation, and metallization, and combinations thereof, may be performed.

Advantageously, the dried layers of the inventive compositions possess a number of highly desirable properties. For example, the layers will exhibit low outgassing for vacuum etch processes. That is, if a 15- μm thick film of the composition is baked at 80-250°C for 2-60 minutes (more preferably 2-4 minutes), the solvents will be driven from the composition so that subsequent baking at 140-300°C for 2-4 minutes results in a film thickness change of less than about 5%, preferably less than about 2%, and even more preferably less than about 1.0% or even 0% (referred to as the "Film Shrinkage Test"). Thus, the dried layers can be heated to temperatures of up to about 350°C, preferably up to about 320°C, more preferably up to about 300°C, without chemical reactions occurring in the layer. In some embodiments, the layers can also be exposed to polar solvents (e.g., *N*-methyl-2-pyrrolidone) at a temperature of 80°C for 15 minutes without reacting.

The bond integrity of the dried layers can be maintained even upon exposure to an acid or base. That is, a dried layer of the composition having a thickness of about 15 μm can be submerged in an acidic media (e.g., concentrated sulfuric acid) or base (e.g., 30 wt. % KOH) at 85°C for about 45 minutes while maintaining bond integrity. Bond integrity can be evaluated by using a glass carrier substrate and visually observing the bonding composition layer through the glass carrier substrate to check for bubbles, voids, etc. Also, bond integrity is maintained if the active wafer and carrier substrate cannot be separated by hand.

After the desired processing has occurred, the active wafer or substrate can be separated from the carrier substrate. In one embodiment, the active wafer and substrate are separated by heating to a temperature sufficient to soften the bonding layer. More specifically, the stack is heated to temperatures of at least about 100°C, preferably at least about 120°C, and more preferably from about 150°C to about 300°C. These temperature ranges represent the preferred

debonding temperatures of the bonding composition layer. This heating will cause the bonding composition layer to soften and form softened bonding composition layer 24' as shown in Fig. 1(c), at which point the two substrates can be separated, for example, by sliding apart. Fig. 1(c) shows an axis 28, which passes through both of wafer 12 and substrate 14, and the sliding forces would be applied in a direction generally transverse to axis 28. Instead of sliding, wafer 12 or substrate 14 can be separated by lifting upward (i.e., in a direction that is generally away from the other of wafer 12 or substrate 14) to separate the wafer 12 from the substrate 14.

Alternatively, instead of heating to soften the layer, the bonding composition can be dissolved using a solvent. Once the layer is dissolved, the active wafer and substrate can be thereafter separated. Suitable solvents for use in dissolving the bonding layer can be any solvent that was part of the composition prior to drying, such as those selected from the group consisting of MEK and cyclopentanone, as well as hydrocarbon solvents selected from the group consisting of limonene, mesitylene, dipentene, pinene, bicyclohexyl, cyclododecene, 1-tert-butyl-3,5-dimethylbenzene, butylcyclohexane, cyclooctane, cycloheptane, cyclohexane, methylcyclohexane, and mixtures thereof.

Whether the bonding composition is softened or dissolved, it will be appreciated that separation can be accomplished by simply applying force to slide and/or lift one of wafer 12 or substrate 14 while maintaining the other in a substantially stationary position so as to resist the sliding or lifting force (e.g., by applying simultaneous opposing sliding or lifting forces to wafer 12 and substrate 14). This can all be accomplished via conventional equipment.

Any bonding composition remaining in the device areas can be easily removed by rinsing with a suitable solvent followed by spin-drying. Suitable solvents include the original solvent that was part of the composition prior to drying as well as those solvents listed above suitable for dissolving the composition during debonding. Any composition remaining behind will be completely dissolved (at least about 98%, preferably at least about 99%, and more preferably about 100%) after 5-15 minutes of exposure to the solvent. It is also acceptable to remove any remaining bonding composition using a plasma etch, either alone or in combination with a solvent removal process. After this step, a clean, bonding composition-free wafer 12' and carrier substrate 14 (not shown in their clean state) will remain.

EXAMPLES

The following examples set forth preferred methods in accordance with the invention. It is to be understood, however, that these examples are provided by way of illustration and nothing therein should be taken as a limitation upon the overall scope of the invention.

5

EXAMPLE 1

Cycloolefin Copolymer Resin and
Low Molecular Weight COC Resin Blends

In this Example, formulations containing cycloolefin copolymers and a low molecular weight COC resin were made. Antioxidants were added to some of the formulations.

10

1. *Sample 1.1*

In this procedure, 1.2 grams of an ethene-norbornene copolymer (TOPAS® 5010, T_g 110°C; obtained from TOPAS Advanced Polymers, Florence, KY) were dissolved in 6 grams of D-limonene (Florida Chemical Co.), along with 2.8 grams of a low molecular weight cycloolefin copolymer (TOPAS® Toner TM, M_w 8,000, M_w/M_n 2.0). The solution was allowed to stir at room temperature until the ingredients were in solution. The solution had about 40% solids.

15

2. *Sample 1.2*

In this procedure, 0.75 grams of an ethene-norbornene copolymer (TOPAS® 8007, T_g 78°C) and 3.25 grams of low molecular weight COC (TOPAS® Toner TM) were dissolved in 6 grams of D-limonene. The solution was allowed to stir at room temperature until the ingredients were in solution. The solution had about 40% solids.

20

3. *Sample 1.3*

For this procedure, 1.519 grams of an ethene-norbornene copolymer (TOPAS® 5013, T_g 134°C) were dissolved in 5.92 grams of D-limonene along with 2.481 grams of a low molecular weight cycloolefin copolymer (TOPAS® Toner TM), 0.04 grams of a phenolic antioxidant (IRGANOX® 1010), and 0.04 grams of a phosphonite antioxidant (IRGAFOX® P-

25

30

EPQ). The solution was allowed to stir at room temperature until the ingredients were in solution. The solution had about 40% solids.

4. *Sample 1.4*

5 In this procedure, 1.2 grams of an ethene-norbornene copolymer (TOPAS® 8007) were dissolved in 5.92 grams of D-limonene along with 2.8 grams of a low molecular weight cycloolefin copolymer (TOPAS® Toner TM), 0.04 grams of a phenolic antioxidant (IRGANOX® 1010), and 0.04 grams of a phosphonite antioxidant (IRGAFOX® P-EPQ). The solution was allowed to stir at room temperature until the ingredients were in solution. The
10 solution had about 40% solids.

5. *Sample 1.5*

For this procedure, 2.365 grams of an ethene-norbornene copolymer (TOPAS® 5013) were dissolved in 5.92 grams of D-limonene along with 1.635 grams of a low molecular weight
15 cycloolefin copolymer (TOPAS® Toner TM), 0.04 grams of a phenolic antioxidant (IRGANOX® 1010), and 0.04 grams of a phosphonite antioxidant (IRGAFOX® P-EPQ). The solution was allowed to stir at room temperature until the ingredients were in solution. The solution had about 40% solids.

20 6. *Sample 1.6*

In this procedure, 2.2 grams of a hydrogenated norbornene-based copolymer prepared by ring-opening polymerization (ZEONOR® 1060, T_g 100°C; obtained from Zeon Chemicals, Louisville, KY) and 1.8 grams of a low molecular weight cycloolefin copolymer (TOPAS® Toner TM) were dissolved in 5.92 grams of cyclooctane (Aldrich, Milwaukee, WI). The solution
25 was allowed to stir at room temperature until the ingredients were in solution. The solution had about 40% solids.

EXAMPLE 2

Cycloolefin Copolymer Resins and Tackifier Blends

In this Example, formulations were made containing cycloolefin copolymers blended with various tackifiers. As in Example 1, antioxidants were added to some of the formulations.

5

1. *Sample 2.1*

In this procedure, 0.83 grams of an ethene-norbornene copolymer (TOPAS® 8007) were dissolved in 5.92 grams of D-limonene, along with 3.17 grams of a hydrogenated hydrocarbon resin (REGALITE® R1125; obtained from Eastman Chemical Co., Kingsport TN), 0.04 grams
10 of a phenolic antioxidant (IRGANOX® 1010), and 0.04 grams of a phosphonite antioxidant (IRGAFOX® P-EPQ). The solution was allowed to stir at room temperature until the ingredients were in solution. The solution had about 40% solids.

2. *Sample 2.2*

For this procedure, 0.7 grams of an ethene-norbornene copolymer (TOPAS® 8007) and
15 3.3 grams of a styrenated terpene resin (ZONATAC® NG98; obtained from Arizona Chemical, Jacksonville, FL) were dissolved in 5.92 grams of D-limonene, along with 0.04 grams of a phenolic antioxidant (IRGANOX® 1010), and 0.04 grams of a phosphonite antioxidant (IRGAFOX® P-EPQ). The solution was allowed to stir at room temperature until the ingredients
20 were in solution. The solution had about 40% solids.

3. *Sample 2.3*

In this formulation, 1.9 grams of an ethene-norbornene copolymer (TOPAS® 5013) were dissolved in 5.92 grams of D-limonene, along with 2.1 grams of a cyclo-aliphatic hydrocarbon
25 resin (ARKON® P-140; obtained from Arakawa Chemical USA Inc., Chicago, IL), 0.04 grams of a phenolic antioxidant (IRGANOX® 1010), and 0.04 grams of a phosphonite antioxidant (IRGAFOX® P-EPQ). The solution was allowed to stir at room temperature until the ingredients were in solution.

4. *Sample 2.4*

For this procedure, 2.42 grams of an ethene-norbornene copolymer (TOPAS® 5013) were dissolved in 5.92 grams of D-limonene, along with 1.58 grams of a cyclo-aliphatic hydrocarbon resin (PLASTOLYN® R-1140; obtained from Arakawa Chemical USA Inc., Chicago, IL), 0.04 grams of a phenolic antioxidant (IRGANOX® 1010), and 0.04 grams of a phosphonite antioxidant (IRGAFOX® P-EPQ). The solution was allowed to stir at room temperature until the ingredients were in solution. The solution had about 40% solids.

EXAMPLE 3

Application, Bonding and Debonding, and Analysis

The formulations prepared in Examples 1 and 2 above were spin-coated onto various substrate wafers. After baking to evaporate the solvent and allowing the bonding composition to reflow, a second wafer was bonded to each coated wafer by applying pressure. A typical procedure for temporary wafer bonding using the bonding compositions is illustrated in Fig. 2. The bonded wafers were tested for mechanical strength, thermal stability, and chemical resistance. The wafers were tested for debonding by manually sliding them apart at acceptable temperatures. After debonding, the bonding composition residue was cleaned using a solvent rinse and spinning.

The rheological properties of each formulation from Examples 1 and 2 were tested. All of these materials were successfully tested for debonding. It was determined that the preferred debonding temperature for samples 1.1, 1.2, 2.1, and 2.2 was 150°C. The preferred debonding temperature for samples 1.3, 1.4, and 2.3 was 200°C, and the preferred debonding temperature for samples 1.5, 1.6, and 2.4 was 250°C. The storage modulus (G') and melt viscosity (η^* , complex coefficient of viscosity) for each sample at their preferred debonding temperatures are reported below. The rheological data is also illustrated in Figs. 3-5 for each debonding temperature.

Table 1

Sample number	Storage Modulus, G' (Pa)	Viscosity, η^* (Pa·s)	Debonding Temperature (°C)
1.1	25.5	35.0	150
1.2	9.6	16.7	150

1.3	8.0	13.9	200
1.4	3.5	5.1	200
1.5	16.3	20.1	250
1.6	8.1	15.1	250
2.1	24.9	14.5	150
2.2	1.3	2.4	150
2.3	21.7	28.7	200
2.4	5.5	14.1	250

Further studies on thermal stability and chemical resistance were also carried out on these compositions. Thermogravimetric analysis (TGA) was carried out on a TA Instruments thermogravimetric analyzer. The TGA samples were obtained by scraping off the spin-coated and baked bonding composition samples from Examples 1 and 2. For the isothermal TGA measurement, the samples were heated in nitrogen at a rate of 10°C/min., up to their preferred debonding temperature plus 50°C, and kept constant at that temperature for 1 hour to determine the thermal stability of the particular bonding composition. The isothermal measurements for each sample formulation are reported below in Table 2. For the scanning TGA measurement, the samples were heated in nitrogen at a rate of 10°C/min. from room temperature to 650°C.

Table 2 - Isothermal thermogravimetric results - thermal stability (in N₂)

Sample	Weight Loss (%) (Isothermal for 1 hour)	Isothermal temperature (°C)
1.1	0.123	200
1.2	0.847	200
1.3	1.268	250
1.4	0.764	250
1.5	0.752	300
1.6	0.596	300
2.1	5.496	200
2.2	4.650	200
2.3	5.737	250
2.4	5.191	300

As can be seen from the Table above, all of the COC-low molecular weight COC resin blends (Example 1) possessed the required thermal stability at least up to 300°C and exhibited minimal weight loss (< 1.5-wt%). The COC-tackifier blends (Example 2) had an average weight loss of about 5-wt% when maintained at the testing temperature. However, as shown in Table 3, below, the 1-wt% weight loss temperatures were higher than their respective bonding/debonding temperatures, suggesting sufficient thermal resistance for wafer-bonding applications.

Table 3 - Scanning thermogravimetric results

Sample	Temperature at 1.0% weight loss (°C)	Debonding Temperature (°C)
2.1	214	150
2.2	223	150
2.3	228	200
2.4	252	250

To determine chemical resistance, two silicon wafers were bonded using the particular bonding composition to be tested. The bonded wafers were put into chemical baths of *N*-Methyl-2-Pyrrolidone (NMP) or 30% by weight KOH at 85°C, and concentrated sulfuric acid at room temperature to determine chemical resistance. The bond integrity was visually observed after 45 minutes, and the stability of the bonding composition against the respective chemical was determined. All bonding compositions retained the bond integrity.

We Claim:

1. A wafer bonding method comprising:

providing a stack comprising first and second substrates bonded together via a bonding
5 layer, said layer being formed from a composition comprising a cycloolefin
copolymer dissolved or dispersed in a solvent system; and
separating said first and second substrates.

2. The method of claim 1, further comprising subjecting said stack to a temperature
10 at least about 100°C to soften said bonding layer prior to said separating.

3. The method of claim 1, further comprising dissolving said bonding layer with a
solvent prior to said separating.

15 4. The method of claim 3, said solvent being selected from the group consisting of
limonene, mesitylene, dipentene, pinene, bicyclohexyl, cyclododecene,
1-tert-butyl-3,5-dimethylbenzene, butylcyclohexane, cyclooctane, cycloheptane, cyclohexane,
methylcyclohexane, methyl ethyl ketone, cyclopentanone, and mixtures thereof.

20 5. The method of claim 1, wherein said separating comprises applying a force to at
least one of said first and second substrates while causing the other of said first and second
substrates to resist said force, said force being applied in a sufficient amount so as to separate
said first and second substrates.

25 6. The method of claim 1, further comprising subjecting said stack to a process
selected from the group consisting of backgrinding, metallizing, patterning, passivation, and
combinations thereof, prior to said separating.

30 7. The method of claim 1, wherein said providing a stack comprises:
applying a bonding composition to at least one of the first and second substrates; and
contacting the substrates with one another so as to bond the substrates together.

8. The method of claim 1, wherein:

said first substrate has a first surface and a second surface remote from said first surface and comprising at least one active site and plurality of topographical features, and said bonding composition layer is bonded to said second surface; and

5 said second substrate comprises a bonding surface that is bonded to said bonding composition layer.

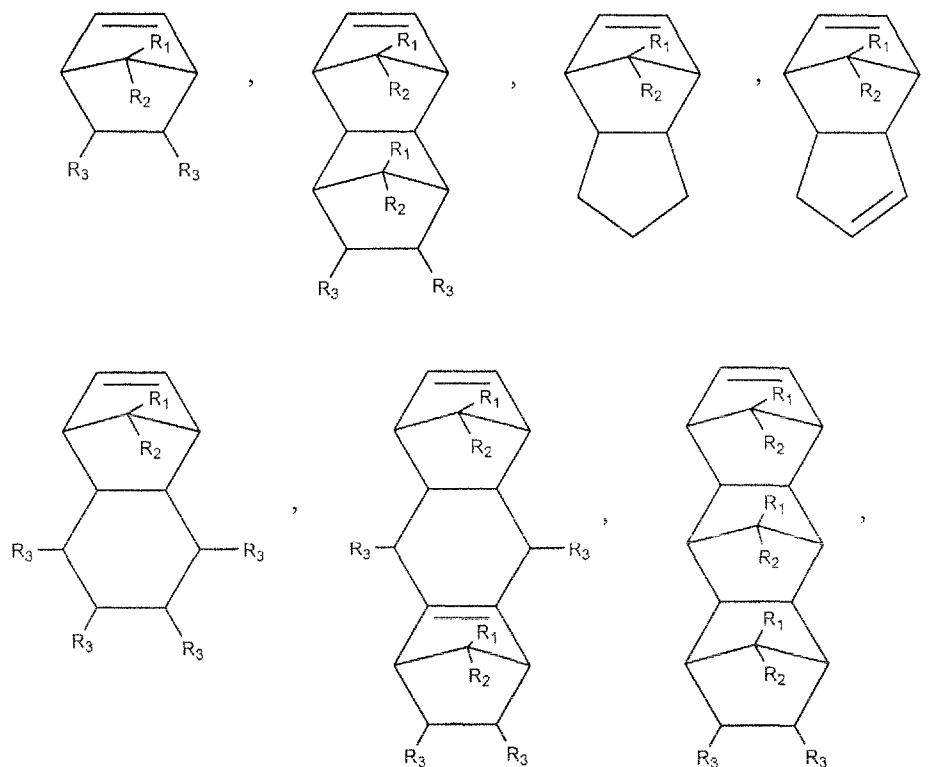
9. The method of claim 8, wherein:

10 said topographical features present respective end surfaces remote from the first surface of said first substrate, and at least one of the end surfaces is further from the first surface of the first substrate than the other of said end surfaces, said further end surface defining a plane that is substantially parallel to said first surface; and the distance from said plane to the bonding surface on said second substrate varying by less than about 20% across said plane and second substrate bonding surface.

15

10. The method of claim 1, said composition further comprising an ingredient selected from the group consisting of tackifiers, low molecular weight cycloolefin copolymers, plasticizers, antioxidants, and mixtures thereof.

11. The method of claim 1, wherein said copolymer is formed from the polymerization of a cyclic olefin selected from the group consisting of



and combinations of the foregoing,

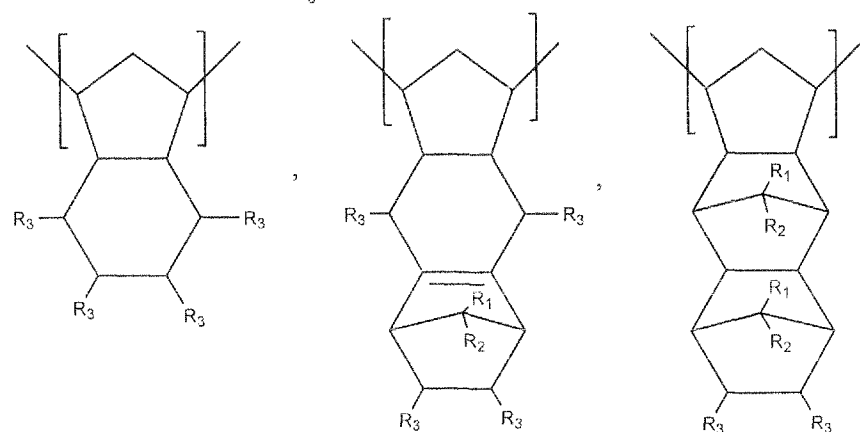
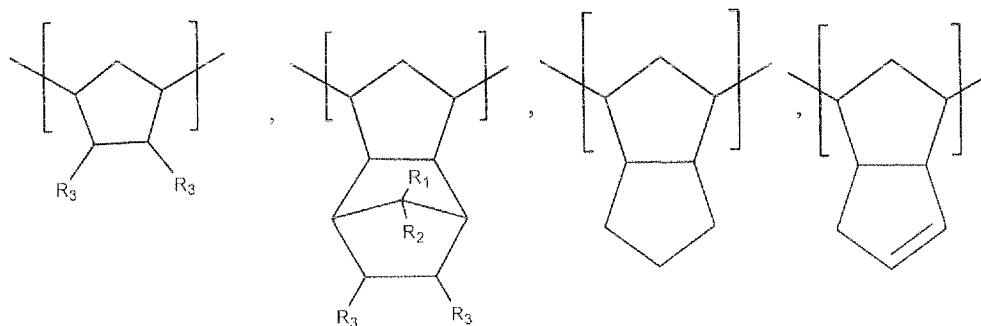
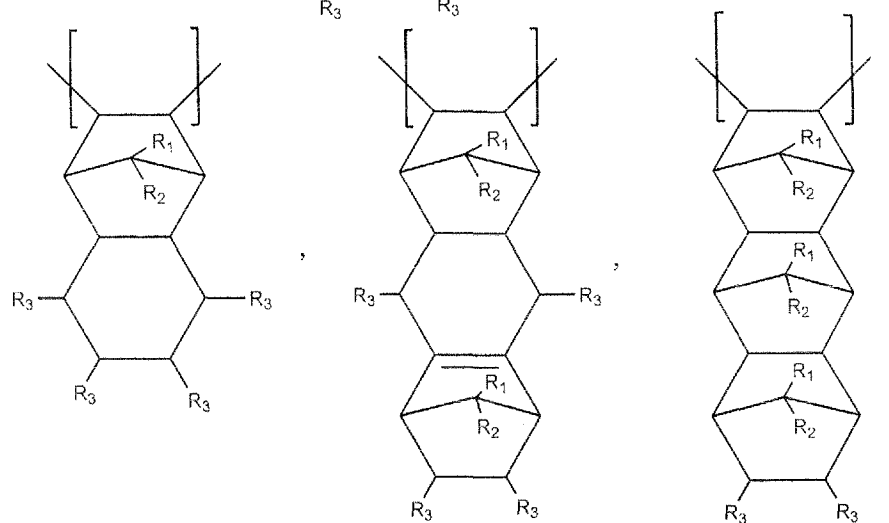
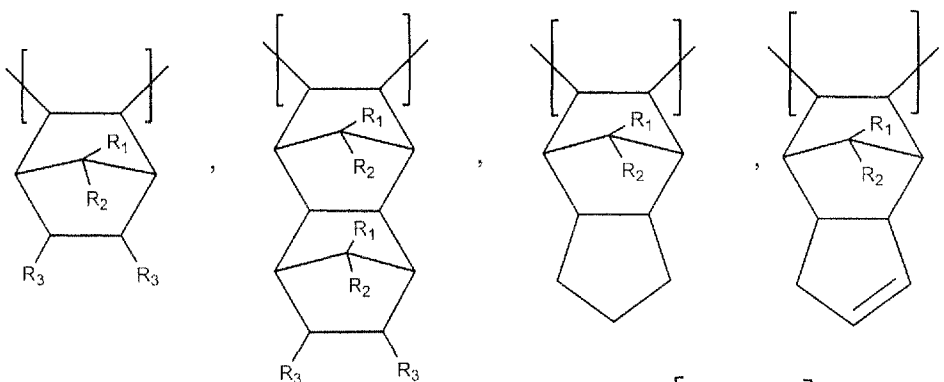
where:

each R_1 and R_2 is individually selected from the group consisting of -H, and alkyl groups; and

each R_3 is individually selected from the group consisting of -H, substituted and unsubstituted aryl groups, alkyl groups, cycloalkyl groups, aralkyl groups, ester groups, ether groups, acetyl groups, alcohols, aldehyde groups, ketones, nitriles, and combinations thereof.

12. The method of claim 1, wherein said copolymer comprises recurring monomers of:

(I):



and combinations of the foregoing,

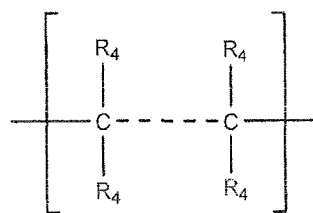
where:

each R_1 and R_2 is individually selected from the group consisting of -H, and alkyl groups; and

each R_3 is individually selected from the group consisting of -H, substituted and unsubstituted aryl groups, alkyl groups, cycloalkyl groups, aralkyl groups, ester groups, ether groups, acetyl groups, alcohols, aldehyde groups, ketones, nitriles, and combinations thereof;

and

(II):



where:

--- is a single or double-bond; and

each R_4 is individually selected from the group consisting of -H and alkyl groups.

13. An article comprising:

a first substrate having a back surface and an active surface, said active surface comprising at least one active site and a plurality of topography features;

a second substrate having a bonding surface; and

a bonding layer bonded to said active surface and to said bonding surface, wherein said bonding layer is formed from a composition comprising a cycloolefin copolymer dissolved or dispersed in a solvent system.

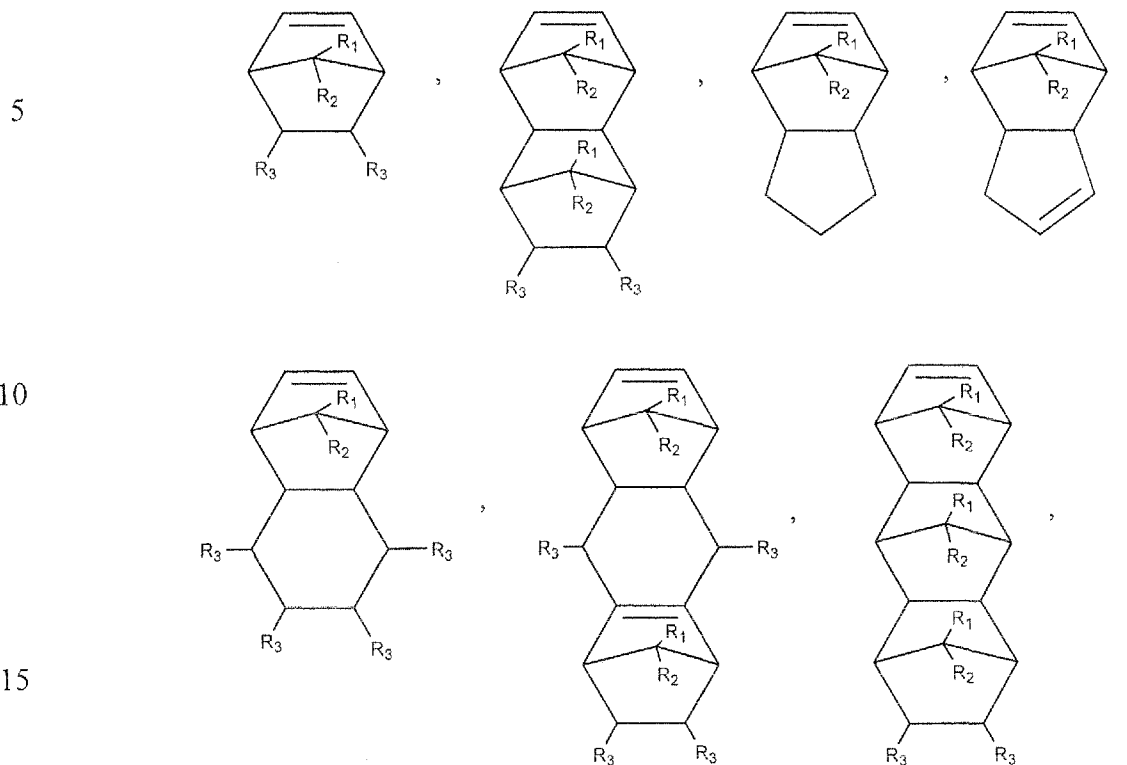
14. The article of claim 13, wherein:

said topographical features present respective end surfaces remote from the back surface of said first substrate, and at least one of the end surfaces is further from the back surface of the first substrate than the other of said end surfaces, said further end surface defining a plane that is substantially parallel to said first surface; and the distance from said plane to the bonding surface on said second substrate varying by less than about 20% along said plane and second substrate bonding surface.

15. The article of claim 13, wherein said first substrate is selected from the group consisting of microelectromechanical system devices, display devices, flexible substrates, compound semiconductors, low k dielectric layers, dielectric layers, ion implant layers, and substrates comprising silicon, aluminum, tungsten, tungsten silicide, gallium arsenide, germanium, tantalum, tantalum nitride, SiGe, and mixtures of the foregoing.

16. The article of claim 13, wherein said second substrate comprises a material selected from the group consisting of sapphire, ceramic, glass, quartz, aluminum, silver, silicon, glass-ceramic composites, and combinations thereof.

17. The article of claim 13, wherein said copolymer is formed from the polymerization of a cyclic olefin selected from the group consisting of



and combinations of the foregoing,

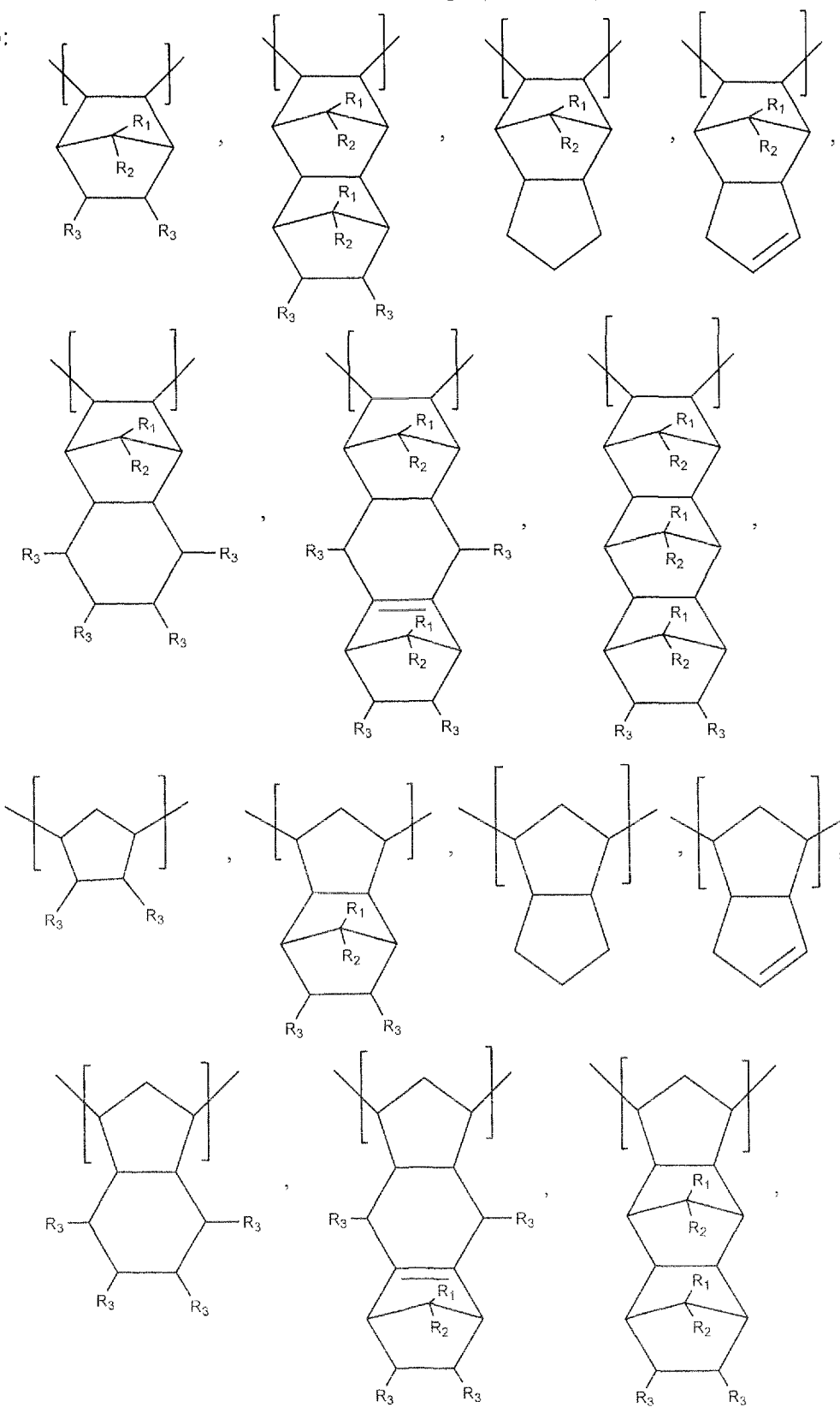
where:

20 each R_1 and R_2 is individually selected from the group consisting of -H, and alkyl groups; and

25 R_3 is individually selected from the group consisting of -H, substituted and unsubstituted aryl groups, alkyl groups, cycloalkyl groups, aralkyl groups, ester groups, ether groups, acetyl groups, alcohols, aldehyde groups, ketones, nitriles, and combinations thereof.

18. The article of claim 13, wherein copolymer comprises recurring monomers of:

(I):



and combinations of the foregoing,

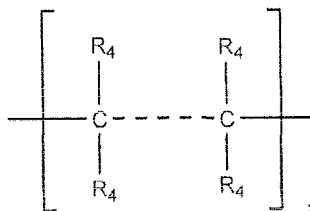
where:

each R_1 and R_2 is individually selected from the group consisting of -H, and alkyl groups; and

5 each R_3 is individually selected from the group consisting of -H, substituted and unsubstituted aryl groups, alkyl groups, cycloalkyl groups, aralkyl groups, ester groups, ether groups, acetyl groups, alcohols, aldehyde groups, ketones, nitriles, and combinations thereof;

10 and

(II):



15 where:

--- is a single or double-bond; and

each R_4 is individually selected from the group consisting of -H and alkyl groups.

20 19. The article of claim 13, said composition further comprising an ingredient selected from the group consisting of tackifiers, low molecular weight cycloolefin copolymers, plasticizers, antioxidants, and mixtures thereof.

25 20. The article of claim 19, wherein said ingredient is a tackifier selected from the group consisting of polyterpene resins, beta-polyterpene resins, styrenated terpene resins, polymerized rosin resins, rosin ester resins, cyclo-aliphatic hydrocarbon resins, C5 aliphatic hydrocarbon resins, hydrogenated hydrocarbon resins, and mixtures thereof.

30 21. The article of claim 19, wherein said ingredient is a low molecular weight cycloolefin copolymer having a weight average molecular weight of less than about 50,000 Daltons.

22. A composition useful for bonding two substrates together, said composition comprising a cycloolefin copolymer and an ingredient dissolved or dispersed in a solvent system, said ingredient being selected from the group consisting of tackifier resins, low molecular weight cycloolefin copolymers, and mixtures thereof.

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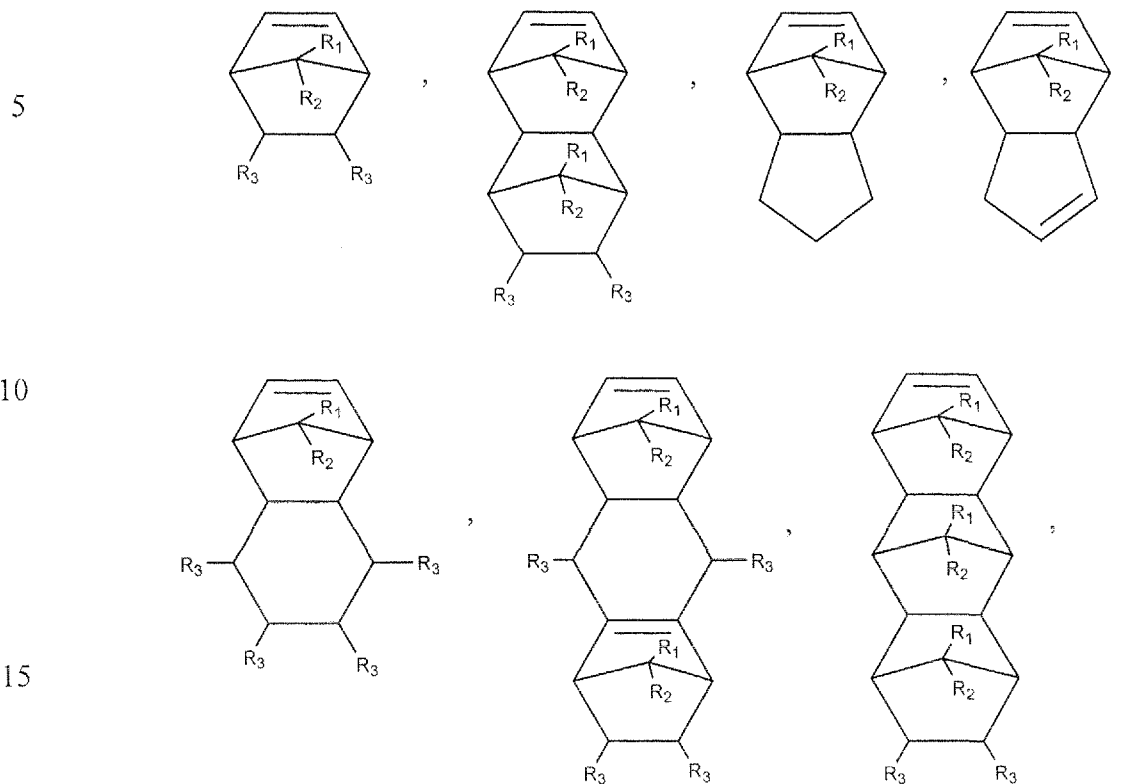
23. The composition of claim 22, said composition comprising from about 5% to about 85% by weight of said copolymer, based upon the total weight of the composition taken as 100% by weight.

10

24. The composition of claim 22, wherein said solvent system is selected from the group consisting of limonene, mesitylene, dipentene, pinene, bicyclohexyl, cyclododecene, 1-tert-butyl-3,5-dimethylbenzene, butylcyclohexane, cyclooctane, cycloheptane, cyclohexane, methylcyclohexane, methyl ethyl ketone, cyclopentanone, and mixtures thereof.

15

25. The composition of claim 22, said copolymer being formed from the polymerization of a cyclic olefin selected from the group consisting of



and combinations of the foregoing,

where:

- 20 each R_1 and R_2 is individually selected from the group consisting of -H, and alkyl groups; and
- each R_3 is individually selected from the group consisting of -H, substituted and unsubstituted aryl groups, alkyl groups, cycloalkyl groups, aralkyl groups, ester groups, ether groups, acetyl groups, alcohols, aldehyde groups, ketones, nitriles, and combinations thereof.
- 25

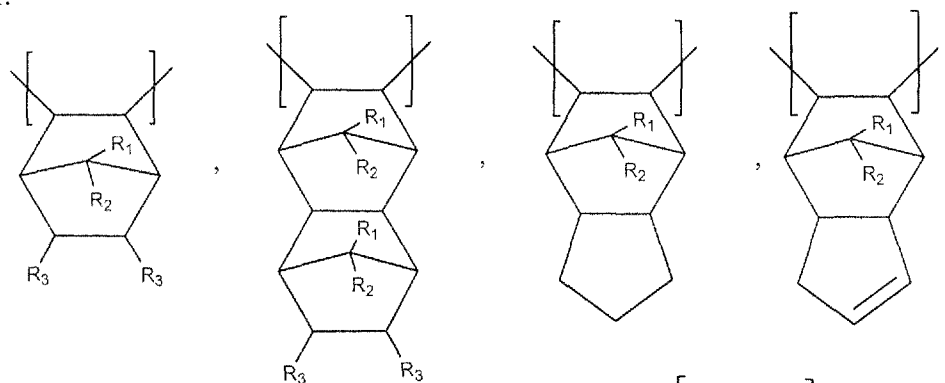
26. The composition of claim 25, wherein said cyclic olefin is polymerized with an acyclic olefin selected from the group consisting of branched and unbranched C_2 - C_{20} alkenes.

30

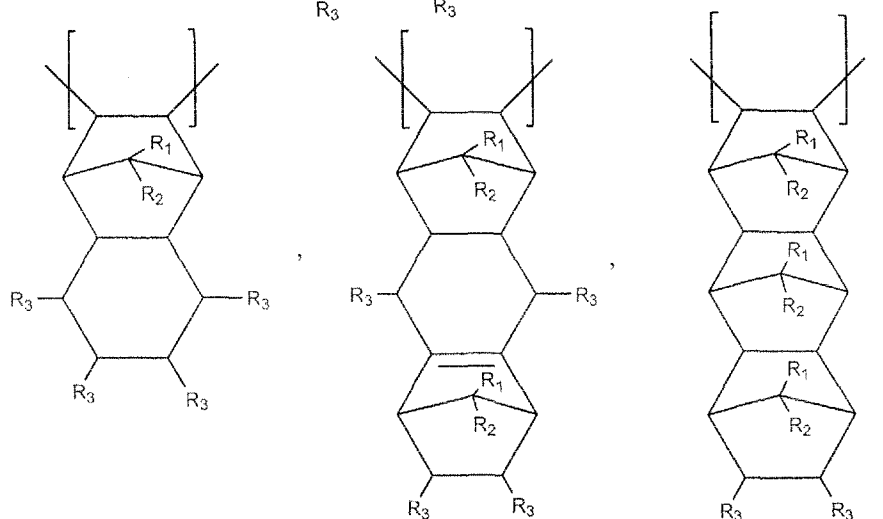
27. The composition of claim 22, wherein said copolymer comprises recurring monomers of:

(I):

5

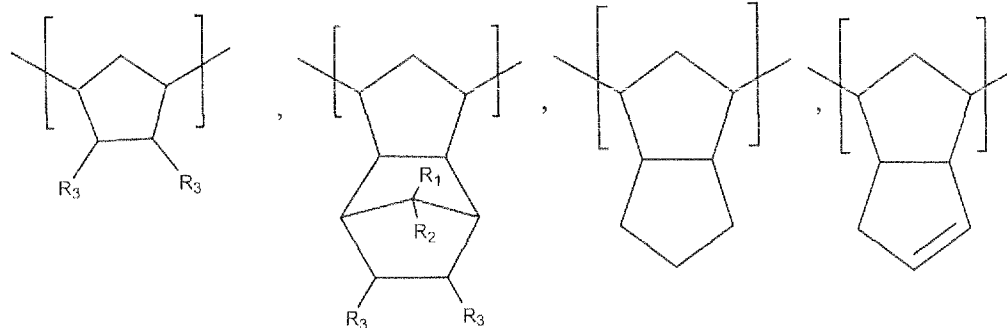


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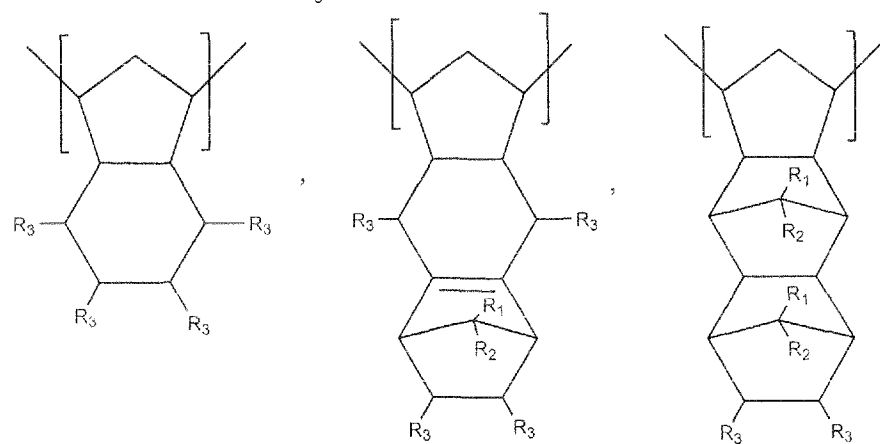


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25



30

and combinations of the foregoing,

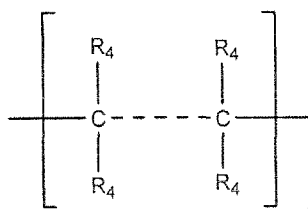
where:

each R_1 and R_2 is individually selected from the group consisting of -H, and alkyl groups; and

5 each R_3 is individually selected from the group consisting of -H, substituted and unsubstituted aryl groups, alkyl groups, cycloalkyl groups, aralkyl groups, ester groups, ether groups, acetyl groups, alcohols, aldehyde groups, ketones, nitriles, and combinations thereof;

10 and

(II):



15 where:

----- is a single or double-bond; and

each R_4 is individually selected from the group consisting of -H and alkyl groups.

20 28. The composition of claim 22, where said ingredient is present in said composition at a level of from about 2% to about 80% by weight, based upon the total weight of the composition taken as 100% by weight.

25 29. The composition of claim 22, wherein said ingredient is a tackifier resin selected from the group consisting of polyterpene resins, beta-polyterpene resins, styrenated terpene resins, polymerized rosin resins, rosin ester resins, cyclo-aliphatic hydrocarbon resins, C5 aliphatic hydrocarbon resins, hydrogenated hydrocarbon resins, and mixtures thereof.

30 30. The composition of claim 22, wherein said ingredient is a low molecular weight cycloolefin copolymer having a weight average molecular weight of less than about 50,000 Daltons.

31. The composition of claim 22, wherein said composition is essentially free of adhesion promoting agents.

32. The composition of claim 22, said composition having thermal stability up to a
5 temperature of about 350°C.

33. The composition of claim 22, said composition being essentially free of crosslinking agents.

10 34. The composition of claim 22, said composition further comprising an antioxidant selected from the group consisting of phenolic antioxidants, phosphite antioxidants, phosphonite antioxidants, and mixtures thereof.

15 35. The composition of claim 22, said composition having a melt viscosity of less than about 100 Pa•S at the debonding temperature of said composition.

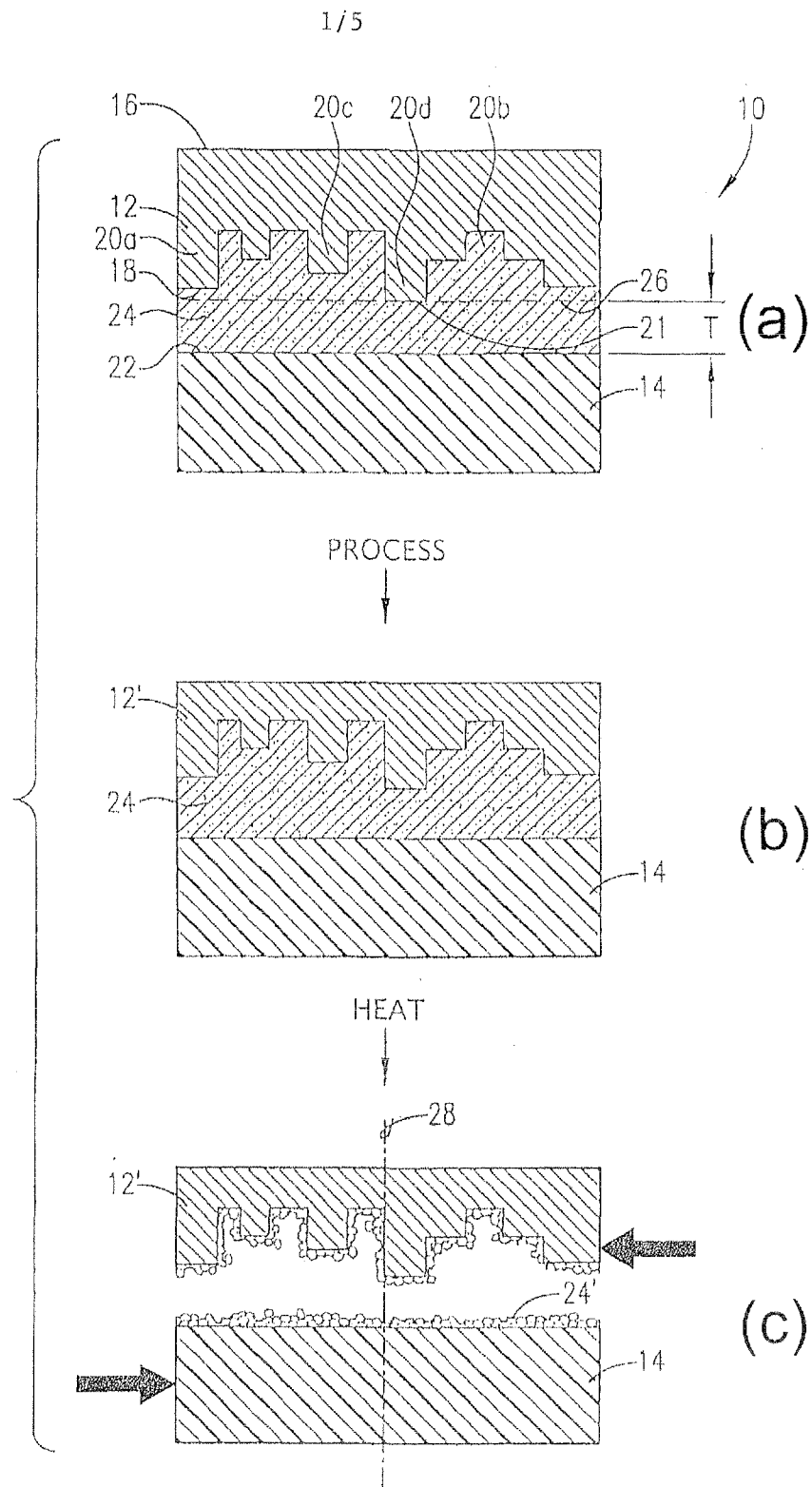


Fig. 1

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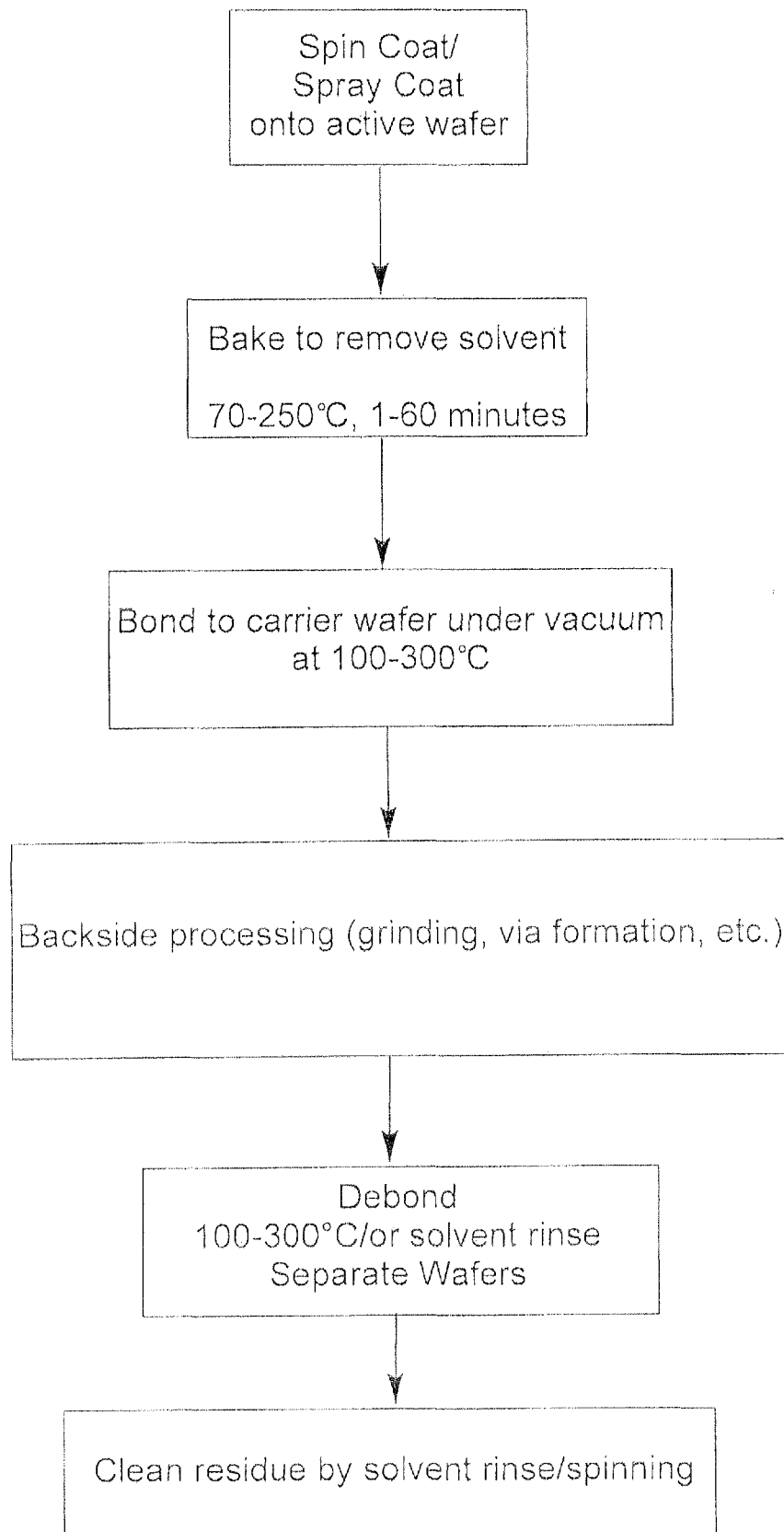


Fig. 2

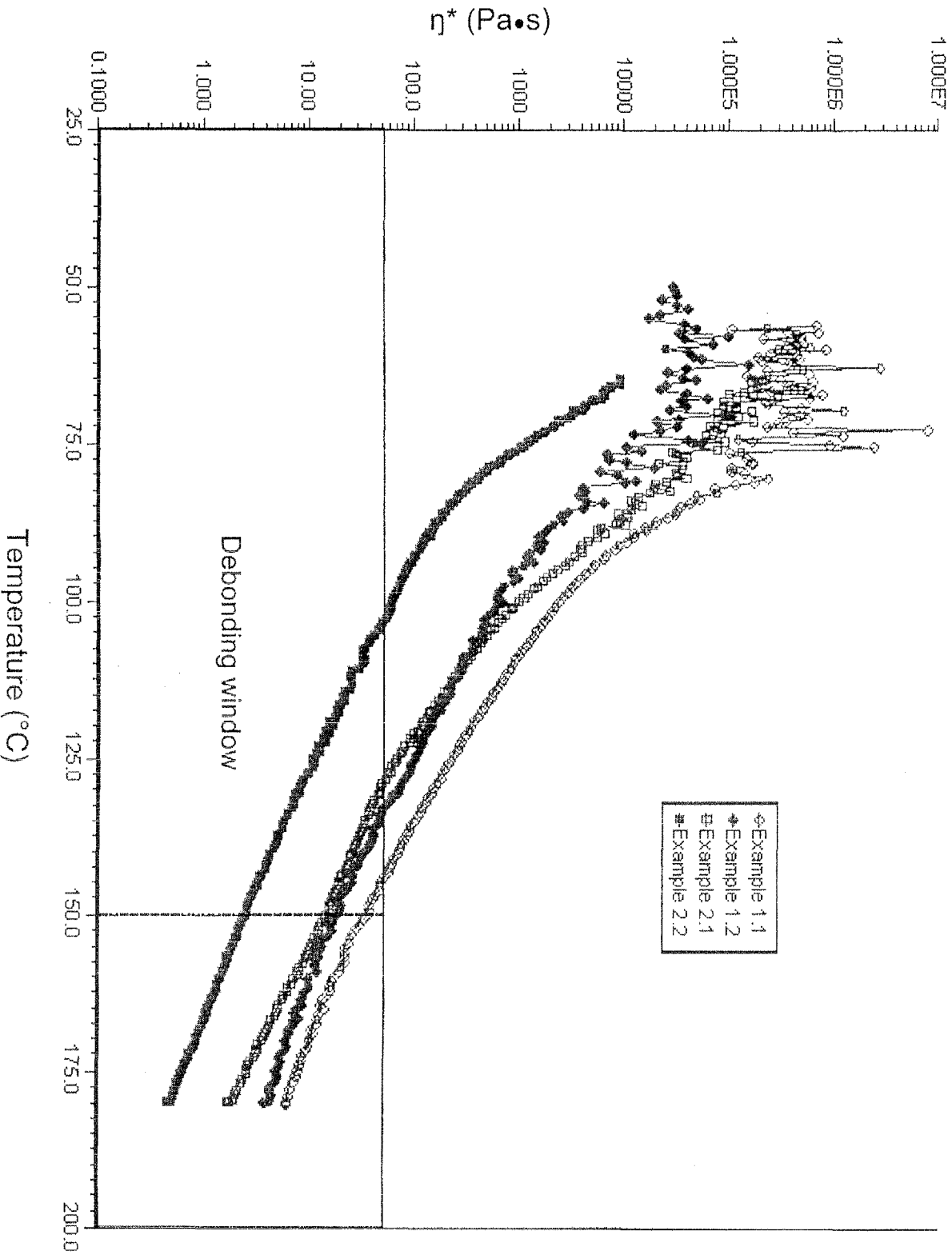


Fig. 3

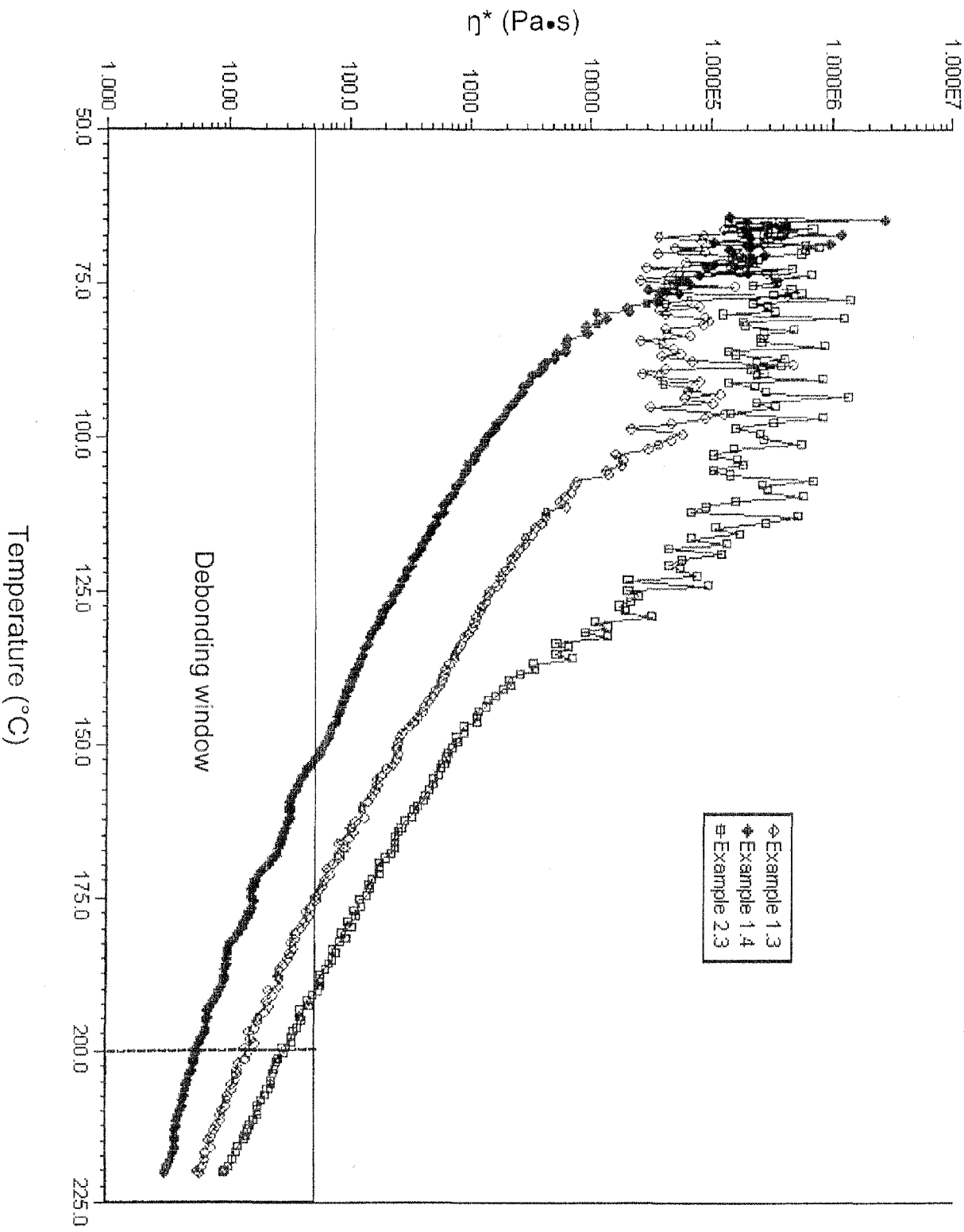


Fig. 4

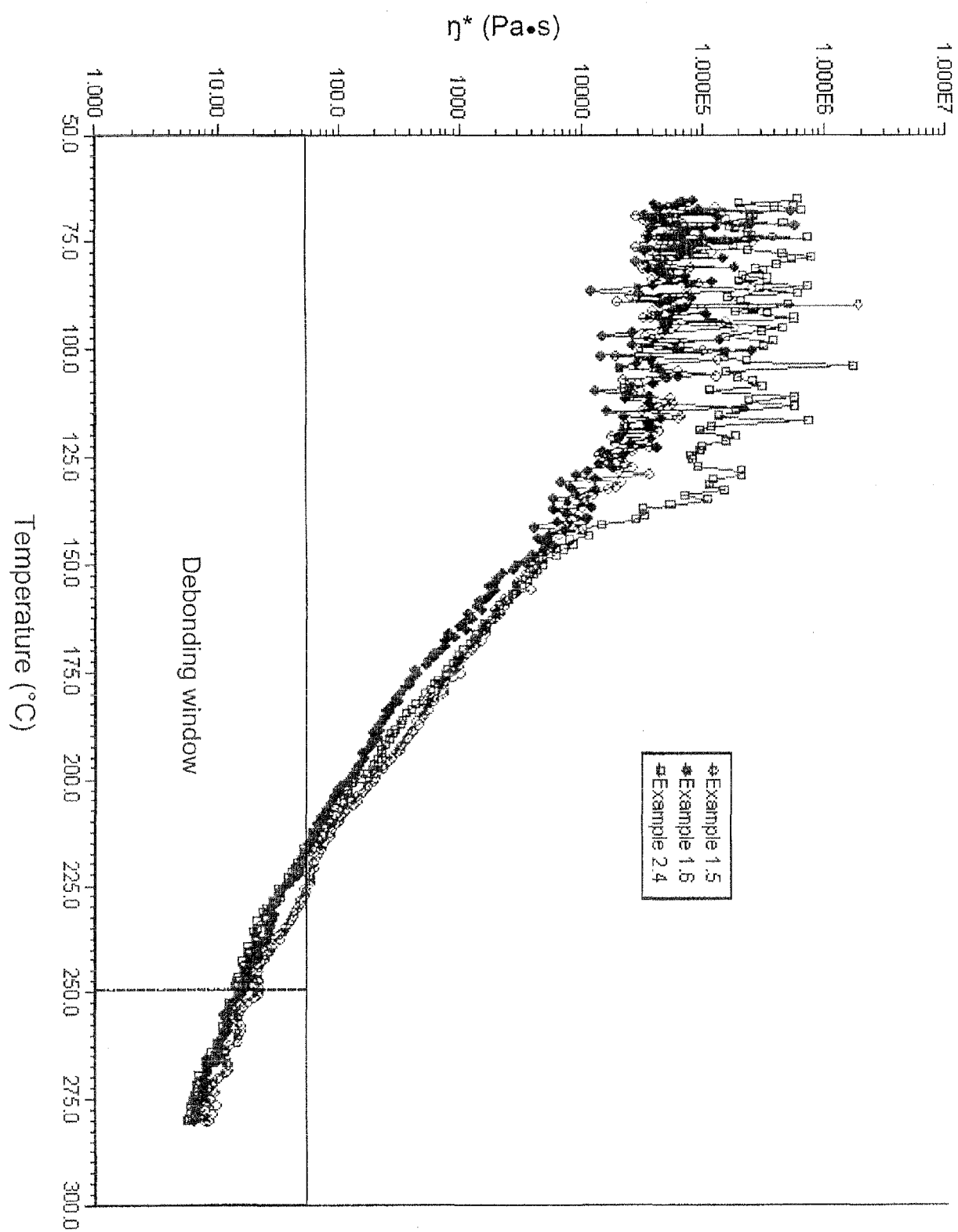


Fig. 5