

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
15 February 2001 (15.02.2001)

PCT

(10) International Publication Number
WO 01/10937 A1

(51) International Patent Classification⁷: C08J 3/22,
C08L 27/12 // C08F 265/06

(21) International Application Number: PCT/US99/27506

(22) International Filing Date:
19 November 1999 (19.11.1999)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
09/371,070 9 August 1999 (09.08.1999) US

(71) Applicant: 3M INNOVATIVE PROPERTIES COMPANY [US/US]; 3m Center, Post Office Box 33427, Saint Paul, MN 55133-3427 (US).

(72) Inventors: EIAN, Gilbert, L.; Post Office Box 33427, Saint Paul, MN 55133-3427 (US). LUDWIG, Bret, W.; Post Office Box 33427, Saint Paul, MN 55133-3427 (US). ANDRUS, Milton, H., Jr.; Post Office Box 33427, Saint Paul, MN 55133-3427 (US).

(74) Agents: SCHULTE, Daniel, C. et al.; 3 M Innovative Properting Company, Office of Intellectual Property Counsel, Post Office Box 33427, Saint Paul, MN 55133-3427 (US).

(81) Designated States (*national*): AE, AL, AM, AT, AT (utility model), AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, CZ (utility model), DE, DE (utility model), DK, DK (utility model), DM, EE, EE (utility model), ES, FI, FI (utility model), GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK (utility model), SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

— With international search report.

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: POLYMERIC COMPOSTIONS

(57) Abstract: Described are compositions comprising core/shell particles and fluoropolymer, wherein the core comprises an ultraviolet absorber, as well as multiphase polymeric compositions comprising a polymeric core phase and a polymeric shell phase, wherein the polymeric core phase comprises an ultraviolet absorber.

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POLYMERIC COMPOSITIONS

Field of the Invention

The invention contemplates UV-absorbing compositions including latices,
5 agglomerate particles, powders, films, coatings, multilayer articles, and other
materials prepared from or containing polymeric particles or derivatives thereof,
and associated methods.

Background

10 Ultraviolet radiation (UV) can cause degradation of certain materials if
exposed. Chemical materials known as ultraviolet absorbers, or UVAs, can be
used to protect materials from the damaging effects of UV radiation. A UVA can
be incorporated into a material to protect that material from UV radiation, or, a
composition that contains UVA can be applied to a UV-sensitive substrate to
15 protect the substrate.

Protective coating compositions, sometimes referred to as "topcoats," can
be applied to outdoor-durable materials such as signs based on flexible substrates
and optionally having applied graphics, where the coating functions to inhibit dirt
buildup or dirt penetration, as a barrier to water, to prevent plasticizers or other
20 ingredients from migrating out of the substrate, or to allow ease of cleaning. A
topcoat can include polymeric materials (e.g., a fluoropolymer to provide dirt
resistance or cleanability), as well as stabilizers to protect the topcoat or the
substrate from degradation, e.g., due to UV radiation. Degradation may involve
yellowing, embrittlement, or loss of clarity, gloss, or water resistance.

25 Unfortunately, while it can be desirable to incorporate a UVA into a
protective coating, UVAs can cause some difficult problems. One problem is the
relative impermanence of UVAs in many chemical compositions. Non-reactive
UVAs can be included in a chemical composition as a dispersed compound, not
chemically attached to any other component. These UVAs can be lost from a
30 composition by volatilization during processing (e.g., drying), or by otherwise
migrating to the surface of a composition followed by removal as dust or wash off.
Loss of the UVA leaves the composition and its substrate less protected from
ultraviolet radiation, allowing UV radiation to degrade the composition or

substrate. One imperfect remedy to this problem is to include larger amounts of UVA in a composition.

A further problem with UVAs is that they can be incompatible with different polymeric materials (e.g., fluoropolymers). This incompatibility can lead to instability (e.g., thermodynamic instability) or water sensitivity of the composition, which can cause a loss of physical or optical properties, including loss of clarity or increased fogginess. Incompatibility can also cause increased or accelerated loss of UVA by migration, bleeding, or blooming.

Attempts to incorporate UVAs into chemical compositions such as topcoats have been met with a variety of frustrating results, especially when the composition includes an ingredient that is incompatible with the UVA, as are many fluoropolymers. There is a general need to identify ultraviolet absorbing materials and compositions, and also to identify materials that can be used to prepare UV-absorbing compositions such as films and coatings. There is a further need to incorporate UVAs into chemical compositions that contain other materials with which the UVA may not be compatible, wherein the UVA becomes a relatively permanent component of the composition, and wherein the composition is relatively thermodynamically stable, to provide long-term protection from ultraviolet radiation.

20

Summary of the Invention

The invention provides ultraviolet radiation-absorbing compositions including latices of core/shell particles wherein the core comprises a core polymer and a UVA, as well as multiphase polymeric compositions comprising a polymeric core phase and a polymeric shell phase, wherein the polymeric core phase includes ultraviolet absorber.

Preferred multiphase polymeric compositions comprising a core phase, a shell phase, and a phase of an additional polymeric material can exhibit improved thermodynamic stability and UVA retention, thereby exhibiting time-stable protection from ultraviolet radiation, with lasting physical and mechanical properties. This can be true even if the UVA is used in combination with an additional material that is not compatible with the UVA.

30

An exemplary composition is a latex comprising core/shell particles and a polymeric film-forming material, wherein the latex may be coated and dried to form a film, or which may be spray dried to form a powder which can be further processed. Such a latex can be formed into a film or coating e.g., by coating and drying, or, dried to form a powder, preferably spray-dried to form a powder comprising agglomerate particles which can be coated onto a substrate and fused to form a film or coating.

Multiphase polymeric compositions of the invention can comprise phase domains of a polymeric core phase, a polymeric shell phase, and a polymeric film-forming phase, wherein the morphology is such that the polymeric core phase does not substantially contact the polymeric film-forming phase, but both of these phases contact the polymeric shell phase, which separates the other two phases. It has been found that compositions having this preferred morphology, particularly if the core particle (or a component thereof) is not compatible with the polymeric film-forming material, can exhibit improved initial physical properties, which can be maintained with aging, as compared to chemical compositions containing chemically identical ingredients, in identical amounts, but exhibiting a different morphology. Such properties can include one or more of: flexibility, lasting protection from ultraviolet radiation due to reduced migration loss (e.g., bleeding or blooming) of the UVA (especially if the UVA is not chemically attached to the core polymer); thermodynamic stability; and improved resistance to water (e.g., reduced water sensitivity).

A particular advantage of this preferred morphology is that it allows the use of a core particle or a component thereof, e.g., a core polymer or a UVA, in a composition that contains another material (e.g., a polymeric film-forming material) that is incompatible with the core particle or core particle component. If a core particle or its component is incompatible with a polymeric film-forming material, the shell material shields the core particle phase from the incompatible phase, thus preventing the consequences otherwise associated with including the core particle with an incompatible material, and thereby preferably achieving a thermodynamically stable composition.

Multiphase polymeric compositions having the described morphology can be prepared by various methods. For instance, a multiphase composition can be prepared from a latex that includes polymeric particles having a core/shell structure, wherein the core comprises UVA, and particles of polymeric film-forming material. This can be accomplished by forming the latex into a film or coating, and allowing the latex to dry. Alternatively, such a multiphase composition can be prepared by spray-drying the latex to prepare a powder of agglomerate particles, powder coating the agglomerate particles, and fusing the coated powder.

10 An aspect of the invention relates to compositions comprising fluoropolymer and polymeric core/shell particles wherein the core comprises a UVA. The composition can be in the form of one or more of the following: a latex; an agglomerate particle of the core/shell particles fused to fluoropolymer particles; a powder comprising core/shell particles and fluoropolymer particles, or
15 their agglomerates; a polymeric composition such as a multiphase polymeric composition, e.g., in the form of a film, coating, or multilayer article; or a product or composition derived from or containing any of these compositions. Such compositions can block or absorb ultraviolet radiation for any desired reason and can exhibit desirable cleanability properties, and desired mechanical or physical
20 properties such as flexibility. The core/shell particles can be included in a composition to protect that composition itself from UV radiation, or, a UVA-containing film or coating can be disposed onto another material to protect that material from UV radiation.

 Another aspect of the invention relates to a multiphase polymeric
25 composition comprising a polymeric core phase, a polymeric shell phase, and a fluoropolymer phase, wherein the polymeric core phase comprises an ultraviolet absorber and wherein the polymeric core phase and the fluoropolymer phase do not substantially contact one another, but both the polymeric core phase and the fluoropolymer phase contact the polymeric shell phase.

30 Another aspect of the invention relates to a method for preparing a multiphase polymeric composition comprising a polymeric core phase, a polymeric shell phase, and a polymeric film-forming phase, wherein the polymeric core phase

comprises an ultraviolet absorber. The method includes steps comprising: powder coating onto a substrate polymeric materials comprising the polymeric core phase, the polymeric shell phase, and the polymeric film-forming phase, and fusing the polymeric materials to form a multiphase polymeric composition wherein the polymeric core phase and the polymeric film-forming phase do not substantially contact one another, but both the polymeric core phase and the polymeric film-forming phase contact the polymeric shell phase.

As used herein, the following terms shall be given the recited meanings:

The term "thermoplastic" means materials that soften or flow upon exposure to heat and/or pressure. Thermoplastic is contrasted with "thermoset," which describes materials that react irreversibly upon heating so that subsequent applications of heat and pressure do not cause them to soften or flow.

"(Meth)acrylate" means either acrylate or methacrylate.

"Phase" is used in a manner not inconsistent with its generally accepted meaning in the chemical art, for instance to refer to a discrete, typically homogeneous component of a chemical composition.

"Domain," when referring to a phase of a multiphase composition, refers to individual, continuous or discontinuous, microscopic or macroscopic portions of a phase within the multiphase composition. Examples of discontinuous domains include the following, as illustrated in Figure 3: individual core particle domains; polymeric shell domains; and fluorochemical domains.

In a multiphase polymeric composition, if individual domains of one phase have an affinity for each other (there is an affinity between separate, individual domains of the same phase) that is significantly greater than the affinity between domains of that phase and domains of a chemically different phase of the composition which the first phase contacts, individual domains of the like phases will be thermodynamically driven to combine with each other, causing polymer segregation and growth in domain size of like phase domains. Phases of a multiphase polymeric composition are referred to as "incompatible" if, while both separate phases are contained in a multiphase composition, individual domains of a like phase tend to combine with each other within the composition to form larger phase domains. If domains of a like phase of a multiphase polymeric composition

do in fact form larger-sized phase domains, i.e., the composition suffers gross symptoms of polymer segregation, the multiphase polymeric composition is considered to be "thermodynamically unstable."

5 Gross symptoms of polymer segregation can include decreased optical clarity, loss of flexibility, embrittlement, and other effects. Polymer segregation within a multiphase polymeric composition can be considered to occur upon a significant loss of clarity or flexibility, or a significant increase in brittleness. These properties can be measured by known methods.

10 Brittleness can be measured by measuring elongation, with a 20% decrease in elongation being considered significant and indicating polymer segregation and thermodynamic instability.

Clarity can be measured using a Hazemeter. If the result of a haze measurement according to ASTM D1003, upon aging, increases by 10 percentage points, e.g., 5 percentage points or more (e.g., from a reading of 0% to 5%, see the
15 Examples Section and Table 1), it is considered to have lost a significant amount of clarity, and is considered to be thermodynamically unstable.

Flexibility of a multiphase polymeric film composition applied to a flexible substrate can be detected by the following test: samples of the film (approximately 20 to 50 micrometers in thickness) are coated onto a substrate (approximately 500
20 to 600 micrometers in thickness, e.g., 3M PANAFLEX™ 930 Sign and Awning Substrate) (and optionally aged, e.g., for 1 week at 150F). Two samples of the coated film can be cooled to room temperature (about 25°C), and then bent and creased, one in a direction to cause compression of the coating, and another to cause extension. Creasing can be a fold to 180 degrees, followed by heavy finger
25 pressure along the fold. If the coating visibly cracks, the coating is considered to have lost flexibility. To more easily see cracks, a permanent marker can be applied to the creased coating, and the marker can be washed off with isopropyl alcohol. If a crack is present, the marker will stain the crack and remain visible after washing; if no cracks are present, the marker will wash clean.

30

Brief Description of the Drawings

Figure 1 is an SEM photomicrograph at 2.0kv x500 magnification of agglomerate particles prepared by spray drying a latex containing core/shell particles and fluoropolymer particles.

5 Figure 2 is an SEM photo at 3.0kv x25.0K magnification containing the surface of agglomerate particles prepared by spray drying a latex of core/shell particles and fluoropolymer particles.

Figure 3 is a transmission electron micrograph (TEM) taken at 60,000 times magnification of a multiphase film prepared according to example. The
10 smaller dark, circular regions are domains of the polymeric core phase of the core/polymeric shell (dyed with RuO₄); the core phase domains are surrounded by lighter domains of polymeric shell. The larger domains, generally relatively light, and generally not containing domains of the darker core polymer phase are domains of a fluoropolymer phase.

15 Figure 4 is a transmission electron micrograph (TEM) taken at 60,000 times magnification of a multiphase film prepared according to Comparative Example 2. The small phase domains are of the core particle phase; the larger, nondistinct, essentially continuous phase comprises a mixture of polymeric shell and fluoropolymer.

20 Figure 5 is a schematic view of a method of applying a multiphase polymeric coating onto a substrate according to this invention.

Figure 6 is a schematic cross-sectional view illustrating a preferred embodiment of the invention.

Detailed Description

25 Polymeric core/shell particles used in the practice of the invention exhibit a core/shell structure, meaning the particles comprise a polymeric core and a polymeric shell. The polymeric core comprises core polymer and ultraviolet absorber (UVA), which may or may not be chemically incorporated into the core
30 polymer. (The polymeric core can include non-polymeric materials, and may be referred to herein as the "core" or "core particle.") These particles are described in Assignee's copending United States Patent Application entitled UV-Absorbing

Particles, having Serial No. 09/371,064, Attorney's Docket No. 54644USA6A, which was filed on even date herewith.

The core polymer can be chosen to be useful for a specific application of the core/shell particle to provide various physical, chemical, or mechanical properties. The core polymer can be chosen to be a relatively soft or rubbery material, e.g., having a glass transition temperature (T_g) of 40 degrees Celsius (40C) or lower. A soft or rubbery core polymer may impart flexibility to a core/shell particle or a composition containing or derived from the core/shell particle. Alternatively, the core polymer may be relatively hard, or even thermosetting or thermoset. A relatively hard core polymer can impart dirt resistance or cleanability properties to a core particle so that if a portion of a core particle is exposed at the surface of a composition, it will not cause dirt buildup, or will at least be cleanable. The core polymer can be crosslinked to prevent breakage or disruption of the core particle during processing, e.g., to a film or coating, which could cause core material to be present at the surface of the film or coating.

The core polymer can comprise monomeric units derived from one or more reactive monomers or comonomers (referred to collectively herein as the "core monomer" or "core monomers"). The core monomer can include any reactive compound (e.g., monomer, dimer, trimer, oligomer, prepolymer, polymer, etc.) capable of forming a useful core polymer (meaning a homopolymer or a copolymer). Examples of useful core monomers include monofunctional reactive compounds comprising unsaturated moieties such as vinyls, e.g., (meth)acrylates, with lower (meth)acrylates being preferred, or other reactive compounds such as epoxies, alcohols, or isocyanates. Specific examples include butyl acrylate, hexyl acrylate, octyl acrylate, decyl acrylate, and butyl methacrylate, with ethyl acrylate and methyl methacrylate being preferred.

The core monomer can include multi-functional reactive compounds having suitable reactive moieties, e.g., two, three, or more reactive moieties such as vinyls, (meth)acrylates, epoxies, alcohols, isocyanates, etc. Such multi-functional compounds (sometimes referred to or used as crosslinkers) are known in the art of polymer chemistry, and useful examples include but are not limited to multi-

functional vinyl compounds such as multifunctional (meth)acrylate compounds, multifunctional styrenes, and multifunctional allyl compounds, specifically including allyl acrylate, allyl methacrylate (AMA), butanediol diacrylate (BDDA), and hexanediol diacrylate (HDDA).

5 Ultraviolet radiation absorbers, UVAs, are known and commercially available chemical materials which absorb ultraviolet radiation. *See, e.g., Rabek, J. F., Photostabilization of Polymers, 203-42 (1990).* A variety of UVAs are known and commercially available, and can be prepared by known methods. *See id.* A UVA can preferably be soluble in a core polymer, and can be chosen to have
10 desired UV-absorption properties for a given application of the core/shell particles. Examples of UVAs include benzophenones, benzotriazoles, triazines, cinnamates, cyanoacrylates, dicyano ethylenes, and para-aminobenzoates. For protecting a PVC substrate, preferred UVAs include benzotriazoles and benzophenones.

 The UVA can be included in, contained by, or attached to the core particle
15 in any configuration, and in any chemical or physical manner. A UVA can be a relatively low molecular weight compound dispersed in, not chemically attached to, the core polymer. Such dispersible UVAs are commercially available, with a single example being TINUVIN 1130, from Ciba Specialties Corp.

 Optionally, a UVA can be functionalized with one or more reactive
20 moieties to provide a reactive, monomeric UVA which can be included in and reacted with the core monomer to become chemically incorporated into the core polymer as a monomeric unit of the backbone or as a group pendant from the core polymer. Suitable reactive moieties include unsaturated moieties such as vinyls, e.g., (meth)acrylate and styrene, or other useful reactive moieties such as alcohols,
25 isocyanates, epoxies, etc. Specific examples of functionalized UVAs include (meth)acrylate-functionalized UVAs such as (meth)acrylate-functional benzotriazoles and benzophenones. These compounds are well known, can be prepared by known methods, and are commercially available, e.g., NORBLOC UVAs such as NORBLOC 7966.

30 A core particle can be prepared from any useful amounts of various core monomers, reactive (monomeric) UVA, non-reactive (dispersible) UVA, and crosslinker. Useful amounts of these different ingredients can be from about 50 to

98 parts by weight core monomer based on the total weight of the core particle, preferably from about 65 to 95 parts by weight, and more preferably from about 75 to 90 parts by weight core monomer (for purposes of these ranges the monomer does not include reactive (monomeric) UVA), and up to about 5 weight percent crosslinker, preferably from about 0.1 to 3 weight percent crosslinker, e.g., from about 0.2 to about 2 weight percent crosslinker, based on the total weight of the core particle. The amount of UVA included in a core particle can be any useful amount, depending on factors such as the chemistry of the UVA, the substrate, the core monomer, etc. In general, UVA can be included in a core particle in an amount in the range from about 2 to 50 parts by weight UVA based on the total weight of the core particle, preferably from about 5 to 35 parts by weight UVA, and more preferably from about 10 to 25 parts by weight; these ranges are the same whether the UVA is monomeric, i.e., reactive with the core monomer, or non-reactive. A preferred core polymer may also be prepared from a single acrylate or (meth)acrylate monomer, and optionally a reactive UVA and crosslinker, e.g., ethyl acrylate and a (meth)acrylate-functional UVA, and crosslinker.

The polymeric shell takes the form of a polymeric material disposed on the core, preferably completely surrounding (e.g., encapsulating) the core. Still, it is possible for production processes to result in particles wherein the polymeric shell does not completely surround the core, but only partially covers the core, leaving a portion of the core exposed. These particles, if produced, will typically be present in relatively small amounts compared to core/shell particles where the polymeric shell does completely surround or encapsulate the core.

The polymeric shell comprises a polymeric material (shell polymer) useful for a chosen core/shell particle and application, to provide desired physical, mechanical, or chemical properties. The shell polymer may be chosen to be a thermoplastic polymer such as one having a T_g sufficiently high to yield a powder composition that flows freely, without particles substantially gumming or clinging together, but still low enough that the core/shell particles and compositions or products containing or derived from the core/shell particles can be suitable for processing and still exhibit useful chemical, physical, and mechanical properties. The shell polymer can also be sufficiently hard (e.g., have a sufficiently high T_g)

to exhibit dirt resistance and cleanability properties. It may be desirable to crosslink the shell polymer in order to provide desired physical properties. For instance, the shell polymer could be crosslinked to increase solvent resistance, so the core/shell particles may be added to solvent-based systems while still
5 maintaining their core/shell structure.

The polymeric shell, or components thereof, may be either compatible or incompatible with the core particle, while preferably exhibiting a useful level of adherence to the core particle. The ingredients of the polymeric shell may be chosen based on compatibility or incompatibility with a polymeric film-forming
10 material with which the core/shell particles may be intended for use (e.g., mixed with, coated on, or otherwise associated with). If the core/shell particles are designed for use in a composition comprising a polymeric film-forming material, e.g., a fluoropolymer, the polymeric shell can preferably be, but is not required to be, compatible with that polymeric film-forming material.

15 Monomers used to prepare the shell polymer (shell monomers) can be chosen to provide a polymeric shell according to the above considerations, e.g., to provide desired physical properties such as hardness or softness, or compatibility or cleanability properties, and can be chosen to be thermoplastic, thermosetting or crosslinked. Preferred shell monomers can include (meth)acrylate monomers such
20 as methyl methacrylate (MMA), methyl acrylate, ethyl methacrylate, ethyl acrylate (EA), and mixtures of these. Particularly preferred shell monomers include MMA and EA.

Selected shell monomers can be included in a variety of useful amounts, with preferred amounts of methyl methacrylate and ethyl acrylate being in the
25 range from 70:30 to 95:5 (MMA:EA) (by weight), more preferably in the range from about 80:20 to 90:10, MMA:EA.

The amounts (by weight) of core particle and polymeric shell in a core/shell particle can be any amounts of each which are found to be useful for a particular application, and to provide desired physical or mechanical properties such as
30 flexibility or cleanability, with an exemplary range being from about 1:1 to 1:9 parts by weight core particle per parts by weight polymeric shell (core:shell).

Preferred amounts of core particle to polymeric shell can be in the range from about 30:70 to 15:85.

Polymeric core/shell particles can be prepared from the above-described materials, by methods known in the polymer art. The chosen method can depend
5 on a number of factors including the identity of the core and shell monomers, whether the UVA is non-reactive and therefore will be dispersed in the core polymer or reactive and will be an attached chemical component of the core polymer, or whether the core polymer or polymeric shell is thermoplastic, thermosetting, or crosslinked. Examples of suitable methods include those
10 described, e.g., in United States Patent Number 5,461,125 (Lu et al.), and Segall et al., *Core-Shell Structured Latex Particles. II. Synthesis and Characterization of Poly(n-butyl acrylate)/Poly(benzyl methacrylate-styrene) Structured Latex Particles*, J. Applied Poly Sci. 58, 401-417 (1995). Specifically, the core/shell particles can be prepared by semi-continuous or two-stage emulsion
15 polymerization methods, wherein a first polymerization produces a core particle comprising the UVA, and by a second polymerization a polymeric shell is formed on the core.

Preferred methods for preparing core/shell particles can produce an aqueous latex comprising a dispersion of core/shell particles in water. The
20 core/shell particles can be present in a range of differently shaped and sized particles, typically having an average (mean) size (diameter) in the range from about 40 to 200 nanometers. The size of core/shell particles can be measured by known analytical methods, for example by light scattering methods using a light scattering apparatus, such as a COULTER N4 MD submicron particle analyzer.

25 The core/shell particles can be combined with a polymeric film-forming material, and processed to film-forming, UV-absorbing chemical compositions.

Polymeric film-forming materials are well known, and are described throughout the patent and scientific literature; See e.g., *Organic Coatings: Science and Technology*, Wicks, Jones, and Pappas 35-48 (1992), and *Encyclopedia of*
30 *Polymer Science and Engineering*, Supplement Volume, pp. 53-59 (1989). Polymeric film-forming materials can typically be considered to be polymeric materials that can be formed into a continuous coating or film (e.g., a solid), e.g.,

by drying with removal of solvent, by chemical reaction, or by melting or fusing. The polymeric film-forming material can be in any useful form, such as a solid (e.g., a powder), a liquid (e.g., a high solids liquid such as a polymeric resin or a low solids liquid such as an aqueous latex), or any combination of these or other forms. The polymeric film-forming material can be reactive or non-reactive, thermoplastic, thermosetting, or otherwise polymerizable, and can be either compatible or incompatible with either the polymeric shell or the core particle. The polymeric film-forming material can have any useful chemistry, including chemistries similar to or identical to a component of the core or polymeric shells. Examples of some suitable polymeric film-forming materials include but are not limited to polyvinyl chloride, polyamide, polyester, polyacrylate, polymethacrylate, polyethylene, polypropylene, fluoropolymers, and mixtures of one or more of these.

The core/shell particles can be particularly useful with a fluoropolymer. Fluoropolymers and their preparation are well known. Examples of fluoropolymers include thermoplastic fluoropolymers such as those available in the form of fluoropolymer powders and fluoropolymer latices. Latices can for some applications be preferred over powders, because latices can be combined with a core/shell particle latex and then processed further, e.g., spray dried to form agglomerate particles as described *infra*.

Specific examples of useful thermoplastic fluoropolymers include homopolymers and copolymers comprising monomeric units derived from fluorinated monomers such as vinylidene fluoride (VF₂), hexafluoropropene (HFP), chlorotrifluoroethylene (CTFE), vinyl fluoride (VF), trifluoroethylene (TrFE), and tetrafluoroethylene (TFE), among others, optionally in combination with one or more other non-fluorinated monomer. In general, any amounts of these and other fluorinated and non-fluorinated comonomers can be used to prepare a fluoropolymer. Particularly preferred fluoropolymers can include fluoropolymer prepared from vinylidene fluoride and HFP, with the amounts of VF₂ and HFP being approximately 80 to 100 weight percent VF₂ and 0 to 20 weight percent HFP, with preferred fluoropolymers being prepared from 95 to 100 wt% VF₂ and 0 to 5 wt% HFP. Such fluoropolymers are available in the form of

latices, typically having mean particle sizes in the range from 200 to 400 nanometers.

Commercially available fluoropolymer powders and latices include THV-500P fluoroterpolymer powder and KEL-F 3700 from Dyneon, of St. Paul, MN, 5 KYNAR PVDF homo- and copolymer powders, and KYNAR 32 PVDF latex from Elf Atochem.

The polymeric film-forming material may be combined with the core/shell particles while either material is in any physical form or during any stage of processing, for example when the core/shell particles exist within a latex, or as a 10 powder of core/shell particles or their agglomerates, etc. Alternatively, the core/shell particles may be added to a liquid or solid polymeric film-forming material, followed by extrusion.

The relative amounts of core/shell particles and polymeric film-forming material in a given composition can be any amounts of each that are found to be 15 useful, with amounts for any particular application depending on factors such as the desired application or product construction, e.g., the physical or mechanical properties desired of a composition or material prepared from the core/shell particles, the chemistries of the core and the polymeric shell, the chemistry of the polymeric film-forming material, the choice of processing methods used, etc.

20 An exemplary composition of the invention is an aqueous latex which contains core/shell particles and particles of polymeric film-forming material e.g., in the form of dispersed polymeric particles. In a preferred embodiment, a latex which contains core/shell particles can be combined with a fluoropolymer latex. It can be preferable for the latices to be miscible with one another, and to be suitably 25 stable in the form of a combined latex. "Miscible" with respect to latices means that in combining the latices, a dispersion is maintained and the combination of the two separate latices does not cause coagulation. (Coagulation of the latices can sometimes be prevented by pH adjustment prior to mixing or by adding one latex to another very slowly.)

30 Aqueous latices can have solids contents as known in the polymer and latex arts, and can be prepared, combined, and processed into mixtures or other compositions by known methods. The latex may be processed as desired, for

example dried by known methods to form a powder, or applied as a water-based coating to form a film upon drying. If applied as a coating and dried, the latex coating can be dried to remove water at one temperature, e.g., 66°C, followed by another heating step where the polymeric components are melted or fused together to form a continuous film or coating.

It will be appreciated by a skilled artisan that a latex composition comprising core/shell particles and polymeric film-forming material can include other ingredients or additives useful in the production or processing of latices and films. The chosen ingredients or additives can depend on the desired use of the latex, but can include foam control agents, coalescing aids, anti-fungal agents, rheology control materials such as thickeners, pigments, leveling agents, etc.

Preferred methods for drying a latex which contains core/shell particles (alone or with another material) include spray drying methods. During spray-drying, latex particles tend to fuse together to form larger agglomerate particles composed of the individual latex particles. A latex of core/shell particles and particles of a polymeric film-forming material, if spray-dried, will typically produce agglomerate particles composed of particles of the polymeric film-forming material fused with core/shell particles.

Agglomerate particles can preferably exist as a free-flowing powder having a combination of particle size, thermoplastic properties (e.g., melt viscosity), and thermal stability, which allow the particles to be processed into a desired product construction. Spray-drying can produce agglomerate particles having a range of sizes, and the size of the agglomerate particles can be any size capable of being processed into a useful product. After spray drying, agglomerate particles can be graded to obtain a desired, e.g., narrow, size distribution. An example of a useful size range for agglomerate particles can be in the range from about 10 to 200 micrometers (μm), more preferably from about 10 to 70 μm . Although particles outside of these size ranges may also be useful, particles smaller than 10 μm may present explosion hazards during powder coating, and particles larger than 200 μm may be difficult to electrostatically charge for powder coating, and, if powder coated, may produce an overly thick particle layer that may be difficult to fuse.

Preferred agglomerate particles are shown in figures 1 and 2. Figure 1 shows a number of agglomerate particles. The photomicrograph of figure 2 is a closeup of a single agglomerate particle from figure 1. These photomicrographs show agglomerate particles comprised of core/shell particles and fluoropolymer particles fused together. In figure 2, the fluoropolymer particles are the larger spherical particles, and the smaller spherical particles attached to and surrounding the larger fluoropolymer particles are the core/shell particles.

A skilled artisan will appreciate that a powder composition comprising the core/shell particles can additionally include, in understood amounts, other ingredients known to facilitate the production or processing of a powder composition, such as plasticizers, stabilizers, flow aids to improve handling and processability (e.g., silica), pigments, and extenders.

Another exemplary composition of the invention is a multiphase polymeric composition which contains a polymeric core phase, a polymeric shell phase, and a polymeric film-forming phase, e.g., in the form of a film or a coating, and wherein the core phase comprises a UVA. Preferably, the composition exhibits a multiphase morphology wherein core phase domains and polymeric film-forming material phase domains do not substantially contact one another, but are substantially or entirely separated from each other by polymeric shell phase domains, e.g., both polymeric film-forming material phase domains and core phase domains contact polymeric shell phase domains, but do not significantly contact each other. Of course in practice there may be some degree of contact between domains of the core particle phase and the polymeric film-forming material phase, e.g., due to imperfect processing of the core/shell particles or due to breakage during processing, but this type of inadvertent contact will typically be kept to insubstantial levels.

The core/shell particles used in this preferred morphology can be particularly useful for allowing incorporation of a core particle material into a polymeric film-forming material with which the core particle material (e.g., a core particle, core polymer, or a UVA contained in a core particle) is not compatible. The described morphology provides a polymeric composition that is relatively thermodynamically stable, because incompatible phases, e.g., the core phase and

the polymeric film-forming material phase, are separated by the polymeric shell phase; the incompatible phases do not contact each other, thus resulting in thermodynamic stability and reducing the tendency of the multiphase composition to change morphology over time.

5 In these compositions, the core phase, the polymeric shell phase, and the polymeric film-forming material phase can be any of those materials described above, and can be chosen to provide particular properties for a desired application. Preferred multiphase compositions can include ingredients having appropriate chemistries for a particular application, and in appropriate amounts, to provide
10 desired properties such as flexibility, weather resistance, thermodynamic stability, and adhesion to a substrate. In a preferred multiphase composition, the amounts of polymeric core phase, polymeric shell phase, and polymeric film-forming material phase can be sufficient to exhibit one or more of the following: flexibility, ultraviolet radiation absorption, dirt and water resistance, prevention of plasticizer
15 blooming or bleeding from a substrate, and minimization of blooming, bleeding, or other migration of UVA from the composition. Because these properties can be dependent on a number of factors, such as the particular chemistries of the different components, a broad range of amounts of the different ingredients may be useful. An exemplary range can be from about 80:20 to 10:90 (by weight)
20 combined polymeric core and polymeric shell phases, per polymeric film-forming material phase. Preferred amounts can be in the range from about 20:80 to 60:40 (combined polymeric core and shell phase: polymeric film-forming material phase).

 As one example, a multiphase polymeric composition may be used as a
25 topcoat to be applied to a flexible substrate exposed to outdoor conditions. Such a topcoat can include ingredients including polymeric core phase, polymeric shell phase, and polymeric film-forming material phase, chosen to provide properties such as weatherability, durability, dirt resistance, water resistance, flexibility, toughness, adhesion to a desired substrate, barrier properties (e.g., resistance to
30 plasticizers), compatibility between different phases, thermodynamic stability, etc. A preferred polymeric core phase for use in a topcoat can be of a rubbery crosslinked material which provides flexibility to the final construction, without

brittleness. A polymeric shell phase for a topcoat can be chosen, e.g., to provide chosen properties such as cleanability, and to enhance adhesion between the multiphase polymeric composition and a chosen substrate such as PVC. The polymeric film-forming material can be chosen to add other specifically desired properties, e.g., a fluoropolymer may be chosen to provide weatherability, dirt and water resistance, and cleanability. The polymeric shell phase can preferably be, but is not required to be, compatible with the film-forming polymeric material phase. Preferred polymeric shell materials that are compatible with many fluoropolymers can have a T_g below about 105C, with T_gs in the range from about 40C to 90C being particularly preferred.

A specific example of a preferred multiphase composition is a film comprising separate phase domains of a poly(ethyl acrylate) core phase, a poly(meth)acrylate shell phase, and fluoropolymer. Both the fluoropolymer and the core polymeric phases can be compatible with the polymeric shell phase, but the fluoropolymer phase may or may not be compatible with the polymeric core phase.

It has been found that such multiphase polymeric compositions comprising a polymeric core phase separated from a polymeric film-forming phase by a polymeric shell phase can exhibit lasting UVA protection due to reduced migration loss (e.g., bleeding or blooming) of the UVA; thermodynamic stability; and improved resistance to water (e.g., reduced water sensitivity), compared to chemical compositions containing the same amounts of chemically identical ingredients but exhibiting a different morphology. The improved properties are believed to be due, at least in part, to the fact that the UVA in the polymeric core phase is surrounded and contained by polymeric shell phase (and is optionally and preferably chemically bound to the core polymer). Further, separation of incompatible phases can improve thermodynamic stability, which leads to improved water resistance (e.g., less water sensitivity). See, e.g., Comparative Example 2, *infra*.

Multiphase polymeric compositions having the preferred morphology, optionally including a polymeric core phase that is incompatible with a polymeric film-forming material phase, can be prepared by any successful method. As an

example, the morphology can be achieved by providing a film or coating comprising core/shell particles and particles of the polymeric film-forming material, and processing the film or coating such that polymeric components of the composition fuse together or solidify into a preferably continuous, cohesive film or coating. The process can be practiced with care to maintain the core/shell structure, e.g., prevent dissolution of the core/shell particles, to achieve the preferred multiphase morphology.

More specifically, by one example of such a method, a multiphase composition having the preferred morphology can be achieved by providing a film or coating of a latex comprising core/shell particles and particles of the polymeric film-forming material, followed by drying to form a film.

The preferred morphology can also be achieved by powder coating particles comprising core/shell particles and particles of the polymeric film-forming material, (e.g., agglomerate particles obtained by spray-drying), followed by fusing. Fusing should take place at a temperature that is above the melting point of any crystalline polymeric phase of the powder, and above the Tg of any amorphous polymeric phase, a temperature sufficient to melt and thereby fuse the particles into a preferably continuous, multiphase polymeric film or coating, to an extent that the core phase does not come into substantial contact with the film-forming polymeric material phase.

Figure 5 schematically illustrates a method of coating a powder onto a flexible substrate. Substrate 10 moves through powder cloud 12 coming from electrostatic fluidized bed powder coater 14, so that a particle layer 16 is formed on a surface of the substrate. The particles of powder cloud 12 are shown much larger than actual size for the purpose of illustration. Substrate 10 may be in the form of a long continuous web (as shown), or a smaller piece of material laid on a carrier web. In a another technique (see for example "Powder Coating," edited by Nicholas P. Liberto, published by the Powder Coating Institute, Chapter 10 (1994)), powder cloud 12 can be generated by placing a powder in the chamber of the coater and passing ionized air through the powder until it fluidizes. Preferably, the powder is pre-dried in a conditioning chamber (not shown) before use. The substrate passing through powder cloud 12 can be electrostatically charged to

cause the powder to be attracted to the substrate. To facilitate deposition of a uniform coating, a charge can be applied to the substrate by known methods, e.g., by corona treatment methods or by contacting the web to a material which appears higher or lower on the triboelectric series, depending on the desired polarity of the charge. A grounding plate 17 made of aluminum or another like material can be placed behind the substrate to provide a ground potential to attract the charged powder to the surface of the substrate. The coating weight of particle layer 16 can be controlled by one or more of the line speed, the voltage applied to the air supply, and the particle size of the powder. Optionally, both surfaces of a substrate may be coated by passing the substrate between two powder coaters, or by making two passes over the same coater and inverting the substrate between passes.

Although electrostatic fluidized bed powder coating is a preferred method for continuous powder coating, other types of powder coating methods such as electrostatic spray coating may also be used. Powder coating equipment is well known and complete systems are readily available commercially. A nonlimiting example of a powder coating equipment manufacturer is Electrostatic Technology Incorporated (ETI), Branford CT, USA.

Again referring to Figure 5, to fuse the coated powder into a continuous film or coating, the coated substrate passes through a nip configuration defined by heated roll 20 and backup roll 18. The nip configuration applies heat and pressure simultaneously to fuse the powder particles of particle layer 16 into a continuous, multiphase polymeric layer 22, and bond the layer to substrate 10, thereby forming a composite sheet material 30. No preheating stage is required prior to the nip, but such a stage may be useful if desired, e.g., to achieve a higher line speed. Heated roll 20 is typically made of metal, and its outer surface is preferably covered with a material having release properties such as poly(tetrafluoroethylene) to prevent transfer of either melted thermoplastic powder or fused thermoplastic layer from the substrate to the roll. Backup roll 18 preferably has a resilient surface, such as rubber.

The temperature and pressure of the heated roll can be chosen to be sufficient enough to fuse the coated powder into a continuous layer, yet not so high as to distort or degrade the substrate. If substrate 10 is a thermoplastic film or

other material likely to soften or distort at the elevated temperatures in the nip, support can be provided to the substrate in the form of a carrier web, liner, or belt system (not shown). The backup roll may be at ambient temperature, it may be heated to facilitate fusing, or it may be chilled to provide further thermal protection for the substrate.

As an alternative to the nip configuration shown in Figure 5, simultaneous heat and pressure may be applied by other suitable means, such as a heated press, or by a belt configuration.

While the above-detailed processes can be used to prepare a multilayer polymeric composition having the described preferred morphology, this morphology may not be achieved by just any coating or film-forming process or method. For instance, the described preferred morphology may not necessarily result from the same composition (i.e., identical chemical ingredients in identical amounts) being prepared into a film, if prepared by other processing methods. If, as described, a latex of the core/shell particles is blended with a latex of particles comprising the polymeric film-forming material, followed by spray drying to produce agglomerate particles, the resulting agglomerate particles can be powder coated and fused to achieve a composition with the desired morphology. If, on the other hand, the latices are spray dried separately, and then blended into a powder mixture and powder coated, the desired morphology may not necessarily be achieved, but a different morphology might be achieved which although possibly useful, may not exhibit the same physical properties of the described preferred morphology. Furthermore, if the core/shell particles become dissolved or are otherwise caused to lose their core/shell structure during processing, e.g., by mixture with a solvent, sufficient heating for an extended period of time, or by exposure to any other processing condition that would cause the loss of the core/shell structure, a composition prepared by such a method may not exhibit the described preferred morphology or resultant properties. In particular, if the core/shell particles are solvent coated, it is possible that the desired morphology may not be achieved. See Comparative Example 3.

The core/shell particles can be used to prepare various products having UV radiation absorption or blocking properties. An example is composite sheet

material 33, shown in figure 6. Multiphase polymeric layer 22, (e.g., as a topcoat), in the form of a continuous film prepared from or containing polymeric core/shell particles, is bonded to substrate 10. The multiphase polymeric layer can be transparent or translucent in appearance, and can generally have a thickness in the range from about 12.7 μm to about 254 μm (0.5 mil to 10 mils). An example of a protective coating for an outdoor sign substrate (topcoat) can be translucent and can have a thickness in the range from 10 μm to 25.4 μm (0.4 mil to 1 mil).

Substrates that can be suitable include most any material that can benefit from protection from ultraviolet radiation, such as plastics, or materials that might otherwise be coated with a polymeric material, wherein the polymeric material can be UV-radiation resistant. Preferred substrates can be flexible two-dimensional materials capable of receiving a powder coating and capable of withstanding heat and pressure conditions used to process the powder into a coating or film. The substrate can be used in conjunction with a supporting liner, or can be internally reinforced in order to meet process requirements. The thickness of the substrate can be any useful thickness, and can preferably be in the range from about 12.7 to about 1270 μm (0.5 to 50 mil), more preferably from about 50.8 to about 762 μm (2 to 30 mil).

Preferred substrates include flexible thermoplastic films such as those made from polyester, polyamide, polyimide, polycarbonate, plasticized polymers such as PVC, polyacrylates such as polymethyl methacrylate, polypropylene, polyethylene, and copolymers or combinations thereof. The thermoplastic film may optionally contain additional components such as pigments, fillers, reinforcing fibers or mesh, and ultraviolet absorbing agents, to provide properties important for some applications. An example of a suitable substrate for use in outdoor backlit signs and awnings is a polyester mesh scrim that has been coated on both sides with a plastisol-containing PVC resin, a phthalate ester plasticizer, and pigment, and then fused. Such a substrate is commercially available under the trade name PANAFLEX, from 3M of St. Paul, MN.

The powder may also be coated onto a release liner and optionally fused into a film. The coat liner could then be used as a casting liner for another film or

coating such as a PVC plastisol or organosol. Optionally, the pre-fused film could be laminated to a previously fused organosol or plastisol film.

The substrate surface may optionally be treated to increase adhesion between a topcoat and a substrate, e.g., by etching or other treatment such as a non-polymeric, chemical agent applied to chemically modify the surface of the substrate to allow the topcoat to bond, or by application of an adhesion-promoting primer or tie-layer. Such treatment is not required, and is preferably avoided if adhesion between the substrate and the topcoat is otherwise adequate. These and other embodiments and applications of the compositions of the invention will be apparent to a skilled artisan. See also Assignee's copending United States Patent Application, entitled Multi-layer Articles Including UV-Absorbing Polymeric Compositions, having U.S. Serial No. 09/371,060, Attorney's Docket No. 54645USA4A, which was filed on even date herewith.

Examples

Example 1 – Preparation of a polymeric UV absorber (UVA) having a core/shell structure

This example details preparation of a latex particle having a 30/70 core/shell ratio with the core composition being 17/83 Norbloc™ 7966/ ethyl acrylate and the shell composition being 80/20 methyl methacrylate/ethyl acrylate. Water (331.4g), sodium lauryl sulfate (1.5g), isooctyl thioglycolate (0.45g), and a mixture (premix) of ethyl acrylate (37.5g) and Norbloc™ 7966 (7.5g) a polymerizable UV absorber (available from Janssen Pharmaceutica, Titusville, NJ) were charged to a reaction flask, stirred and purged with nitrogen while heating to 85-90°C. When the system was well dispersed, the batch was cooled to 75°C and a premix of potassium persulfate initiator (0.45g) in water (17.55g) was added. After an initial induction period, the polymerization started and the batch temperature rose to about 80°C. After the peak temperature had been reached (about 5 to 10 minutes), the shell monomer premix consisting of methyl methacrylate (84g), ethyl acrylate (21g) and isooctyl thioglycolate (0.53g) was added over about 90 minutes while the batch temperature was held at 80°C. When addition was complete, the

batch was held at 80°C for 90 minutes then cooled and filtered through cheesecloth.

The product had inherent viscosity of 0.36 deciliter per gram (dl/g) measured in tetrahydrofuran solvent. Analysis of the dried polymer by differential scanning calorimetry showed two separate glass transitions at -6°C and 70°C.

Example 2 – Preparation of a polymeric UV absorber (UVA) having a core/shell structure and a crosslinked core.

A latex with 20/80 core/shell ratio was prepared by the procedure of Example 1 above, modified as follows. The core composition was prepared using 0.45 g allyl methacrylate, 5.01g NORBLOC™ 7966, 24.54g ethyl acrylate, and 0.20g isooctyl thioglycolate. The shell monomers included 102g methyl methacrylate, 18g ethyl acrylate and 0.78g isooctyl thioglycolate. The shell monomer mixture was added over a period of 90 minutes. The product had IV = 0.30 dl/g and a single glass transition temperature at 82°C. The addition of allyl methacrylate to the core resulted in crosslinking of the core and disappearance of the lower Tg that was noted in Example 1.

Examples 3 and 4 – Preparation of Core/Shell Latices with Alternative Polymerizable UV Absorbers

Latices with a 30/70 core/shell ratio were prepared by the procedure outlined in Example 1, modified as follows. The core compositions for these examples were prepared from polymerizable UVA and ethyl acrylate in a weight ratio of 16.7/83.3, and the shell polymer was prepared from methyl methacrylate/ethyl acrylate in a weight ratio of 80/20. The shell monomer mixture for both Example 3 and 4 contained 84g methyl methacrylate, 21g ethyl acrylate and 0.53g isooctyl thioglycolate.

The core monomer mixture of Example 3 contained 37.5g ethyl acrylate, 7.5g CGL 104, a polymerizable benzotriazole available from Ciba Specialty Chemicals, and 0.45g isooctyl thioglycolate. The product had IV = 0.38 dl/g and showed two glass transition temperatures at -2°C and 73°C.

In example 4, Cyasorb™ 416, a polymerizable benzophenone UV absorber available from Cytec, was substituted for the CGL 104 used in Example 3. The product had IV = 0.18 dl/g and showed two glass transition temperatures at 1°C and 73°C.

5

Example 5 – Non Polymerizable UVA in core/shell particle having a crosslinked core.

A latex with a crosslinked core containing a non polymerizable UVA was prepared using the procedure of Example 2 except Tinuvin™ 1130, a non-polymerizable benzotriazole UV absorber available from Ciba Specialty Chemicals was substituted for NORBLOC™ 7966. The product had IV = 0.30 dl/g and a single Tg at 74°C.

15 **Comparative Example 1 – Preparation of a polymeric UV absorber without a core/shell structure**

Homogeneous latex particles were prepared, having a composition of 80/15/5 methyl methacrylate/ethyl acrylate/ Norbloc™ 7966. These particles were prepared using a two step (core/shell) process wherein the composition of the core and the shell were identical. The procedure used was similar to that used in Example 1 except that the core monomer mixture consisted of 36g methyl methacrylate, 6.75g ethyl acrylate, 2.25g NORBLOC™ 7966, and 0.45g isooctyl thioglycolate, and the shell monomer contained 84g methyl methacrylate, 15.75g ethyl acrylate, 5.25g Norbloc™ 7966 and 0.53g isooctyl thioglycolate. The product had IV of 0.31 dl/g and a single glass transition temperature at 87°C.

Preparation of Powder Coating Compositions

Example 6

30 A latex was prepared by blending the core/shell UVA latex of Example 1 with a fluoropolymer latex consisting of Hylar™ FXH-6, a 95/5 vinylidene fluoride/hexafluoropropylene copolymer from Ausimont USA, in ratios of 80% fluoropolymer solids to 20% acrylic solids. The resulting latex mixture was then spray dried to yield agglomerate powder particles in which the two polymeric components were blended and fused on submicron scale. (See Figures 1 and 2).

35

Spray drying was accomplished under the following conditions on a Büchi B-191 Mini Spray Dryer (available from Brinkman Instruments, Westbury, N.Y):

5	Inlet Temperature: 140°C	Outlet Temperature: 40-55°C
	Pressurized air flow: 550 liters/hr.	Pump Speed: 100%
	Aspirator: 100%	Vacuum 40 - 50 mbar

Comparative Example 2

A similar powder coating composition was prepared using the
10 homogeneous latex particles prepared in Comparative Example 1 and the fluoropolymer described in Example 6.

Preparation of Powder Coated Articles

The powder coating compositions prepared as described in Example 6 and
15 Comparative Example 2 above were coated onto 10.1 cm x 15.2 cm pieces of PANAFLEX™ 930 vinyl outdoor sign substrate available from 3M Company, St. Paul, MN. A coating weight of 33grams per square meter of the 80/20 fluoropolymer/acrylic powder blend was applied to the substrate using a laboratory-scale C-30 electrostatic fluidized bed powder coater (Electrostatic
20 Technology, Inc., Bradford, CT). The voltage in the coater was set to give the desired coating weight (settings varied with the powder used), and the substrate was held over the coater for approximately 5 seconds. The coated substrate was then hand fed through a nip configuration to fuse the particle layer. The nip configuration consisted of a TEFLON-coated roll heated to 200°C and an unheated
25 backup roll. A piece of aluminum foil was threaded over the heated roll. The coated substrate was placed powder side down on the foil and fed through the nip at an applied air pressure of about 276 kPa (40 psi) and a line speed of 1 meter per minute. The resulting fused protective layer had a thickness of approximately 25 µm.

30 The same compositions were coated onto a TELFON-coated release liner and removed as a free film, which had the following UV-absorbing properties.

The ultraviolet spectra of the films were essentially identical to those of monomeric or polymeric benzotriazole UV absorbers and showed absorption maxima at 240, 298 and 336 nm due to the benzotriazole chromophore.

Example 7 – Preparation of Cast Films Using a Core/Shell Latex

Latices were prepared by blending the core/shell UVA latex of Example 1 with a fluoropolymer emulsion of HYLAR™ FXH-6, a 95/5 vinylidene fluoride/hexafluoropropylene copolymer from Ausimont USA, in ratios of 80% fluoropolymer solids to 20% acrylic solids. This latex blend was bar-coated onto a hydrophylically-primed polyester liner at a wet thickness of 100 microns. The coating was then dried in a 67°C oven for 3 minutes before transfer to a 180°C oven for 1 minute. The resultant continuous, dry film was 10-15 microns thick.

10

Comparative Example 3 – Preparation of Solvent Cast Films

The powder coating composition prepared according to Example 6 was dissolved in acetone (44°C) in an amount sufficient to prepare a 10% solids solution. The resultant solution was bar-coated onto a polyester liner at a wet thickness of ~100 microns. The coating was then heated for 1 minute at 190°C to yield a dry film having a thickness of approximately 10-15 microns.

15

Example 8 - Resistance of Coated Films to Hot Water

The water resistance of fused films was evaluated by immersion of fused film samples into deionized water at 66°C for 16 hours. This test simulates long term exposure to hot and humid conditions. The haze or clarity of fused films was measured before and after immersion using a Gardner PG 5500 Digital Photometric Unit and ASTM D1003 as a standard test method.

20

Samples were prepared by powder coating the blended test materials prepared according to Example 6 and Comparative Example 2 onto a TEFLON-coated flexible substrate and then fusing the coatings using the methods described previously (see Preparation of Powder Coated Articles). The fused films (25 µm thick) were then removed from the TEFLON-coated substrate and evaluated for water resistance. The results for the Example 6 composition containing core/shell UVA particles from (Example 1) and, the Comparative Example 2 composition, containing the homogeneous UVA-containing particles (from Comparative Example 1) are shown in the Table 1 below. It is worth noting that the absolute

25
30

haze is not particularly relevant, because a film may be useful with various levels of haze or translucence; this test instead measures stability by tracking a change, e.g., increase, in haze.

5

TABLE 1

Powder Coating Composition	Haze Before Immersion	Haze After Immersion	Difference
Example 6	25.6%	28.1%	2.5%
Comp. Ex. 2	22.0%	51.6%	29.6%

The results in Table 1 indicate that the film containing the core/shell UVA (Ex. 6) exhibited much less change in haze caused by the immersion challenge. This indicates that when powder coated and fused, the core/shell particle
10 containing a UV absorber in the core, can be included in a fluoropolymer to produce a more thermodynamically stable multiphase composition, than homogeneous UV absorbers.

Additionally, the core/shell particles showed improved flexibility in qualitative comparison to the homogeneous particles in coated films made with the
15 fluoropolymer/acrylic blends. Folding (using finger pressure) the coated substrate 180 degrees to form a sharp crease tested the coating flexibility. The substrates were creased with the coating on the inside to test the coating versus compressive forces and with the coating on the outside to test versus extensional forces. The film containing the homogeneous acrylic cracked under both compression and
20 extension, the film containing the core/shell did not crack under either stress.

The water resistance of the latex cast films (Example 7) and the solvent cast films (Comparative Example 3) are shown in Table 2 below.

TABLE 2

Composition	Haze Before Immersion	Haze After Immersion	Difference
Example 7	1.1%	11.2%	10.1%
Comp. Ex. 3	1.5%	13.5%	12.0%

Claims:

1. A chemical composition comprising:
a polymeric particle having a core/shell structure, the core comprising an
ultraviolet absorber; and
5 fluoropolymer.
2. The composition of claim 1 wherein the core is incompatible with
the fluoropolymer.
- 10 3. The composition of claim 1 wherein the core comprises a polymer
derived from monomers comprising ethyl acrylate.
4. The composition of claim 3 wherein the core polymer is derived
from monomers further comprising a (meth)acrylate-functional ultraviolet
15 absorber.
5. The composition of claim 3 wherein the core comprises a polymer
prepared from monomers consisting essentially of ethyl acrylate, (meth)acrylate-
functional ultraviolet absorber, and crosslinker.
20
6. The composition of claim 2 wherein the shell comprises a
polymeric shell that is compatible with the fluoropolymer.
7. The composition of claim 2 wherein the polymeric shell comprises
25 a poly(meth)acrylate.
8. The composition of claim 7 wherein the polymeric shell is prepared
from monomers chosen from the group consisting of methyl acrylate, methyl
methacrylate, ethyl acrylate, ethyl methacrylate, and mixtures thereof.

30

9. The composition of claim 1 wherein the fluoropolymer is prepared from fluorinated monomers comprising vinylidene fluoride, hexafluoropropene, and mixtures thereof.
- 5 10. The composition of claim 1 wherein the composition is an aqueous latex.
11. The composition of claim 1 wherein the composition is a powder.
- 10 12. The composition of claim 11 wherein the composition comprises agglomerate particles comprising core/shell particles and fluoropolymer particles fused together.
13. The composition of claim 1 wherein the core/shell particles
15 comprise from about 10 to 50 parts by weight core particle, and from about 50 to 90 parts by weight polymeric shell.
14. The composition of claim 13 wherein the composition comprises from about 10 to 80 parts by weight core/shell particles and from about 20 to 90
20 parts by weight fluoropolymer.
15. A multiphase polymeric composition comprising a polymeric core phase, a polymeric shell phase, and a fluoropolymer phase,
wherein the polymeric core phase comprises an ultraviolet absorber, and
25 wherein the polymeric core phase and the fluoropolymer phase do not substantially contact one another, but both the polymeric core phase and the fluoropolymer phase contact the polymeric shell phase.
16. The composition of claim 15 wherein domains of the polymeric
30 shell phase surround domains of the polymeric core phase.

17. The composition of claim 15 wherein the polymeric core phase is incompatible with the fluoropolymer phase, and wherein the polymeric shell phase is compatible with both the polymeric core phase and the fluoropolymer phase.

5 18. The composition of claim 17 wherein the composition is thermodynamically stable.

19. The composition of claim 15 wherein the polymeric core phase comprises a core polymer derived from monomers comprising ethyl acrylate.
10

20. The composition of claim 19 wherein the core phase is a polymer prepared from monomers consisting essentially of ethyl acrylate and a (meth)acrylate-functional ultraviolet absorber.

15 21. The composition of claim 15 wherein the composition is a flexible film.

22. The composition of claim 15 wherein the polymeric shell phase comprises a shell polymer prepared from monomers chosen from the group
20 consisting of methyl acrylate, methyl methacrylate, ethyl acrylate, ethyl methacrylate, and mixtures thereof.

23. The composition of claim 15 wherein the fluoropolymer is prepared from fluorinated monomers comprising vinylidene fluoride, hexafluoropropene,
25 and mixtures thereof.

24. A method of preparing a multiphase polymeric composition comprising a polymeric core phase, a polymeric shell phase, and a polymeric film-forming phase, wherein the polymeric core phase comprises an ultraviolet
30 absorber, the composition being prepared according to steps comprising:
powder coating onto a substrate polymeric materials comprising the polymeric core phase, the polymeric shell phase, and the polymeric film-forming

phase, and fusing the polymeric materials to form a multiphase polymeric composition wherein the polymeric core phase and the polymeric film-forming phase do not substantially contact one another, but both the polymeric core phase and the polymeric film-forming phase contact the polymeric shell phase.

5

25. The method of claim 24 wherein the polymeric materials include core/shell particles comprising a core particle surrounded by a polymeric shell.

26. The method of claim 25 wherein the polymeric core phase
10 comprises a core polymer prepared from monomers comprising ethyl acrylate.

27. The method of claim 25 wherein the polymeric materials are coated as a powder comprising agglomerate particles of core/shell particles and particles of the polymeric film-forming material.

15

28. The method of claim 27 wherein the agglomerate particles are prepared by spray drying a latex comprising core/shell particles and particles of polymeric film-forming material.

29. The method of claim 24 wherein the polymeric film-forming
20 material is a fluoropolymer.

30. The method of claim 24 wherein the multiphase polymeric composition is thermodynamically stable.

25

31. The method of claim 24 wherein the multiphase polymeric composition is flexible.

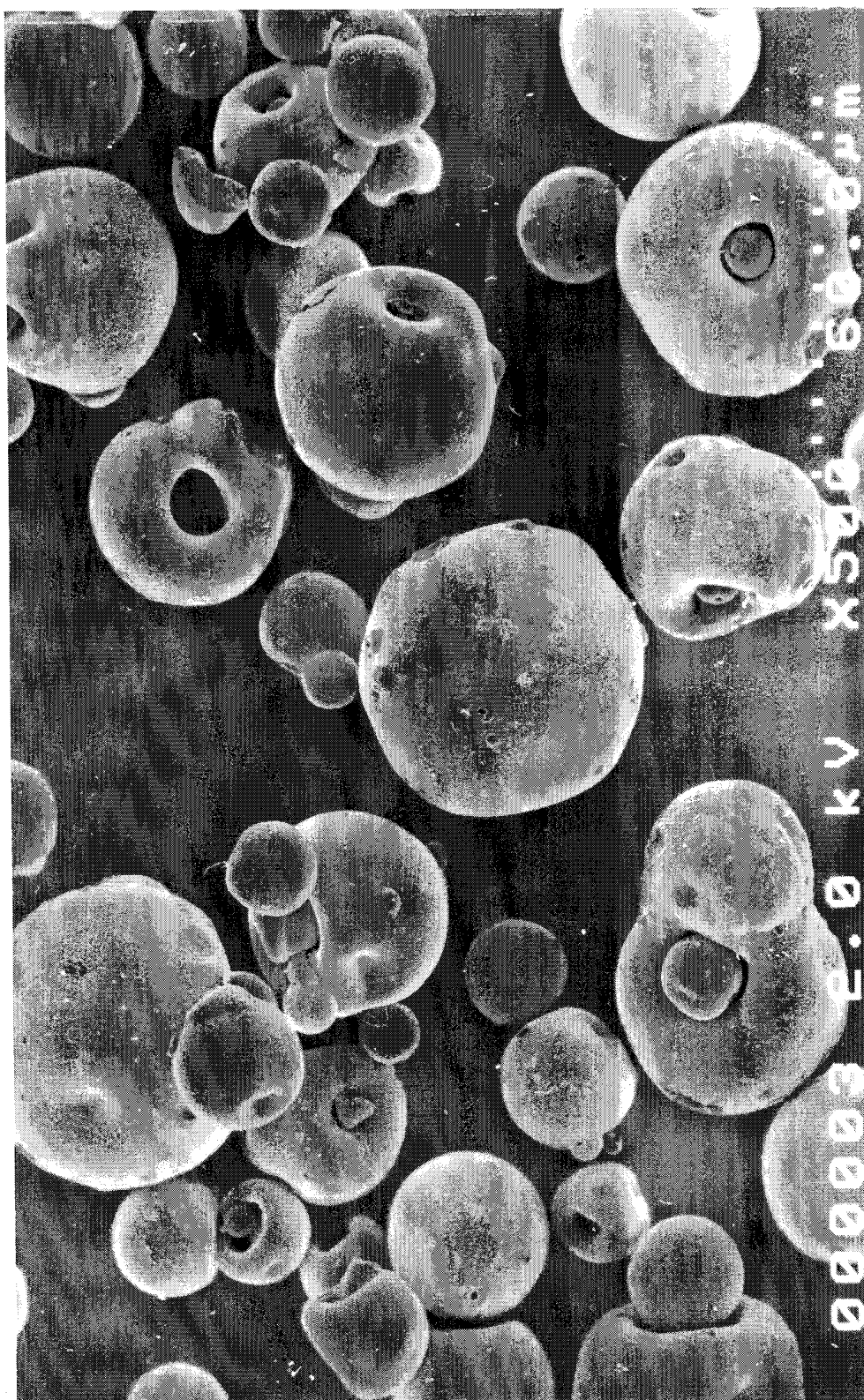


FIG. 1

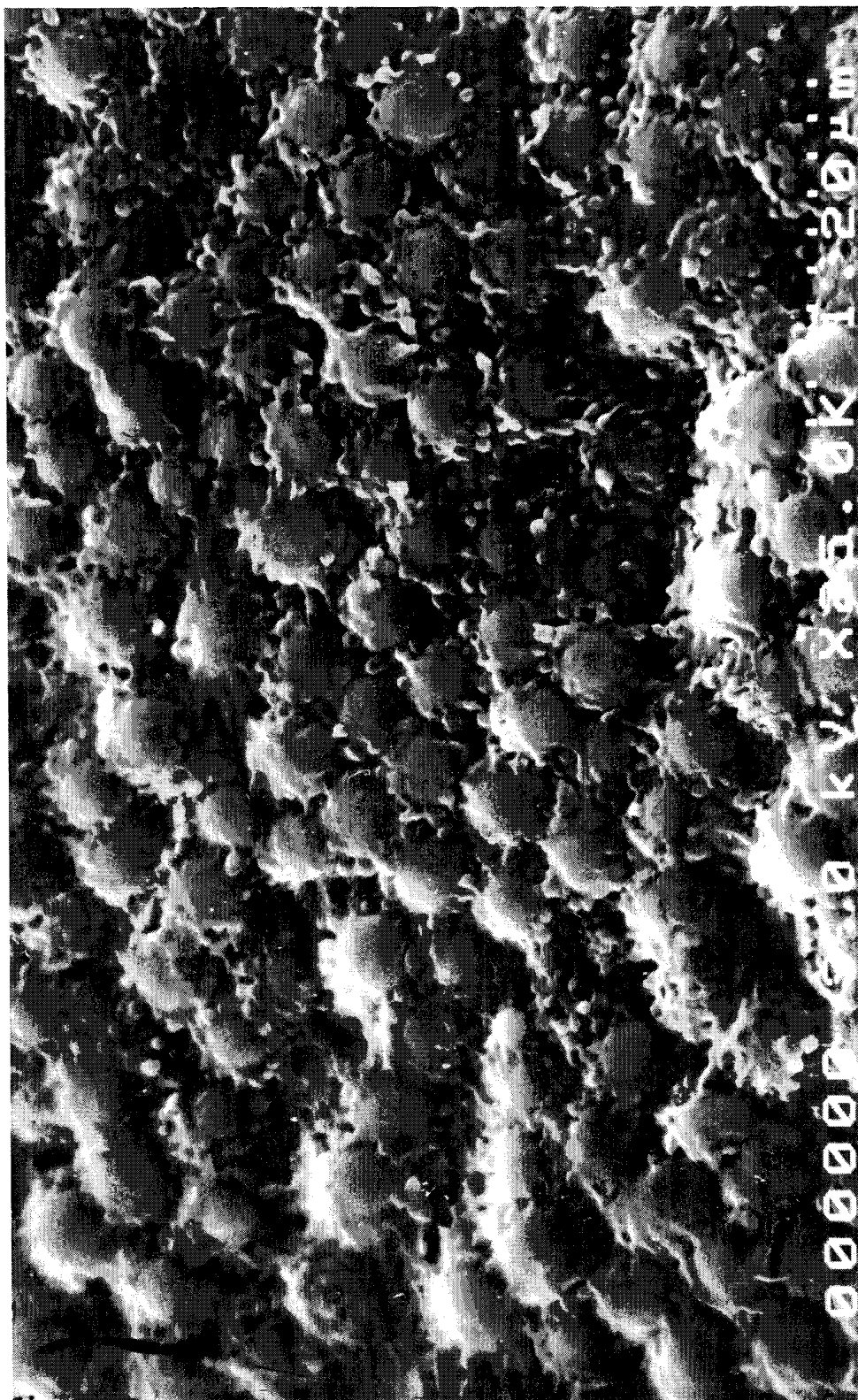


FIG. 2

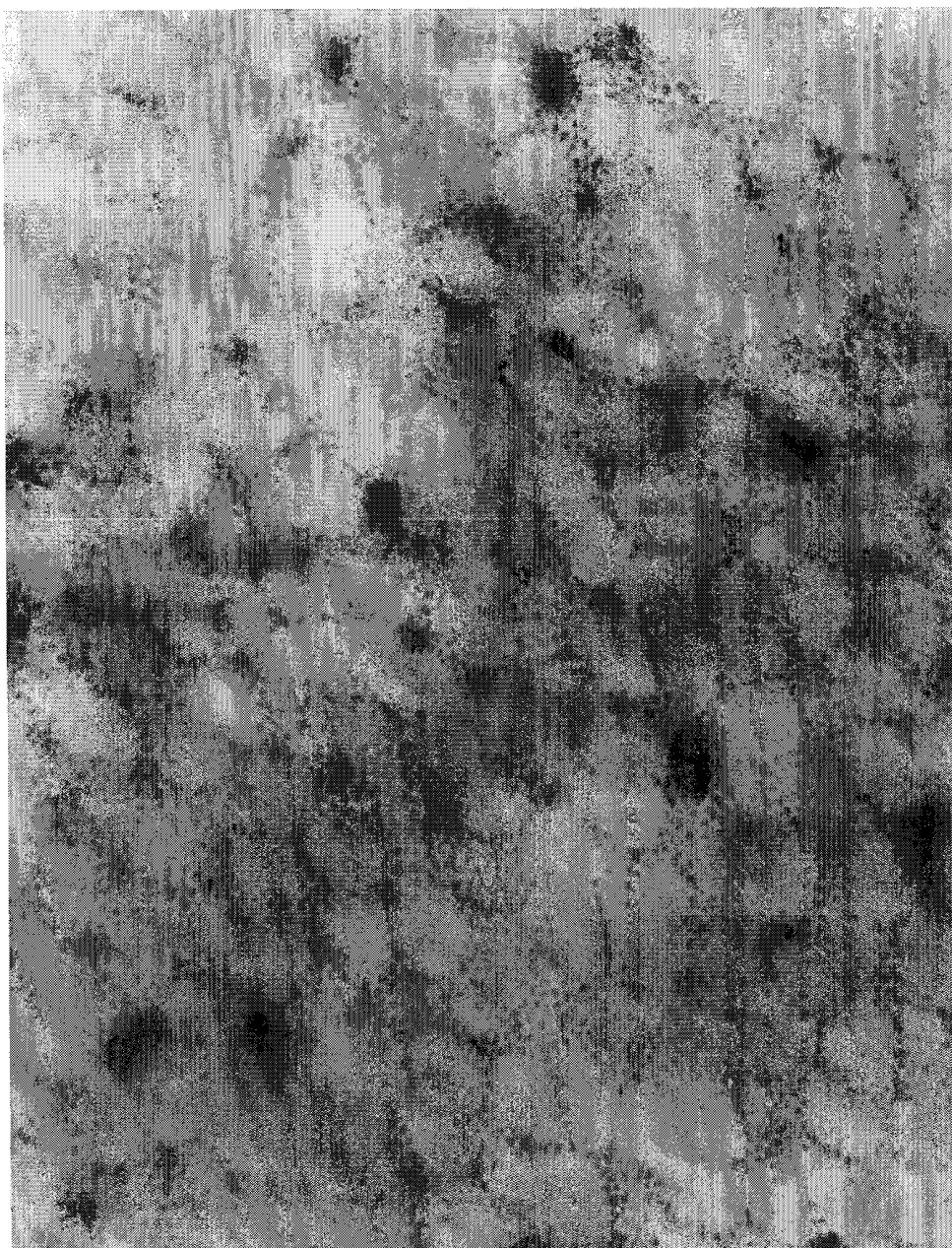


FIG. 3

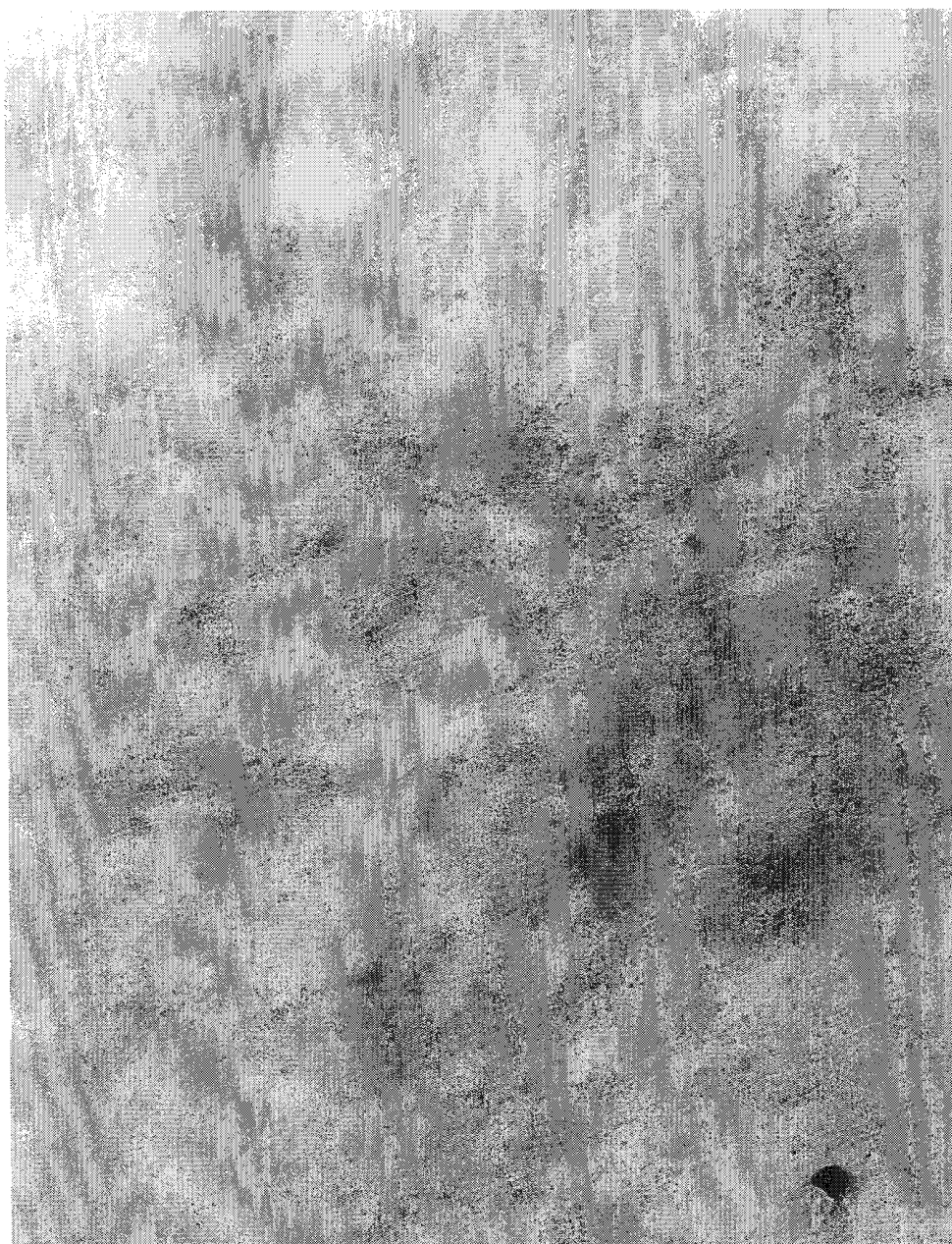


FIG. 4

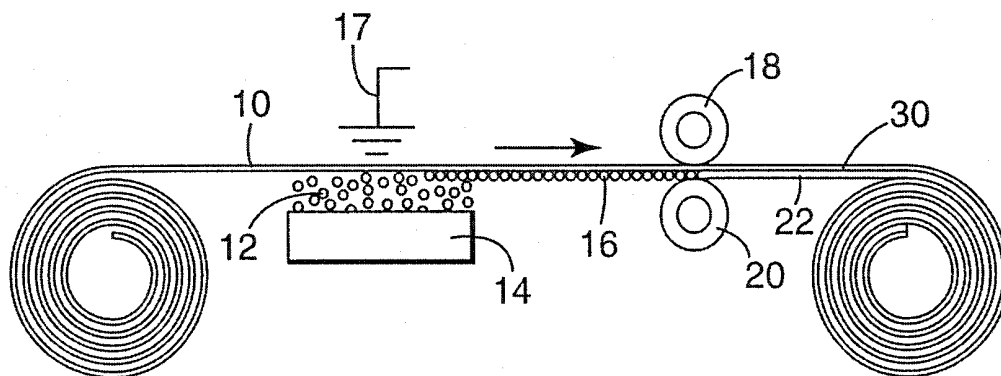


Fig. 5

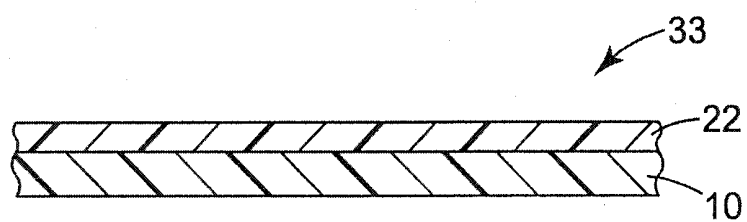


Fig. 6

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/27506

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C08J3/22 C08L27/12 //C08F265/06

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 C08J C08L

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)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 4 413 095 A (MIZUNO TOSHIYA ET AL) 1 November 1983 (1983-11-01) claim 1 column 1, line 42 - line 59 column 4, line 27 - line 40 ---	1,4,11, 15,24
A	DATABASE WPI Derwent Publications Ltd., London, GB; AN 1982-07046E XP002137973 & JP 56 163140 A (KUREHA CHEM IND CO LTD), 15 December 1981 (1981-12-15) abstract --- -/--	1,2,4,7



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents :

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

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"&" document member of the same patent family

Date of the actual completion of the international search

18 May 2000

Date of mailing of the international search report

29/05/2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Hallemesch, A

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/27506

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>DATABASE WPI Derwent Publications Ltd., London, GB; AN 1996-056019 XP002137974 & JP 07 316242 A (TAISEI KAKO CO), 5 December 1995 (1995-12-05) abstract</p> <p style="text-align: center;">---</p>	1
A	<p>US 5 587 424 A (LANGSTEIN GERHARD ET AL) 24 December 1996 (1996-12-24) claim 1</p> <p style="text-align: center;">-----</p>	1

INTERNATIONAL SEARCH REPORT

Information on patent family members

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