

Description

Background of the invention

This invention relates to polymeric compositions and processes particularly useful in surface coating compositions, and more particularly pertains to an aqueous blend of certain reactive self-curing water dispersed polymers, water-dispersed epoxy-acrylic copolymers, and optionally containing a polymeric phosphate additive, to provide water-based can coating compositions for interior coatings for food and beverage containers and similar sanitary coating applications.

Water-based coating compositions for use as internal sanitary liners for metal containers are suggested in U.S. 3,991,216. Such polymers are based on interpolymers of copolymerized acrylamide monomer, carboxylic monomer, and other ethylenically unsaturated monomers. However, such polymers are difficult to spray and often exhibit deficient film properties such as lack of resistance to ethanol and hence are undesirable for containers for alcoholic beverages.

Epoxy resins are particularly desirable for use in surface coating materials as a vehicle or polymeric binder for pigments, fillers, and other additives whereby epoxy resins advantageously provide toughness, flexibility, adhesion, and chemical resistance and water-dispersed coating compositions containing epoxy resins are highly desirable for can coating compositions. Coatings for soft drink and beer cans, for instance, are critical due to taste sensitivity wherein such sanitary can coatings must not alter the product taste of canned beverages. Taste problems can occur in a variety of ways such as by leaching of coating components into the beverage, or by adsorption of flavor by the coating, or sometimes by chemical reaction, or by some combination thereof. In commonly assigned U.S. 4,212,781, a process is disclosed for modifying epoxy resin by reacting the epoxy resin with addition polymerizable monomer in the presence of at least 3% by weight of benzoyl peroxide (or the free radical initiating equivalent thereof) based on monomer at a suitable reaction temperature. This reaction produces a reaction mixture containing an in-situ formed blend of resinous materials comprising an epoxy-acrylic copolymer mixture containing epoxy resin, graft epoxy-acrylic polymer, and associatively-formed ungrafted addition polymer. The in-situ polymerized monomers include acid functional monomers to provide acid functionality in the reaction mixture sufficiently high to effect stable dispersion of the resulting reaction product in a basic aqueous medium. In a preferred embodiment of U.S. 4,212,781, a polyglycidyl ether of bisphenol-A is reacted with a mixture of addition polymerizable monomers containing an acrylic acid such as methacrylic acid. The epoxy resin has a molecular weight above 4,000 and provides from 50% to 90% of the initial reaction mixture. The reaction takes place in the presence of benzoyl peroxide at an elevated temperature above 80°C and preferably between about 110°C and 130°C to effect addition polymerization of the monomer and product addition polymer grafted to the epoxy resin. The reaction product can be dispersed in a basic aqueous medium to form a water-reducible epoxy-acrylic copolymer mixture.

United States Patent 4,289,674 describes a solution technique for copolymerizing an alpha beta ethylenically unsaturated carboxylic acid such as acrylic and methacrylic acids, with N-ethoxymethylacrylamide or N-ethoxymethylmethacrylamide and optionally a further alpha beta ethylenically unsaturated monomer such as styrene and methacrylic acid esters. Such copolymers may be used with epoxy resins to form coating compositions.

U.S. Patent 4,289,674 relates to the production of base-solubilized dispersions of acrylic polymers in combination with epoxy resins. The acrylic polymer is one formed by reaction of alpha, beta-unsaturated carboxylic acid with N-ethoxymethylacrylamide or methacrylamide and optionally a further ethylenically-unsaturated monomer. These are apparently components which correspond to the components of the self-curing water dispersed polymer of the patentee's claims. However, the method of production used in U.S. Patent 4,289,674 is a conventional solution polymerization technique in which all of the components are premixed to form a monomer charge which is fed into a reactor. This contrasts with the patentee's invention in which the alkylol acrylamide is employed in a second stage of an emulsion polymerization, this resulting in the advantages in the product noted above. The patentee's invention as claimed does therefore represent a patentable advance over U.S. Patent 4,289,674 in view of the improved properties of the product.

In the light of this, it has been appreciated that the real invention in the present case lies in the fact that by using a two-stage polymerization process for the production of the copolymer of the alkylol acrylamide, the functional monomer and the "non-functional monomer" one can obtain a water dispersed polymer having surprisingly good rheological properties such as viscosity, stability and spray application apparently as a result of the two-stage polymerization forming a relatively rigid or hard polymer particle surface which stabilizes the viscosity of the water dispersed blends.

It also has been found that certain reactive self-curing water dispersed emulsion copolymers blended with epoxy-acrylic copolymer and polymeric phosphate provides an excellent coating suitable for an interior coating for beverage and food containers. The reactive self-curing water dispersed polymer contains emulsion copolymerized monomers including functional carboxyl, hydroxyl, amine or amide monomers preferably in combination with alkylol acrylamide monomers. The monomers can be copolymerized together in a step-wise reaction to concentrate the alkylol acrylamide on the surface of the polymer particles which advantageously provides an improved water dispersed polymer exhibiting surprisingly good rheological properties including viscosity, stability, and spray application. By contacting

alkylol acrylamide on the polymer surface, it is believed that a minor amount of alkylol acrylamide reacts with a minor amount of functional monomer during the addition polymerization of ethylenic monomers to provide a relatively rigid or hard polymer particle surface, which apparently stabilizes the viscosity of the water dispersed blend as well as provide considerable shear resistance during subsequent spray application of the blend. Upon ultimately heat curing of the composition of this invention, the water dispersed polymer becomes self-curing by the alkylol acrylamide reacting with the functional monomer groups in the polymer. By blending the reactive self-curing water dispersed polymer with a water-dispersed epoxy-acrylic copolymer, the blend provides an excellent sprayable interior coating particularly suitable for beverage cans. A preferred composition in accordance with this invention further contains a phosphated polymer such as an epoxy phosphate polymer. The inclusion of phosphated polymer provides improved coating characteristics such as solvent resistance and improved porosity properties. The reactive self-curing water dispersed polymers are high molecular weight polymers and advantageously provide good film integrity properties, high solids content, good spray application, and minimal use of solvents. These and other advantages of this invention will become more apparent by referring to the detailed description of the invention and the illustrative examples.

Summary of the invention

Briefly, the invention is based on a certain self-curing water dispersed polymer produced by a two stage emulsion polymerizing ethylenic monomers, including functional carboxyl, hydroxyl, amine or amide monomers in combination with alkylol acrylamide monomers and other ethylenically unsaturated monomers not containing functional groups, and preferably thereafter blend the water dispersed polymer with an epoxy-acrylic copolymer and optionally a phosphated polymer. The composition of this invention comprises a water-dispersed polymeric blend of (a) the reactive self-curing water dispersed polymer, and (b) an epoxy-acrylic copolymer. The preferred composition further contains (c) a phosphated polymer. An aminoplast cross-linking resin can be added to the water-dispersed polymeric blend to provide improved curing properties.

Detailed description of the invention

In accordance with this invention, the aqueous composition comprises certain reactive self-curing water dispersed polymers blended with epoxy-acrylic copolymer, phosphated polymer, and an aminoplast cross-linking resin.

Referring first to the reactive self-curing water dispersed polymer, the reactive self-curing polymer is a polymer of copolymerized ethylenically unsaturated monomers comprising (i) alkylol acrylamide and preferably alkylated alkylol acrylamide monomers, (ii) functional monomers consisting of reactive carboxy, hydroxyl, amine, or amide monomers, and (iii) other ethylenically unsaturated monomers to produce a self-reactive alkylol acrylamide water dispersed polymer. The self-curing water dispersed polymer is synthesized by step-wise polymerization of monomers in water whereby the alkylol acrylamide monomers are polymerized in the second polymerization step. The two stage monomers can be proportioned on a weight basis from 25/75 to 75/25 first stage/second stage monomers.

The alkylated alkylol acrylamide monomers can be derivatives of acrylamide, methacrylamide, methylol acrylamide, or similar alkyl modified acrylamide monomer as shown for example in U.S. 3,991,216; U.S. 4,097,438; and U.S. 4,305,859. The acrylamide monomers preferably are alkylated with an alkyl group such as methyl, ethyl, propyl, n-butyl, or iso-butyl, and similar alkylated alkylol acrylamide monomers, wherein the butylated monomers are preferred. Functional monomers, include carboxyl, hydroxyl, amino, and amido functional group containing monomers. Carboxy containing monomers include acrylic acid and lower alkyl substituted acrylic acids wherein the preferred carboxylic monomers are acrylic and methacrylic acids. Hydroxyl containing monomers are hydroxy containing ethylenically unsaturated monomers including hydroxy alkyl acrylates such as 2-hydroxy ethyl acrylate and methacrylate, 2-hydroxypropyl acrylate and methacrylate, and similar hydroxy alkyl acrylates. Amido containing monomers include acrylamide and methacrylamide or similar alkyl alkylol acrylamide monomers. Other reactive monomers include N-methylol acrylamide or methacrylamide monomers. The remaining monomers that can be copolymerized with the alkylol acrylamide monomer and functional monomers to form a reactive self-curing polymer are ethylenically unsaturated monomers such as vinyl monomers, including for example, vinyl esters, such as vinyl acetate, vinyl propionate, vinyl butyrates, vinyl benzoate, isopropenyl acetate and similar vinyl esters; and vinyl halides such as vinyl chloride. Other ethylenically unsaturated monomers can include, for example, those monomeric materials exhibiting ethylenic double bond unsaturation such as polymerizable allylic acrylic, fumaric, maleic, or like ethylenically unsaturated double bond functionality. Further ethylenically unsaturated monomers can include, for example, styrene, methyl styrenes and similar alkyl styrenes, chlorostyrene, vinyl toluene, vinyl naphthalene, divinyl benzene, diallyl phthalate and similar diallyl derivatives, butadiene, alkyl esters of acrylic and methacrylic acid and similar ethylenically unsaturated monomers. Further suitable ethylenically unsaturated monomers include acrylic unsaturated monomers such as lower alkyl esters of acrylic or methacrylic acid having an alkyl ester portion containing between 1 to 12 carbon atoms as well as aromatic derivatives of acrylic and methacrylic acid. Useful acrylic monomers include, for example, acrylic and methacrylic acid, methyl acrylate and methacrylate, ethyl acrylate and methacrylate, butyl acrylate and

methacrylate, propyl acrylate and methacrylate, 2-ethyl hexyl acrylate and methacrylate, cyclohexyl acrylate and methacrylate, decyl acrylate and methacrylate, isodecylacrylate and methacrylate, benzyl acrylate and methacrylate and various reaction products such as butyl, phenyl, and cresyl glycidyl ethers reacted with acrylic and methacrylic acids, hydroxyl alkyl acrylates and methacrylates such as hydroxyethyl and hydroxypropyl acrylates and methacrylates, as well as amino acrylates and methacrylates.

A further improved self-curing water-dispersed polymer can be produced by polymerizing the ethylenic monomer mixture including reactive functional monomer and alkylol acrylamide monomer with a hydroxy alkyl phosphate ester monomer containing at least one unsaturated double bond. The hydroxy alkyl phosphate ester monomer can be prepared, for instance, by reacting an acrylic glycol with phosphorus pentoxide or reacting an acrylic monoepoxide with superphosphoric acid to produce a hydroxy acrylic phosphate ester monomer. The mixture of ethylenic monomers described above in accordance with this invention can contain by weight between 1% and 10% hydroxy alkyl phosphate ester monomer to be copolymerized and produce a phosphated self-curing water dispersed polymer. As a further variation, the water dispersed self-curing polymer can be produced by polymerizing the ethylenic monomer mixture including reactive functional monomer and alkylol acrylamide monomer in the presence of a high molecular weight epoxy phosphate ester surfactant to provide a self-curing water-dispersed polymer emulsified or otherwise dispersed in water by the epoxy phosphate surfactant. The epoxy phosphate esters comprise epoxy-phosphate produced by reacting between 0.05% and 5% phosphoric acid with a relatively high molecular weight adducts of bisphenol A reacted with low molecular weight epoxy resin such as Dow DER 333. During the polymerization process, the epoxy phosphate surfactant acts as a primary surfactant and controls the particle size of the self-curing copolymer and further stabilizes the polymer particles as the particle size increases. The foregoing self-curing water-dispersed polymers containing phosphates provide improved self-curing reactivity wherein the phosphate functions as a cross-linking promoter. Further advantages are the phosphate can further participate in the cross-linking reaction and the film properties exhibit improved corrosion resistance.

The reactive self-curing alkylol acrylamide water dispersed polymer preferably is a copolymer prepared by copolymerizing the ethylenically unsaturated monomers in water by two step polymerization described above through free radical induced polymerization using peroxy or azo catalysts, common redox catalysts, ultraviolet radiation or the like. Free radical initiators also include various peroxygen compounds such as persulfates, benzoyl peroxide, t-butyl hydroperoxide, cumene hydroperoxide, and similar peroxide catalysts; and azo compounds such as azo bis-isobutyronitrile and dimethylazobis-isobutyrate. Initiating systems further include alkali metal persulfate or ammonium persulfate with or without a reducing substance adapted to activate the persulfate. The initiators or catalysts ordinarily are used at a level of about 0.1% to 1% by weight of monomers. The resulting self-curing water dispersed polymer contains by weight between 1% and 20% reactive carboxyl, hydroxyl, amine or amide monomers, and the remaining being other ethylenically unsaturated monomers. Suitable water dispersed polymers can be produced by various aqueous polymerization methods. These polymers can include emulsion polymers produced by emulsion polymerization of monomers with water soluble initiator and in the presence of emulsifier; suspension polymers produced by aqueous suspension polymerization using a suspending agent; or microemulsion polymers produced by aqueous microemulsion polymerization in the monomer droplets and stabilized by long chain aliphatic alcohols; or azeotropic emulsion polymers produced in a water-solvent azeotropic mixture using soap emulsifiers. Other useful water dispersed polymers can be produced by copolymerization of the monomers in water containing an ionizing agent and a minor amount of organic cosolvent as disclosed in U.S. 4,218,356.

Referring next to the epoxy-acrylic copolymers, the copolymer comprises epoxy resin coreacted with monomers including acrylic monomers, or coreacted with preformed acrylic copolymer, to produce an epoxy-acrylic copolymer comprising coreacted epoxy resin and acrylic polymer. The preferred epoxy-acrylic copolymer comprises an epoxy-acrylic graft copolymer mixture containing epoxy resin, epoxy-acrylic graft copolymer, and ungrafted addition polymer produced by polymerizing ethylenically unsaturated monomers in-situ with epoxy resin in the presence of at least 3% benzoyl peroxide (or equivalent) as more particularly set forth in U.S. 4,212,781 and said patent is incorporated herein by reference. The in-situ polymerization of monomers generally comprises reacting the ethylenically unsaturated monomers in the presence of epoxy resin with at least 3% of benzoyl peroxide by weight of the monomer. The ethylenically unsaturated monomers include carboxyl functional monomers including acrylic acid monomer such as acrylic acid and lower alkyl substituted acrylic acids such as methacrylic acid to provide carboxyl functionality means for dispersing the epoxy-acrylic copolymer mixture into water. The preferred acrylic acid is methacrylic acid. The balance of the monomers preferably are nonreactive under the contemplated conditions of polymerization, although small amounts of other reactive monomers may be used such as hydroxy monomers illustrated by 2-hydroxy ethyl methacrylate, amide monomers illustrated by acrylamide, or N-methylol monomers illustrated by N-methylol acrylamide. The remaining monomers are nonreactive monomers illustrated by acrylate and methacrylate esters, such as ethyl acrylate, methyl methacrylate, or isobutyl methacrylate, styrene, or vinyl toluene, vinyl acetate, vinyl chloride, vinylidene chloride, acrylonitrile, and generally alkyl esters of an acrylic acid, generally the lower alkyl esters, that is, those esters in which the esterifying group contains from 1 to 4 carbon atoms, and particularly ethyl acrylate. Other useful monomers in this class include other C₁₋₁₅ alkyl acrylate esters and

methacrylate esters such as, for example, propyl acrylate, isopropyl acrylate, butyl acrylate, isobutyl acrylate, tertiary butyl acrylate, pentyl acrylate, decyl acrylate, lauryl acrylate, isobornyl acrylate, methyl methacrylate, butyl methacrylate, isobutyl methacrylate, hexyl methacrylate, 2-ethyl hexyl methacrylate, octyl methacrylate, and nonyl methacrylate. Other useful monomers are those readily commercially available monomers having vinyl unsaturation and include styrenic monomers such as styrene, vinyl toluene, divinyl benzene, isoprene and butadiene. The in-situ formed carboxyl-functional polymer can have a molecular weight between 5,000 and 20,000 and preferably between 7,000 and 15,000. The carboxyl content ($-\text{COOH}$) should comprise at least 2% by weight of the monomer mixture and preferably should be above 5%. The epoxy resin portion of the epoxy-acrylic copolymer can be either aliphatic or aromatic, although the aromatic epoxy resins are preferred. The most preferred epoxy resins are polyglycidyl ethers of bisphenol-A, especially those having 1,2-epoxy equivalency of from about 1.3 to about 2. The molecular weight should be from about 350 to about 20,000, and preferably, for sanitary coating compositions, from about 4,000 to about 10,000. When the in-situ polymeric blend contains from 50% to 90% by weight of epoxy resin based on total polymer solids, the epoxy resin selected should have a molecular weight in the range from about 4,000 to about 10,000, particularly for the preparation of sanitary coating compositions. Mixtures of monoepoxides and diepoxides are desirable. Another procedural variation is the introduction of the aromatic polyether which is devoid of oxirane functionality by reacting epoxide groups with benzoic acid, phenol or similar monoreactive epoxide blocking agent. Still another variation comprises providing an epoxy phosphate as the epoxy resin component of the epoxy-acrylic copolymer wherein the epoxy phosphate preferably contains up to about 5% by weight coreacted phosphate. The epoxy phosphate is then coreacted with acrylic monomers including carboxyl monomers by in-situ copolymerization induced by at least 3% peroxide catalyst by weight based on monomers to produce an epoxy-acrylic composition containing epoxy phosphate-acrylic graft copolymer. In preferred practice, the epoxy resin is a mixture including aromatic polyether devoid of oxirane functionality, aromatic polyether having a single oxirane group, and aromatic polyether having two oxirane groups. This mixture of epoxy functionality maximizes compatibility, although the aromatic polyether devoid of oxirane functionality can be added later, and the mixture can be heated and agitated to enhance the intimacy of the association between the various components.

The in-situ formed epoxy-acrylic copolymer can be prepared by in-situ polymerization of the ethylenic monomer with epoxy resin. The epoxy resin can be heated in a reactor wherein the polymerizable monomer can be added slowly over a period of at least two or three hours along with a solvent and a free radical initiator. Although the reaction may be conducted in the absence of solvent, a solvent system is preferred for the in-situ polymerization of monomers in the presence of epoxy resin. A preferred solvent system comprises two miscible solvents, one of which dissolves the epoxy resin and the other of which dissolves the monomers. The particular solvents satisfactory for the epoxy resin are solvents such as xylene, benzene, ethyl benzene, toluene, and the alkoxy alkanols. For the monomer, alcohols such as methanol, ethanol, propanol, butanol, and the like, are suitable, with butanol being preferred. Ethylene glycol monobutyl ether, ethylene glycol monobutyl ether acetate, and the like, hexane, mineral spirits, and the like, are also suitable. For subsequent dispersion into water, then the solvents selected should be water-soluble materials, as are acetone, butanol, ethanol, propanol, ethylene glycol monoethyl ether, and the like. Ordinarily the amount of solvent may be in the range from about 5% to 30% by weight of the sum of the other components. In practice, the epoxy resin and the mixture of polymerizable monomers are reacted together in the presence of a free radical initiator, preferably of the peroxide type, and benzoyl peroxide is most preferred. Typical and useful free radical initiators include cumene hydroperoxide, benzoyl peroxide, t-butyl perbenzoate, t-butyl peroxide, lauroyl peroxide, methyl ethyl ketone peroxide, chlorobenzoyl peroxide, and the like. Benzoyl peroxide is preferred as the free radical initiator for use in the practice of the present invention. The amount of free radical catalyst is expressed in terms of percentage by weight of benzoyl peroxide based on the total weight of the polymerizable monomer, or equivalent, at the temperature of use. The amount of catalyst should be at least 3%, and preferably more than 4% by weight of benzoyl peroxide or equivalent based on monomer weight. The reaction temperature preferably is maintained in the range from about 80°C to about 130°C, although the temperature may be adjusted within a relatively wide range to accommodate the reactivity of the mixture. Thus, operating temperatures in the range from about 30°C to about 200°C are feasible, depending upon the end results and operating conditions selected. After the monomers are added, the reaction mixture is normally held for up to three hours at reaction temperature to complete the monomer conversion. The in-situ polymerization of the monomers produces an in-situ formed carboxyl functional polymer containing at least about 20% of polymerized monoethylenically unsaturated carboxylic acid monomer based on the total weight of monomers as well as epoxy-acrylic graft polymer and ungrafted epoxy resin, as more particularly set forth in U.S. 4,212,781.

The epoxy-acrylic copolymer also can be an epoxy-acrylic ester copolymer produced from previously described epoxy resin and copolymerized monomers including carboxyl monomers wherein the epoxy-acrylic copolymer comprises acidic or carboxylic copolymer esterified with epoxy resin. The epoxy-acrylic ester can be produced by esterifying a solvent-soluble carboxy functional polymer with an epoxy resin wherein the esterification reaction is carried out preferably in an organic solvent and in the presence of sufficiently high amounts of amine catalyst to produce a nongelled epoxy ester copolymer. The

esterification reaction is carried out in the presence of amine catalyst in sufficient amounts greater than catalytic amounts of 0.3% to avoid gellation and preferably in the presence of greater than 2% amine esterification catalyst based on the weight of reactants subject to esterification to produce an epoxy-acrylic ester. The preformed acrylic polymer preferably comprises copolymerized ethylenically unsaturated monomers including at least about 20% by weight of copolymerized monoethylenically unsaturated carboxylic acid monomer based on the total weight of copolymer to produce a carboxyl functional prepolymer. The epoxy resin portion comprises at least about 40% of the epoxy-acrylic ester polymer and provides an oxirane functionality in a stoichiometric deficiency with respect to carboxyl functionality in the carboxyl preformed polymer in a ratio of about 1:2 to 1:20 whereby the excess carboxyl functionality in the epoxy-acrylic ester provides a means for dispersing the polymer into water by reacting with a base to render the reaction product self-emulsifiable into water, as further described in detail in EPO publication Serial No. 0 006 334 published January 9, 1980.

Still a further alternative for producing the epoxy-acrylic copolymer comprises reacting the epoxy resin with a preformed carboxyl polymer of polymerized ethylenically unsaturated monomers in the presence of amine and considerable amounts of water to produce a complex quaternary ammonium-amine salt linkage in the epoxy-acrylic copolymer such as disclosed in U.S. 4,247,439. Still a further variation of the epoxy-acrylic copolymer can comprise forming a preformed carboxylic polymer of polymerized ethylenic monomers including carboxyl monomers in the presence of melamine resin followed by reacting the preformed carboxylic polymer with epoxy resin in the presence of high amounts of amine as suggested in U.S. 4,289,811.

The epoxy-acrylic copolymer can further consist of an epoxy phosphate ester and an acrylic copolymer to produce the epoxy-acrylic portion of the mixture containing the self-curing water-dispersed polymer. The epoxy phosphate ester can be produced by reacting a high molecular weight epoxy resin with 0.05% to 5% phosphoric acid. The acrylic copolymer preferably is a resinous polyelectrolyte polymer produced by polymerizing ethylenic monomers including acrylic monomers and ionizable functional monomers, such as acrylic or methacrylic acid, in water containing an ionizing agent for ionizing the functional groups of the functional monomers.

Referring next to an optional but a preferred inclusion of a phosphated polymer in the polymeric blend of this invention, the phosphated polymer can include mono- and di-alkyl esters of phosphoric acid, phosphated epoxidized oil, phosphated epoxidized polybutadiene copolymers, phosphated acrylic copolymers, phosphated polyesters, phosphated copolymers containing copolymerized phosphate monomer, epoxy phosphate, and phosphated epoxy-acrylic copolymer.

Phosphated esters of alkyl alcohol can comprise a mixture of mono- and di-alkyl esters of phosphoric acid produced by reacting phosphoric acid with aliphatic alcohols containing between 2 and 20 carbon atoms and preferably between 4 and 8 carbon atoms as suggested in U.S. 2,005,619. Other water dispersed phosphated copolymers can be produced by polymerizing monomers in water containing an ionizing agent and a minor amount of organic cosolvent such as suggested in U.S. 4,218,356 wherein the polymers are phosphated. The water dispersed phosphated copolymers can comprise an aqueous dispersion of a resinous polyelectrolyte formed by the addition polymerization in water of ethylenically unsaturated monomers, including at least a portion of the monomers having functional groups ionizable in water such as carboxyl groups, wherein the water contains an ionizing agent for neutralizing at least a portion of the functional groups in the resulting copolymer, as set forth in U.S. 4,218,356. The phosphated resinous polyelectrolyte can be produced by providing an aqueous medium containing a cosolvent, adding the ionizing agent in advance of or concurrently with the monomers, polymerizing the monomers by free-radical polymerization until the water-dispersed resinous polyelectrolyte is formed, and then phosphating the remaining oxirane groups to produce a phosphated water-dispersed polymer.

Phosphated epoxidized oil can comprise a phosphated ester of epoxidized oil or epoxidized polybutadiene copolymer by reacting the epoxidized polymer with up to about 5% phosphoric acid. Phosphated acrylic copolymers can be produced by reacting an acrylic copolymer containing oxirane or hydroxy functionality with phosphoric acid. The acrylic copolymer can be produced by reacting ethylenically unsaturated monomers, particularly including glycidyl acrylate or methacrylate or hydroxy functional monomers as well as other ethylenically unsaturated monomers such as methyl methacrylate, butyl methacrylate, hexyl methacrylate, octyl methacrylate, isodecyl methacrylate, stearyl methacrylate, methyl acrylate, ethyl acrylate, butyl acrylate, hexyl acrylate, 2-ethylhexyl acrylate, octyl acrylate, decyl acrylate and dodecyl acrylate, substituted alkyl esters, for example, hydroxyalkyl esters such as hydroxyethyl and hydroxypropyl acrylate and methacrylate, styrene, alpha-methyl styrene, alpha-chloro styrene and vinyl toluene. Phosphated acrylic copolymers comprise phosphoric acid reacted with acrylic copolymer containing oxirane or hydroxy functionality wherein the acrylic copolymer comprises copolymerized ethylenic monomers including glycidyl acrylate or methacrylate or hydroxyl functional monomer.

Phosphated polyesters can be hydroxy terminated polyesters reacted with phosphoric acid wherein the polyesters ordinarily are dicarboxylic saturated or unsaturated acid esterified with simple glycols. Glycols include ethylene glycol, propylene glycol, diethylene glycol, dipropylene glycol 1,3 or 1,4 butanediol, hexanediols as well as minor amounts of polyols such as pentaerythritol, triethylene glycol, trimethylol propane, or glycerol. The unsaturated dicarboxylic acid component can be alpha-beta

3A

unsaturated dicarboxylic acids or anhydrides such as maleic, fumaric, itaconic or the like. Saturated dicarboxylic acids or anhydrides include phthalic anhydride or acid, terephthalic acid, isophthalic acid, adipic acid, sebacic acid, succinic acid or similar saturated acids or anhydrides. Monocarboxylic acids or fatty acids can be added to a minor amount and can include lower alkyl acids as well as lauric acid, palmitic acid, myristic acid and stearic acids. The hydroxyl terminated polyester is produced by reacting excess equivalents of glycol and polyol with lesser equivalents of carboxylic acid whereupon the excess hydroxyl groups are phosphated with phosphoric acid. Similarly, polyester containing terminal glycidyl groups can be reacted with phosphoric acid to produce a phosphated polyester.

Phosphated copolymers can be produced by copolymerizing ethylenic monomers with a phosphated hydroxy ethyl acrylate or similar phosphated monomers. Phosphated copolymer based on copolymerization of ethylenically unsaturated monomers in combination with phosphated ethylenically unsaturated monomer can be produced by polymerizing the ethylenic monomers previously described with the phosphate monomer in an aqueous polymerization process such as emulsion, suspension, aqueous ionic resinous polyelectrolyte, dispersion, microemulsion azeotropic and similar process previously described herein.

Phosphated epoxy-acrylic copolymers can be produced by reacting ethylenic monomers including ethylenic carboxyl monomers in the presence of epoxy phosphate resin by in-situ polymerization to produce phosphated epoxy graft copolymer. Other phosphated epoxy-acrylic copolymers can be produced by reacting epoxy resin with preformed acrylic carboxyl copolymer in the presence of high amounts of amine to produce an epoxy-acrylic ester copolymer or in the presence of amine and water to produce an epoxy-acrylic quaternary ammonium copolymer wherein the phosphoric acid is coreacted either before or after the epoxy acrylic copolymer formation. Another phosphated epoxy polymer comprises an epoxy polymer coreacted with phosphoric acid to produce a phosphated epoxy ester polymer. The phosphated epoxy polymer contains on an equivalent basis at least about 0.1 equivalent of phosphate per equivalent of epoxide and preferably about 0.1 to 1 equivalent of phosphate per epoxide equivalent on the epoxy polymer. Thus, a mole of monoepoxide polymer can contain up to about 1 mole of phosphate whereas a mole of a diepoxide polymer can contain up to 2 moles of phosphate. On a weight basis, the phosphated epoxy polymer contains between about 0.05% and 5%, and preferably between 0.1% and 3% coreacted phosphoric acid, although excess molar amounts of phosphate reactant in any appreciable amount is undesirable. The phosphated epoxide resin can be synthesized by heating a solution of epoxy polymer dispersed in a suitable solvent such as methyl ethyl ketone or 2-butoxy ethanol and then reacted with phosphoric acid or preferably polyphosphoric acid at reflux temperatures of 120°C to 145°C for sufficient time, typically 2 to 8 hours, to completely react the phosphoric acid with available epoxide groups. The preparation of epoxy phosphate resins are shown in U.S. 4,289,812 and U.S. 4,316,922. In a further variation of this aspect of the invention, the epoxy phosphate can be produced by reacting a non-phosphated epoxy resin with an epoxy resin containing higher levels of phosphate, provided that the resulting epoxy resin mixture contains between about 0.05% and 5% by weight coreacted phosphoric acid. For instance, a weight ratio of 9 parts non-phosphated epoxy mixed with 1 part phosphated (10%) epoxy resin reacted together at temperatures about 120°C for at least 2 hours provides an epoxy phosphate containing 1% coreacted phosphoric acid in accordance with this invention. In practice, the non-phosphated epoxy resin is reacted with phosphated (10%) epoxy resin at sufficiently high temperatures for time sufficient to convert the mixture to predominately a monophosphate, which can be enhanced by the addition of about 2% water based on solids to hydrolyze the phosphated mixture and preferably convert the mixture to a predominately monophosphated epoxy resin. The epoxy phosphate containing between 0.05% and 5% coreacted phosphoric acid in accordance with this invention then can be dispersed into an amine and water mixture by mechanical mixing. The resulting aqueous dispersed epoxy phosphate can be added to the epoxy-acrylic copolymer mixture and the self-curing water dispersed polymer in accordance with this invention.

A highly desirable sprayable coating composition can be produced based on the reactive self-curing polymer blended with amounts of epoxy-acrylic copolymer and phosphated polymer as the matrix polymer. Broadly the composition can comprise between about 5% and 99% self-curing polymer, 1% and 70% epoxy-acrylic copolymer, and 0% and 50% phosphated polymer. Preferred coating compositions of this invention contain a polymer blend on a weight basis comprising between 40% and 90% reactive self-curing polymer, between about 1% and 30% epoxy-acrylic copolymer, and between 0% and 30% and preferably 3% and 20% and most preferably between 3% and 15% phosphated polymer. The foregoing binder matrix polymer compositions are then mixed with a water-dispersible cross-linking component generally referred to as aminoplast resins adapted to heat cure and cross-link with the carboxyl functionality of the epoxy-acrylic copolymer mixture. On a polymer weight basis, the coating composition binder contains between 0% and 15% but preferably between 1% and 10% aminoplast cross-linking resin mixed with between 85% and 100% of the above indicated matrix polymer compositions.

Referring next to the aminoplast cross-linking resins, aminoplasts are melamine or melamine derivatives such as methylol melamine or similar alkylated melamine formaldehyde resins. Aminoplasts further include benzoguanamine, acetoguanimine, as well as ureaformaldehyde. Commercially available aminoplasts which are water-soluble or water-dispersible for the instant purpose include Cymel 301, Cymel

38

303, Cymel 370, and Cymel 373 (all being products of American Cyanamid, Stamford, Conn. Cymel is a Trade mark, said aminoplasts being melamine based, e.g., hexamethoxy-methyl melamine for Cymel 301), and Beetle 80 (products of American Cyanamid which are methylated or butylated ureas Beetle is a Trade Mark). Other suitable aminoplast resins are of the type produced by the reaction of aldehyde and formal guanamine; ammeline; 2 - chloro - 4,6 - diamine - 1,3,5 - triazine; 2 - phenyl - p - oxy - 4,6 - diamino - 1,3,5 - triazine; and 2,4,6 - triethyltriamino - 1,3,5 - triazine. The mono, di-, or tri-aryl melamines, such as 2,4,6 - triphenyl - triamino - 1,3,5 - triazine, are preferred.

The water-dispersed coating compositions can be produced by mixing together the various water-dispersed polymers. The self-curing polymer is prepared in aqueous medium. The aminoplast polymer can be dispersed into water by mechanical mixing. The epoxy-acrylic copolymer and phosphated polymer can be prepared in solvent and then subsequently dispersed into water using a fugitive base such as primary secondary, and tertiary alkyl, alkanol, aromatic amines, or alkanolalkyl mixed amines such as mono-ethanol amine, dimethyl ethanol amine, diethanol amine, triethyl amine, dimethyl aniline, ammonium hydroxide, and the like, as more particularly described in U.S. 4,212,781. Alternatively, one of the polymers of the blend can be produced in the presence of one of the other polymers in the polymeric blend of this invention. For instance, the water dispersed self-curing polymer comprising polymerized monomers including alkyl alkylol acrylamide monomer can be polymerized in the presence of either the epoxy-acrylic copolymer or the phosphated polymer or in a mixture thereof. Similarly, the epoxy-acrylic copolymer or the phosphated polymer can be prepared by in-situ polymerization of the monomers in the presence of the other preformed polymer or by in-situ polymerization in the presence of the self-curing polymer. The amount of water contained in the coating composition containing the epoxy-acrylic copolymer, the reactive self-curing polymer, the phosphated polymer, and the aminoplast resin depends on the viscosity desired, which in turn, relates to the method of application. For spraying, preferably the coating composition contains between about 10% and 30% by weight polymeric solids relative to 70% to 90% water including other volatiles such as minor amounts of solvent. For applications other than spraying, the aqueous polymeric dispersions can contain between about 10% and 40% by weight water. Organic solvents can be utilized to facilitate spray or other applications methods and such solvents include n-butanol, 2-butoxy-ethanol-1, xylene, toluene, and preferably n-butanol is used in combination with 2-butoxy-ethanol-1. The coating composition of the present invention can be pigmented and/or opacified with known pigments and opacifiers. For many uses, including food uses, the preferred pigment is titanium dioxide. The resulting aqueous coating composition can be applied satisfactorily by conventional methods known in the coating industry. Thus, spraying, rolling, dipping, and flow coating application methods can be used for both clear and pigmented films, although spraying is preferred. After application onto the metal substrate, the coating is cured thermally at temperatures in the range from about 95°C to about 235°C or higher for time sufficient to effect complete curing as well as volatilizing of any fugitive component therein. Upon heat curing, the reactive self-curing polymer is believed to become reactive and self-curing wherein the alkyl chain of the alkoxy acrylamine splits from the alkylol acrylamide chain whereby the acrylamide chain of the polymer reacts with the functional monomer groups of carboxyl, hydroxyl, or amido groups, or can react with the carboxyl or hydroxyl functionality in the epoxy-acrylic copolymer mixture as well as the epoxy phosphate and the aminoplast resin.

For metal sheet substrates intended as beverage containers and particularly for carbonated beverages such as beer, the coating should be applied at a rate in the range from 0.5 to 15 milligrams of polymer coating per square inch (775—23,250 mg/m²) of exposed metal surface. To attain the foregoing, the water-dispersible coating is applied can be as thick as 0.1 to 1 mil (0.0025—0.025 mm).

For a better understanding of the present invention, the following Examples are provided. In this application, all parts are parts by weight, all percentages are weight percentages, and temperatures are degrees Centigrade unless otherwise expressly noted.

Example 1 Preparation of self-curing copolymer

A two stage thermosetting acrylic emulsion copolymer containing styrene/ethyl acrylate/methacrylic acid/N-isobutoxy methylol acrylamide was prepared by loading 100 parts of deionized water and 0.80 parts sodium dihexyl sulfosuccinate into the reactor and heating under nitrogen sparge with agitation to 76—77°C. At equilibrated reaction temperature the nitrogen sparge was removed and 0.10 parts ammonium bicarbonate added. A mixture of 3.0 parts styrene and 2.0 parts ethyl acrylate was then added to the reactor and emulsified 10 minutes, whereupon 0.25 parts ammonium persulfate were added and allowed to react 20 minutes before starting monomer stage number one.

Monomer stage number one contained 33.0 parts styrene, 26.5 parts ethyl acrylate and 3.50 parts methacrylic acid. This monomer stage was added at a constant rate to the reactor so that addition was complete after 3.0 hours. Monomer stage number two consisted of 14.0 parts styrene, 11.5 parts ethyl acrylate, 1.5 parts methacrylic acid and 5.0 parts N-isobutoxy methylol acrylamide, and was added continuously to the reactor during 1.25 hours. After monomer mix number two was completed, the batch was held for 2 hours at the reaction temperature before cooling and filtration.

Monomer stage sizes are not necessarily confined to those described in this example. Satisfactory

39

spray characteristics were obtained when the first stage/second stage ratio was maintained at a weight ratio of between 25/75 and 75/25.

Examples 2—10

- 5 In a manner similar to Example 1, additional self-curing copolymers were prepared with the alkylol acrylamide in the second monomer stage.

10

15

20

25

30

35

40

45

50

55

60

65

Examples 2—10

	Ex. 2	Ex. 3	Ex. 4	Ex. 5	Ex. 6	Ex. 7	Ex. 8	Ex. 9	Ex. 10
Alkylol acrylamide	NMA ¹	NMA	NMA/NIBMA ²	NIBMA	NIBMA	NIBMA	NIBMA	NIBMA	NIBMA
Alkylol acrylamide in second stage	2.0	3.0	1.0/5.0	5.0	10.0	10.0	10.0	10.0	15.0
Carboxyl monomer level	4.0	5.0	5.0	5.0	3.0	5.0	4.0	10.0	5.0
Additional functional monomer	HEA ³	none	none	none	none	HEA	none	none	none
Additional functional monomer level	3.0					5.0			
Polymer Tg	20°	30°	40°	40°	30°	20°	30°	50°	30°
Other monomers	91.0	92.0	89.0	90.0	87.0	80.0	86.0	80.0	80.0

¹N-methylol acrylamide²N-isobutoxy methylol acrylamide³Hydroxy ethyl acrylate

Examples 11—13

In contrast, to Examples 1—10, emulsion polymers were prepared containing an equal quantity of alkylol acrylamide in both monomer stages.

TABLE 2

	Ex. 11	Ex. 12	Ex. 13
Alkylol acrylamide	NMA	NIBMA	NIBMA
Alkylol acrylamide level	3.0	5.0	10.0
Carboxyl monomer level	5.0	5.0	5.0
Polymer Tg	30	40	50

Example 14

A) Preparation of epoxy-acrylic copolymer

An epoxy-acrylic graft copolymer was prepared according to the following procedure: 2.31 pounds (1.05 kg) of epoxy resin (DER-333, DER is a trade mark) were heated in an agitated reactor to about 82°; 1.17 pounds (0.53 kg) of bisphenol-A were added with agitation. The reactor then was heated to about 191° over a period of about 2 hours and held there for an additional 2 hours. Periodic testing for viscosity and percent oxirane was made. Target oxirane value was about 0.6% and viscosity at 25°C between Z and Z₁ (Gardner-Holt). When these values were obtained, 1.35 pounds of 2 - butoxy - ethanol - 1 were added, followed by 2.03 pounds (9.2 kg) of N-butanol. The molecular weight of the epoxy resin at this point was about 5,500 based on oxirane content. In a separate vessel, there was loaded and mixed the following: 0.64 pounds (0.29 kg) of methacrylic acid, 0.4 pounds (0.18 kg) of styrene, 0.44 pounds (0.2 kg) of ethyl acrylate, and 0.1 pounds (0.045 kg) of benzoyl peroxide. This monomer mixture was added gradually to the reactor containing the epoxy resin over a period of 2 hours at uniform rate. The reaction temperature was maintained at 118°. Viscosity was checked periodically on samples. The batch was cooled to 85°; its Acid Number on solids was 85.

The resin batch then was fed into an agitated reducing vessel containing a mixture of 10.95 pounds (4.97 kg) of deionized water (resistivity at least 50,000 ohm-cm) and 0.57 pounds (0.26 kg) of dimethylethanolamine. The temperature of the resulting blend was 50°. It was held there for about an hour, then the blend was cooled to below 32° by adding 5 pounds (2.3 kg) of the cool deionized water.

B) The effect of variation in composition

In the above example, the amount of benzoyl peroxide employed during the reaction was about 6.8% by weight based upon the monomer mixture. To demonstrate the effect of changes in composition with respect to proportions of the epoxy resin and the several monomers in the monomer mixture, several variations were made in Table 3 below.

TABLE 3

	Visc. ¹	Epoxy resin	MAA ²	ST ³	EA ⁴	Acid number
45	U	70	13	8	9	85
50	Z—Z ₁	30	13	34	23	85
	Z—Z ₁	70	13	7.7	0.3	85
	Z—Z ₁	80	13	6.8	0.2	85
55	Z	50	32.5	17.0	0.5	210
	Y	60	25.75	13.85	0.4	168
	Z ₅	80	13	6.8	0.2	85

¹Viscosity of epoxy resin at 40% NV in 2-butoxy ethanol

²MAA=Methacrylic acid

³ST=Styrene

⁴EA=Ethyl acrylate

Example 15
Epoxy phosphate synthesis
Procedure A

The epoxy phosphate was prepared according to the following procedure: 1005 gms. of epoxy resin (DER-333) and 340.5 gms. of bisphenol-A and 178 gms. of butyl cellosolve were heated to 140°C in a 5L round bottom flask equipped with stirrer, condenser and thermometer. When the temperature was at 140°C, the heat was turned off and the exotherm raised the temperature to 188°C. After the exotherm was peak, the batch was kept at 175°C for an additional 5 hours. Periodic testing for viscosity and percent oxirane was made. Target oxirane value was about 2.28% and viscosity at 40% non-volatile between I—J. When these values were obtained, 227 gms. of butyl cellosolve were added, and the batch was cooled to 120°C. When the batch temperature was at 120°C, a mixture of 63.64 gms. of 85% phosphoric acid and 20 gms. of butyl cellosolve was added dropwise. The exotherm raised the batch temperature to 145°C. The batch was cooled to 120°C and held for a half hour. Twenty-seven gms. of water were added to the reaction mixture and the batch was held at 120°C for an additional 4 hours. After the hold, 241 gms. of butanol, 78 gms. of butyl cellosolve, 122.5 gms. of dimethylethanolamine and 2,500 gms. of deionized water were added respectively. The final mixture was agitated for 2 hours to obtain a stable emulsion.

Procedure B

Epoxy phosphate was produced by adding 816 gms. of epoxy resin (DER-333), 384 gms. of bisphenol-A, and 163 gms. of butyl cellosolve in a 5L round bottom flask equipped with a stirrer, condenser, and thermometer and heated to 140°C. When the temperature was at 140°C, the heat was turned off and the exotherm raised the temperature to 155°C. After the exotherm was peak, the heat was turned back on to keep the temperature at 175°C for an additional 2 hours. Periodic testing for viscosity and percent oxirane were made. Oxirane value was about 0.87% and the viscosity was X—Y at 40% NV in butyl cellosolve. When these values were obtained, 163 grams of butyl cellosolve were added, and the batch was cooled to 125°C. Mixture of 14.2 gms. of polyphosphoric acid (FMC) and 50 gms. of butyl cellosolve was added over 45 minutes, 30 gms. of extra butyl cellosolve was added as line rinse. The batch was held at 120°C for 1 hour. Then 23 gms. of DI water were added to the reaction mixture and the held batch was at 120°C for an additional 2 hours. After the hold time, the heat was turned off and 203 gms. of butanol were added over 8 minutes. 1550 gms. of DI water and 17.4 gms. of dimethyl ethanolamine were heated in a letdown container to 60°C. The above resin was dropped slowly into water amine mixture to form a stable emulsion. The resulting emulsion was adjusted to 25% NV by adding 1000 grams of DI water and stirring continued for 2 hours to insure an homogeneous mixture.

In accordance with the Procedure A or B various epoxy phosphates were produced with the following molecular weight variation.

TABLE 4

	Molecular weight	% Oxirane	% Acid	% NV	Emulsion
	9410	0.34	0.35	30.3	Good
	6150	0.52	0.39	34.4	Good
	4000	0.80	0.92	34.9	Good
	3080	1.04	0.92	32.8	Good
	2500	1.26	2.30	23.0	Good
	1400	2.28	4.20	22.4	Good

Example 16

Polymeric blend

About 860 gms. of emulsion copolymer from Example 1 were added to a 5L reaction flask along with additional deionized water to provide a NVM of about 23%. This mixture was agitated slowly to avoid foaming for about 15 minutes. Thereafter, a premix of 15.5 grams of DMEA and 82.8 gms. of water were added with continued agitation for about 30 minutes. About 72.8 grams of butoxy ethanol were added and agitated for 30 minutes. Thereafter, about 220.8 grams of epoxy phosphate from Example 15 and 815 grams of epoxy acrylic copolymers from Example 14 were added slowly over a time period of about 1 hour. The 259.3 grams of n-butanol were added and blended with increased agitation for about 15 minutes. Then about 31.3 grams of aminoplast resin Cymel 303 were added and mixed in for about 15 minutes. About 68.1 grams of deionized water were added followed by additional mixing of about 1 hour and then filtered.

The resulting polymeric mixture provided an excellent spray applied coating on the interior of cans.

To demonstrate the effect of changing the compositions of epoxy phosphate, water dispersed copolymer, and epoxy acrylic copolymers, several blend compositions are shown in Table 5. The epoxy phosphate was Example 15, the epoxy acrylic was Example 14(a), and the water dispersed copolymers were from Examples 1—10.

TABLE 5

	% by solid content			Coating properties			
	Epoxy phosphate	Water dispersed copolymer	Epoxy acrylic	Spray result	MEK rubs	Water resistance	Porosity
5	0	75	25	Good	Pass	Fair	Fair
10	5	80	15	Good	Pass	Good	Good
15	10	80	10	Good	Pass	Good	Good
20	15	80	5	Good	Pass	Good	Good
25	20	80	0	Poor	Pass	Good	Good
30	5	90	5	Fair	Pass	Good	Good
35	10	75	15	Good	Pass	Good	Good
40	0	90	10	Fair	Fail	Poor	Poor
45	10	60	30	Good	Pass	Fair	Fair
50	10	40	50	Good	Pass	Good	Good
55	8	84	8	Good	Pass	Good	Good
60	5	85	10	Fair	Pass	Good	Good
65	5	95	0	Poor	Pass	Fair	Fair
70	30	10	60	Good	Pass	Good	Good
75	40	20	40	Fair	Pass	Good	Good
80	50	30	20	Fair	Pass	Good	Good
85	28	30	42	Good	Pass	Good	Good

Example 17**Polymeric blend for coating**

In a manner similar to Example 16, a coating composition was prepared as follows. About 1031 gms. of emulsion from Example 13 were added to a 5L reaction flask along with 922 gms. of deionized water. This mixture was agitated slowly to avoid foaming. A premix of 18.7 gms. of DMEA and 40.0 gms. of water were added with continued agitation for about 30 minutes. Then 73.6 gms. of butoxy ethanol were added and agitated for 30 minutes. Thereafter, 263.7 gms. of n-Butanol were added and blended with increased agitation for about 15 minutes. Then 354.0 gms. of epoxy acrylic copolymers from Example 14 and 183.0 gms. of epoxy phosphate from Example 15 were added slowly over a time period of about 1 hour. To the resulting polymeric mixture, 31.3 gms. of amino-plast resin Cymel 303 was then added to provide an excellent sprayable can coating composition particularly useful for spray applications to interiors of beverages cans. Coating spray applied to cans were heat cured and provided excellent cured interior can coatings.

Example 18**Polymeric blend for coating**

In a manner similar to Example 16 a coating composition was prepared as follows. About 1000 gms. of emulsion copolymer from Example 1 were added into a 5L reaction flask with slight agitation. A premix of

18.1 gms. of DMEA and 37 gms. of water were added slowly to the latex with continued agitation. About 226 gms. of BuOH were added over a period of 1 hour. Thereafter, about 711 gms. of epoxy acrylic copolymers from Example 14 were added over 15 minutes. Then about 72.5 gms. of butoxy ethanol were added and followed by 900 gms. of deionized water. The mixture was agitated continuously for an additional 1 hour. The resulting polymeric mixture was mixed with an aminoplast crosslinking agent to obtain an excellent spray applied coating for the interior of cans. Spray applied coatings to cans were cured and provided excellent interior can coatings.

Example 19

10 Polymeric blend for coating

In a manner similar to Example 16, a coating composition was prepared as follows. About 1000 gms. of water dispersed copolymer from Example 13 were added to 770 gms. of deionized water in a 5L reaction flask with slow agitation. A premix of 18.4 gms. of DMEA and 100 gms. of water were added with continued agitation for about 30 minutes. Then a premix of 72 gms of butoxy ethanol, 226 gms. of n-butanol, and 130 gms. of water were added and blended for 30 minutes. About 716.8 gms. of epoxy acrylic copolymers from Example 15 were added over 30 minutes and followed by 32.8 gms. of aminoplast resin Cymel 303 and 80 gms. of water. The resulting polymeric mixture was agitated for another 2 hrs. for complete mixing. The resulting coating was spray applied to interiors of cans and heat cured to produce an excellent interior can coating.

20 Example 20

Polymeric blend for coating

In a manner similar to Example 16, a coating composition was prepared as follows. A mixture of 500 gms. of emulsion copolymer from Example 1 and 500 gms. of deionized water were loaded into a 5L reaction flask. A premix solution of 10 gms. DMEA and 100 gms. of deionized water were added to emulsion over 30 minutes. Then 243.0 gms. of epoxy phosphate from Example 15 and 1326.1 gms. of epoxy acrylic copolymers from Example 14 were added with increased agitation over 1 hour. About 87 gms. of n-butanol and 30.5 gms. of Cymel 303 were added and followed by 340 gms. of deionized water. The final polymeric mixture was agitated for another hour. The coatings from this process provided excellent spray applied coatings. The cured films exhibited excellent film integrity properties.

Example 21

Preparation of epoxy-acrylic ester copolymer

(a) Epoxy resin. To a 12-liter round-bottomed, 4-necked flask was charged 4144 g of a liquid spray resin, DER 333 (from Dow Chemical Company, Midland, Michigan DER is a trade mark), 2135 g of bisphenol A, and 409 g of 2-butoxy-ethanol-1. A water-cooled condenser was placed on one neck of the flask, an air driven mechanical stirrer was placed through the second neck, and a nitrogen feed was placed through the third, and a thermometer was placed through the fourth neck of the flask. Heating was started and the mixture in the flask was stirred under a vacuum of 15 inches. The temperature was raised to 140°C whereupon the heating and vacuum were discontinued. About 40 g of a volatile was collected comprising water, xylene, (from DER 333) and 2-butoxy-ethanol-1. The exotherm carried the reaction temperature to about 175°C, where it was held for 5 hours. The viscosity of the epoxy resin after one hour was Y Gardner-Holt (40% solution in 2-butoxy-ethanol-1), Y+ after 3 hours, and Z after 5 hours. At the end of the 5-hour hold, 1012 g of 2-butoxy-ethanol-1 was added, and the epoxy solution was left to cool to room temperature.

(b) Acrylic terpolymer. To a 12 liter 4-necked round-bottomed flask was charged 5503 g of n-butanol. Through the 4 necks were placed a water cooled condenser, an air-driven mechanical stirrer, a thermometer-nitrogen inlet line and monomer addition funnel. The solvent was heated to 113°C. A monomer mixture of 1764 g of methacrylic acid, 924 g of styrene, 26 g of ethyl acrylate, 182 g of benzoyl peroxide was slowly added to the refluxing solvent. The addition took 2 hours, and the reaction mixture was held at 114°C for another hour before cooling to room temperature. The non-volatile of the reaction mixture was 347.

Epoxy-acrylic ester copolymer

Into a 5-liter round-bottomed, 4-necked flask was charged 1306 g of epoxy resin (a), and 694 g of the acrylic resin (b). The reaction mixture was stirred under nitrogen sparge and heated to 105°C and held at that temperature for 4 hours. At the end of the hold, 23.6 g of dimethylethanol amine and 144 g of 2-butoxy-ethanol-1 were added in 4 minutes, and the reaction mixture was held at 102—105°C for ½ hour. The viscosity of the resin is U ½, (from 1 part resin, 1 part pyrol). Into a 2 gallon stainless steel container was charged 2043 g of deionized water and 86.1 g of dimethylethanol amine. The container was placed on a hot plate so that the temperature of the water was 50°C before 1500 g of the resin from the 5-liter flask was dropped into the agitated water. An emulsion formed easily and the non-volatile of the emulsion was 23.5% viscosity Ford #4 cup, 38 seconds, acid number 34.1 and base number 76.4.

Binder composition

A binder composition was produced by mixing the following:

- 1200 grams of emulsion copolymer of Example 1
- 893 grams of deionized water
- 5 21.7 grams dimethyl ethanolamine
- 86.8 grams of deionized water
- 106.7 grams of 2-butoxy-ethanol-1
- 225.5 grams of epoxy phosphate
- 284.3 grams Epoxy-acrylic Ester from above
- 10 363.6 grams of n-butanol
- 32.7 grams of Cymel 303
- 400.5 grams of deionized water

The foregoing blend was evaluated as an interior can coating. NVM was 18.8%; viscosity was 15 seconds #4 Ford cup; Acid No.=35.5; base No.=29; percent neutralization was 81.7%; organic solvent/solids ratio was 0.9; 2-butoxy-ethanol-1/n-butanol ratio was 30/70; and Cymel was 5% based on solids.

The coating formed a clear continuous coating upon air drying as well as upon baking. The coating had a wedge bend failure of 24%; pasteurization at 170°F (77°C) for 10 minutes produced a slight blush; blush adhesion was excellent; porosity ratio less than 1; MEK rubs of 24 overall; and general film properties were excellent.

Example 22

Preparation of epoxy-acrylic graft quaternary ammonium copolymer

High molecular weight epoxy resin was prepared by reacting 438 g of Epon 828 (liquid epoxy resin, Bisphenol type from Shell Chemical Co., Houston, Texas Epon is a trade mark), 248 grams (g) of Bisphenol A in 125 g of 2-butoxy-ethanol-1 and 23 g of xylene. 0.32 g of sodium acetate trihydrate in 3.6 g of water was now added. The reaction mixture was charged into a 12-liter 4 necked round bottom flask. Through one neck was placed an air driven mechanical stirrer, through another neck a water cooled condenser, through the third neck a thermometer and a nitrogen inlet, and the fourth neck an addition funnel, heating was provided by a heating mantle. When the temperature reached 140°C, heating was stopped and the exotherm carried the temperature of the reaction mixture to 175°C. The temperature was held at 175°C for five hours. At the end of the hold, the viscosity of the epoxy resin (40% solution in 2-butoxy-ethanol-1) was A1—Z2 (Gardner Holt viscosity). Heating was stopped and 484 g of n-butanol was carefully added. The temperature at the end of the butanol addition was about 115°C. A monomer mixture consisting of methacrylic acid 114 g, styrene 60 g, ethyl acrylate 1.6 g benzoyl peroxide (78% active in water) and 45 g of 2-butoxy-ethanol-1 was added to the epoxy resin in two hours, the temperature was held at 115°C. After the monomer mixture was added, 25 g of n-butanol was used to rinse the addition funnel and then added to the reaction mixture which was held at 115°C for another hour. A second monomer mixture consisting of 888 g of styrene, 350 g of ethyl acrylate, 186 g of methacrylic acid, 36.4 g of benzoyl peroxide (78% active in water), 289 g of 2-butoxy-ethanol-1 and 148 g of n-butanol was slowly added to the epoxy-acrylic graft copolymer at 115°C. The addition took 3 hours, and the reaction mixture was held at 115°C for another half hour before 719 g of n-butanol was added. Mixture was cooled from 115°C to 85°C whereupon 450 g of epoxy resin Epon 828 was slowly added to the reaction mixture and then held for ½ hour at 90°C. Then 80 g of dimethylethanol amine and 50 g of water were added and the reaction mixture was held at 80°C for 1 hour. Then 29 g of dimethylethanol was added to obtain a dispersion of 20% NVM. The resultant copolymer was believed to be an epoxy-acrylic graft containing quaternary ammonium functionalities. The boiling Joy and porosity of the film was drastically improved.

Example 23

N-isobutoxy methyl acrylamide (NiBMA) self-curing copolymer by suspension polymerization.

(a) To a 5 liter round-bottom 4-necked flask was charged 2211 g of deionized water. The flask was fitted with nitrogen inlet, thermometer, water cooled condenser and air driven mechanical stirrer. The water was heated to 60°C before 110 g of Vinol 523 (polyvinyl alcohol from Air Products and Chemical, Wayne, Pa. Vinol is a trade mark) was added and let dissolved for about 20 minutes. A monomer mixture consisted of 544 g of styrene, 435 g of ethyl acrylate, 54.5 g of methacrylic acid and 64 g of NiBMA monomer (85% in isobutanol) and 13 g of lauroyl peroxide was batch added to the aqueous solution at about 70°C and the polymerization was carried out at 70°C for about 5 hours. At the end of that time, the polymerization mixture was let cool to room temperature. The nonvolatile of the suspension was determined to be 29.2% and the viscosity by Ford #4 cup was 25 seconds.

A blend of suspension polymer (a), epoxy-acrylic graft copolymer (Example 14a), and epoxy phosphate (Example 15) was prepared as follows.

Into a 5-liter 4-necked round-bottomed flask was charged 1000 g of the suspension polymer (a) and 754.5 g of deionized water. The flask was fitted with nitrogen inlet, thermometer, water cooled condenser and an air driven-mechanical stirrer. The mixture was heated to 38°C, and 18 g of dimethylamino ethanol and 83.6 g of deionized water were added. The temperature was increased to 48°C and mixing was continued for 15 minutes. Then 55.9 g of 2-ethoxy-butanol-1 was added followed by 2.8 g of deionized

water. The mixture was stirred for 40 minutes at 43°C whereupon 200.3 g of an epoxy-phosphate (Example 15) was added in 30 minutes followed by 68.2 g of deionized water over 15 minutes. Then 378.1 g of epoxy-acrylic graft copolymer was added over 30 minutes followed by 77.1 g of deionized water over 5 minutes. Then 168.4 g of n-butanol was added in 10 minutes followed by addition of 29.1 g of Cymel 303. Additional n-butanol of 55.2 g was added over 15 minutes followed by the addition of 205.4 g of deionized water. The non-volatile of the blend was 15.1%, and viscosity Ford #4 cup was 13 seconds. A film drawn down on a glass plate followed by baking over a hot plate at 380°F (193°C) for 5 minutes provided a smooth coating.

10 Example 24

Blend containing epoxy graft acrylic-containing phosphated monomer

A 5L flask equipped with a stirrer, a thermometer, a condenser, and a N₂ inlet was charged with 434.5 gms. of 2-butoxy ethanol and heated to 150°C. When the temperature was at 150°C, 1000 gms. of solid epoxy resin (Epon 1009F) was added slowly to dissolve in the solvent. When all the epoxy resin was dissolved, the temperature was lowered to 115°C and 507.7 gms. of n-butanol was slowly added to maintain the temperature at 115°C. A mixture of 159.1 gms. of methacrylic acid, 83.2 gms. of styrene 2.3 gms. of ethyl acrylate, 25.6 gms. of glycidyl methacrylate phosphate, 21.7 gms. of benzoyl peroxide and 62.4 gms. of 2-butoxy ethanol was added to the epoxy solution over 1-1/2 hours. After all the monomer mixture was added, 34.9 gms. of n-butanol was used to rinse the addition funnel. The reaction mixture was kept at 115°C for 3 hours to complete the polymerization. The resin was then fed into an agitated reducing vessel containing a mixture of 2550 gms. of deionized water and 103 gms. of DMEA. The temperature of the resulting blend was 50°C. It was held there for about an hour, then the blend was cooled to below 32°C by adding 100 gms. of the cool deionized water. The phosphated epoxy-acrylic graft copolymer was blended with emulsion copolymer from Example 1 to produce an excellent spray applied interior can coating.

25 Example 25

Blend containing epoxy acrylic melamine graft

About 125 gms. of emulsion copolymer from Example 1 were added to a 5L flask along with 112.5 gms. of deionized water. This mixture was agitated slowly for about 15 minutes. A premix of 1.3 gms. of dimethyl amino ethanol amine and 12.5 gms. of deionized water were added with continued agitation. The 9.5 gms. of 2-butoxy ethanol was added and followed by 25.2 gms. of epoxy phosphate from Example 15 and 56.4 gms. of epoxy acrylic melamine graft produced according to Example 1 in U.S. 4,289,811. Thereafter, 28.5 gms. of n-butanol and 3.7 gms. of aminoplast resin, Cymel 303, were added and mixed very well. A mixture of 0.5 gms. of DMEA and 3.7 gms. of deionized water was then added and mixed for 1 hour. The resulting polymeric mixture provided an excellent spray applied coating on the interior of cans.

35 Example 26

Blend containing phosphorylated hydroxy terminated polybutadiene

To a suitable reactor 184 parts of hydroxy terminated polybutadiene (Hycar 1300×29 by B.F. Goodrich, Hycar is a trade mark) were added and 79 parts of 2-butoxy ethanol. Heating and agitation were begun with N₂ sparge and at 50°C, a mixture of 1.84 parts of super phosphoric acid and 44 parts of 2-butoxy ethanol was added slowly over 11 minutes. The temperature of the reactants was retained at 125°C for 25 minutes. At the end of this hold period, 3.7 parts of water was added and the temperature of the reaction mixture was retained at 125°C for 2 hours. At the end of the hold period, the temperature was lowered to 75°C and 3.7 parts of dimethyl amino ethanol amine was added and followed by 303.2 parts of water. One hundred and ten parts of 2-butoxy ethanol was then added and followed by 25 parts of butanol. Twenty six parts of DMEA was added and 200 parts of water was then added to adjust to the required solid.

Blend composition:

50 To a 5L reaction flask, 1000 gms. of water dispersed copolymer from Example 13 were added along with 754 gms. of water. This mixture was agitated slowly for 15 minutes. A premix of 18 gms. of DMEA and 200 gms. of water were added with continued agitation. Then 341.7 gms. of polybutadiene phosphate from the example above was added and rinsed by 17 gms. of water. Thereafter, 379.1 gms. of epoxy acrylic copolymer from Example 14 were added and rinsed by 19 gms. of water. Next, 228.7 gms. of n-butanol was added slowly and followed by a mixture of 8.73 gms. of butanol and 29.1 gms. of Cymel 303. To adjust the mixture to required solid content, 63.4 gms. of water was added. The resulting mixture formed good film for spray applied coatings.

Example 27

60 Blend containing phosphorylated polyester

A conventional polymerization vessel equipped with a stirrer, condenser and water separator, thermometer, and a nitrogen inlet was charged with a tall oil fatty acid (Sylfat V-18, Sylfat is a trade mark) and heated up to 80°C. When the temperature was at 80°C., 5120 gms. of a resinous polyol (RJ-101 from Monsanto) was added and continued to heat-up to 125°C. A vacuum of 18" was needed to pull out water which was trapped in RJ-101. The reaction mixture was heated to 240°C and xylene was added, as needed,

to aid the removal of water. The temperature was maintained at 240°C for 2½ hours and the acid number was 3.9 on solid. The heat was turned off and the reaction mixture was cooled to 120°C. When the temperature was at 120°C, 1567 gms. of acetone was added very slowly. When all the addition was finished, the temperature of the batch was 51°C. The resulting polyester was phosphorylated as follows. A conventional vessel equipped with a stirrer, condenser, and a thermometer was charged with 386.1 gms. of resin from above. At 25°C 15 gms. of P₂O₅ solid was added with stirring. The reaction mixture was heated up to 125°C and 79 gms. of 2-butoxy ethanol was added to dilute to 77.9% solid. This phosphorylated polyester was then added to a water dispersed copolymer (Example 13) and epoxy-acrylic copolymer (Example 14) to produce a good coating for spray application to interior of cans.

Example 28

Synthesis of phosphated epoxy-acrylic graft

An epoxy phosphate acrylic copolymer was made as follows. 1141 gms. of 95% Epon 828 in xylene, 614 gms. of bisphenol A, and 310 gms. of 2-butoxy ethanol was charged into a 5 l flask equipped with a stirrer, a thermometer, a N₂ inlet, a condenser and additional funnel. The mixture was heated to 90°C with N₂ sparge, when the temperature was at 90°C, a mixture of 0.52 gm. of sodium acetate and 3 gms. of water was added, and the heat was continued to 140°C. When the temperature reached 140°C, the heat was turned off. The exotherm raised the temperature to 160°C. The heat was turned back on to raise the temperature to 175°C. The reaction mixture was held at 175°C for five hours and cooled down to 125°C by adding 360 gms. of 2-butoxy ethanol. When the temperature was at 125°C, a mixture of 7.32 gms. of polyphosphoric acid and 20 gms. 2-butoxy ethanol was added slowly and held for 30 minutes. Thirty-four gms. of water was then added slowly to the reaction mixture and held at 125°C for two hours. After the two hours hold, 888 gms. of BuOH was added to the reaction mixture and the temperature was dropped to 113°C. A mixture of 283 gms. of methacrylic acid, 148 gms. of styrene, and 4 gms. of ethyl acrylate, 38.5 gms. of benzoyl peroxide and 32.4 gms. of 2-butoxy ethanol was added over two hours and then the reaction mixture was held at 114°C for additional two hours. The resulting resin was then dropped into the mixture of 3867 gms. of DI water and 162 gms. of dimethyl ethanolamine. The phosphated epoxy-acrylic graft copolymer was mixed with a latex (Example 13). The resulting emulsion was a stable opalescence emulsion which gave good protective coating for steel can in softdrink application when cured with 5% Cymel 303.

Example 29

Thermosetting copolymers containing alkyl hydroxy phosphate surfactant

Alkyl hydroxy phosphates were prepared by the reaction of superphosphoric acid with alkyl epoxides by the method disclosed in U.S. 3,487,130. By using 1,2-epoxyhexadecane, 240 parts of 1,2-epoxyhexadecane were added dropwise to a stirred solution of 105 parts superphosphoric acid in 200 parts tetrahydrofuran. During addition of epoxyhexadecane, the reaction mixture was maintained at 10–15°C, after which time it was slowly raised to room temperature. 300 parts of diethyl ether were then added to the batch, and the solution washed with two 200 ml portions of saturated sodium chloride solution. The organic layer was dried with sodium sulfate and solvent evaporated to yield about 300 parts hexadecane hydroxy phosphate. Under the preparative conditions, mono and diesters comprise the majority of reaction products. Separation of hydroxy phosphate mono and diesters was not required for use in water dispersed copolymers nor in epoxy/acrylic blends.

A thermosetting emulsion copolymer containing the foregoing alkyl hydroxy phosphate as a surfactant was prepared by loading 115 parts deionized water and 0.10 parts hexadecane hydroxy phosphate into the reactor and heated to 50–52°C. At 50°C the pH of the solution was adjusted to 8.5–9.0 by the addition of dimethyl ethanolamine and then heated under nitrogen sparge with agitation to 76–78°C. At equilibrated reaction temperature, the nitrogen sparge was discontinued, and a mixture of 1.2 parts styrene and 0.80 parts ethyl acrylate was added to the reactor and emulsified 10 minutes. After this time, 0.30 parts ammonium persulfate were added to the reactor, and allowed to react at least 20 minutes before beginning the monomer feed.

The monomer feed consisted of 55.0 parts styrene, 26.0 parts ethyl acrylate, 5.0 parts methacrylic acid and 12.0 parts N-isobutoxy methyl acrylamide. Monomers were added to the reactor at a constant rate during 4.5–5.0 hours while maintaining the batch temperature at 76–78°. The batch was then held an additional two hours at reaction temperature, before cooling and filtration.

Example 30

Thermosetting copolymers containing phosphate functionality

Monomers containing phosphate functionality were prepared by the reaction of phosphorus pentoxide with hydroxyl ethyl acrylate using the method disclosed in U.S. 3,686,371. In this method, 140 parts of phosphorus pentoxide were added slowly to a vigorously stirred solution of 350 parts hydroxyl ethyl acrylate and 0.05 parts MEHQ inhibitor. Reaction temperature was maintained at 25–35° with external cooling. After addition of phosphorus pentoxide was complete, the reaction was stirred for 2.5 hours at 30°C and then heated for two hours at 45° to insure complete reaction. Both mono and diesters were present in the reaction mixture but did not require separation prior to polymerization.

Phosphate monomers can be emulsion or solution copolymerized using standard semi-continuous methodology. Solution polymerization is best accomplished in a 1:2 mixture of butoxy-ethanol and n-butanol. Emulsion polymerization can proceed in aqueous media without the preserve of additional organic solvents.

A thermosetting emulsion copolymer containing phosphate monomer was prepared by loading 110 parts deionized water and 0.10 parts sodium dihexyl sulfosuccinate into the reactor. The solution was heated under nitrogen sparge to 76—78°C. At equilibrated reaction temperature, the nitrogen sparge was removed and 0.10 parts ammonium bicarbonate added. A mixture of 1.2 parts styrene and 0.80 parts ethyl acrylate was then added to the reactor and emulsified 10 minutes whereupon 0.30 parts ammonium persulfate were added and allowed to react 20 minutes before beginning the monomer feed.

The monomer mix contained 35.5 parts styrene, 50.0 parts ethyl acrylate, 4.0 parts methacrylic acid, 10.0 parts N-isobutoxy methyl acrylamide, and 0.50 parts 2-phosphate ethyl methacrylate. Monomers were added to the reactor at constant rate during 4.5—5.0 hours while maintaining the reaction temperature at 76—78°C. The batch was then held an additional two hours at reaction temperature, during which time an additional 400 parts deionized water was added to reduce viscosity.

Example 31

Thermosetting copolymers containing alkyl phosphates

Alkyl phosphates were prepared by the reaction of superphosphoric acid with aliphatic alcohols. The procedure employed is identical to that used to phosphate hydroxyl alkyl acrylates, as illustrated in Example 29.

A thermosetting emulsion copolymer containing alkyl phosphate was prepared by loading 115.0 parts deionized water and 0.10 parts octyl phosphate into the reactor and heated to 50—52°C. At 50—52°, pH of the aqueous phase was adjusted to pH 7—7.5 with dimethyl ethanolamine, and then heated under nitrogen sparge with agitation to 76—78°C. At equilibrated reaction temperature, the nitrogen sparge was discontinued, and a mixture of 1.2 parts styrene and 0.80 parts ethyl acrylate was added to the reactor. This monomer portion was emulsified 10 minutes and then 0.30 parts ammonium persulfate were added to the reactor and allowed to react 20 minutes before beginning the monomer feed.

The monomer mix contained 45.0 parts styrene, 38.0 parts ethyl acrylate, 5.0 parts methacrylic acid, and 10.0 parts N-isobutoxy methyl acrylamide. Monomers were added to the reactor during 3.5—4.0 hours while maintaining the batch temperature at 76—78°C. During the monomer feed, an additional 900 parts of deionized water were added to the batch to reduce viscosity. The batch was held an additional three hours at reaction temperature before cooling and filtration.

The foregoing emulsions produced in Examples 29—31 were each mixed with an epoxy-acrylic copolymer in accordance with this invention to produce an excellent interior can coating.

Example 32

Preparation of water solution dispersion

A 5L flask equipped with a thermometer, a condenser, a stirrer, and a N₂ inlet was charged with a mixture of 1000 gms. of DI water and 114.6 gms. of dimethyl aminoethanol. The mixture was heated to 85°C. A monomer mixture containing 548 gms. of styrene, 487.1 gms. of ethyl acrylate, 121.7 gms. of methacrylic acid, 24.3 gms. of benzoin, 379.5 gms. of butyl cellosolve, 379.5 gms. of n-butanol, 60.9 gms. of N(isobutoxy methyl)acrylamide, and 34.8 gms. of tert-butyl hydroperoxide was added over 4 hrs. When the addition of monomer mixture was completed, the reaction mixture was held at 85°C for 1 hour. Then, 900 gms. of DI water was added and followed by a mixture of 100 gms. of DI water and 34.8 gms. of tert-butyl hydroperoxide. The reaction mixture was held at 85°C for another 2 hrs. to complete the polymerization.

Blend composition

To a 5L flask equipped with a condenser and a stirrer, 1514 gms. of water soluble dispersion from the foregoing water soluble dispersion were added along with 463 gms. of DI water. The mixture was mixed for 15 minutes. Next, 200.3 gms. of epoxy phosphate from Example 15 was added and rinsed with 200 gms. of DI water. The resulting mixture was mixed for 1 hr., then 379.1 gms. of epoxy acrylic copolymers from Example 14 was added and rinsed with 200 gms. of DI water. The mixture was stirred for 1 hr. A mixture of 13 gms. of n-butanol and 29.1 gms. of Cymel 303 was then added and followed by 100 gms. of DI water. The resulting mixture was mixed for 2 hrs. and then filtered.

The resulting polymeric mixture was cured and provided good interior can coating.

Example 33

Blend with ammonium phosphate

To a 5L flask equipped with a condenser and a stirrer, 1000 gms. of latex from Example 1 were added along with 1000 gms. of DI water. A mixture of 18.1 gms. of dimethyl ethanol amine and 50 gms. water was then added slowly. Next, 76.6 gms. of butyl cellosolve and 222.8 gms. of n-butanol were added to the mixture. Thereafter, 715 gms. of epoxy acrylic copolymer from Example 14 was added and continued to stir for 2 hrs. Then, 1.33 gms. of ammonium phosphate was added and continued to stir for an additional 1 hr.

The resulting mixture was then mixed with 5% Cymel 303. The coating was cured into a clear film with good film properties.

Claims

1. A coating composition comprising an aqueous dispersion of a self curing copolymer of ethylenically unsaturated alkylol acrylamide monomer, ethylenically unsaturated functional monomer containing at least one of carboxyl, hydroxy, amino or amido groups and a further ethylenically unsaturated monomer lacking such functional groups, said polymer having been prepared by a two stage emulsion polymerization wherein the alkylol acrylamide monomers are polymerized in the second polymerization step so as to concentrate alkylol acrylamide on the polymer surface and effect crosslinking with a minor amount of said functional monomer to result in a relatively rigid polymer particle surface.
2. A water dispersed coating composition according to Claim 1, comprising on a polymer solids weight basis:
 - between 5% and 99% of said self-curing water dispersed polymer comprising copolymerized ethylenically unsaturated alkylol acrylamide monomer, ethylenically unsaturated functional monomer containing reactive carboxyl, hydroxy, amino or amido functional groups, and other ethylenically unsaturated monomer;
 - between 1% and 70% epoxy-acrylic copolymer; and
 - between 0% and 50% of a phosphated polymer.
3. A composition as claimed in Claim 2, characterized in that said phosphated polymer is present in an amount of from 3 to 20% on a polymer solids weight basis.
4. A composition as claimed in any one of the preceding claims, wherein the composition further comprises from 1 to 10% of an aminoplast resin on a polymer solids weight basis.
5. A composition as claimed in any one of the preceding claims, wherein the alkylol acrylamide component of the self curing water dispersed polymer is an alkylated alkylol acrylamide.
6. A composition according to any one of the preceding claims, wherein the epoxy acrylic copolymer is a graft copolymer, in admixture with ungrafted polymer, epoxy resin and ungrafted acrylic admixture polymer.
7. A composition according to any one of the preceding claims, wherein said epoxy acrylic is a phosphated epoxy-acrylic polymer.
8. A composition according to any one of the preceding claims, wherein said epoxy acrylic copolymer is an epoxy acrylic ester copolymer.
9. A composition according to any one of the preceding claims, wherein said self-curing water dispersed polymer was formed by copolymerization of its constituent unit in the presence of a phosphate or phosphoric acid.
10. A composition according to any one of the preceding claims, wherein a phosphated polymer is present and said phosphated polymer comprises an epoxy phosphate.
11. A composition according to any one of the preceding claims, wherein a phosphated polymer is present and is selected from a) phosphated resinous polyelectrolyte polymers produced by reacting ethylenically unsaturated monomers including carboxyl or hydroxyl monomer in water containing an ionizing agent, b) phosphated epoxidized oil c) phosphated epoxidized butadiene copolymer, d) polymer containing copolymerized phosphate monomer, e) phosphated polyester.
12. A water dispersed coating composition according to Claim 1, containing said water dispersed self-curing polymer comprising copolymerized ethylenically unsaturated alkylol acrylamide monomer, ethylenically unsaturated functional monomer containing reactive functional groups, and other ethylenically unsaturated monomer, wherein the self-curing polymer contains phosphate.
13. The composition in Claim 12, wherein the phosphate is an alkyl phosphate ester monomer copolymerized with the other monomers.
14. A process for preparing a self-curing water dispersed polymer capable of use in the composition of Claim 1, characterized in that (a) alkylol acrylamide, (b) functional monomer selected from carboxyl, hydroxyl, amine and amide monomers and (c) other ethylenic monomer is emulsion polymerized to two steps, polymerization of alkylol acrylamide being effected in the second step, said polymerization being effected in such manner that in the second step a minor amount of said alkylol acrylamide monomer reacts with a minor amount of said functional monomer to result in a relatively rigid polymer particle surface.
15. A process according to Claim 14, wherein said acrylamide monomer is an alkylated alkylol acrylamide monomer.
16. A process according to Claim 15, wherein said alkylated alkylol acrylamide monomer is a butylated alkylol acrylamide monomer.
17. A process according to any one of Claims 14—16, wherein said ethylenically unsaturated monomers comprise by weight between 1% and 20% alkylol acrylamide monomer, between 1% and 20% functional monomer, the remainder being other ethylenic monomer.
18. A process according to any one of Claims 14—17, wherein the monomers for the self-curing polymers are polymerized in the presence of ionizing agent to produce a resinous polyelectrolyte water dispersed self-curing polymer.

The resulting mixture was then mixed with 5% Cymel 303. The coating was cured into a clear film with good film properties.

Claims

5

1. A coating composition comprising an aqueous dispersion of a self curing copolymer of ethylenically unsaturated alkylol acrylamide monomer, ethylenically unsaturated functional monomer containing at least one of carboxyl, hydroxy, amino or amido groups and a further ethylenically unsaturated monomer lacking such functional groups, said polymer having been prepared by a two stage emulsion polymerization wherein the alkylol acrylamide monomers are polymerized in the second polymerization step so as to concentrate alkylol acrylamide on the polymer surface and effect crosslinking with a minor amount of said functional monomer to result in a relatively rigid polymer particle surface.

10

2. A water dispersed coating composition according to Claim 1, comprising on a polymer solids weight basis:

15

between 5% and 99% of said self-curing water dispersed polymer comprising copolymerized ethylenically unsaturated alkylol acrylamide monomer, ethylenically unsaturated functional monomer containing reactive carboxyl, hydroxy, amino or amido functional groups, and other ethylenically unsaturated monomer;

20

between 1% and 70% epoxy-acrylic copolymer; and

between 0% and 50% of a phosphated polymer.

3. A composition as claimed in Claim 2, characterized in that said phosphated polymer is present in an amount of from 3 to 20% on a polymer solids weight basis.

4. A composition as claimed in any one of the preceding claims, wherein the composition further comprises from 1 to 10% of an aminoplast resin on a polymer solids weight basis.

25

5. A composition as claimed in any one of the preceding claims, wherein the alkylol acrylamide component of the self curing water dispersed polymer is an alkylated alkylol acrylamide.

6. A composition according to any one of the preceding claims, wherein the epoxy acrylic copolymer is a graft copolymer, in admixture with ungrafted polymer, epoxy resin and ungrafted acrylic admixture polymer.

30

7. A composition according to any one of the preceding claims, wherein said epoxy acrylic is a phosphated epoxy-acrylic polymer.

8. A composition according to any one of the preceding claims, wherein said epoxy acrylic copolymer is an epoxy acrylic ester copolymer.

35

9. A composition according to any one of the preceding claims, wherein said self-curing water dispersed polymer was formed by copolymerization of its constituent unit in the presence of a phosphate or phosphoric acid.

10. A composition according to any one of the preceding claims, wherein a phosphated polymer is present and said phosphated polymer comprises an epoxy phosphate.

40

11. A composition according to any one of the preceding claims, wherein a phosphated polymer is present and is selected from a) phosphated resinous polyelectrolyte polymers produced by reacting ethylenically unsaturated monomers including carboxyl or hydroxyl monomer in water containing an ionizing agent, b) phosphated epoxidized oil c) phosphated epoxidized butadiene copolymer, d) polymer containing copolymerized phosphate monomer, e) phosphated polyester.

45

12. A water dispersed coating composition according to Claim 1, containing said water dispersed self-curing polymer comprising copolymerized ethylenically unsaturated alkylol acrylamide monomer, ethylenically unsaturated functional monomer containing reactive functional groups, and other ethylenically unsaturated monomer, wherein the self-curing polymer contains phosphate.

13. The composition in Claim 12, wherein the phosphate is an alkyl phosphate ester monomer copolymerized with the other monomers.

50

14. A process for preparing a self-curing water dispersed polymer capable of use in the composition of Claim 1, characterized in that (a) alkylol acrylamide, (b) functional monomer selected from carboxyl, hydroxyl, amine and amide monomers and (c) other ethylenic monomer is emulsion polymerized to two steps, polymerization of alkylol acrylamide being effected in the second step, said polymerization being effected in such manner that in the second step a minor amount of said alkylol acrylamide monomer reacts with a minor amount of said functional monomer to result in a relatively rigid polymer particle surface.

55

15. A process according to Claim 14, wherein said acrylamide monomer is an alkylated alkylol acrylamide monomer.

16. A process according to Claim 15, wherein said alkylated alkylol acrylamide monomer is a butylated alkylol acrylamide monomer.

60

17. A process according to any one of Claims 14—16, wherein said ethylenically unsaturated monomers comprise by weight between 1% and 20% alkylol acrylamide monomer, between 1% and 20% functional monomer, the remainder being other ethylenic monomer.

18. A process according to any one of Claims 14—17, wherein the monomers for the self-curing polymers are polymerized in the presence of ionizing agent to produce a resinous polyelectrolyte water dispersed self-curing polymer.

65



19. A process for preparing a copolymer as claimed in any one of Claims 2—10, which comprises first forming a self-curing water dispersed polymer according to any one of Claims 14—18 and then mixing said self-curing water dispersed polymer with an epoxy-acrylic copolymer and optionally phosphated polymer to provide a water dispersed composition comprising between 1% and 99% of said self-curing polymer, between 1% and 70% of said epoxy-acrylic copolymer, and 0% to 50% phosphated polymer.

20. A process according to any one of Claims 19, wherever the monomers are phosphated in the presence of a phosphate surfactant.

21. The use of a composition according to any one of Claims 1—13, which comprises spraying an application of said composition to a substrate and then heat curing to cross-link said functional groups with said alkylated acrylamide groups.

22. A composition according to any one of Claims 2—6, wherein said epoxy acrylic copolymer is quaternary polymer.

23. A composition according to any one of Claims 2—6, wherein said epoxy acrylic copolymer is produced by reaction epoxy resin with a melamine graft copolymer mixture of melamine and acrylic copolymer in the presence of sufficient amine to produce a non-gelled epoxy-acrylic ester copolymer.

Patentansprüche

1. Beschichtungszusammensetzung umfassend eine wässrige Dispersion eines selbsthärtenden Copolymers von äthylenisch ungesättigtem Alkylolacrylamidmonomer, äthylenisch ungesättigtem funktionellen Monomer mit mindestens einer Carboxyl-, Hydroxyl-, Amino- oder Amidogruppe und einem weiteren äthylenisch ungesättigten Monomer, dem solche funktionelle Gruppen fehlen, welches Polymer durch eine zweistufige Emulsionspolymerisation hergestellt wurde, in der die Alkylolacrylamidmonomere im zweiten Polymerisationsschritt polymerisiert werden, um das Alkylolacrylamid an der Polymeroberfläche zu konzentrieren.

2. In Wasser dispergierte Beschichtungszusammensetzung nach Anspruch 1, umfassend, bezogen auf das Polymerfeststoffgewicht,

5 bis 99% des genannten selbsthärtenden in Wasser dispergierten Polymers umfassend copolymerisiert äthylenisch ungesättigtes Alkylolacrylamidmonomer, äthylenisch ungesättigtes funktionelles Monomer mit reaktiven funktionellen Carboxyl-, Hydroxyl-, Amino- oder Amidogruppen und ein anderes äthylenisch ungesättigtes Monomer;

1 bis 70% Epoxyacrylcopolymer und
0 bis 50% eines phosphatierten Polymers.

3. Zusammensetzung, wie in Anspruch 2 beansprucht, dadurch gekennzeichnet, daß das genannte phosphatierte Polymer in einem Anteil von 3 bis 20%, bezogen auf das Polymerfeststoffgewicht, vorhanden ist.

4. Zusammensetzung, wie in einem der vorhergehenden Ansprüche, beansprucht, wobei die Zusammensetzung auch 1 bis 10% eines Aminoplastharzes, bezogen auf das Polymerfeststoffgewicht, aufweist.

5. Zusammensetzung, wie in einem der vorhergehenden Ansprüche beansprucht, worin die Alkylolacrylamidkomponente des selbsthärtenden, in Wasser dispergierten Polymers ein alkyliertes Alkylolacrylamid ist.

6. Zusammensetzung nach einem der vorhergehenden Ansprüche, worin das Epoxyacrylcopolymer ein Pfropfcopolymer in Mischung mit einem nicht-gepfropften Polymer, einem Epoxyharz und einem nichtgepfropften Acrylzusatzpolymer ist.

7. Zusammensetzung nach einem der vorhergehenden Ansprüche, worin das Epoxyacrylcopolymer ein phosphatiertes Epoxyacrylpolymer ist.

8. Zusammensetzung nach einem der vorhergehenden Ansprüche, worin das Epoxyacrylcopolymer ein Epoxyacrylestercopolymer ist.

9. Zusammensetzung nach einem der vorhergehenden Ansprüche, worin das selbsthärtende, in Wasser dispergierte Polymer durch Copolymerisation seiner Konstituenteneinheit in Anwesenheit eines Phosphats oder von Phosphorsäure gebildet wurde.

10. Zusammensetzung nach einem der vorhergehenden Ansprüche, worin ein phosphatiertes Polymer vorhanden ist und dieses phosphatierte Polymer ein Epoxyphosphat umfaßt.

11. Zusammensetzung nach einem der vorhergehenden Ansprüche, worin ein phosphatiertes Polymer vorhanden ist, das ausgewählt ist aus a) phosphatierten harzartigen Polyelektrolytpolymeren, hergestellt durch Umsetzen von äthylenisch ungesättigten Monomeren einschließlich Carboxyl- oder Hydroxylmonomer in ein Ionisierungsmittel enthaltendem Wasser, b) phosphatiertem epoxidierten Öl, c) phosphatiertem epoxidierten Butadiencopolymer, d) copolymerisiertes Phosphatmonomer enthaltendem Polymer und e) phosphatiertem Polyester.

12. In Wasser dispergierte Beschichtungszusammensetzung nach Anspruch 1, die das genannte, in Wasser dispergierte selbsthärtende Polymer enthält, das copolymerisiert äthylenisch ungesättigtes Alkylolacrylamidmonomer, äthylenisch ungesättigtes funktionelles Monomer mit reaktiven funktionellen Gruppe und ein anderes äthylenisch ungesättigtes Monomer umfaßt, wobei das selbsthärtende Polymer Phosphat enthält.



13. Zusammensetzung nach Anspruch 12, worin das Phosphat, ein Alkylphosphatestermonomer ist, das mit den anderen Monomeren copolymerisiert ist.

14. Verfahren zum Herstellen eines selbsthärtenden, in Wasser dispergierten Polymers, das in der Zusammensetzung nach Anspruch 1 verwendet werden kann, dadurch gekennzeichnet, daß (a) Alkylolacrylamid, (b) funktionelles Monomer ausgewählt aus Carboxyl-, Hydroxyl-, Amin- und Amidmonomeren und (c) ein anderes äthylenisches Monomer in zwei Schritten emulsionspolymerisiert werden, wobei die Polymerisation des Alkylolacrylamids im zweiten Schritt bewirkt wird und die Polymerisation auf solche Weise durchgeführt wird, daß im zweiten Schritt ein geringer Anteil des Alkylolacrylamidmonomers mit einem geringen Anteil des funktionellen Monomers reagiert, wobei eine relativ starre Polymerteilchenoberfläche erhalten wird.

15. Verfahren nach Anspruch 14, in dem das Acrylamidmonomer ein alkyliertes Alkylolacrylamidmonomer ist.

16. Verfahren nach Anspruch 15, in dem das alkylierte Alkylolacrylamidmonomer ein butyliertes Alkylolacrylamidmonomer ist.

17. Verfahren nach einem der Ansprüche 14 bis 16, in dem die Äthylenisch ungesättigten Monomer 1 bis 20 Gew.% Alkylolacrylamid und 1 bis 20 Gew.% funktionelles Monomer umfassen, wobei der Rest ein anderes äthylenisches Monomer ist.

18. Verfahren nach einem der Ansprüche 14 bis 17, in dem die Monomer für die selbsthärtenden Polymere in Anwesenheit des Ionisierungsmittels polymerisiert werden, wobei ein harzartiges in Wasser dispergiertes selbsthärtendes Polyelektrolytpolymer erhalten wird.

19. Verfahren zum Herstellen eines Copolymers, wie in einem der Ansprüche 2 bis 10 beansprucht, in dem zuerst ein selbsthärtendes, in Wasser dispergiertes Polymer nach einem der Ansprüche 14 bis 18 gebildet wird und dann dieses selbsthärtende, in Wasser dispergierte Polymer mit einem Epoxyacrylcopolymer und gegebenenfalls phosphatiertem Polymer unter Bildung einer in Wasser dispergierten Zusammensetzung gemischt wird, die 1 bis 99% des selbsthärtenden Polymers, 1 bis 70% des Epoxyacrylcopolymers und 0 bis 50% phosphatiertes Polymer umfaßt.

20. Verfahren nach Anspruch 19, in dem die Monomere in Anwesenheit eines oberflächenaktiven Phosphats phosphatiert werden.

21. Verwendung einer Zusammensetzung nach einem der Ansprüche 1 bis 13, die das Aufsprühen der genannten Zusammensetzung auf ein Substrat und das anschließende Wärmehärten zum Vernetzen der funktionellen Gruppen mit den alkylierten Acrylamidgruppen umfaßt.

22. Zusammensetzung nach einem der Ansprüche 2 bis 6, worin das Epoxyacrylcopolymer ein quaternäres Polymer ist.

23. Zusammensetzung nach einem der Ansprüche 2 bis 6, worin das Epoxyacrylcopolymer durch Umsetzen von Epoxyharz mit einer Melaminpfropfcopolymermischung von Melamin und Acrylcopolymer in Anwesenheit von ausreichende Amin zur Bildung eines nicht-gelierten Epoxyacrylestercopolymers hergestellt ist.

40 Revendications

1. Composition d'enduction comprenant une dispersion aqueuse d'un copolymère autoréticulable d'un monomère alkylolacrylamide à insaturation éthylénique, d'un monomère fonctionnel à insaturation éthylénique renfermant au moins un groupe carboxyle, hydroxy, amino ou amido et un autre monomère à insaturation éthylénique dépourvu d'un tel groupe fonctionnel, ledit polymère ayant été préparé par polymérisation en émulsion à deux stades dans laquelle les alkylolacrylamides monomères sont polymérisés au second stade de polymérisation de manière à concentrer l'alkylolacrylamidesur la surface du polymère.

2. Composition d'enduction dispersée dans l'eau selon la revendication 1, qui comprend par rapport au poids des solides du polymère:

de 5 à 99% dudit polymère autoréticulable dispersé dans l'eau comprenant un alkylolacrylamide à insaturation éthylénique copolymérisé, un monomère fonctionnel à insaturation éthylénique contenant des groupes fonctionnels carboxyle, hydroxy, amino ou amido et un autre monomère à insaturation éthylénique;

de 1 à 70% d'un copolymère époxy-acrylique; et
de 0 à 50% d'un polymère phosphaté.

3. Composition selon la revendication 2, caractérisé en ce que ledit polymère phosphaté est présent à raison de 3 à 20% par rapport au poids des solides du polymère.

4. Composition selon l'une quelconque des revendications précédentes, dans laquelle la composition comprend également de 1 à 10% d'une résine aminoplaste par rapport au poids des solides du polymère.

5. Composition selon l'une quelconque des revendications précédentes, dans laquelle le composant alkylolacrylamide du polymère autoréticulable dispersé dans l'eau est un alkylolacrylamide alkylé.

6. Composition selon l'une quelconque des revendications précédentes, dans laquelle le copolymère époxy-acrylique est un copolymère greffé, en mélange avec un polymère non greffé, une résine époxy et un polymère mélangé acrylique non greffé.



7. Composition selon l'une quelconque des revendications précédentes, dans laquelle ledit époxy-acrylique est un polymère époxy acrylique phosphaté.

8. Composition selon l'une quelconque des revendications précédentes, dans laquelle ledit copolymère époxy-acrylique est un polymère époxy-ester acrylique.

9. Copolymère selon l'une quelconque des revendications précédentes, dans lequel ledit polymère autoréticulable dispersé dans l'eau est formé par copolymérisation de son motif constituant en présence d'un phosphate ou d'un acide phosphorique.

10. Composition selon l'une quelconque des revendications précédentes, dans laquelle est présent un polymère phosphaté et ledit polymère phosphaté comprend un époxy phosphate.

11. Composition selon l'une quelconque des revendications précédentes, dans laquelle un polymère phosphaté est présent et il est choisi parmi (a) des polymères de polyélectrolytes résineux phosphatés qu'on obtient en faisant réagir des monomères à insaturation éthylénique, notamment un monomère carboxyle ou hydroxyle dans l'eau contenant un agent ionisant, (b) une huile époxydée phosphatée, (c) un copolymère de butadiène époxydé phosphaté, (d) un polymère contenant un phosphate monomère copolymérisé (e) un polyester phosphaté.

12. Composition d'enduction dispersée dans l'eau selon la revendication 1, contenant ledit polymère autoréticulable dispersé dans l'eau comprenant un alkylolacrylamide copolymérisé à insaturation éthylénique, un monomère fonctionnel à insaturation éthylénique contenant des groupes fonctionnels réactifs et un autre monomère à insaturation éthylénique, le polymère autoréticulable contenant un phosphate.

13. Composition selon la revendication 12, dans laquelle le phosphate est un alkylphosphate monomère copolymérisé avec les autres monomères.

14. Procédé de préparation d'un polymère autoréticulable dispersé dans l'eau pouvant servir dans la composition selon la revendication 1, caractérisé en ce qu'on polymérise en émulsion en deux stades (a) un alkylolacrylamide, (b) un monomère fonctionnel choisi parmi les monomères carboxyle, hydroxyle, amine et amide et (c) un autre monomère éthylénique, la polymérisation de l'alkylolacrylamide étant effectuée au second stade, ladite polymérisation étant effectuée de manière qu'au second stade, une petite quantité dudit alkylolacrylamide monomère réagisse avec une petite quantité dudit monomère fonctionnel pour donner une surface relativement rigide aux particules du polymère.

15. Procédé selon la revendication 14, dans lequel ledit acrylamide monomère est un alkylolacrylamide monomère alkylé.

16. Procédé selon la revendication 15, dans lequel l'alkylolacrylamide alkylé est un alkylolacrylamide butylé.

17. Procédé selon l'une quelconque des revendications 14 à 16, dans lequel lesdits monomères à insaturation éthylénique comprennent, en poids, de 1 à 20% d'alkylolacrylamide monomère, de 1 à 20% d'un monomère fonctionnel, le complément étant l'autre monomère éthylénique.

18. Procédé selon l'une quelconque des revendications 14 à 17, dans lequel on polymérise les monomères pour former des polymères autoréticulables en présence d'un agent ionisant pour obtenir un polymère autoréticulable du type polyélectrolyte résineux dispersé dans l'eau.

19. Procédé de préparation d'un copolymère tel que défini dans l'une quelconque des revendications 2 à 10, qui consiste à former d'abord un polymère autoréticulable dispersé dans l'eau selon l'une quelconque des revendications 14 à 18 et ensuite à mélanger ledit polymères autoréticulables dispersé dans l'eau avec un copolymère époxy-acrylique et, facultativement, un polymère phosphaté pour former une composition dispersée dans l'eau comprenant de 1 à 99% dudit polymère autoréticulable, de 1 à 70% dudit copolymère époxy-acrylique et 0 à 50% du polymère phosphaté.

20. Procédé selon la revendication 19, dans lequel les monomères sont phosphatés en présence d'un phosphate surfactif.

21. Utilisation d'une composition selon l'une quelconque des revendications 1 à 13, qui consiste à pulvériser une dose de ladite composition sur un substrat et ensuite à durcir thermiquement afin de réticuler lesdits groupes fonctionnels avec lesdits acrylamides alkylés.

22. Composition selon l'une quelconque des revendications 2 à 6, dans laquelle ledit copolymère époxy-acrylique est un polymère quaternaire.

23. Composition selon l'une quelconque des revendications 2 à 6, dans laquelle ledit copolymère époxy-acrylique est produit par réaction d'une résine époxy avec un mélange copolymère greffé de mélamine et de copolymère acrylique en présence d'une quantité suffisante d'amine pour obtenir un copolymère non gélifié époxy/ester acrylique.

