

IMPACT MODIFIED VINYL CHLORIDE
RESIN COMPOSITION, AND
GRAFT COPOLYMER COMPOSITIONS

BACKGROUND OF THE INVENTION

Field of the Invention

The present invention relates to vinyl chloride resin compositions and impact modifiers, and more particularly relates to impact modified vinyl chloride resin compositions and graft copolymer compositions.

Description of the Related Art

Polyvinyl chloride resins and rigid vinyl aromatic-vinyl cyanide resins are generally relatively brittle, and consequently are typically impact modified with graft copolymers such as acrylonitrile-styrene-butadiene graft copolymers (ABS) comprising a butadiene rubber substrate and an acrylonitrile-styrene superstrate. Traditional surfactants based on linear fatty acids, used in emulsion processes for making the ABS resin tend to fog or volatilize from the surface of ABS resin or blends thereof either in the molten state and/or in high temperature end-use environments such as is experienced in the interior of an automobile during the summer months. Individuals purchasing a new automobile have typically experienced the deposit of organic material on the interior of a windshield after 6 months to a year.



Accordingly, there is a need and desire for impact modified resin compositions which exhibit reduced levels of fogging.

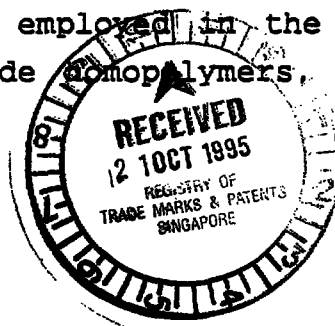
Summary of the Invention

5 An impact modified resin composition is provided comprising (a) an impact modifier containing a residual amount of a C_{30} to C_{100} polyacid surfactant. The compositions exhibit reduced levels of fogging, and are useful for making automotive dash components.
10 The impact modified composition further contains a vinyl chloride resin and/or a vinyl aromatic-unsaturated nitrile resin.

Detailed Description of the Invention

 A thermoplastic composition is provided
15 comprising (a) a vinyl chloride resin or vinyl aromatic-vinyl cyanide resin present at a level of from 20 to 95 percent by weight based on the total weight of the thermoplastic composition, more preferably from 60 to 92 percent by weight thereof,
20 and most preferably from 80 to 90 percent by weight thereof, and (b) an impact modifier composition present at a level of from 5 to 80 percent by weight based on the total weight of the thermoplastic composition, more preferably from 8 to 40 percent by
25 weight thereof, and most preferably 10 to 20 percent by weight thereof. Optionally, the thermoplastic resin compositions containing the vinyl chloride resin may further an amount of (c) a non-graft non-rubber vinyl polymer at a level of from, for example,
30 5 to 40 percent by weight and for further example from 10 to 20 percent by weight based on the total weight of the thermoplastic composition.

Vinyl chloride resin employed in the present invention is vinyl chloride homopolymers, such as



polyvinyl chloride, and copolymers prepared by copolymerizing more than 50 percent by weight of vinyl chloride and at least one monoolefinic monomer, such as vinyl acetate, vinyl stearate, acrylic ester, methacrylic ester, styrene, acrylonitrile, ethylene, propylene and the like.

Vinyl aromatic-vinyl cyanide resin comprises from 60 to 90 percent by weight vinyl aromatic monomer based on the total weight of the vinyl aromatic-vinyl cyanide resin, more preferably from 65 to 80 percent by weight thereof, and most preferably from 72 to 75 percent by weight thereof; a vinyl cyanide (unsaturated nitrile) monomer is present at a level of from 10 to 40 percent by weight, more preferably from 20 to 35 percent by weight and most preferably from 25 to 28 percent by weight.

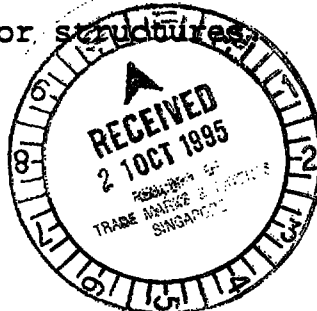
The rubber graft copolymer useful in the impact modifier composition (graft copolymer composition) comprises (i) the rubber substrate, and (ii) a rigid polymeric superstrate portion grafted to the rubber substrate, and optionally a free rigid component. The rubber substrate is preferably present in the impact modifier composition at a level of from 10 to 90 percent by weight based on the total weight of the graft copolymer, more preferably from 40 to 85 percent by weight thereof, and more preferably 50 to 85 percent by weight thereof and most preferably from 65 to 85 percent by weight thereof; and the rigid superstrate (and optional free rigid in combination) is preferably present at a level of from 10 to 90 percent by weight based on the total weight of the graft copolymer, preferably from 15 to 60 percent by weight thereof, more preferably from 15 to 50 percent by weight thereof, and most preferably from 15 to 35



percent by weight thereof. The free rigid portion, if present, is at about 0-50% by weight based on the total weight of the composition. The rubber substrate is a rubber having a T_g (glass transition temperature) of less than 0°C.

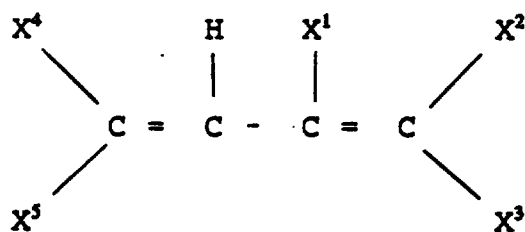
Examples of rubbery polymers for the substrate include: conjugated dienes, copolymers of a diene with styrene, acrylonitrile, methacrylonitrile or C₁ to C₈ alkyl acrylate which contain at least 50% (preferably at least 65% by weight) conjugated dienes, polyisoprene or mixtures thereof; olefin rubbers i.e. ethylene propylene copolymer (EPR) or ethylene propylene non-conjugated diene (EPDM); silicone rubbers; or C₁ or C₈ alkyl acrylate homopolymers or copolymers with butadiene and/or styrene. The acrylic polymer may also contain up to 5% of one or more polyfunctional crosslinking agents such as alkylenediol di(meth)acrylates, alkyleneetriol tri(meth)acrylates, polyester di(meth)acrylates, divinylbenzene, trivinylbenzene, butadiene, isoprene and optionally graftable monomers such as, triallyl cyanurate, triallyl isocyanurate, allyl (meth)acrylate, diallyl maleate, diallyl fumarate, diallyl adipate, triallyl esters of citric acid or mixtures of these agents.

The diene rubbers may preferably be polybutadiene, polyisoprene and copolymers of butadiene with up to 35% by weight of comonomers such as styrene, acrylonitrile, methylmethacrylate or C₁-C₈-alkylacrylate which are produced by aqueous radical emulsion polymerisation and may optionally be cross-linked with di or trivinyl or allyl monomers or mixtures of such monomers or structures.



The rubbers may be cross-linked, particulate emulsion copolymers substantially of C₁-C₈-alkylacrylate, in particular C₂-C₆-alkylacrylate, optionally in admixture with up to 15% by weight of comonomers such as styrene, methylmethacrylate, butadiene, vinyl methyl ether or acrylonitrile and optionally up to 5% by weight of a polyfunctional crosslinking comonomer, e.g. divinylbenzene, glycol-bis-acrylates, bisacrylamides, phosphoric acid triallylester, citric acid triallylester, allylesters of acrylic acid or methacrylic acid, triallylcyanurate, triallylisocyanurate. Also suitable are mixtures of diene- and alkylacrylate rubbers and rubbers which have a so-called core/sheath structure, e.g. a core of diene rubber and a sheath of acrylate or vice versa.

Specific conjugated diene monomers normally utilized in preparing the rubber substrate of the graft polymer are generically described by the following formula:



wherein each X¹, X², X³, X⁴ and X⁵ is individually selected from the group consisting of hydrogen, alkyl groups containing from one to five carbon atoms, chlorine or bromine. Examples of dienes that may be used are butadiene, isoprene, 1,3-heptadiene, methyl-1,3-pentadiene, 2,3-dimethylbutadiene, 2-ethyl-1,3-pentadiene 1,3- and 2,4-hexadienes, chloro and bromo

substituted butadienes such as dichlorobutadiene, bromobutadiene, dibromobutadiene, mixtures thereof, and the like. A preferred conjugated diene is 1,3 butadiene.

5 The substrate polymer, as mentioned, is preferably a conjugated diene polymer such as polybutadiene, polyisoprene, or a copolymer, such as butadiene-styrene, butadiene-acrylonitrile, or the like. The rubbery polymeric substrate portion must
10 exhibit a glass transition temperature (T_g) of less than about 0°C .

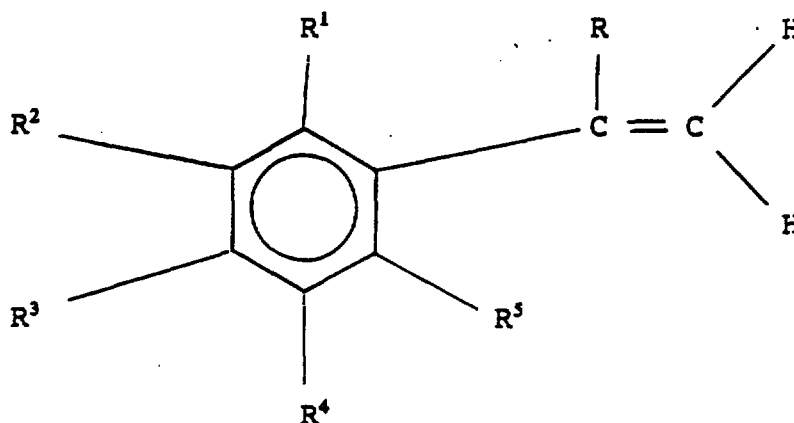
Mixtures of one or more rubbery polymers previously described for preparing the graft polymers, or mixtures of one or more graft polymers
15 disclosed herein may also be employed. Furthermore, the rubber may comprise either a block or random copolymer.

The rubber particle size used in this invention as measured by simple light transmission methods or
20 capillary hydrodynamic chromatography (CHDF) may be described as having an average particle size by weight of: 0.05 to 1.2 microns, preferably 0.2 to 0.8 microns, for the emulsion based polymerized rubber latices. The rubber substrate is preferably
25 a particulate, crosslinked diene or alkyl acrylate rubber, and preferably has a gel content greater than 50%.

Suitable superstrates include vinylidene polymers including homopolymers and copolymers.

30 Monovinylidene aromatic monomers (vinyl aromatic monomers) which may be employed in making of the superstrate include styrene, alpha-methyl styrene, halostyrenes i.e. dibromostyrene, mono or di alkyl, alkoxy or hydroxy substitute groups on the nuclear

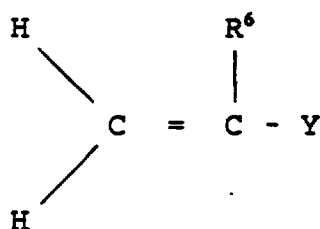
ring of the monovinylidene aromatic monomer i.e. vinyl toluene, vinylxylene, butylstyrene, para-hydroxystyrene or methoxystyrene or mixtures thereof. The monovinylidene aromatic monomers utilized are
 5 generically described by the following formula:



wherein each R^1 , R^2 , R^3 , R^4 and R^5 is individually selected from the group consisting of hydrogen, alkyl groups of 1 to 5 carbon atoms, cycloalkyl, aryl, alkaryl, aralkyl, alkoxy, aryloxy, and halogens. R
 10 is selected from the group consisting of hydrogen, alkyl groups of 1 to 5 carbon atoms and halogens such as bromine and chlorine. Examples of substituted vinylaromatic compounds include styrene, 4-methylstyrene, 3,5-diethylstyrene, 4-n-propylstyrene,
 15 α -methylstyrene, α -methyl vinyltoluene, α -chlorostyrene, α -bromostyrene, dichlorostyrene, dibromostyrene, tetrachlorostyrene, mixtures thereof and the like. The preferred monovinylidene aromatic monomers used are styrene and/or α -methylstyrene.

20 Monomers which may be used alone or in combination with the monovinylidene aromatic monomer includes acrylonitrile, methacrylonitrile, C_1 to C_8 alkyl or aryl substituted acrylate, C_1 to C_8 alkyl,

aryl or haloaryl substituted methacrylate, acrylic acid, methacrylic acid, itaconic acid, acrylamide, N-substituted acrylamide or methacrylamide, maleic anhydride, maleimide, N-alkyl, aryl or haloaryl substituted maleimide, glycidyl (meth)acrylates, hydroxy alkyl (meth)acrylates or mixtures thereof. The unsaturated nitrile (acrylonitrile or substituted acrylonitrile) or acrylic acid esters are described generically by the following formula:



wherein R^6 may be selected from the same group set out for R as previously defined and Y is selected from the group consisting of cyano and carbalkoxy groups wherein the alkoxy group of the carbalkoxy contains from one or about twelve carbon atoms. Examples of such monomers include acrylonitrile, ethacrylonitrile, methacrylonitrile, α -chloroacrylonitrile, α -bromoacrylonitrile, methyl acrylate, methyl methacrylate, ethyl acrylate, butyl acrylate, propylacrylate, isopropyl acrylate and mixtures thereof. The preferred unsaturate nitrile monomer is acrylonitrile and the preferred acrylic acid esters are ethyl acrylate and methyl methacrylate. It is also preferred that the acrylic acid esters, when included, are employed in combination with styrene or acrylonitrile.

The number average molecular weight of the grafted rigid superstrate of the monovinylidene

aromatic resin is designed to be in the range of 10,000 to 350,000. The ratio of monovinylidene aromatic monomer to the second and optionally third monomer may range from 100/0 to 50/50 preferably 85/15 to 60/40. Additional monomer(s) may optionally replace 0 to 50% of one or both of the first and second monomers. Preferred graft superstrates include copolymers of styrene and acrylonitrile, copolymers of α -methylstyrene and acrylonitrile, and methylmethacrylate polymers or copolymers with up to 50% by weight of C_1 - C_6 alkylacrylates, acrylonitrile or styrene.

These graft copolymers may be polymerized either by emulsion or combined processes such as emulsion-suspension or other emulsion techniques well known in the art. Furthermore, these rubber modified monovinylidene aromatic graft copolymers may be produced either by continuous, semibatch or batch processes.

Optionally the composition may include ungrafted (free) rigid phase portion which is a reaction product of the same monomers as set out for the superstrate, specifically vinyl aromatic monomers, unsaturated nitrile monomers, alkyl (alkyl) acrylate monomers and mixtures thereof. The free rigid phase is a term for the ungrafted rigid polymer. Preferably the free rigid phase and the grafted superstrate of the impact modifier composition are in respective ratios of 50:50 to 0:100, optionally from 30:70 to 5:95. In other words, it is preferred that the grafted superstrate be present in amounts much greater than the free rigid phase.

Alternatively, the monovinylidene aromatic monomers may be replaced in whole or in part with (or

be free of) acrylic acid esters such as methymethacrylate and/or ethylacrylate

The preferred impact modifier compositions (graft copolymer compositions) are compositions which
5 comprise amounts of a graft copolymer selected from the group consisting of acrylonitrile-butadiene-styrene graft copolymers, methymethacrylate-butadiene-styrene graft copolymers
10 methymethacrylate-alkyl acrylate grafted to butadiene or styrene-butadiene rubber, methymethacrylate-acrylonitrile-butadiene-styrene graft copolymers, methymethacrylate-ethylacrylate
15 onto polybutylacrylate butylacrylate-styrene-acrylonitrile graft copolymers, butylacrylate-ethylacrylate-styrene graft copolymers, and acrylonitrile-ethylacrylate-styrene graft copolymers.

Specific examples of monovinylidene aromatic graft copolymers include but are not limited to the following: acrylonitrile-butadiene-styrene (ABS),
20 acrylonitrile-styrene-butyl acrylate (ASA), methymethacrylate-acrylonitrile-butadiene-styrene (MABS), acrylonitrile-ethylene-propylene-non-conjugated diene-styrene (AES), methymethacrylate-butadiene-styrene and methymethacrylate-butadiene.

25 Suitable vinylidene rigid polymers including homopolymers and copolymers used in this invention and prepared independently from the vinylidene graft copolymer include monovinylidene aromatic monomers (first monomer) as described previously and other
30 comonomers (second monomer) also described earlier. The compositional ratio of the first monomer to the second may range from 100/0 to 50/50 preferably 85/15 to 60/40. Additional monomer(s) may optionally

replace 0 to 50% of one or both of the first and second monomers.

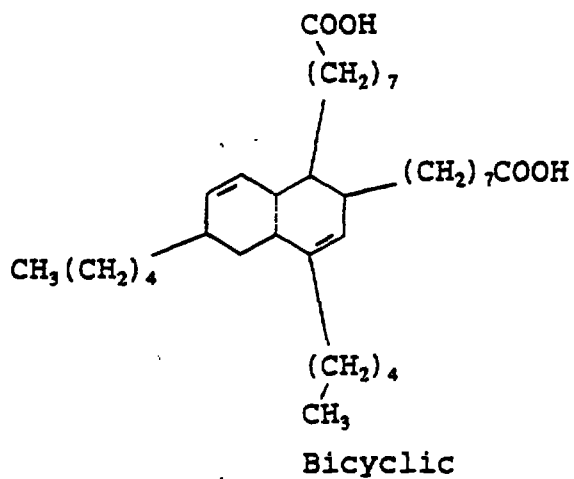
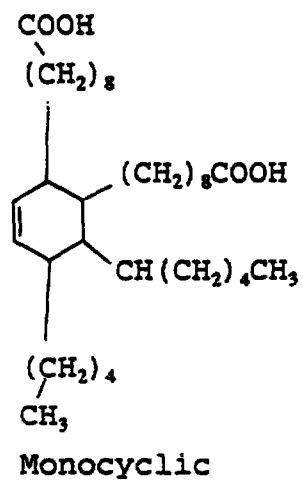
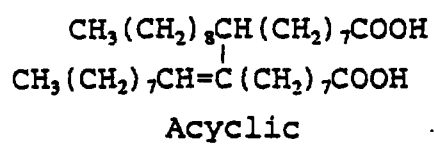
5 The number average molecular weight of the vinylidene rigid polymer is designed to be in the range of 10,000 to 350,000. Preferred vinylidene rigid polymers include copolymers of styrene and acrylonitrile, and homopolymers or copolymers of methylmethacrylate with up to 50% by weight of C_1 to C_8 alkylacrylates, acrylonitrile or styrene.

10 These vinylidene rigid polymers may be polymerized either by emulsion, emulsion-suspension, suspension or bulk techniques well known in the art. Furthermore, the rigid polymers may be produced either by continuous, semibatch or batch processes.

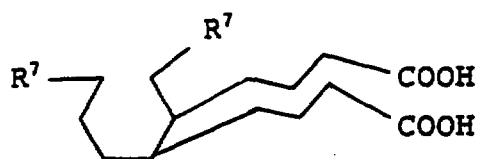
15 Antioxidants such as thio ester antioxidants and hindered phenols may be added to the impact modifier. The present invention preferably utilizes (adds to the emulsion) an antioxidant composition which contains an amount of polyacid surfactant. The antioxidant may be in the form of an antioxidant emulsion.

20 The polyacid is preferably present in the graft copolymer composition (impact modifier composition) at a level of from 0.25 to 5 percent by weight based on the total weight of the graft copolymer composition, more preferably present at a level of from 0.75 to 3.5 percent by weight thereof, more preferably is present at a level of from 1.5 to 2.75 percent by weight thereof. The polyacids have from 30 to 108 carbon atoms (C_{30} to C_{108}), more preferably from C_{32} to C_{60} . The polyacid component may be in the form of an admixture of polyacids, and is preferably substantially (>50 percent by weight) di and/or triacids, and more preferably substantially dimerized

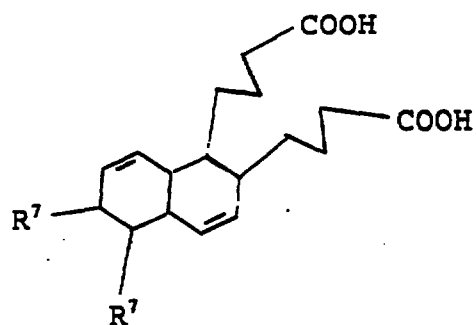
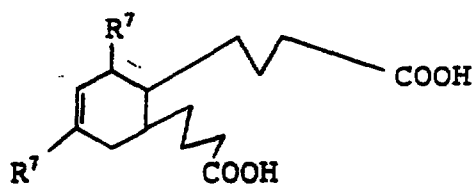
fatty acids and/or trimerized fatty acids. Preferably the polyacid in the form of carboxylate salts of the polyacid function are surfactants, emulsifiers or soaps. The dimerized fatty acid soaps that are useful in the emulsifying agents are soaps of polycarboxylic acids resulting from the polymerization of drying oil or semi-drying oil fatty acids. Suitable fatty acids include polyunsaturated fatty acids, such as linoleic acid, as well as mixtures of fatty acids that contain a substantial amount of polyunsaturated acids, such as soybean oil fatty acids and tall oil fatty acids. Such acids may be dimerized, for example, by heating them under pressure in the presence of an alkaline catalyst. The term "dimerized fatty acids" as used herein is intended to include the mixture of acids resulting from the dimerization of drying oil or semi-drying oil fatty acids. This mixture ordinarily includes a major amount of dimeric acids, trimeric acids and higher polymeric acids (and smaller amounts of unpolymerized (monomeric) acids). Examples include the following but are not limited to these structures:



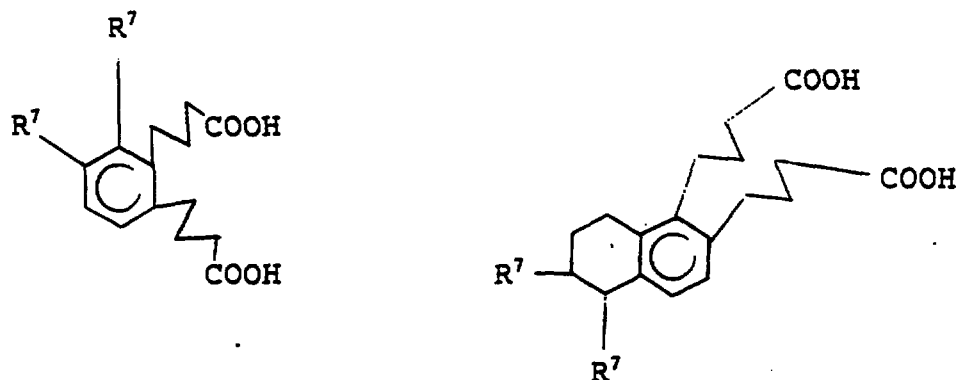
linear dimer acids:



cyclic dimer acids:



aromatic dimer acids:



Each R⁷ is preferably a hydrogen or aliphatic hydrocarbon radical (alkyl) having from 1 to 18 carbon atoms and is more preferably of the formula (CH₂)_x, wherein x is from 1 to 22. Dimerized fatty acids may be formed by dimerizing fatty acids containing ethylenic linkages and from about 14 to about 22 carbon atoms in length. Monobasic acids which dimerize to form products as illustrated above include: oleic, elaidic, palmitoleic, linoleic, linolenic, licanic, arachidonic, erucic, clupanodonic, elaeostearic, etc. In commercial practice naturally occurring mixtures of these acids are generally employed for production of dimerized fatty acids. Such acids may be derived from sources such as tall oil fatty acids, tallow (animal grease) fatty acids and vegetable oils, e.g. soya, linseed, cottonseed, and other oils comprised of unsaturated fatty acid glycerides. In general, the dimerization is carried out by heating the monomeric acid at an elevated temperature, with or without a catalyst, while avoiding cracking and decarboxylation. U.S. patents 2,482,761, 2,664,429, 2,793,219, 2,793,220, 2,955,121, 3,076,003, 3,507,890 and 3,925,342

describe dimerization in further detail. Various types of dimerized fatty acids are available and the commercial products of dimerized fatty acids frequently contain trimerized fatty acids, which are

5 tribasic acids formed as a by-product through the polymerization of three molecules of the fatty acids. In addition, the commercial products may contain small percentages of isomerized monobasic acids or unreacted or fully saturated monomeric monobasic

10 fatty acids which did not polymerize or which were not removed after the polymerization was carried out. The dimerized fatty acids employed to prepare the novel compositions of this invention preferably comprise a product having not more than about 70%

15 tribasic acid and for very low non juicing characteristics not more than 20% monobasic acids. And preferably not more than 10% by weight monobasic acid based on the total weight of the polyacid component. The dimerized acid content is preferably

20 at least 75% by weight based on the total weight of the surfactant. For improved color, the dimerized fatty acids may be partially or fully saturated by hydrogenation in a subsequent reaction step as described in Chapter 5 entitled "Hydrogenation of

25 Fatty Acids" in the book "Fatty Acids in Industry" edited by Robert Johnson and Earle Fritz and published by Marcel Dekker, Inc. NY, NY. All dimer acids are liquid at 25°C even though their number average molecular weight is typically above 500 (for

30 example, about 560). The dimerization process may produce a mixture of monoacids, diacids, triacids, and higher polymeric acids. Preferably more than about half of the dimerized acid is comprised of isomers with cyclic aliphatic or aromatic structures.

The cyclic structures include monocyclic and bicyclic aliphatic and aromatic structures. The cyclic structures generally form during the dimerization process, but their mode of formation is not critical to this invention. It is further important to note that there is no rosin acid, abietic acid or derivatives thereof present in these compositions in part because such materials as abietic acid can be undesirably volatile and can result in juicing and fogging.

Preferred polyacids are cyclic polyacids and cyclic aromatic polyacids. Preferably the polyacid is soluble and/or miscible in the thermoplastic compositions. The polyacid is preferably a C_{30} to C_{100} polyacid in order to minimize fogging, and are more preferably C_{34} to C_{36} polyacids.

The process of this invention involves emulsion polymerizing a diene rubber substrate with an ethylenically unsaturated compound selected from the group of monovinyl aromatic monomers, unsaturated nitrile monomers, alkyl (alka) acrylate monomers, and mixtures thereof, the polymerization being in the presence of a polyacid surfactant to deter formation of coagulum, the polymerization forming an impact modifier composition containing the polyacid surfactant. Preferably the thermoplastic compositions obtained pursuant to the present invention when injection molded pursuant to the following conditions: a 5 oz. capacity molding machine with a clamp force of 150 tons; a 4" x 6" x 0.125" removable mold cavity insert; a mold temperature of 80°F; a melt stock temperature of 575°F; an injection speed of 1.0 inch/second; a screw speed of 75 RPM; a screw back pressure of 100 psi; a

cycle time of approximately 32 seconds; a short shot equaling 95% of cavity capacity; an accumulation of mold deposit resulting from 150 molding cycles; removing the cavity from the molding machine and
5 washing the removable mold cavity with chloroform to collect the residue in a 20 mil scintillation vial; weighing a tarred scintillation vial to determine the amount of accumulated residue in milligrams after carefully removing the chloroform; analyzing the
10 contents by Fourier Transform InfraRed (FTIR), high pressure liquid chromatography (HPLC), etc. yields a fogging level of less than 12 mgs, more preferably less than 9 mgs, and most preferably less than 7 mgs, after 150 short shots. The process further involves
15 injection molding a thermoplastic composition containing the impact modifier composition to form a molded article. The thermoplastic compositions made by this present process exhibit reduced levels of fogging compared to those which contain a monoacid
20 surfactant. The injection molding process typically occurs at a temperature of at least 500°F, more typically at temperatures of between 515 and 575°F, and most preferably at temperatures between 525 and 550°F. These relatively high injection molding
25 temperatures contribute to the problems associated with juicing, fogging and plate out of linear monoacid surfactants, and the present polyacid surfactants effectively resist such high levels of juicing, fogging and plate out.

30 The final composition may also contain ordinary additives, such as coloring agent, heat stabilizer, light stabilizer, molding aid, etc.

The compositions are useful for making automotive dash components such as dash boards, and

thus are useful for automobiles having glass windshields and thermoplastic dash boards.

ExamplesTable 1A

Series I - Rigid PVC/ABS Compounds

Experimental #	A	1	B	2	3	C
Emulsifier	LFA	Dimer	LFA	Dimer	Dimer	LFA
% PBd in ABS	66.2	65.0	66.3	65.1	64.4	69.5
% Emulsifier in ABS	2.40	2.40	2.40	2.40	2.40	2.55
Fogging						
% Gloss Retention	90.8	93.6	87.9	94.0	94.6	86.7
Fogging of ABS resin alone						
% Gloss Retention	43.2	92.0	36.2	50.9	49.8	34.1
• PVC	100					
ABS Resin	12					
Acrylic Process Aid	2.5					
Tin mercaptide						
stabilizer	2.5					
Lubricants	0.70					

All ABS were coagulated with acid.

LFA is a linear mono fatty acid.

Dimer is a dimerized fatty acid.

Examples 1-4 and comparative examples A-D had the formulation "*" set.

Series II - 30% PBd Substrate Based ABS resin

Neat (Pure resin as is)

Conditions: 100°C in a circulating oil heated bath for 3 hours - Glass plate over bowl head to condense the volatile components.

Table 1B

<u>Experiment</u>	<u>Fatty Acid</u>	<u>% Gloss Retention</u>
		60° Gloss
4	2.36% Dimer acid Based ABS	89.8
D	2.36% LFA Based ABS	52.9
Dimer = dimerized fatty acid		

This is the replacement sheet

Date: - 4 APR 1996

Table II

Experiment #	5	E	
SAN resin	0	0	0 to 75%
2.36% Dimer acid based ABS**	80		0 to 100%
2.36 LFA Based ABS**		80	
Nitrile rubber	20	20	0 to 50%
Monomeric or polymeric plasticizer	0	0	0 to 60%
Fogging			
% Gloss Retention	36	31	

The chemical nature of the condensate was not analyzed to identify components.

** 30% PBd rubber based ABS resin (Rubber level in ABS can range from 10 to 85%.)

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Table IIIABS/SAN

Experiment #	F	6	7	8	G
SAN -1	60	60	60	60	
ABS -1					100
ABS -2	40				
ABS -3		40			
ABS -4			40		
ABS -5				40	
EBS wax	1.0	----->			
Mg Stearate	0.5	----->			
Silicon Oil	0.1	----->			
Juice, mgs	19.2u	21.2v	17.5w	14.0x	22.7y

SAN-1 A bulk polymerized SAN containing 1.5% of a low molecular weight SAN oligomer.

ABS-1 An all emulsion grade ABS resin with 20% rubber containing 2.0% linear monofatty acid (LFA) added in the rubber substrate, graft stage and AOE #1 make-up.

ABS-2 A 50% PBd rubber grade ABS containing 1.54 pphr of a linear fatty monoacid used in the rubber substrate and AOE #2 make-up.

ABS-3 A 50% PBd rubber grade ABS containing 1.54 pphr of a dimer acid used in the rubber substrate and AOE #2 make-up.

ABS-4 A 50% PBd rubber grade ABS containing 1.54 pphr of a dimer acid used in the rubber substrate and AOE #3 make-up.

ABS-5 A 50% PBd rubber grade ABS containing 1.54 pphr of a dimer acid used in the rubber substrate and AOE #4 make-up.

Composition of juice as analyzed by FTIR & HPLC - the organic acids were derivatized for analysis by HPLC.

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% SAN oligomer		% organic acid					% AO	% Unknown
		Palmetic	Oleic	Elaidic	Stearic	Dimer		
Saturated		yes	no	no	yes	mixed		
acid		mono	mono	mono	mono	di		
u	25.0	19.4	3.6	1.9	13.8	0.0	0.0	63.7
v	30.2	14.7	0.26	0.29	24.1	0.0	0.0	69.6
w	27.4	12.6	0.19	0.25	21.4	0.0	0.0	61.8
x	29.2	9.9	0.14	0.20	16.4	0.0	1.0	55.8
y	0.0	37.5	3.8	6.9	21.3	0.0	0.0	69.5

With regards to detection of organic saturated mono acids in all samples of Table III, it is believed to be derived from the magnesium stearate and/or the ethylene bis stearamide wax (EBS).

The majority of oleic and elaidic acids in samples u and y are due to the linear fatty acid emulsifier and to a lesser extent from the magnesium stearate and/or the ethylene bis stearamide wax (EBS).

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Claims

- 1 A thermoplastic resin composition comprising:
 - (a) a vinyl chloride resin; and
 - (b) a graft copolymer containing a C_{30} - C_{108} polyacid surfactant.
- 2 The composition of Claim 1 wherein said vinyl chloride resin is present at a level of from 20 to 95 percent by weight based on the total weight of the composition.
- 3 The composition of Claim 1 wherein said graft copolymer is present at a level of from 5 to 80 percent by weight based on the total weight of the composition.
- 4 The composition of Claim 1 wherein said polyacid is a cyclic dimer acid.
- 5 An article made from the composition of Claim 1.
- 6 A low juicing impact modifier composition comprising:
 - (a) a rubber substrate with a T_g of less than 0°C present at a level of from 10 to 90 weight percent based on the total weight of the impact modifier composition;
 - (b) a superstrate grafted to said substrate, said superstrate with a glass transition temperature greater than 20°C being derived from a compound selected from the group consisting of vinyl aromatics, unsaturated nitriles, alkyl (alkyl) acrylates and mixtures thereof; and
 - (c) a polyacid selected from the group consisting of C_{30} to C_{108} polyacids.
- 7 The modifier composition of Claim 6 wherein said polyacid is present at a level from 0.25 to 5 percent by weight based on the total weight of the composition.

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- 8 The modifier composition of Claim 7 wherein said polyacid is selected from the group consisting of aliphatic or cyclic aromatic.
- 9 The composition of Claim 8 wherein said surfactant is present at a level of from 0.75 to 3.5 weight percent based on the total weight of the composition.
- 10 The composition of Claim 6 wherein said substrate is present at a level of between 50 and 85 weight percent based on the total weight of the composition.

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Claims

- 1 A thermoplastic resin composition comprising:
 - (a) a vinyl chloride resin; and
 - (b) a graft copolymer containing a C_{30} - C_{108} polyacid surfactant.
- 2 The composition of Claim 1 wherein said vinyl chloride resin is present at a level of from 20 to 95 percent by weight based on the total weight of the composition.
- 3 The composition of Claim 1 wherein said graft copolymer is present at a level of from 5 to 80 percent by weight based on the total weight of the composition.
- 4 The composition of Claim 1 wherein said polyacid is a cyclic dimer acid.
- 5 An article made from the composition of Claim 1.
- 6 A low juicing impact modifier composition comprising:
 - (a) a rubber substrate with a T_g of less than 0°C present at a level of from 10 to 90 weight percent based on the total weight of the impact modifier composition;
 - (b) a superstrate grafted to said substrate, said superstrate with a glass transition temperature greater than 20°C being derived from a compound selected from the group consisting of vinyl aromatics, unsaturated nitriles, alkyl (alkyl) acrylates and mixtures thereof; and
 - (c) a polyacid selected from the group consisting of C_{30} to C_{108} polyacids.
- 7 The modifier composition of Claim 6 wherein said polyacid is present at a level from 0.25 to 5 percent by weight based on the total weight of the composition.



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- 8 The modifier composition of Claim 7 wherein said polyacid is selected from the group consisting of aliphatic or cyclic aromatic.
- 9 The composition of Claim 8 wherein said surfactant is present at a level of from 0.75 to 3.5 weight percent based on the total weight of the composition.
- 10 The composition of Claim 6 wherein said substrate is present at a level of between 50 and 85 weight percent based on the total weight of the composition.