

COMMONWEALTH of AUSTRALIA

PATENTS ACT 1952

APPLICATION FOR A STANDARD PATENT

†
We ROHM AND HAAS COMPANY of
Independence Mall West,
Philadelphia, Pennsylvania 19105,
United States of America.

hereby apply for the grant of a Standard Patent for an invention entitled:

"POLYIMIDES, PREPARATION OF POLYIMIDES AND BLENDS OF
POLYIMIDES"

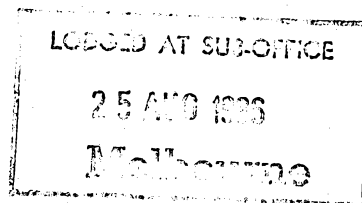
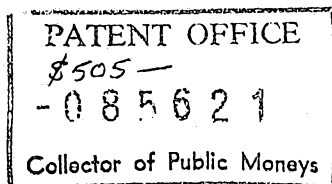
which is described in the accompanying ~~provisional~~
complete specification.

Details of basic application(s):—

<u>Number</u>	<u>Convention Country</u>	<u>Date</u>
770,034	United States of America	27th August, 1985

APPLICATION ACCEPTED AND AMENDMENTS

ALLOWED 30.11.89



The address for service is care of DAVIES & COLLISON, Patent Attorneys, of 1 Little Collins Street, Melbourne, in the State of Victoria, Commonwealth of Australia.

Dated this 25th day of August, 1986

To: THE COMMISSIONER OF PATENTS

H. M. Dimington

(a member of the firm of DAVIES &
COLLISON for and on behalf of the Applicant).

Davies & Collison, Melbourne and Canberra.

COMMONWEALTH OF AUSTRALIA

PATENTS ACT 1952

DECLARATION IN SUPPORT OF CONVENTION OR
NON-CONVENTION APPLICATION FOR A PATENT

Insert title of invention.

In support of the Application made for a patent for an invention
entitled: "POLYIMIDES, PREPARATION OF POLYIMIDES AND BLENDS OF
POLYIMIDES"

Insert full name(s) and address(es)
of Declarant(s) being the appli-
cant(s) or person(s) authorized to
sign on behalf of an applicant
company.

I Kenneth Norman Rimington
We Patent Attorney of,
Davies & Collison
of 1 Little Collins Street,
Melbourne 3000,
Australia

Cross out whichever of paragraphs
1(a) or 1(b) does not apply.

1(a) relates to application made
by individual(s).

1(b) relates to application made
by company; insert name of
applicant company.

do solemnly and sincerely declare as follows:—

1. (a) ~~I am~~ the applicant..... for the patent
~~We are~~

or (b) I am authorized by

Rohm and Haas Company

Cross out whichever of paragraphs
2(a) or 2(b) does not apply.

2(a) relates to application made
by inventor(s).

2(b) relates to application made
by company(s) or person(s) who
are not inventor(s); insert full
name(s) and address(es) of inven-
tors.

the applicant..... for the patent to make this declaration on ^{its}
~~their~~ behalf.

2. (a) ~~I am~~ the actual inventor..... of the invention
~~We are~~

or (b) Michael Paul Hallden-Abberton of 964 Whitney Lane,
Maple Glen, Pennsylvania 19002, United States of America;
Newman Mayer Bortnick of 509 Orelan Mill Road, Orelan,
Pennsylvania 19075, United States of America; Leslie Alan
Cohen of 2550 Greek Road, Langhorne, Pennsylvania 19047,
United States of America; Harry Charles Fromuth, Jr. of 904
Sterner Mill Road, Trevoise, Pennsylvania 19047, United
States of America; and William Thayer Freed of RD #1 Box
E323, Stockton, New Jersey 08559, United States of America

~~is~~ the actual inventor(s)..... of the invention and the facts upon which the applicant.....
~~are~~

~~is~~ entitled to make the application are as follows:—
~~are~~

The actual inventors have assigned the invention to the
said applicant.

State manner in which appli-
cant(s) derive title from inven-
tor(s)

Cross out paragraphs 3 and 4
for non-convention applications.
For convention applications
insert basic country(s) followed
by date(s) and basic applicant(s).

3. The basic application..... as defined by Section 141 of the Act ^{was}
~~were~~ made
in United States of America on the 27th August, 1985
by Michael Paul Hallden-Abberton, Newman Mayer Bortnick, Leslie
Alan Cohen, Harry Charles ~~on the~~ Fromuth, Jr. and William
~~by~~ Thayer Freed
in on the
by

4. The basic application..... referred to in paragraph 3 of this Declaration ^{was}
~~were~~ the first application..... made in a Convention country in respect of the invention the subject
of the application.

Insert place and date of signature.

Declared at Melbourne this 25th day of August, 1986

Signature of Declarant(s) (no
attestation required).

K. N. Rimington

Note: Initial all alterations.

DAVIES & COLLISON, MELBOURNE and CANBERRA.

(12) PATENT ABRIDGMENT (11) Document No. AU-B-61803/86
(19) AUSTRALIAN PATENT OFFICE (10) Acceptance No. 593835

(54) Title
POLYIMIDES WITH BLOCKED ACID GROUPS

International Patent Classification(s)
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(56) Prior Art Documents
US 4129619
US 3639357
US 3284425

(57) Claim

1. A thermoplastic polymer comprising imide groups and optionally acid and/or anhydride functional groups and having a degree of imidization of at least 1% characterized in that the degree of imidization on said polymer and the acid and/or anhydride functionality on said polymer are as set forth in the Table below;

<u>when the degree of imidization on said polymer is</u>	<u>the milliequivalents of acid and/or anhydride per gram of imidizable polymer component</u>
--	---

from 1% to 10%	0 to 0.20
above 10% to 20%	0 to 0.28
above 20% to 30%	0 to 0.32
above 30% to 40%	0 to 0.31
above 40% to 50%	0 to 0.28
above 50% to 60%	0 to 0.28
above 60% to 70%	0 to 0.27

above 70% to 80%	0 to 0.28
above 80% to 90%	0 to 0.33
and above 90% to less than 95%	0 to 0.40

7. A process for the preparation of a thermoplastic polymer comprising imide groups and optionally acid and/or anhydride functional groups and having a degree of imidization of at least 1%, said process comprising reacting a ~~first~~ polymer comprising imide groups and active acidic and/or anhydride functional groups with a predetermined amount of acidic and/or anhydride functional group blocking agent to reduce or substantially eliminate the active acid and/or anhydride functional groups in the polymer.

14. A blend of at least one thermoplastic polymer and a second polymer, said second polymer being a polymer as claimed in any one of claims 1 to 6 or a polymer prepared by the process as claimed in any one of claims 7 to 13.

COMMONWEALTH OF AUSTRALIA

PATENT ACT 1952

COMPLETE SPECIFICATION

(Original)

FOR OFFICE USE

Class

Int. Class

Application Number:
Lodged:

61803/86

Complete Specification Lodged:
Accepted:
Published:

Priority:

Related Art:

This document contains the
amendments made under
Section 49.

and is correct for printing.

Name of Applicant: ROHM AND HAAS COMPANY

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Actual Inventor(s): Michael Paul HALIDEN-AEBERTON
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1 Little Collins Street, Melbourne, 3000.

Complete Specification for the invention entitled:

"POLYIMIDES, PREPARATION OF POLYIMIDES AND BLENDS OF
POLYIMIDES"

The following statement is a full description of this invention,
including the best method of performing it known to us :-

POLYIMIDES, PREPARATION OF POLYIMIDES AND BLENDS OF
POLYIMIDES

This invention is concerned with polymers comprising imide groups and optionally acid and/or anhydride functional groups, a process for the
5 preparation of such polymers, and blends of said imide polymers with other thermoplastic materials.

US-A-4246374 describes and claims polyglutarimide polymers and their preparation. This patent teaches that the degree of imidization of an
10 acrylic polymer may vary from as low as 1% imidization up to about 100% imidization and that imidization is accomplished with anhydrous ammonia or an anhydrous primary amine.

US-A-3284425 is directed to the preparation of polymethacrylimide products. This patent discusses the
15 possibility that acid or ester groups may become isolated between adjacent imide rings. US-A-3284425 also teaches the formation of polymethacrylimide products which "...are substantially free of carboxy or ester groups...in which the degree of imidization is at
20 least about 95%..." Thus, while recognizing the presence of acid groups in the polyimides, US-A-3284425 teaches imidizing to at least about 95% imidization in order to lower or remove the acid functionality of a
25 polymethacrylimide.

Further, there is no recognition in US-A-3284425 concerning adverse effects of the normal amount of acid and anhydride functionality on a polyimide polymer.

US-A-4518717 describes a process for making
30 imide polymers by reacting a monomeric dianhydride with a lactam or oxime to form a bis-imide having an acid-group-carrying radical attached to the imide nitrogen.

The bis-imide monomer is then esterified with a reactive solvent that acts as an esterifying agent; methanol is typical. In the esterification process, the aromatic imide functionality is retained. The esterified bis-imide monomer is subsequently reacted with an aromatic diamine monomer in a polycondensation reaction, to form an imide or amide-imide polymer.

EP-A-0076691 describes the preparation of polyimides which are free from methyl substitution on the nitrogen atom. EP-A-0076691 teaches the preparation of imide polymers free from methyl substitution on the nitrogen atom starting from a non-crosslinked polymer which contains glutaric anhydride units. The polyglutaric anhydrides are reacted with ammonia to yield the desired polyimides free from methyl substitution on the nitrogen atom. As the degree of imidization exceeds 95% and approaches complete imidization, the amount of residual acid and anhydride units remaining in the polymer chain becomes less significant, although the degree of imidization, if above 95%, is a factor which limits the utility of the resultant polyimide. When the degree of imidization is less than 95%, then the residual acid and anhydride functionality on the polymer chain results in less desirable properties than could be obtained when the acid and anhydride functionality is reduced or eliminated.

Makromol. Chem., 184, pages 1705-1726 (1983) discloses an analytical technique for confirming the chemical structure of the reaction products of polymethacrylic acid and formamide. The technique encompasses treating copolymers of methacrylic acid and methacrylimide with diazomethane. There is no disclosure in the reference procedure concerning the degree of imidization. Further, the imide groups are reacted along with the acid groups.

Japanese published patent application No.
60-20904 (A) describes a process for preparing a
polyglutarimide by condensing a polyamide or a
copolymer of a methacrylamide with a methacrylate
5 ester.

US-A-4458046 discloses blends of vinyl chloride
polymers and copolymers of the styrene-maleimide type.
US-A-4458046 teaches that the imide used be prepared
from a dicarboxylic acid moiety and that complete
10 conversion of the dicarboxylic acid moiety to the
desired imide be obtained by using an excess of the
stoichiometric amount of ammonia or amine required for
such conversion.

As a by-product of imidization of (meth)acrylic
15 polymers by reaction with ammonia or primary amines,
anhydride and acid groups are formed on the polymer
chain. The acid and anhydride groups are intermediate
in the formation of imide units. When the degree of
imidization exceeds 95% and approaches 100%, the amount
20 of acid and anhydride units present on the resultant
imide product decreases and poses less of a problem.
However, when the degree of imidization exceeds 95%,
particularly when an autoclave process is used, then a
high degree of discoloration of the resultant polymer
25 and degradation of the polymer itself is often
obtained. The degradation of the polymer may also lead
to alteration of the desirable properties of the
resultant imide polymer.

When the degree of imidization of a polymer is
30 95% or less, then the amount of acid and anhydride
functionality normally present on the polymer chain as
a natural incidence of the reaction involved becomes
objectionable in that the presence of acid and
anhydride functionality generally adversely affects the
35

properties of the polyimide. For example, the presence of the normal amount of acid and anhydride functionality on an imide or polyglutarimide polymer will alter the miscibility of such polymer with other thermoplastic polymers. Additionally, acid and anhydride functionality affects the weatherability of a molded article prepared from such polymer in that the molded article will absorb more water as a result of the presence of the normal amount of acid and anhydride functionality. Further, high acid and anhydride functionality can result in higher melt viscosity which would translate into the polymer being more difficult to process into various molded articles.

It is an object of this invention therefore, to provide a polyimide wherein the amount of acid and/or anhydride functionality is reduced from the amount normally present, or is substantially eliminated.

A further object is to provide a process for reducing or substantially eliminating active acid and/or anhydride functional groups existing on a polyimide.

A still further object of this invention is to prepare blends of polyimides with other thermoplastic polymers wherein the acid and/or anhydride functionality of the polyimide in the blend has either been reduced from that which was originally present or has been substantially eliminated.

In accordance with one aspect of the invention, a polymer comprises imide groups, preferably glutarimide groups, and optionally acid and/or anhydride functional groups and has a degree of imidization of at least 1%, characterized in that the degree of imidization on said polymer, and the acid and/or anhydride functionality on said polymer are as set forth in the Table below:

	<u>when the degree of imidization on said polymer is</u>	<u>the milliequivalents of acid and/or anhydride per gram of imidizable polymer component are</u>
	from 1% to 10%	0 to 0.20
5	above 10% to 20%	0 to 0.28
	above 20% to 30%	0 to 0.32
	above 30% to 40%	0 to 0.31
	above 40% to 50%	0 to 0.28
	above 50% to 60%	0 to 0.28
10	above 60% to 70%	0 to 0.27
	above 70% to 80%	0 to 0.28
	above 80% to 90%	0 to 0.33
	and above 90% to less than 95%	0 to 0.40

15 In accordance with another aspect of the present invention, a process is provided for the preparation of a thermoplastic polymer comprising imide groups and optionally acid and/or anhydride functional groups and having a degree of imidization of at least 1%, said process comprising reacting a first polymer comprising

20 imide groups and active acidic and/or anhydride functional groups with a predetermined amount of acidic and/or anhydride functional group blocking agent to reduce or substantially eliminate the active acid and/or anhydride functional groups on the polymer.

25 Preferably the process is carried out at a temperature of from 50°C to 450°C. Preferably the process is carried out at a pressure of from 1 atmosphere (101.3 KiloPascals) to 1000 atmospheres (101300 KiloPascals). Preferably said polymer is a glutarimide.

30 This invention also contemplates a blend of at least one other thermoplastic polymer and a second polymer, said second polymer being a polyimide polymer as hereinbefore defined or a polyimide polymer as prepared by the process as hereinbefore defined.

It is surprising to find that when acid and/or anhydride functionality of a polyimide is reduced or substantially eliminated, the resultant polyimide may exhibit altered miscibility and compatibility with other thermoplastic polymers, improved weatherability characteristics, and a lower melt viscosity thereby enhancing the processability of the polyimide.

The polyimides of the present invention may also possess superior thermal stability. Preferred polyimides of this invention are those which have a thermal stability indicated by a 1% weight loss, by thermogravimetric analysis (TGA), at a temperature above 285°C in air and above 300°C in nitrogen.

As used in the specification and claims, the term "degree of imidization" refers to the weight percentage of imide groups on a polymer in relation to the groups of the polymer capable of being reacted to form imide groups. For example, one may have a copolymer of styrene and methacrylic acid. Because the styrene is not capable of being imidized, the degree of imidization is based on the percentage of imide groups which are subsequently formed from the methacrylic acid portion of the copolymer.

Any thermoplastic polyimide, containing acid and/or anhydride groups, may be used in practicing the process of this invention. For example, one may use the polyglutarimides disclosed in US-A-4246374. Additionally, one may use a polymethacrylimide of the type set forth in US-A-3284425. The polyglutarimide product of this invention may be prepared from a polyimide such as those derived from esters of acrylic or methacrylic acid wherein the ester moiety contains from 1 to 20 carbon atoms and which has been reacted with ammonia or a primary amine wherein the hydrocarbon

portion of the amine is hydrogen or a substituted or unsubstituted alkyl or aryl group having up to 20 carbon atoms, may also be used. Such polyimides are disclosed in US-A-4246374. A preferred polyimide is derived from methyl methacrylate. Preferably the imide units are derived from a primary amine having substituted or unsubstituted alkyl or aryl groups of up to 20 carbon atoms, and more preferably are derived from methylamine, ammonia, aniline or cyclohexylamine.

The composition of the thermoplastic polyimide may be varied depending upon the properties desired from the final product. A preferred polyimide contains up to about 50%, and preferably from about 10% to about 50%, by weight, of acrylic or methacrylic ester units, preferably wherein the ester moiety contains from 1 to 20 carbon atoms. Another preferred polyimide contains from 50 to 99%, by weight, of acrylic or methacrylic ester units, with the preferred ester moiety as stated above. Among the ester moieties, a more preferred one is methyl.

Among the polyimides which may be used to practice the process of this invention are the polyglutarimides described above and copolymers thereof with, for example, styrenic materials such as styrene, ring substituted styrenes and/or alpha-methyl styrene, all optionally containing acrylonitrile and/or methacrylonitrile; polyimides containing succinimide functionality such as those prepared by the copolymerization of an alkyl methacrylate with a maleimide, N-alkylmaleimides and/or N-arylmaleimide; polyimides prepared by the copolymerization of alkyl methacrylates, maleimide and/or N-alkyl and/or N-arylmaleimides, styrene and/or ring substituted styrenes or alpha-methylstyrene, all optionally

containing acrylonitrile and/or methacrylonitrile;
copolymers of styrene, ring substituted styrenes, or
alpha-methylstyrene with maleimide, N-alkylmaleimides
and/or N-arylmaleimides; copolymers of ethylene and
5 propylene containing glutarimide functionality; graft
polymers of methyl methacrylate or methacrylic acid
onto polyethylene, polypropylene, and/or ethylene-
propylene diene rubber; copolymers of maleic anhydride
and olefins, followed by imidization, and graft
10 copolymers of maleic anhydride onto polyolefins
followed by imidization; and copolymers of
alkylmethacrylate, styrene, itaconimide, citraconimide,
alpha-methylmaleimide or N-methyl alpha-
methylmaleimide, all optionally with acrylonitrile
15 and/or methacrylonitrile. Whichever polyimide is used
to practice the process of this invention, such
polyimide must have residual acid and/or anhydride
groups.

Particularly preferred polyimides are those set
20 forth in US-A-4246374 and, even more particularly
preferred, are the polyglutarimides prepared by the
reaction of (1) poly(methyl methacrylate) with ammonia
or methylamine and (2) a copolymer of poly(methyl
methacrylate) and ethyl acrylate with ammonia or
25 methylamine and (3) a copolymer of (methyl)styrene and
methyl methacrylate with ammonia or methylamine,
because these polyimides are simple to prepare and easy
to work with.

The first polymer may be prepared in any
30 suitable manner and the invention is not to be
construed as being limited to the method of preparation
of the polyimide. For example, one may prepare a
polyimide, for example a polyglutarimide, by heating,
with stirring or mixing, the selected polymer such as

poly(methyl methacrylate), preferably in an autoclave, in the presence of a primary amine, such as butylamine or methylamine. Other suitable amines include aniline and cyclohexylamine; ammonia is also suitable for
5 preparing the polyimide. The reaction may be conducted in the presence of water and at an elevated pressure such as between 8 and 85 atmospheres (810.4 and 8610.5 KiloPascals). The temperature which may be used may vary from 180°C to 350°C, with the understanding that
10 the pressure used will vary with the temperature used.

One may also prepare the first polymer as described in US-A-4246374 wherein an extruder is used in the preparation. Generally, when using an extruder, the polymer, such as poly(methacrylic acid) may be
15 continuously fed to an extruder and ammonia or a primary amine, such as methylamine, may be introduced continuously through an injection port. Unwanted by-products and excess ammonia or primary amine are removed by progressively reducing the pressure at
20 downstream extruder vents. The temperature in the extruder may be varied and such variation will be dependent on the nature of the starting materials, pressure being used, residence time, melt viscosity of the polymer being extruded, etc. Generally, a
25 temperature of from about 200°C to about 450°C may be used in preparing the polyimide and a pressure may be used which is from about atmospheric to about 1000 atmospheres (101300 KiloPascals), although pressures of less than 500 atmospheres (50650 KiloPascals) are
30 generally suitable. The reaction time of the acrylic polymer in the reaction zone of the extruder will generally vary from about 0.1 to about 1000 seconds and preferably from about 30 to about 300 seconds. The desired degree of imidization is obtained by suitable

adjustment of various reaction parameters such as residence time. The extruder is generally equipped with a screw which advances the materials present through the extruder. In a particularly preferred embodiment, the extruder may be equipped with at least one pair of screws. The number of pairs of screws which is present in the extruder is dictated solely by economic and practical process conditions. In one embodiment, each screw member of each pair of screws which is present in the extruder may rotate in a direction counter to the rotational direction of the other screw member of the pair. In this manner mixing may be enhanced, as may removal of undesired materials.

The first polymer may also be prepared in a tubular reaction vessel such as a baffled in-line mixer, wherein the baffles provide the necessary mixing. The baffled in-line mixer may also be equipped with various ports for the introduction of necessary reactants or materials.

It is preferred that an extruder be used and that the extruder which is used for preparing the polyimide, be equipped with at least one pair of screws wherein each member of the pair rotates in a direction counter to the rotational direction of the other screw member of the pair.

Because the preparation of the first polymer may involve amide and/or acid intermediates which may give rise to imides or anhydrides, if an extruder apparatus is not used to prepare the polyimide, the final product may contain imide, amide, acid, anhydride and ester moieties. The preference for the reaction to take place in the extruder is for the reason that the proper use of the extruder tends to cause all significant amide groups to further react to imide

although some acid and anhydride groups are still present. Thus, polyimides prepared using an extruder are usually substantially free of amide functionality and contain less acid and anhydride functionality than polyimides prepared using other processes. Preferably, regardless of the method or apparatus used, the polyimide is substantially free of amide functionality.

The invention is not to be construed as limited to any particular method for preparing polyimides.

Regardless of which apparatus or method is used for preparing the polyimides, for example a polyglutaramide, the degree of imidization should be controlled so that the final product preferably has a degree of imidization of up to 99%, and preferably from 10% to no more than 95%. It is more preferred however that the degree of imidization of the polyimide be from 25% to 95%, even more preferred that the degree of imidization be from 40% to 88%, and still more preferred that the degree of imidization be from 50% to 85%.

In a preferred embodiment of the present invention, the acid and anhydride functionality of the first polymer having up to 88% degree of imidization is reduced to no more than 0.25 milliequivalents per gram of imidizable polymer component, and in a more preferred embodiment the acid and anhydride functionality of a polyimide having up to 85% degree of imidization is reduced to no more than 0.25 milliequivalents per gram of imidizable polymer component.

If the degree of imidization exceeds 95%, then, although the acid and anhydride level of such a polyimide may be low, such polyimides may suffer from a high degree of discoloration, a partially degraded polymer, and undesirable alteration of the properties of the polyimide.

Although the final polyimide product of this invention preferably has no more than about a 95% degree of imidization, the process of this invention may be used to remove acid and anhydride functionality from a polyimide containing such functionality wherein the degree of imidization of such polyimide exceeds 95% and is less than 100%.

The agent which is used to block the acid and/or anhydride functional groups may be any agent which is capable of reacting with the acid and anhydride groups and incapable of reacting with the imide functionality and which will not adversely affect the properties of the polyimide, either during the time the reaction with the acid and anhydride functionality is being conducted or in the final polymer.

The term acid and/or anhydride functional group means any carboxylic acid, carboxylic acid salt, and/or anhydride group present on the polyimide.

Among the types of agents which may be used to partially or completely remove acid and anhydride functionality which is present on the polyimide are alkylating agents such as orthoesters, ketals, carbonates and/or sulfoxides, preferably alkyl, aryl, alkaryl and/or aralkyl carbonate and/or dialkyl sulfoxide. Other agents, including esterification agents, may also be used such as siloxanes, silyl ethers, silylenol ethers, trialkyl phosphates, trialkyl phosphites, dialkyl sulfates, alkyl alkylsulfonates, alkyl arylsulfonates, dialkyl carbonates, diaryl carbonates, aryl isocyanates, carbodiimides, trialkylsilyl halides, enol esters, alcohols, alkyl esters, alkyl isocyanates, quaternary ammonium salts, ureas and/or guanidines.

Among the specific agents which are preferably used to remove acid and anhydride functionality are one or more of dimethyl carbonate, 2,2-dimethoxypropane, dimethyl sulfoxide, diethyl sulfoxide, triethyl
5 orthoformate, trimethyl orthoacetate, trimethyl orthoformate, diphenyl carbonate, dimethyl sulfate, methyl toluenesulfonate, methyl trifluoromethyl-sulfonate, methyl acetate, methanol, ethanol, methyl isocyanate, p-chlorophenyl isocyanate, dimethyl
10 carbodiimide, dimethyl t-butylsilyl chloride, isopropenyl acetate, dimethyl urea, tetramethyl-ammonium hydroxide, dimethyl diethoxysilane, tetra-n-butoxysilane, dimethyl(trimethylsilyl) phosphite, trimethyl phosphite, trimethyl phosphate, and tricresyl
15 phosphate.

The amount of agent used to block the acid and/or anhydride functional groups will vary depending upon the amount of acid and anhydride present. This may be readily determined by the titration method
20 described hereafter. The amount of agent used will generally be at least the stoichiometric amount of agent needed to substantially eliminate the acid and anhydride functionality if it is desired to remove substantially all acid and anhydride functionality.
25 Generally, when substantially eliminating all acid and anhydride functionality from the polyimide, it is desirable to use from 10% to 50% excess above stoichiometry of the agent.

When it is desired to only partially reduce the
30 acid and anhydride functionality, then it is preferred to use less than a stoichiometric amount of agent to avoid removing more acid and anhydride functionality than is desired.

In some instances, the agent used may react with water which is associated with the first polymer. In such instances, procedures may be used to assure that the desired amount of agent is present to react with the acid and anhydride groups present on the polyimide. One procedure is to dry the polyimide, for example, under vacuum at a temperature of from 100°C to 135°C for 8 to 20 hours. A second procedure is to use an excess, above the amount of agent required, so that the excess amount of agent will react with the water present and the desired amount of agent will react with the acid and anhydride groups present on the polyimide.

The reaction wherein the acid and anhydride functionality is removed or reduced may be conducted in a single or multi-screw extruder, in an autoclave, in a continuously recirculated pipe reactor, in a baffled in-line mixer which may be used in series with an extruder, in a wiping-film evaporator, a falling film reactor or evaporator, in a disc extruder or reactor, or other melt-mixing device.

If the reaction is conducted using an extruder, then it may be conducted as a continuous process wherein a polyimide is prepared in an extruder and is then passed, in the extruder, into a zone wherein the agent is introduced and the polyimide is contacted with the agent for a period of time of from 15 second to 20 minutes at a temperature of from 150°C to 400°C and at a pressure of from about atmospheric to 15000 KiloPascals (2200 psi), in the absence of any solvent and/or catalyst.

Alternatively, a polyimide, which has been previously prepared, may be introduced into an extruder at an elevated temperature so as to form a molten mass, and may then be contacted with the agent at the aforesaid temperature and pressure.

If the reaction of the agent and the acid and anhydride groups is to be conducted in an autoclave, then the reaction may take place, with or without a solvent for the polyimide, in the autoclave for a period of time of from 10 minutes to 8 hours at temperatures of from 100°C to 180°C and in an unpressurized vessel or a pressurized vessel at pressures up to 1000 atmospheres (101300 KiloPascals). In some instances, the agent, if a liquid, may also act as the solvent for the polyimide. In other instances a separate solvent may be used.

The solvent used may vary widely depending on the nature of the polyimide solute to be reacted. Generally, the solvent should be at least somewhat polar in nature and should be non-reactive with the agents to be used. If the agent is not to also act as the solvent for the polyimide then the solvent should also be non-reactive with the polyimide.

Among the solvents which may be used are diglyme, triglyme, sulfolane, tetrahydrofuran and/or tetrahydropyran.

Yet another method for conducting the reaction of the polyimide with the agent is to melt the polyimide in the autoclave and to conduct the reaction between the acid and anhydride function groups of the polyimide and the agent in a molten state without the addition of any solvent.

Still another method is to prepare the polyimide in a reaction vessel, such as an autoclave, and, as the very last step in the preparation, to add a suitable agent for the removal of the acid and anhydride groups.

If one is to conduct a continuous reaction to prepare a polyimide and remove or reduce acid and anhydride functionality, whether in an extruder,

autoclave or other reaction vessel, then the amount of acid and anhydride groups present on the polyimide may be determined by first conducting the imidization reaction under the conditions to be used in the extruder, autoclave or other reaction vessel and measuring the amount of acid and/or anhydride functional groups present on the resultant polyimide. In this manner, one may predetermine the amount of agent to be added, if a continuous process is to be utilized, to reduce or substantially eliminate the desired amount of acid and anhydride groups. In some cases by-products or unreacted materials from the imidization reaction may enhance or impede the subsequent reaction of the polyimide with the agent.

If desired, the reaction of the agent with the polyimide may be conducted in the presence of a catalyst. The catalyst preferably catalyzes the reaction of the agent with the polyimide, but does not catalyze degradation of the polyimide nor cause other undesirable reactions. Among the catalysts which may be used are base catalysts such as trimethylamine, triethylamine, benzyldimethylamine and/or tetramethylammonium hydroxide. One may also use acid catalysts or Lewis acid catalysts such as p-toluenesulfonic acid, tetrabutyl titanate and/or manganese tetraacetate.

Optionally, the polyimide may be present in the final or outermost stage of a multistage polymer and the reaction is conducted with the entire multistage polymer. Further, the polyimide may be one of the elements of a block or graft copolymer, and the reaction is conducted in the presence of the entire block or graft copolymer.

In another embodiment, the polyimide may be in the core of a multistage polymer or in the second or other intermediate stage of a multistage polymer and the acid and anhydride functionality may be removed as aforesaid. The amount of acid and anhydride functionality on any of the polyimides of the stages of the multistage polymer may be adjusted to that which is desired.

The polyimides of this invention may be blended with one or more other thermoplastic polymers and may even be blended with another polyimide which may or may not have had its acid and/or anhydride functionality level reduced or substantially eliminated. The polyimide according to this invention may be blended, as a multistage polymer, wherein at least one of the stages is a polyimide according to this invention, with another thermoplastic polymer. When the thermoplastic polymer itself is a multistage polymer, the second polymer may also be a multistage polymer and preferably the outermost stage of the second polymer contains imide units.

Generally, when blending polyimides prepared according to this invention, with other thermoplastic polymers, the amount of polyimide present may vary widely from 1% to 99%, by weight or the entire composition. In a preferred blend, the second polymer is present in an amount of up to 50%, based on the total weight of said thermoplastic polymer plus said second polymer. In another preferred blend the second polymer is present in an amount from 50% to 99%, based on the total weight of said thermoplastic polymer plus said second polymer.

The advantage of blending a polyimide of this invention with at least one other thermoplastic polymer is that the resultant blends generally exhibit a better combination of properties which may include better melt viscosity, greater compatibility and/or miscibility, greater weatherability, greater impact resistance, better ductility, and better solvent resistance than the combination of properties of blends of polyimides which have not undergone diminution or substantial removal of acid and anhydride functionality.

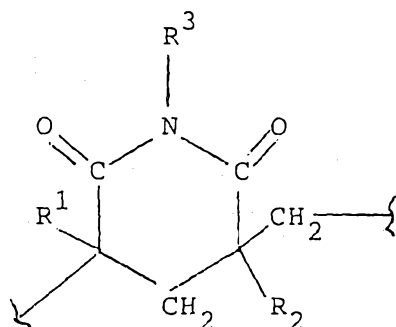
The polyimides of this invention, particularly the polyglutarimides, may be blended with one or more other thermoplastic polymers (as used herein "-" indicates blended polymers, "/" copolymers, and "/" graft or block polymers) such as butadiene/styrene/(meth)acrylic, styrene/(meth)acrylic, and (meth)acrylic multistage polymers; butadiene/-styrene rubbers, ethylene/propylene/diene rubbers, polyamides, polyamide-multistage polymer blends, ethylene/vinyl acetate, styrene/acrylonitrile, methyl styrene/acrylonitrile, styrene/acrylonitrile-multistage polymer blends, styrene/acrylonitrile-ethylene/propylene/diene rubber blends, alpha-methyl-styrene/acrylonitrile, alpha-methyl-styrene/styrene/acrylonitrile, alpha-methyl-styrene/methyl methacrylate/ethyl acrylate, butadiene//acrylonitrile/styrene, polycarbonate, polycarbonate-multistage polymer blends, polybutylene terephthalate, polybutylene terephthalate-polycarbonate blends, polybutylene terephthalate-multistage polymer blends, polybutylene terephthalate/polytetrahydrofuran, polyvinyl chloride, polyvinyl chloride-multistage polymer blends, polyvinyl chloride-(meth)acrylate blends, chlorinated polyvinyl chloride, acrylonitrile/-

(meth)acrylate-multistage polymer blends,
acrylonitrile/(meth)acrylate/styrene, epichlorohydrin/-
bisphenol-A, polyethylene terephthalate or other
polyalkylene terephthalate, polyethylene terephthalate-
5 glycol modified, polyethylene terephthalate-
polycarbonate blends, polycaprolactone, polyarylate,
copolyester of bisphenol-A with isophthalic and/or
terephthalic acids, poly(meth)acrylates, polyamide,
polystyrene, polyacetal, polystyrene, high-impact
10 polystyrene, styrene/maleic anhydride,
styrene/maleimide, polyolefins, polyvinylidene
fluoride, polyvinylidene fluoride-multistage polymer
blends, polyvinylidene chloride, cellulose, polyethylene
oxide, polyamideimide, polyetherester,
15 polyetheresteramide and polyetheramide. Blends having
modified performance characteristics can be prepared
from any other polymer systems with polyimides. These
include polyphenylene sulfide, polyphenylene oxide,
polysulfone, polyphenylene oxide-
20 styrene blends, polyphenylene oxide-high impact
polystyrene blends, polyvinylidene chloride,
polyvinylidene chloride/(meth)acrylonitrile,
polyvinylidene chloride/(meth)acrylate, polyvinyl
alcohol, polyvinyl acetate, polyetheretherketone,
25 polyetherimide and thermoplastic polyimides. Random or
block copolymers which combine the different
functionalities can also be used, and ternary or higher
blends can be made from combinations of any of these
polymer types with the imide polymers.

30 The imide polymers and blends with other polymer
systems can be further modified by the incorporation of
glass or other fibers, or particulate fillers or
reinforcing agents such as talc or other minerals,
glass spheres, or metallic particles. In general, the
35 imide polymers exhibit good adhesion to glass and other

inorganic fillers. Further modification of properties of the polyimides and blends can be obtained by incorporating additives such as flame retardants, blowing agents, antioxidants, thermal stabilizers, pigments, flatting agents, lubricants, antistatic agents, conductive materials, toners and/or UV stabilizers.

This invention is preferably concerned with a thermoplastic polymer comprising up to 95% imide units of the structural formula



wherein R¹, R² and R³ independently represent hydrogen or C₁ to about C₂₀ unsubstituted or substituted alkyl, aryl, or mixtures thereof, said polymer being non-crosslinked and soluble in dimethyl formamide, said polymer containing substantially no acid and/or anhydride groups.

In the Examples which follow, certain tests are conducted. These tests are as follows:

The Vicat is determined by ASTM Test Method 1525-65T using 40 mil (0.1 cm) penetration and heating rate B.

The total acid and anhydride functionality was determined by titrating solutions of the polymers in either 1:1 methylene dichloride-methanol or 3.75:1 dimethyl sulfoxide-water. A 0.3 g sample of the

polymer is dissolved in 75 ml (75 cm^3) of methylene dichloride-methanol. A measured volume of 0.1 N sodium hydroxide solution, containing more than enough base to neutralize the acid and anhydride in the sample, is added to the polymer solution, and the excess base is titrated with 0.1 N hydrochloric acid. The total acid and anhydride functionality is the difference, in milliequivalents, between the added base and the acid solution used to reach neutrality, corrected by subtracting the acid titration value of a solvent blank (the blank is usually less than 5% of the total acid and anhydride functionality). If dimethylsulfoxide is used, the polymer is dissolved in 75 ml (75 cm^3) of dimethylsulfoxide, and 20 ml (20 cm^3) of water is added to the solution, which is then titrated as above. In the samples below, the end point of the titration was detected either colorimetrically or potentiometrically, and in some instances other solvents were used to prepare the polymer solution for titration.

The total acid and anhydride functionality of a copolymer or blend containing polymer components that are imidizable is calculated as milliequivalents per gram of the imidizable polymer components.

Molecular weight is determined by gel permeation chromatography using a commercial Waters instrument and a tetrahydrofuran solvent, and is a weight average molecular weight.

Dynamic thermogravimetric analysis (TGA), as used herein, is a standard test conducted using a programmed temperature-increase rate of 20°C per minute and an atmosphere of either air or nitrogen, as indicated. Results given herein for TGA were either obtained using, or are consistent with those obtained using, a DuPont thermogravimetric analyzer in

combination with differential thermal analyzer;
specific operating instructions for the TGA instrument
and for this particular test are described in E. I.
DuPont de Nemours, Inc. Instrument Products Division
5 Preliminary Product Bulletin 950-1(A-36177).

Although compatibility and/or miscibility of
blended polymers is not necessary to the compositions
of the present invention, compatibility and/or
miscibility of such blends may be enhanced by reduction
10 of the acid and anhydride functionality. Measurements
of glass transition temperature (T_g) and visual
examination of polymer samples, as described below,
help to quantify observations of such compatibility
and/or miscibility enhancement.

15 The glass transition temperature (T_g) of polymer
and blend samples is determined by differential
scanning calorimetry (DSC), using a Perkin-Elmer Model
DSC-2 calorimeter. The sample is heated in a nitrogen
atmosphere at 20°C per minute, and the reference
20 material in the calorimeter is 60-80 mg aluminum. The
 T_g value is taken from the DSC chart as the temperature
at the midpoint of the transition. All blend samples,
prior to the DSC run, are preheated for one minute at a
temperature 20°C higher than the higher T_g value for
25 either component in the blend, and then cooled to a
temperature below the lower T_g value of either
component.

The DSC values for T_g are usually sufficient to
indicate whether a blend is miscible. However, if the
30 T_g values of the two components are similar, or if one
of the components undergoes a phase change (e.g.
melting or recrystallization) near the T_g of the other
component, the DSC result may not be adequate to detect
whether the sample is miscible. In such a case other

techniques for detecting miscibility may be used. For example, domains indicating incompatibility in blend samples may be detected by transmission electron microscopy (TEM), and the sensitivity of this technique may, under some conditions, be enhanced, as by staining one of the phases. An example of this is staining phases containing carbon-carbon unsaturation using osmium tetroxide.

Compatibility or miscibility of polymer blends is determined by an examination of the blend of the polyimide with a specified polymer after mastication of the blend in a Brabender mixing device at a temperature of from 193 to 246°C for ten minutes in an air atmosphere. Samples of the resulting blend are removed from the melt, formed into small, flattened "buttons" and examined visually for clarity. With some blends, the refractive indices of the two polymers are similar at room temperature, and incompatibility is difficult to detect by visual examination of the button sample. In such a case the button is observed at several different temperatures; the refractive indices of two immiscible components will change at different rates if one component is heated above its glass transition temperature.

In the examples which follow, the percent nitrogen of the polyimides used is set forth. The degree of imidization, by weight, may be calculated from the percent nitrogen given: divide the percent of nitrogen actually present in the polyimide by the theoretical amount of nitrogen at 100% imidization and multiply by 100.

In order to more fully illustrate the nature of this invention and the manner of practicing the same, the following examples are presented.

Examples 1 through 10

A polyglutarimide having N-methylimide groups, prepared in accordance with the procedure disclosed in US-A-4246374, is used. The imide was prepared from a feedstock of poly(methyl methacrylate) having a weight average molecular weight of 200,000.

The extruder used is a 2.0 cm-diameter (0.8 inch) Welding Engineers non-intermeshing twin screw extruder. The extruder has one section about six diameters long, for introducing and plasticating the polyglutarimide followed by a closed barrel section about 6 diameters long for pumping and pressure generation, followed by the reaction zone. This zone, about 31.5 diameters long, comprises a ported barrel section for introducing blocking agents, closed barrel sections, and a venting section for removing volatile products. Another vented section, operated at substantially atmospheric pressure, follows the reaction zone, for removing additional volatiles.

The extruder is purged with the polyglutarimide, to remove contaminants. The polyglutarimide is then added to the plastication zone of the extruder at a rate of 60 grams per minute. At this time the barrel temperature in the reaction zone is about 300°C, and the blocking agent pressure is about 3500 KiloPascals (500 psig). The polymer resides in the extruder for an average of about 120 seconds.

The polyglutarimide used has a nitrogen content of 7.13 weight percent, as determined by micro Kjeldahl analysis, and a Vicat of 170°C.

The initial, total acid and anhydride of the polyglutarimide used is 0.585 ± 0.020 milliequivalents per gram of imidizable component in the polymer.

In ten separate experiments, the polyglutarimide described above was processed in the extruder. For eight of the runs, trimethyl orthoformate was used for removing anhydride and acid functionality from the polyglutarimide. The extruder screw speed was, for all examples, 400 rpm. The trimethylorthoformate (TMOF) was pumped through a port into the inlet section of the extruder where it reacted with the polyglutarimide at a temperature of from 200°C to 350°C and a pressure of 3450 \pm 70 KiloPascals (500 \pm 10 psig).

In Table I, the letter "d" means that, prior to introduction in the extruder, the polyglutarimide is dried, under vacuum, at a temperature of 135°C. for eight to twelve hours. The letter "u" means that the polyglutarimide is not dried prior to introduction into the extruder. In Examples 3 and 4, no excess TMOF is added to the extruder. Table I sets forth the results obtained.

15 00 01003

TABLE I

	Example No.	Agent fee rate (grams/ minute)	Ratio (equivs. /equivs.) of Agent to Acid and Anhydride	Extruder Reac. Zone Temp. °C.	Vicat of final pro- ducts °C.	Acid & Anhy- dride Content of Final Prod- uct (meq./g/)
5	1 d+	none	0.00	313	170	0.585
	2 u+	none	0.00	304	170	0.569
	3 d	1.9	0.52	298	165	0.264
	4 u	1.9	0.52	290	170	0.545
10	5 d	4.4	1.17	291	160	0.032
	6 u	4.8	1.30	298	160	0.300
	7 d	8.2	2.21	286	160	0.033
	8 u	7.8	2.08	292	160	0.032
	9 d	10.7	2.86	297	160	0.065
15	10 u	10.9	2.92	299	161	0.031

+ Comparative

Example 4 illustrates that when the polyglutarimide is not dried prior to reaction with the agent, there is an effect on the amount of agent available to react with acid and anhydride groups. In Example 5 (dried), a slight excess of agent, over the stoichiometric amount required, is used. The slight excess in Example 6 (undried), which is substantially the same as Example 5, results in a higher acid and anhydride content because of the reaction of water with the agent used, whereas in Example 5, the reaction is substantially quantitative. Example 8 (undried) illustrates that the effect of the reaction of water and TMOF can be overcome by using a sufficient excess of TMOF over the stoichiometric amount required.

15 Examples 11 through 14

The procedure of Examples 1-10 was repeated except that the agent used was triethyl orthoformate (TEOF), to convert the acid and anhydride functionality to an ethyl ester; all polyimides used were undried; the polyimide feed rate was 3 grams/minute; the TEOF feed rate is set forth in Table II; the screw speed of the extruder was 250 rpm; the temperature in the zone for reacting TEOF with the acid and anhydride groups was 302°C. The percent nitrogen in each of the polyimides of the examples was 7.13 wt. %. The initial acid and anhydride level of polyimide used was 0.585 ± 0.020 milliequivalents per gram (meq./g) determined as in Examples 1-10. The results are set forth in Table II. The pressure in the reaction zone was 2900 KiloPascals (420 psig) for Example 12; 2345 KiloPascals (340 psig) for Example 13, and 2070 KiloPascals (300 psig) for Example 14.

TABLE II

Example No.	Agent Feed rate (grams/minute)	Ratio (equivs./equivs.) of agent to acid and anhydride	Vicat of final product (°C.)	Acid & Anhydride Content of Final Product (meq./g.)
11+	none	0.00	170	0.585
12	6.3	2.11	157	0.066
13	13.8	4.62	158	0.081
14	22.4	7.50	158	0.093

+ Comparative

10 Examples 13 and 14 show that the removal of acid and anhydride groups is substantially complete in Example 12 and that the additional agent used in Examples 13 and 14 has little benefit over the amount of agent used in Example 12. The amount of acid and anhydride remaining in Examples 12, 13 and 14 are substantially the same and are within the experimental error of the titration method used, which is the same as is used in Examples 1-10.

Examples 15 through 20

20 The procedure of Examples 11 through 14 were repeated except that the agent used was trimethyl orthoformate (TMOF) and the polyimide used is a 201°C-Vicat ammonia polyimide which has a nitrogen content of 7.72 weight %. The polyimide used had an acid and anhydride content of 0.610 milliequivalents per gram of polyimide, as determined by titrating it in a 3.75/1.0 dimethylsulfoxide/water solution. The polyimide samples used were not dried. The polyimide

feed rate was 30 grams per minute. The pressure in the reaction zone was 2205 KiloPascals (320 psig) for Example 16; 3585 KiloPascals (520 psig) for Example 17; 3655 KiloPascals (530 psig) for Example 18; 3205 KiloPascals (465 psig) for Example 19; and 3035 KiloPascals (440 psig) for Example 20.

The results are set forth in Table III.

TABLE III

Example No.	Agent Feed rate (grams/minute)	Ratio (equivs./equivs.) of agent to acid and anhydride	Vicat of final product (°C.)	Acid & Anhydride Content of Final Product (meq./g.)
15+	none	0.0	201	0.610
16	1.0	0.48	198	0.432
17	1.9	0.97	198	0.311
18	2.9	1.45	192	0.000
19	3.9	1.93	190	0.000
20	7.7	3.81	192	0.000

+ Comparative

20 Examples 21 through 27

The purpose of these examples is to illustrate a continuous process for making polyimides and reducing or substantially eliminating acid and anhydride functionality. In this continuous process, the amount of acid and anhydride of the polyimide prior to treatment with an agent may be determined by removing a sample of polyimide prior to reaction with the agent for reducing acid and anhydride functionality and

titrating the amount of acid and anhydride present. A calculated amount of the agent, based on the desired acid and anhydride level, may then be added.

Alternatively, prior to running the continuous process,

5 one may determine the amount of agent to add by preparing the polyimide under the same conditions to be used in the continuous process, and then measuring the acid and anhydride level. As long as the feed stock is converted to the same polyimide and treated with the
10 agent under the same conditions, the resultant final product should be substantially the same in both cases.

The apparatus of examples 1-10 is used in examples 21 through 27 except that an imidization zone measuring 2.0 cm by 61 cm (0.8 inch by 24 inches) is

15 added to the apparatus as the first zone. The imidization zone also contains a double screw. The feed which is added to the imidization zone is poly(methyl methacrylate) (PMMA) having a weight average molecular weight of 148,000. The PMMA feed
20 rate is 45 grams per minute. To the PMMA in the imidization zone is added methylamine. The barrel temperature of the imidization zone was 302°C. \pm 5°C.

The gauge pressure in the imidization zone for Example 21 was 6137 KiloPascals (890 psig); for Example 22 was
25 6205 KiloPascals (900 psig); for Example 23 was 6275 KiloPascals (910 psig); for Example 24 was 6275 KiloPascals (910 psig); for Example 25 was 6171 KiloPascals (895 psig); and for Example 26 was 7240 KiloPascals (1050 psig). In Example 21 and 26 the
30 polyimides were not treated with any agent after preparation. The procedure of Examples 1-10 for removing acid and anhydride was then repeated. The

agent used was trimethyl orthoformate (TMOF). The temperature of the TMOF/polyimide reaction zone was 304°C. \pm 5°C. The pressure in the TMOF/polyimide reaction zone for Example 22 was 2413 KiloPascals (350 psig); for Example 23 was 3172 KiloPascals (460 psig); for Example 24 was 3103 KiloPascals (450 psig); for Example 25 was 3068 KiloPascals (445 psig); for Example 26 was atmospheric pressure (0.00 psig); and for Example 27 was 3103 KiloPascals (450 psig).

10 Table IV sets forth the results obtained.

25 0 00 01003

TABLE IV

5	Example No.	Rate formation of polyimide (gram/min)	Methyl-amine feed rate (Grams/min)	Agent feed rate (grams/min)	Ratio (equivs./equivs.) of agent to acid and anhydride	Poly-imide Vicat (°C.)	% Nitro-gen of Poly-imide	Poly-imide -wt. av ₃ Mw x 10 ⁻³	Acid & Anhydride Content of final product (meq./g.)
10	21+	38.8	14.6	0.00	0.00	168	6.80	131	0.575
	22	38.8	14.6	1.46	0.61	164	6.63	130	0.251
	23	38.8	14.6	3.10	1.31	157	6.83	127	0.000
	24	38.8	14.6	4.85	2.05	158	6.79	131	0.000
	25	38.8	14.6	8.73	3.69	157	6.77	138	0.000
	26+	38.4	19.9	0.00	0.00	184	7.34	119	0.678
	27	38.4	17.9	8.73	3.16	168	7.33	128	0.000

15 + Comparative

Examples 28 to 34

The procedure of Examples 11 through 14 was repeated with the following changes. In Examples 28, 31, 33 and 37 the imides were not treated with an agent to substantially remove or reduce acid and anhydride functionality. The agent used in Examples 29, 30 and 32 was dimethyl carbonate (DMC). Examples 30 and 32 also utilized triethylamine (TEA) as a catalyst for the reaction between the agent and the acid and anhydride groups. The TEA was added at a feed rate of 0.09 gram per minute in Example 20, and 0.11 gram per minute in Example 32. The agent used for Examples 34-36 was 2,2-dimethoxypropane (DMP). The polyimide used in Examples 28 through 30 was polymethylimide having a Vicat of 172°C and a nitrogen content of 7.04 weight percent. The polyimide used in Examples 31 and 32 was a polymethylimide having a Vicat of 150°C and a nitrogen content of 4.90 weight percent. The polyimide used in Examples 33-36 was a polymethylimide having a Vicat of 170°C and a nitrogen content of 7.24 weight percent. The polyimide used in Examples 37 and 38 was a polymethylimide having a Vicat of 162°C. and a nitrogen content of 3.08 weight percent. The pressure in the agent reaction zone was 2900 KiloPascals (420 psig) for Example 29; 2793 KiloPascals (405 psig) for Example 30; 2137 KiloPascals (310 psig) for Example 32; 2124 KiloPascals (308 psig) for Example 34; 2827 KiloPascals (410 psig) for Examples 35 and 36; and 2137 KiloPascals (310 psig) for Examples 38.

Examples 28-30 used polyimide dried in the manner of Examples 1-10. Examples 31-38 used an undried polyimide. The initial acid and anhydride level of the polyimide of Examples 28-30 was 0.618 milliequivalents per gram (meq/g) of polyimide. The

polyimide of Examples 33-36 had an initial acid and anhydride level of 0.570 meq/g. The polyimide of Examples 31 and 32 was prepared from a 90/10 methyl methacrylate/styrene feedstock and contained 4.90

5 weight percent nitrogen and an initial acid and anhydride level of 0.689 meq/g of imidizable polymer component. The polyimide of Examples 37 and 38 was prepared from a 73.5/24.5/2.0 methyl methacrylate/
10 alpha-methylstyrene/ethyl acrylate feedstock, and had a nitrogen content of 3.08 weight percent and an initial acid and anhydride level of 0.975 meq/g of imidizable polymer component. The results are set forth in Table V.

TABLE V

Example No.	Agent Feed Rate (grams/minute)	Ratio (equivs./equivs.) of agent to acid and anhydride	Agent ¹	Vicat of final product (°C.)	Acid & Anhydride Content of Final Product (meq./g.)
5	28+	none		172	0.618
	29	4.3	DMC	163	0.087
	30	4.2	DMC TEA	162	0.005
	31+	none		150	0.689
	32	5.2	DMC TEA	144	0.020
10	33+	none		170	0.570
	34	2.1	DMP	169	0.478
	35	4.4	DMP	166	0.376
	36	8.6	DMP	166	0.345
	37+	none	none	162	0.975
15	38	9.0	TMOF	150	0.040

¹ DMC = dimethyl carbonate
TMOF = trimethyl orthoformate

TEA = triethylamine

DMP = 2,2-dimethoxypropane

+ Comparative

Examples 35 through 44

These examples illustrate the removal of acid and anhydride functionality from polyimides using a solution reaction. The polyimide used for Examples 35 through 44 was the polyimide of Examples 1-10. The procedure for Examples 35-44 is as follows.

To a 3-neck, 500 ml round-bottom flask equipped with a magnetic stirrer and reflux condenser is added 250 ml of a solvent. A 12.5 g polyimide sample is dissolved in the solvent and the agent is then added to the solution. The resulting solution is then heated to a temperature of from 160°C to 175°C, under nitrogen, for varying periods of time. The solution is then cooled to room temperature and the final product is precipitated by adding it to methanol; it is then filtered and dried. The acid and anhydride level of the final product is determined as in Examples 1-10. The results are given in Table VI.

25 0 06 61113

TABLE VI

5	Example No.	Solvent	Agent ¹ and grams of Agent	Reaction Temp. (°C.)	Reaction Time (hrs.)	Ratio (equivs./equivs.) of agent to acid and anhydride	Acid & Anhydried Content of Product (meq./g.)
	35+	diglyme	none/0.0	160	8.0	0.0	0.508
	36	diglyme	TMOF/7.6	160	4.0	9.8	0.170
	37	diglyme	TMOF/7.6	160	8.0	9.8	0.061
10	38	triglyme	TEOF/10.6	175	2.0	9.8	0.385
	39	triglyme	TEOF/10.6	175	3.5	9.8	0.227
	40	triglyme	TEOF/10.6	175	5.0	9.8	0.128
	41	triglyme	TEOF/10.6	175	7.0	9.8	0.063
	42	triglyme	TEOF/10.6	175	8.2	9.8	0.065
15	43	dimethylsulfoxide	DMSO/275	170	3.0	482	0.263
	44	dimethylsulfoxide	DMSO/275	170	7.0	482	0.217

- 37 -

1
20
TMOF = trimethyl orthoformate
TEOF = triethylorthoformate
DMSO = dimethylsulfoxide
diglyme = diethyleneglycol dimethyl ether
triglyme = triethyleneglycol dimethyl ether
+ Comparative

EXAMPLES 45 through 49

The purpose of these Examples is to demonstrate the change in chemical resistance for the various polyimides which have been prepared. The following
5 Examples utilize the imides prepared in earlier Examples.

Example 45 utilizes the polyimide prepared in Example 1

10 Example 46 utilizes the polyimide prepared in Example 12.

Example 47 utilizes the polyimide prepared in Example 5.

Example 48 utilizes the polyimide prepared in Example 26.

15 Example 49 utilizes the polyimide prepared in Example 27.

The imides used in Examples 45 and 48 were not treated to remove acid and anhydride groups. The polyimides used in Examples 48 and 49 were prepared
20 using a continuous process for the preparation of the polyimide and removal of acid and anhydride groups.

The polyimides used in these Examples were injection molded to form 5.1 x 1.3 x 0.32-cm (2 x 0.5 x 0.125-inch) bars. The weights of the molded bars were
25 determined and each bar was immersed in a different solvent, which was kept at a temperature of 23°C, for seven days. The bars were then removed from the solvent and the percent weight gain or weight loss was then measured by reweighing the bars. For determining

water absorption, a 0.25-mm-thick film of imide was prepared by compression molding pre-dried imide pellets. The film was dried, weighed, immersed in water for fourteen days at a temperature of 50°C, and reweighed.

5

The results are set forth in Table VII

TABLE VII

Example No.	Water Absorption (% Weight Loss or Gain)	Percent Wt. Change-Immersion at 23°C for 7 days					Toluene
		Ammonium Hydroxide (10% aq.)	Methanol	Ethanol	1,1,1-tri-chloro-ethane		
5	45+	4.9	12.3	14.9	1.7	0.37	0.18
	46	3.3	0.98		No Data		
	47	3.8	0.83	7.1	0.64	-13	-35
10	48+	6.0	16.5	19.3	2.52	0.23	0.28
	49	3.9	0.85	8.3	0.82	-11	-33

+ Comparative

As can be seen from Table VII, those polyimides which have had acid and anhydride levels reduced or substantially eliminated absorb less water than polyimides having untreated acid and anhydride levels. This indicates that weatherability would be better for those polyimides having reduced acid and anhydride levels.

Table VII also demonstrates that resistance to ammonium hydroxide, methanol and ethanol is better for those polyimides which have been treated to substantially remove or reduce acid and anhydride levels when compared to untreated polyimides.

Table VII also demonstrates that the treated polyimides are somewhat soluble in 1,1,1-trichloroethylene and toluene whereas the untreated polyimides are not soluble. This indicates that the treated polyimides may be dissolved, to some extent, in some solvents and may therefore be used as coating compositions whereas the untreated polyimides are not soluble in those solvents.

Examples 50 through 57

These examples demonstrate the effect of weathering on various polyimides which have been treated with an agent to remove acid and anhydride groups. The polyglutarimides used in Examples 54 through 57 correspond to the polyglutarimides used in Examples 50 through 53 except that the procedure of Examples 1 through 10 is used to treat the polyglutarimides. The agent used in treating the polyglutarimides is, in each case, trimethyl orthoformate. Example 54 is the treated counterpart of Example 50; Example 55 is the treated counterpart of Example 51; Example 56 is the treated counterpart of

Example 52; and Example 57 is the treated counterpart of Example 53. The milliequivalents of acid and anhydride per gram of imidizable group are set forth in Table VIII. The weight percent nitrogen for each of the polyglutarimides used is also set forth in Table VIII.

The weathering tests employed a Xenon Arc Weather-o-meter in the modified (extreme) test mode, according to ASTM method G26-83. Test samples were molded into 5.1 x 7.6 x 0.32-cm (2 x 3 x 0.125-inch) plaques. The test conditions include a 2-hour cycle: 80 minutes dry (76.7°C black body temperature, 50% relative humidity) and 40 minutes wet (sprayed distilled water). The light level was 6500 watts throughout. Table VIII sets forth the results obtained.

TABLE VIII

5	Example No.	% Nitrogen	Milliequivalents of Acid & Anhydride Per Gram	Total Hrs. of Exposure in Weather-o-meter vs Degree of Surface Crazing ¹					
				500	1000	1500	2000	2500	3000
	50+	5.28	0.349	N	N	N	VL	VL	MH
	51+	6.27	0.460	N	N	N	VL	VL	H
	• 52+	6.87	0.624	N	N	VL	L	H	VH
	53+	7.59	0.798	N	VL	LM	MH	VH	VH
10	54	5.66	0.033	N	N	N	N	N	N
	55	6.39	0.035	N	N	N	N	N	N
	56	7.06	0.002	N	N	N	N	N	N
	57	7.04	0.002	N	N	N	N	N	MH

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- 1
- 15 N = none
VL = very light
L = light
LM = light to medium
M = medium
MH = medium to heavy
- 20 H = heavy
VH = very heavy

+ Comparative

As can be seen from Table VIII, when a polyglutarimide is treated to substantially remove or reduce the acid and anhydride levels, the weatherability of the resultant treated polyglutarimide is much superior to that of the untreated polyglutarimides.

EXAMPLES 58 through 72

A number of N-methyl polyglutarimides were prepared in accordance with the teachings of US-A-4,246,374. Some of the polyglutarimides were then treated with an agent in the manner of Examples 1 through 10, using the agent used in Examples 1 through 10. The polyglutarimides prepared in Examples 58 through 72 were then evaluated for miscibility and compatibility with other polymers. A letter designation is assigned to each example and that letter designation is utilized in later examples in which miscibility and compatibility properties are given. Table IX sets forth the polyglutarimide and its properties. The Tg was determined by differential scanning calorimetry.

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Molecular Weights are determined by gel permeation chromatography in tetrahydrofuran. Milliequivalents of acid and anhydride per gram of imidizable group is determined by titration using methylene dichloride/methanol as set forth in other examples.

EXAMPLES 73 through 81

A number of 50/50 weight/weight blends were prepared with the polyglutarimides of Examples 58 through 72. The polymer blended with the various polyglutarimides was a styrene-acrylonitrile copolymer composed of 75% by weight styrene and 25% by weight acrylonitrile. The Tg for the polyglutarimide, the blend of polyglutarimide and polymer, and the polymer with which the polyglutarimide was blended, was determined. The presence of a single Tg for the blend indicates complete compatibility.

The blends were prepared by melt mixing the polyglutarimide and the styrene-acrylonitrile copolymer in a Brabender melt mixer at a temperature of 232°C. Small (0.9 x 0.15-cm) buttons were formed from the melt.

Table X sets forth the results obtained.

TABLE X

Example No.	Polyglutarimide Used (Table IX)	Observed Tg Values °C	Appearance of Blend
73+	I	165/113	Opaque
74	J	165/113	Opaque
75	K	161/114	Translucent
76	L	154/119	Almost Clear
77	M	128	Clear
78+	E	178/113	Opaque
79	O	131	Clear
80+	E	164/112	Translucent
81	F	130	Clear
82+	C	157/113	Translucent
83	D	125	Clear
84+	A	125	Clear
85	B	124	Clear

+ Comparative

In Examples 73 through 77, the following should be noted. The glutarimide of Example 73 has not been treated with any agent to remove acid and anhydride functionality. The polyglutarimides of Examples 74 through 77 have been treated to remove acid and

anhydride functionality but the amount of acid and anhydride remaining varies so that, as one proceeds from Example 74 through Example 77, the milliequivalents of acid and anhydride in each of these examples changes. Example 73 has 0.585

milliequivalents of acid per gram of imidizable group; Example 74 has 0.512 milliequivalents of acid; Example 75 has 0.354 milliequivalents of acid; Example 76 has 0.191 milliequivalents of acid; and Examples 77 has

0.045 milliequivalents of acid. As the amount of acid and anhydride in the polyglutarimide decreases, the compatibility of the polyglutarimide with the styrene-acrylonitrile copolymer increases as is seen by the appearance or by the single Tg of the blend. In

Examples 78 through 85, all of the even-number examples are of polyglutarimides which have not been treated to remove acid and anhydride functionality. The immediate following, odd-numbered example is of the same

polyglutarimide treated to remove acid and anhydride functionality. It is therefore clear that

compatibility of the treated polyglutarimides with styrene-acrylonitrile copolymer is enhanced. Attention is invited to Example 84, wherein the polyglutarimide is not treated and the blend is clear and shows a

single Tg for the blend. It is known from US-A-4436871 that certain polyglutarimides are compatible with styrene-acrylonitrile copolymers even though no steps have been taken to remove acid and anhydride functionality.

EXAMPLES 86 through 89

The procedure of Examples 73 through 85 was repeated except that the polymer with which the polyglutarimide was blended is changed. The polymer

used in Examples 86 through 89 was Abson 89140 which is an ABS (acrylonitrile/butadiene/styrene terpolymer supplied by Mobay Chemical Company.) All polymer blends were opaque to translucent rather than clear, due to the fact that the initial index of refraction between the rubber and hard phases does not match. However, appearance of a single Tg for the blend indicates compatibility of the polyglutarimide with the ABS polymer (Tg values for the rubber phase are not shown). The results are set forth in Table XI.

TABLE XI

Example Number	Polyglutarimide Used	Observed Tg Values (°C)	Appearance of Blend
86+	N	168/108	Opaque
87	O	129	Semi Opaque
88+	E	155/108	Translucent
89	F	128	Hazy clear

+ Comparative

As is seen from Table XI, Examples 86 and 88 utilize polyglutarimides wherein no steps have been taken to remove acid and anhydride functionality from the polyglutarimide. The results of Examples 87 and 89 show a single Tg for the blend thus demonstrating compatibility of the treated polyglutarimide with the polymer. The material in Example 87 is the material from Example 86, but treated. Similarly, the material in Example 89 is the material from Example 88, but treated.

EXAMPLES 90 through 97

The procedure of Examples 73 through 85 was repeated except that the polymer used was Blendex 586, obtained from Borg-Warner Company. It is believed that Blendex 586 is a terpolymer of alpha methylstyrene/styrene/acrylonitrile. The results are expressed in Table XII.

TABLE XII

Example No.	Polyglutarimide Used	Observed Tg Values °C	Appearance
90+	N	171/129	Opaque
91	O	142	Clear
92+	G	161/128	Opaque
93	H	138	Clear
94+	C	140	Clear
95	D	137	Clear
96+	A	136	Clear
97	B	134	Clear

+ Comparative

Attention is invited to the fact that in Examples 90 and 92, the polyglutarimide is not treated to remove acid and anhydride functionality and the blends are opaque. In Examples 94 and 96, the polyglutarimides are also not treated to remove acid and anhydride functionality and the blends are clear

and exhibit only a single Tg for the blend. The difference resides in the fact that the degree of imidization for Examples 96 and 94 are 67% and 78%, respectively, and the acid and anhydride levels for Example 96 is 0.504 milliequivalents and for Example 94 is 0.452 milliequivalents. When Examples 96 and 94 are compared with Examples 90 and 92, it is seen that Examples 90 and 92 have a higher degree of imidization and a higher acid and anhydride content. The degree of imidization for Example 90 is 91% and the amount of acid and anhydride is 0.798 milliequivalents and the degree of imidization for Example 92 is 82% and the acid and anhydride content is 0.624 milliequivalents. Although the blends for certain non-treated polyglutarimides are clear and exhibit only a single Tg, other properties are altered, such as weatherability.

EXAMPLES 98 to 119

Examples 98 to 119 show that polycarbonate of two different molecular weights and Phenoxy resin (a condensation polymer of epichlorohydrin and bisphenol-A) all show partial to almost complete miscibility with various imides upon melt blending. The alkylated imides uniformly show higher miscibility with these polycarbonate polymers. The similarity of imide and polycarbonate Tg's makes the DSC data in some of the below cases a less reliable indication of miscibility. In these cases the physical appearance is a more important gauge of miscibility. The results of these examples are shown in Table XIII.

TABLE XIII

	Exam- ple No.	Imide Source (Exam- ple)	Imide Vicat (°C) Type ¹	Imide Tg(°C) by ² DSC	Imide %by wt. ³	meq/g total acid ⁴	Imide Polymer Ratio	Observed Tg(°C) by ⁵ DSC	Appear- ance ⁶
5	98+	71	179UC	175	91	0.798	50/50	178/150	OP
	99	72	165/C	163	90	0.002	50/50	165/152	OP/TL
	100+	62	167/UC	164	81	0.513	50/50	166/152	OP
	101	63	157/C	158	84	0.011	50/50	155 (Bd ⁷)	OP/TL
10	102+	60	165/UC	162	78	0.452	50/50	149 (Bd ⁷)	OP
	103	61	157/C	155	76	0.094	50/50	146 (Bd ⁷)	OP/TL
	104+	58	150/UC	147	67	0.504	50/50	143 (Bd ⁷)	OP
	105	59	146/C	143	67	0.101	50/50	142 (Bd ⁷)	OP/TL
	106+	62	167/UC	164	81	0.513	80/20	159	TL
	107	63	157/C	158	84	0.111	80/20	159	TL/TP
	108+	62	167/UC	164	81	0.513	20/80	150	TL
	109	63	157/C	158	84	0.011	20/80	152	TL/TP
	110+	71	179UC	175	91	0.798	50/50	175/147	OP
	111	72	165/C	163	90	0.002	50/50	164/148	OP/TL
	112+	62	167/UC	164	81	0.513	50/50	167/159	OP
	113	63	157/C	158	84	0.011	50/50	152	OP/TL
	114+	60	165/UC	162	78	0.452	50/50	149 (Bd ⁷)	OP
	115	61	157/C	155	76	0.094	50/50	149 (Bd ⁷)	OP/TL
	116+	58	150/UC	147	67	0.504	50/50	149 (Bd ⁷)	OP
25	117	59	146/C	143	67	0.101	50/50	140 (Bd ⁷)	OP/TL
	118+	64	170/UC	168	82	0.624	50/50	162/104	OP/TL
	119	65	160/C	158	84	0.002	50/50	118	Clear

- 1 Imides are all undried methyl imides; UC =
unalkylated, C = alkylated.
- 2 Undried imides.
- 3 Imide weight % imidization level calculated for
5 methyl imides form $100 (\%N/8.377\%) = I$.
- 4 Titrable acid in the imide per gram of sample (0.020
meq/g, titrated in MDC/MeOH - 1/1)
- 5 Since the PC and imide Tgs are similar, appearance
is a better test of miscibility in this case.
- 10 6 OP = Opaque; TL = translucent; TP = transparent;
CL = clear.
- 7 Bd = Broad
- + Comparative

Note: The blended polymer for Examples 98 - 109 is
15 Polycarbonate M50, for Examples 110 - 117 is
Polycarbonate M39, and for Examples 118 and
119 is Phenoxo resin.

EXAMPLES 120 to 125

20 Examples 120 and 121 show that Dylark 332
(approximately 86/14 - styrene/maleic anhydride, by our
analysis) is miscible with a 67% imidized alkylated
methyl imide (imide from Example 59), but not with the
corresponding unalkylated imide (imide from Example
58).

25 An ammonia imide was prepared by treating Dylark
332 at 49 g/m and 260°C in an extruder with 3.9 g/m
anhydrous ammonia at 2826 kPa or 410 psig. according to
US-A-4246374. The resulting imide contained from 0.000
to 0.070 meq/g of acidic functionality and had a
30 nitrogen level of 1.88 weight percent (for 14.1%
maleimide, % N theoretical = 2.01%), indicating about
94 weight % imidization. The Vicat for Dylark 332 was
136°C, (Tg = 130°C) and for the ammonia imide was 151°C
(Tg = 145°C).

When the same unalkylated and alkylated methyl imides used in Examples 120 and 121 were melt blended with this Dylark-based ammonia imide, both blends were observed to be largely to completely miscible, indicating an inherently greater interaction between the acrylic imides and the maleimides versus the maleic anhydride polymer. When the Dylark 332 ammonia derivative was melt blended with a more highly imidized, unalkylated (Example 124) and alkylated (Example 125) acrylic imide, miscibility differences could again be observed between the alkylated and unalkylated blends. Following the previous pattern, the alkylated imide (imide F, Example 125) was completely miscible, while the unalkylated imide (imide E, Example 124) was only partially miscible.

TABLE XIV

	Exam- ple No.	Imide Source (Exam- ple)	Imide Vicat (°C) / Type ¹	Imide Tg (°C) ² DSC	Imide % by wt. ³	meq/g total acid ⁴	Polymer/ Polymer Tg	Poly- mer Ratio	Observed Tg (°C) by DSC	Appear- ance ⁵
5	120+	58	150/UC	147	67	0.504	Dylark- 332/130	50/50	143/130	OP
	121	59	146/C	143	67	0.101	Dylark- 332/130	50/50	134	TL
10	122+	5	150/UC	147	67	0.504	Imide A ⁶ / 145	50/50	147	TL/CL ⁷
	123	59	146/C	143	67	0.101	Imide A ⁶ / 145	50/50	146	CL
	124+	62	167/UC	164	81	0.513	Imide A ⁶ / 145	50/50	150 Bd ⁸	OP
15	125	63	157/C	158	84	0.011	Imide A ⁶ / 145	50/50	153	CL

¹ Imides are all undried methyl imides; UC = unalkylated, C = alkylated.

² For undried imides.

³ Imide weight % imidization level calculated for methyl imides from 100
(%N/8.377%) = % I.

⁴ Milliequivalents of titratable acid in the imide per gram of sample
(± ca. 0.020 meq/g, titrated in MDC/MeOH - 1/1).

⁵ OP = opaque; TL = translucent; TP = transparent; CL = clear.

⁶ Imide A is the ammonia imide prepared from Dylark 332, as described in the text.

⁷ Clear with a trace of turbidity.

⁸ Bd = Broad

+ Comparative

EXAMPLES 126 to 134

Examples 126 to 134 show that polyvinylidene fluoride (Kynar 720) is miscible with alkylated methyl imide but not unalkylated methyl imide (at this imidization level), thus following the same pattern as seen previously. The blends in examples 128 and 129 were prepared on a Brabender melt-mixing device, while examples 130 to 134 were extrusion blended on a 1" (25 cm) Killion extruder. Kynar polyvinylidene fluoride (PVDF) is a crystalline polymer which shows a large crystallization endotherm (CE) by DSC analysis, which obscures any nearby amorphous Tg in polyvinylidene fluoride and any nearby Tg in the blended imide polymer (Example 128). Since it is crystalline, it is also translucent to opaque. The miscibility of polyvinylidene fluoride and the imide from Example 65 (Example 129) suppresses crystallization, giving an amorphous polymer blend with a single Tg. The levels of clarity and Tg's of the blends indicate that while either miscibility or mixing was not complete in all samples, the level of miscibility was quite high. The tensile properties of the imide/Kynar blends also indicate that as little as 5% Kynar polymer gives a ductile material as measured by tensile elongation. The percent elongation to break for Examples 126 and 130-134 are: 5% for Example 126, (no yield), 11% for Example 134, (greater than 6% elongation indicates yielding), 67% for Example 133, 72% for Example 132, 83% for Example 131, and 101% for Example 130. That imides may be used to provide an amorphous polyvinylidene fluoride is also clear from these results.

TABLE XV

	Ex- am- ple No.	Imide Source (Ex- am- ple)	Imide Vicat (°C) / Type ¹	Imide Tg (°C) by ² DSC	Imide % by ³ wt.	meq/g total ⁴ acid	Blended Polymer/ Polymer Tg	Imide/ Poly- mer Ratio	Observed Tg (°C) by ⁵ DSC	Appear- ance ⁸
5	126+	64	170/UC	168	82	0.624	-	100/0	168	CL
	127+	-	-	-	-	-	Kynar ⁵ 720/CE ⁵	0/100	CE ⁵	OP/TL ⁶
10	128+	64	170/UC	168	82	0.624	Kynar ⁵ 720/CE ⁵	50/50	CE ⁵	OP/TL ⁶
	129	65	160/C	158	84	0.002	"	50/50	78 ⁵	CL
	130	65	160/C	158	84	0.002	"	75/25	126	TL
	131	65	160/C	158	84	0.002	"	80/20	130	TL
	132	65	160/C	158	84	0.002	"	85/15	138	T/CL ⁷
15	133	65	160/C	158	84	0.002	"	90/10	145	CL/T ⁷
	134	65	160/C	158	84	0.002	"	95/5	152	CL

¹ Imides are all undried methyl imides; UC = unalkylated, C = alkylated.

² For undried imide.

³ Imide weight % imidization level calculated for methyl imides from $100 (\%N/8.377\%) = \% I$.
⁴ Milliequivalents of titratable acid in the imide per gram of sample (\pm ca. 0.020 meq/g, titrated in MDC/MeOH - 1/1).

⁵ CE indicates crystallization endotherm which obscures any amorphous Tg in polyvinylidene fluoride and any nearby Tg in the blended imide polymer.

⁶ Polyvinylidene fluoride is translucent to opaque due to crystallinity.

⁷ Appearance T/CL was translucent-to-clear (clear with a trace of turbidity), while CL/T was completely clear.

⁸ OP = Opaque; TL = Translucent; TP = Transparent; CL = Clear

+ Comparative.

EXAMPLES 135 to 143

The miscibility of Kynar 720 polyvinylidene fluoride (PVDF) may be exploited to provide enhanced impact resistance in imide blends containing grafted rubber impact modifiers. Example 135 shows typical impact, tensile, and heat-distortion temperature data for an impact-modified blend of unalkylated, 170°C Vicat methylimide, in which the impact modifier (A) was designed to give a high notched Izod impact resistance. Example 136 shows that the same imide, when alkylated, maintains good isotropic impact resistance (Gardener falling dart impact, GVHIT), but loses notched Izod impact resistance. However, when the modifier (B) is specifically designed for the difference in miscibility of the alkylated imide (Example 138), an excellent notched Izod and falling dart impact resistance is maintained. These values diminish in the unalkylated imide (Example 137). Examples 139 and 140 show that as small amounts of PVDF are substituted for the alkylated imide in the modifier B formulation, surprisingly large increases in notched Izod impact value are observed. At 10% PVDF, the impact resistance is high enough to remove 10% rubber modifier and still maintain good impact resistance (Example 141). Fixing the PVDF level at 10% in the formulation in Example 140, and substituting modifier A again shows that lower, but good notched Izod impact values are still obtained.

TABLE XVI

5	Exam- ple No.	Imide Source (Example)	Imide Vicat (°C)/ Type ¹	meq/g total acid	Mod. ² Type	Ratio Imide PVDF/ Mod. (wt. %)	Notched Izod 23°C 0°C Joules/m	Tensile Impact Joules/ cm	GVHIT Joules cm(0.318- cm-thick plaque)	Tensile Prop.		
										% Elong. @ Yield (break)	Modulus x 1000 kPa	DTUFL °C @ 180 kPa (455 kPa)
10	135+	64	170/UC	0.624	A	60/00/40	106.5 38.4	17.4	22.8	8.6 (20)	2220	134 (149)
	136	65	160/C	0.002	A	60/00/40	53.4 22.4	25.2	25.3	5.4 (67)	2503	116 (130)
	137+	64	170/UC	0.624	B	60/00/40	64.6 36.3	10.5	9.38	6.5 (16)	2048	129 (144)
	138	65	160/C	0.002	B	60/00/40	206.7 37.9	29.4	26.2	6.0 (52)	2165	116 (132)
20	139	65	160/C	0.002	B	55/05/40	242.4 61.9	27.1	31.0	5.6 (75)	1896	105 (122)
	140	65	160/C	0.002	B	50/10/40	331.1 138.8	32.4	31.5	5.1 (89)	1751	97 (113)
	141	65	160/C	0.002	B	60/10/30	180.5 38.4	25.6	30.5	5.3 (88)	2179	107 (121)
	142+	64	170/UC	0.624	A	50/10/40	185.3 36.3	21.0	>18	8.6 (36)	1972	107 (127)
25	143	65	160/C	0.002	A	50/10/40	121.2 46.5	46.2	36.2	5.3 (31)	1862	101 (110)

- ¹ Imides are all undried methyl imides. UC = unalkylated, C = alkylated.
² Modifier A = Butylacrylate (65%)/Styrene/Acrylonitrile (26/9%).
 Modifier B = Butylacrylate (65%)/Styrene/Acrylonitrile (30/5%).
 + Comparative

EXAMPLES 144 to 153

As noted in US-A-4255322, methyl imides and polyvinyl chloride (PVC) resins are miscible over a relatively wide range. However, we have found that high imidization levels, high imide acid levels, and changes in the polyvinyl chloride resins can cause immiscibility in certain instances. Examples 144-153 illustrate that reducing the acid and anhydride functionality of the imide can correct such immiscibility. Example 144 shows a 50/50 imide/polyvinyl chloride blend (Geon 103 PVC with a K value of 67) is only partially miscible, giving two sharply defined Tg values, while the same blend, made with alkylated imide (Example 145), is completely miscible.

However, when these same imides are melt blended with a lower molecular weight PVC (Example 146) (Geon 85, K value = 62), both alkylated (Example 147) and unalkylated (Example 146) imides are completely miscible with the PVC. Although the Geon 85 resin appears to be more miscible with imides at lower imidization levels, only partial miscibility is observed with this resin when imides having even higher imidization and acid levels are blended with the PVC resin as in Example 148. This example shows one very broad and uneven Tg by DSC, indicating partial miscibility, whereas the same imide, when alkylated (Example 149), shows complete miscibility and one sharp Tg. The indices of refraction of the imides having 90% imidization or greater are close enough to the PVC refractive index so that appearance is not a conclusive factor.

Examples 150-153 show that the same enhanced miscibility seen with alkylated imides (Examples 144 and 145) is also seen in the two formulations given below, which represent typical polyvinyl chloride siding formulations for twin-screw extruder applications, and which are based on the less-miscible Geon 103 PVC. In the blends and siding formulations, we have also observed that the alkylated imide/PVC materials of the present invention show a better combination of properties than the unalkylated PVC/imide blends. A given, alkylated PVC/imide blend may show less color generation, a lower initial viscosity, a slower viscosity rise (as measured by torque in Brabender melt processing experiments), high impact resistance and heat distortion resistance.

Component:	PVC Formulation I (PVC I)	PVC Formulation II (PVC II)
Geon 103	100.0	100.0
MMA/EA proc. aid ¹	0.5	1.0
BA/S//S/MMA proc. aid ²	1.0	0.5
Me-Tin stabilizer	1.6	1.6
Calcium stearate	1.3	1.3
Paraffin wax	1.0	1.0
TiO ₂	10.0	10.0

- ¹ methyl methacrylate/ethyl acrylate processing aid
² butyl acrylate/styrene//styrene/methyl methacrylate processing aid

Results of these Examples are given in Table XVII.

TABLE XVII

	Exam- ple No.	Imide Source (Exam- ple)	Imide Vicat (°C) Type ¹	Imide Tg (°C) by ² DSC	Imide % by wt. ³	meq/g total acid ⁴	Blended PVC Polymer/ Polymer Tg	Imide/ Poly- mer Ratio	Observed Tg (°C) by DSC	Appear- ance ⁵
5	144+	62	167/UC	164	81	0.513	Geon 103/80	50/50	156/101	OP/TL
	145	63	157/C	158	84	0.011	Geon 103/80	50/50	123	CL ⁶
	146+	62	167/UC	164	81	0.513	Geon 85/80	50/50	106	CL ⁶
	147	63	157/C	158	84	0.011	Geon 85/80	50/50	100	CL
10	148+	71	179/UC	175	91	0.798	Geon 85/80	50/50	115	CL
	149	72	165/C	163	90	0.002	Geon 85/80	50/50	102	CL
	150+	62	167/UC	164	81	0.513	PVC I	50/50	158/96	OP ⁷
	151	63	157/C	158	84	0.011	PVC I	50/50	119	OP ⁷
	152+	62	167/UC	164	81	0.513	PVC II	50/50	156/98	OP ⁷
15	153	63	157/C	158	84	0.011	PVC II	50/50	116	OP ⁷

¹ Imides are all undried methyl imides; UC = unalkylated, C = alkylated.

² Undried Imide.

³ Imide weight-% imidization level for methyl imides = $100 \times (\%N/8.377\%)$.

⁴ Milliequivalents of titratable acid in the imide per gram of sample (\pm ca. 0.020 meq/g titrated in MDC/MeOH - 1/1).

⁵ OP = opaque; TL = translucent; TP = transparent; CL = clear.

⁶ Appearance was clear with a trace of turbidity.

⁷ PVC formulations I and II (see text) contain TiO_2 , and are therefore opaque regardless of miscibility.

+ Comparative

EXAMPLES 154 to 207

The following examples illustrate polyimide blends having 90/10 and 10/90 weight ratios. These blends show that the enhancement of properties can occur for blends containing both small and large amounts of the imides having reduced acid and anhydride functionality. The imides used were prepared in the manner of Examples 52 and 56, respectively; the imide sample treated according to the present invention contained 7.18 weight percent nitrogen and had a weight-average molecular weight of 129,000, and the comparative imide without reduction of the acid and anhydride functionality contained 6.87 weight percent nitrogen and had a weight-average molecular weight of 145,000. Additional properties of the imides, the polymers used for blending, and the blends are given in Table XVIII.

TABLE XVIII

	Exam- ple No.	Imide Vicat (°C) Type ¹	Imide Tg (°C) by ¹ DSC ¹	Imide % by wt. ¹	meq/g total acid ¹	Blended ⁸ Polymer	Imide/ Poly- mer Ratio	Observed Tg (°C) by ¹ DSC ¹	Appear- ance ²
5	154+	170/UC	168	82.01	0.63	-	100/0	168	CL
	155	160/C	158	85.71	0.002	-	100/0	158	CL
						PVDF	0/100	CE ³	OP/TL
	156+	170/UC	168	82.01	0.63	"	90/10	158 ³	OP
10	157	160/C	158	85.71	0.002	"	90/10	146 ³	CL/H-
	158+	170/UC	168	82.01	0.63	"	10/90	92 ³	OP
	159	160/C	158	85.71	0.002	"	10/90	96	TL
						Sty/AN	0/100	113	CL
	160+	170/UC	168	82.01	0.63	"	90/10	163/110	CL/H
15	161	160C	158	85.71	0.002	"	90/10	152	CL
	162+	170/UC	168	82.01	0.63	"	10/90	169/115	TL
	163	160/C	158	85.71	0.002	"	10/90	114	CL
						PETG	0/100	CE ³	CL
	164+	170/UC	168	82.01	0.63	"	90/10	168/89	CL/H-
20	165	160/C	158	85.71	0.002	"	90/10	158/90 ³	CL/H-
	166+	170/UC	168	82.01	0.63	"	10/90	164/95 ³	TL/OP
	167	160/C	158	85.71	0.002	"	10/90	156/96 ³	TL
						PC	0/100	152	CL
	168+	170/UC	168	82.01	0.63		90/10	163	TL
25	169	160/C	158	85.71	0.002		90/10	155	CL/H-

TABLE XVIII (cont'd)

	Exam- ple No.	Imide Vicat (°C) / Type ¹	Imide Tg (°C) by ¹ DSC ¹	Imide % by wt. ¹	meq/g total acid ¹	Blended Polymer ⁸	Imide/ Poly- mer Ratio	Observed Tg (°C) by ¹ DSC ¹	Appear- ance ²
5	170+	170/UC	168	82.01	0.63	PC	10/90	169/148	OP
	171	160/C	158	85.71	0.002		10/90	149	CL/H-
						PET	0/100	84 ³	OP/TL
	172+	170/UC	168	82.01	0.63		90/10	155	TL
10	173	160/C	158	85.71	0.002		90/10	158/77	TL
	174+	170/UC	168	82.01	0.63		10/90	161/87 ³	OP
	175	160/C	158	85.71	0.63		10/90	154/85 ³	OP/H
						PA	0/100	CE ³	OP
	176+	170/UC	168	82.01	0.63		90/10	153 ⁺	TL/OP
15	177	160/C	158	85.71	0.002		90/10	153 ⁴	OP
	178+	170/UC	168	82.01	0.63		10/90	CE ⁴	OP
	179	160/C	158	85.71	0.63		10/90	152 ⁴	OP
						PS	0/100	191	CL
14	180+	170/UC	168	82.01	0.63		90/10	166/189	OP
20	181	160/C	158	85.71	0.002		90/10	159/181	OP
	182+	170/UC	168	82.01	0.63		10/90	164/187	OP
	183	160/C	158	85.71	0.002		10/90	158/188	OP
						Sty/MAH	0/100	130	CL
	184+	170/UC	168	82.01	0.63		90/10	166/129	TP
25	185	160/C	158	85.71	0.002		90/10	157 ⁵	CL/H-

TABLE XVIII (cont'd)

	Exam- ple No.	Imide Vicat (°C) Type ¹	Imide Tg (°C) by ¹ DSC ¹	Imide % by wt. ¹	meq/g total acid ¹	Blended ⁸ Polymer	Imide/ Poly- mer Ratio	Observed Tg (°C) by ¹ DSC ¹	Appear- ance ²
5	186+	170/UC	168	82.01	0.63	Sty/MAH	10/90	164/133	OP/H
	187	160C	158	85.71	0.002		10/90	154/132	OP/H
						AN/MA/BD	0/100	-	TL
	188+	170/UC	168	82.01	0.63		90/10	163/98	TL
	189	160/C	158	85.71	0.002		90/10	153/81	TL
	190+	170/UC	168	82.01	0.63		10/90	83 ⁶	TL
	191	160/C	158	85.71	0.002		10/90	83 ⁶	TL
						EVA	0/100	-	TL
	192+	170/UC	168	82.01	0.63		90/10	168/CE ³	OP
15	193	160/C	158	85.71	0.002		90/10	157/CE ³	OP
	194+	170/UC	168	82.01	0.63		10/90	CE ³	CL/H-
	195	160/C	158	85.71	0.002		10/90	155/CE ³	CL/H-
						PVC	0/100	-	CL
	196+	170/UC	168	82.01	0.63		90/10	161 ⁷	DEC
20	197	160/C	158	85.71	0.002		90/10	152	TP
	198+	170/UC	168	82.01	0.63		10/90	91 ⁷	DEC
	199	160/C	158	85.71	0.002		10/90	102	CL
						Phenoxy	0/100	102	CL
	200+	170/UC	168	82.01	0.63		90/10	163/112	CL
25	201	160/C	158	85.71	0.002		90/10	152	CL/H-

TABLE XVIII (cont'd)

Exam- ple No.	Imide Vicat (°C) / Type ¹	Imide Tg (°C) by ¹ DSC ¹	Imide % by wt. ¹	meg/g total acid ¹	Blended Polymer ³	Imide/ Poly- mer Ratio	Observed Tg (°C) by ¹ DSC ¹	Appear- ance ²
202+	170/UC	168	82.01	0.63	Phenoxy	10/90	160/99	TL/CL
203	160/C	158	85.71	0.002		10/90	99	CL
					Nylon 12	0/100	-	TL
204	170/C	168	82.01	0.63		90/10	162/CE ⁴	CL/TL
205	160/C	158	85.71	0.002		90/10	154/CE ⁴	TL
206+	170/UC	168	82.01	0.63		10/90	CE ⁴ /73	TL
207	160/C	158	85.71	0.002		10/90	CE ⁴ /70	TL

THE FOOTNOTES FOR TABLE NO. XVIII ARE AS FOLLOWS

- 1 As described for Table XVII. C = alkylated and UC = unalkylated.
- 2 CL = Clear; TP Transparent; TL = Translucent; OP = Opaque; DEC = Decomposed; H = Haze (H+ = Heavy Haze; H- = Slight Haze).
- 3 Crystallization endotherm is observed in DSC.
- 4 Crystallization endotherm obscures the Tg.
- 5 Appears as though two Tgs are overlapping.
- 6 The Barex 210/Imide (90/10) blends degrade at a temp. on or around the Tg of the Imide on the Brabender melt mixing device

7 Considerable polymer degradation is observed in all the Geon 85/imide blends, i.e., considerable color formation is observed, but the alkylated imide/Geon 85 blends appear to be more stable.

5 8 Polymers and sources:

PVDF - Polyvinylidene fluoride (Kynar 720 - Pennwalt Corp.)

Sty/AN - 75/25 Styrene/acrylonitrile copolymer (Tyril 1000 - Dow Chemical Co.)

10 PETG - Glycol-modified polyethylene terephthalate (Kodar A-150 Eastman Kodak Co.)

PC - polycarbonate, M_w 20,000 (M39, - Mobay, Inc.)

PET - Polyethylene terephthalate

15 PA - Polyacetal copolymer (M90 - Celanese Corp.)

PS - Polysulfone (Udel P1700 - Union Carbide Corp.)

20 Sty/MAH - Styrene/maleic anhydride copolymer (Dylark 332 Arco Chemical Co.)

AN/MA/BD - Acrylonitrile/methyl acrylate/butadiene (Barex 210 - Vistron Cor.)

EVA - Ethylene/vinyl acetate copolymer (Elvax 750 E. I. DuPont de Nemours & Co., Inc.)

25 PVC - Polyvinyl chloride (Geon 85 - BF Goodrich Co.)

Phenoxy - Epichlorohydrin-bisphenol A condensation polymer (PKHC - Union Carbide Corp.)

30 Nylon 12 - (Grilamid - Emser Industries)

EXAMPLES 208 to 219

The following examples illustrate blends of polyimide copolymers with polystyrene (Sty) and polyphenylene oxide-high impact polystyrene blends

(HIPS/PPO; Noryl, General Electric). The polyimides having acid and anhydride functionality reduced according to the present invention were an imidized 90/10 methyl methacrylate/styrene (MMA/Sty) copolymer having a nitrogen content of 4.95 weight percent (Example 209), and an imidized, 73.5/24.5/2.0 methyl methacrylate/alpha-methylstyrene/ethyl acrylate (MMA/AMS/EA) terpolymer having a nitrogen content of 2.93 weight percent (Example 211). The corresponding polymers not having reduced acid and anhydride functionality (Examples 208 and 210) contained 4.90 and 3.08 weight percent nitrogen, respectively. Additional properties of the polyimides and blends are given in Table XIX.

TABLE XIX

	Exam- ple No.	Imide Source (Exam- ple)	Imide Vicat (°C) / Type ¹	Imide Tg (°C) by ¹ DSC ¹	Imide % by wt. ¹	meq/g total acid ¹	Blended Polymer	Blending Tempera- ture	Observed Tg (°C) by ¹ DSC ¹	Appear- ance ³
5	208+	(2)	150/UC	148	77.8	0.620	(MMA/Sty)	-	148	CL
	209	(2)	144/C	142	78.57	0.018	(MMA/Sty)	-	142	CL
	210+	(2)	162/UC	160	58.0	0.736	(MMA/AMS/EA)	-	160	CL
	211	(2)	150	147	55.49	0.030	(MMA/AMS/EA)	-	147	CL
10	212+	208	150/UC	148	77.8	0.620	Sty	450°F.	144/84	TL
	213	209	144/C	142	78.57	0.018	Sty	450°F.	134/85	TL
	214+	208	150/UC	148	77.8	0.736	HIPS/PPO	450°F.	144 ⁴	OP
	215	209	144/C	142	78.57	0.030	HIPS/PPO	450°F.	142 ⁴	OP
	216+	210	162/UC	160	58.0	0.736	Sty	450°F.	159/141/87	TL
15	217	211	150/C	148	55.49	0.030	Sty	450°F.	139/85	TL
	218+	210	162/UC	160	58.0	0.736	HIPS/PPO	450°F.	162/138 ⁴	OP
	219	211	150/C	148	55.49	0.030	HIPS/PPO	450°F.	145 ⁴	OP

¹ As described for Table XVIII, C = alkylated; UC = unalkylated

² See text

20 ³ CL = Clear; TL = Translucent; OP = Clear

⁴ Small enthalpic peak at 110°C in DSC of Noryl blends is unidentified.
+ Comparative

EXAMPLES 220 to 253

The following examples illustrate a series of 50/50 polyimide/thermoplastic blends that were prepared in a Brabender mixer at the temperature indicated in the accompanying Table; an air atmosphere was used. The polyimide used is the polymethylimide F (Example 63), having a Vicat value of 157°C, a Tg of 158°C, 84% imidization, by weight, and a total acid and anhydride functionality of 0.011 meq/g, reduced according to the present invention. The properties of the blends are given in Table XX.

TABLE XX

Exam- ple No.	Blended Polymer ⁵	Blending Tempera- ture	Observed Tg by DSC (°C)	Appear- ance
15 220	Polyarylate	302°C	155/192	OP
221	Polystyrene	232°C	156/101	OP
222	Polycarbonate (M50)	246°C	155 ²	OP
223	Polysulfone	316°C	155/187	OP
224	Nylon 12 (Grilamid)	232°C	45 ³	TL
225	Nylon 6	232°C	156/53 ⁴	TL
226	Nylon 11	249°C	153/51 ⁴	OP
227	Nylon 66	274°C	157/64 ⁴	OP
228	PETG (6763)	221°C	153/80	OP
229	Polybutylene terephthalate	223°C	150/56 ⁴	OP
25 230	Polyethylene terephthalate	254°C	153/86 ⁴	OP
231	Poly(methyl methacrylate)	232°C	149/116	OP

TABLE XX cont

Exam- ple No.	Blended Polymer ⁵	Blending Tempera- ture	Observed Tg by DSC (°C)	Appear- ance
5				
	232 Styrene/acrylonitrile (75/25)	450°C	130	OP
	233 Aromatic Nylon (Trogamid)	260°C	153	OP
	234 PVC (Geon 85)	199°C	100	CL
50	235 PETG (Kodar A 150)	216°C	149/95	OP
100	236 ABS	232°C	128	TL
150	237 Polycarbonate (M39)	246°C	152 ²	TL
	238 Polyacetal (M90)	210°C	CE ³	OP
	239 Styrene/maleic anhydride	232°C	153/131	OP
	240 Styrene/NH ₃ -Maleimide	246°C	153	CL/H-
15	241 Styrene/N-Methyl Maleimide	246°C	154/133	OP
	242 alpha-Me Sty/Sty/AN copolymer	246°C	138	CL
	243 Polyphenylenesulfide (PPS)	302°C	158/110	OP
	244 Polyetherimide	327°C	158/219	OP
200	245 MA/AN/BD	199°C	142/82	TL
200	246 Phenoxy	193°C	118	TL
	247 Poly(ethylene/8% acrylic acid)	193°C	158 ⁴	OP
	248 Polyvinylidene fluoride	246°C	78	CL
	249 Chlorinated polyvinyl chloride	199°C	149	OP/DEC

TABLE XX cont

Exam- ple No.	Blended Polymer ⁵	Blending Tempera- ture	Observed Tg by DSC (°C)	Appear- ance
250	Polyethylene	199°C	156 ⁴	TL/CL
251	Polypropylene	204°C	CE ⁴	OP
252	Ethylene/vinyl acetate	193°C	156 ⁴	OP
253	Polyvinylidene chloride	193°C	99	TL/CL

- 10 1 OP = opaque, TL = translucent; CL = clear; H = haze
(- is slight); DEC = decomposition.
- 2 Polycarbonate Tg overlaps imide Tg.
- 3 Crystallization endotherm obscures imide Tg.
- 4 Crystallization endotherm observed in DSC.
- 15 5 Polymers and Sources:
- Polyarylate is Ardel D-100 (Union Carbide Corp.)
Polystyrene is PS 656D (Dow Chemical Co.)
- Polycarbonate M39 is M_w 20,000 (Mobay Inc.)
Polycarbonate M50 is M_w 24,600 (Mobay Inc.)
- Polysulfone is UDEL P1700 (Union Carbide Corp.)
- Nylon 12 is Grilamid (Emser Industries)
- Nylon 11 is BMNO (Rilsan Corp.)
- Nylon 66 is Zytel 101 (E. I. DuPont de Nemours & Co.)
- Nylon 6 is Fosta 446 (American Hoechst Corp.)
- 25 PETG 6763 is Glycol modified polyethylene terephthalate
(Eastman Kodak Co.)

Polybutylene terephthalate is Celanex 2002 (Celanese Corp.)

Aromatic Nylon is Trogamid T (Kay-Fries Co.)

PVC is polyvinyl chloride Geon 85 (BF Goodrich Co.)

5 PETG Kodar A-150 is glycol-modified polyethylene terephthalate (Eastman Kodak Co.)

ABS is butadiene//acrylonitrile/styrene Abson 89140 (Mobay Inc.)

Polyacetal M90 is a product of Celanese Corp.

10 Polyvinylidene chloride is Experimental PVDC Resin XU-32009.00 (Dow Chemical Co.)

Other polymer descriptions may be found in the footnotes to Table XVIII.

EXAMPLE 254

15 In a manner similar to that of Examples 242-253, a 50/50 blend of the polyimide from Example 20 with alpha-methylstyrene/acrylonitrile copolymer was prepared at 246°C; the resulting clear blend had a single Tg of 138°C.

20 EXAMPLE 255

In a manner similar to that of Example 233 a 50/50 blend of polyimide (having a Vicat value of 160°C, a Tg of 158°C; 82% imidization and 0.002 meq/g total acid and anhydride functionality, reduced according to the present invention) with Noryl (polyphenylene oxide-high impact polystyrene blend) was prepared. The resulting opaque blend showed Tg values at 158°C and 138°C.

25 In a manner similar to the above examples, 30 copolymers of styrene and maleimide, containing residual maleic anhydride functionality, such as Dylark DKB-162, in which approximately 14% of the anhydride

functionality remains (see US-A-4458046 - Example 1), may be treated to decrease or substantially eliminate the anhydride functionality using the conditions of Examples 7 and 30.

5 Polymeric imides resulting from condensation of dianhydrides and diamines, including the generic class of polyimides referred to as polyetherimides, which contain residual acid functionality, may also be reacted with the agent disclosed above, using the
10 conditions of Examples 7 and 30, where the extrusion temperature is about 350-400°C, to give polyimides with reduced levels of acid functionality.

 Further, polymeric imides resulting from copolymerization of maleimides with other vinyl
15 monomers, such as copolymers of maleimide or N-phenyl maleimide and styrene and/or MMA, in which partial hydrolysis has occurred to give maleamic acids, diacids, or anhydrides, may be treated with the agents disclosed above, using the conditions in Examples 7 or
20 30, to give decreased levels of acid and anhydride functionality.

 Adhesion-enhanced impact resistance is expected in blends of polycarbonates with impact modifiers, and the imides of the present invention; such blends using
25 untreated imides are disclosed in US-A-4254232. Improved miscibility leads to better bonding, and thus to improved impact resistance.

 In general, all imides disclosed in US-A-4246374, especially those polymeric imides
30 containing up to 20% of the lower alkyl acrylates, which contain any residual acid or anhydride functionality, may be effectively treated by the methods disclosed above, such as the method described in Examples 7 and 30, to give polymeric imides with
35 reduced or substantially eliminated acid functionality.

The claims defining the invention are as follows:-

1. A thermoplastic polymer comprising imide groups and optionally acid and/or anhydride functional groups and having a degree of imidization of at least 1% characterized in that the degree of imidization on said polymer and the acid and/or anhydride functionality on said polymer are as set forth in the Table below;

when the degree of imidization on said polymer is	the milliequivalents of acid and/or anhydride per gram of imidizable polymer component
from 1% to 10%	0 to 0.20
above 10% to 20%	0 to 0.28
above 20% to 30%	0 to 0.32
above 30% to 40%	0 to 0.31
above 40% to 50%	0 to 0.28
above 50% to 60%	0 to 0.28
above 60% to 70%	0 to 0.27
above 70% to 80%	0 to 0.28
above 80% to 90%	0 to 0.33
and above 90% to less than 95%	0 to 0.40

2. A polymer according to claim 1 wherein the degree of imidization on said polymer is from 40% to 88%, preferably from 50% to 85%, and the milliequivalents of acid and/or anhydride present is from 0 up to 0.25, preferably less than 0.25.

3. A polymer as claimed in claim 1 or claim 2, wherein said polymer contains up to 50%, preferably from 10% to 50%, by weight, of acrylic or methacrylic acid ester units.

4. A polymer as claimed in claim 1 or claim 2, wherein said polymer contains from 50% to 99%, by weight, of ester units derived from acrylic or methacrylic acid.

5 5. A polymer as claimed in claim 3 or claim 4, wherein the ester moiety contains from 1 to 20 carbon atoms, preferably the ester moiety is methyl.

10 6. A polymer according to any one of the preceding claims, wherein said polymer is a multistage polymer and either the outermost stage and/or the core and/or any other intermediate stage if present of said polymer comprises imide groups.

15 7. A process for the preparation of a thermoplastic polymer comprising imide groups and optionally acid and/or anhydride functional groups and having a degree of imidization of at least 1%, said process comprising reacting a ~~first~~ polymer comprising imide groups and active acidic and/or anhydride functional groups with a predetermined amount of acidic and/or anhydride functional group blocking agent to reduce or
20 substantially eliminate the active acid and/or anhydride functional groups in the polymer.

25 8. A process as claimed in claim 7, wherein said process is carried out at a temperature of from 50°C to 450°C.

9. A process as claimed in claim 7 or claim 8, wherein said process is carried out at a pressure of from 1 atmosphere (101.3 KiloPascals) to 1000 atmospheres (101300 KiloPascals).



10. A process as claimed in any one of claims 7 to 9, wherein said blocking agent is an alkylating agent and/or esterification agent.

5 11. A process as claimed in claim 10, wherein said blocking agent is one or more of orthoesters, ketals, carbonates, sulfoxides, siloxanes, silyl ethers, silylenol ethers, trialkyl phosphates, trialkyl phosphites, dialkyl sulfates, alkyl alkylsulfonates, alkyl arylsulfonates, carbodiimides, trialkylsilyl
10 halides, enol esters, alcohols, alkyl esters, alkyl isocyanates, quaternary ammonium salts, ureas and/or guanidines; and is preferably one or more of dimethyl carbonate, 2,2-dimethoxypropane, dimethyl sulfoxide, diethyl sulfoxide, triethyl orthoformate, trimethyl
15 orthoacetate, trimethyl orthoformate, diphenyl carbonate, dimethyl sulfate, methyl toluenesulfonate, methyl trifluoromethylsulfonate, methyl acetate, methanol, ethanol, methyl isocyanate, p-chlorophenyl isocyanate, dimethyl carbodiimide, dimethyl
20 t-butylsilyl chloride, isopropenyl acetate, dimethyl urea, tetramethyl ammonium hydroxide, dimethyl diethoxysilane, tetra-n-butoxysilane, dimethyl- (trimethylsilyl) phosphite, trimethyl phosphite, trimethyl phosphate, and tricresyl phosphate.

25 12. A process as claimed in any one of claims 7 to 11, wherein said reaction is conducted in a single or multi-screw extruder, in an autoclave, in a continuously recirculated pipe reactor, in a baffled in-line mixer which may be used in series with an
30 extruder, in a wiping-film evaporator, a falling film reactor or evaporator or in a disc extruder or reactor.

13. A process as claimed in any one of claims 7 to 12 wherein said polymer has a degree of imidization of up to 99%, preferably from 10% to 95%.

5 14. A blend of at least one thermoplastic polymer and a second polymer, said second polymer being a polymer as claimed in any one of claims 1 to 6 or a polymer prepared by the process as claimed in any one of claims 7 to 13.

10 15. A blend according to claim 14, wherein said second polymer is present in an amount of from about 1% to about 99%, based on the total weight of said thermoplastic polymer plus said second polymer.

16. A thermoplastic polymer according to claim 1, or a blend thereof, or a process for the preparation thereof, substantially as hereinbefore described with reference to the Examples.

~~17. The steps, features, compositions and compounds referred to or indicated in the specification and/or claims of this application, individually or collectively, and any and all combinations of any two or more of said steps or features.~~

Dated this 25th day of AUGUST, 1986

ROHM AND HAAS COMPANY
By Its Patent Attorneys
DAVIES & COLLISON

