POLY-PHENYLATED DIAMINES AND THEIR USE AS POLYCONDENSATION MONOMERS IN THE SYNTHESIS OF POLYAMIDE, POLY(AMIDE-IMIDE), AND POLYIMIDE POLYMERS

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Abstract

New polyphenylated polynuclear aromatic diamines, such as 1,3-bis[4-aminophenyl]-2,3,5-triphenylbenzene, a process for their manufacture and their use as polycondensation components for the manufacture of polyamide, polyamide-imide and polyimide polymers are described. The polymers obtained with the aromatic diamines according to the invention are readily soluble, rigid-rod polymers and are distinguished by outstanding modulus, tensile compression strength, energy absorption, coefficient of expansion and electrical properties.

15 Claims, No Drawings
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This invention was made with Government support under contract NAG-1-448 awarded by NASA. The Government has certain rights in this invention.

TECHNICAL FIELD OF THE INVENTION

The present invention relates to new poly phenylated aromatic diamines, a process for their manufacture and their use as polycondensation components for the manufacture of polyamide, polyamide-imide and polyimide polymers.

BACKGROUND OF THE INVENTION

Generally, aromatic polyamides, polyamide-imides and polypeptides are difficult to process and generally have low solubility. Some improvement in solubility has been achieved by preparing the polymers from bulky diamines or from dianhydrides containing 3 or more aromatic rings. (compare, for example, DT-AS No. 1,595,733 and DT-OS Nos. 2,009,739, 2,153,829, 2,257,996 and 2,321,513). However, these polymers either cannot be processed from the melt, or are difficult to process from the melt and, in some cases, have inadequate heat and/or chemical stability.

More recently Dams in U.S. Pat. No. 4,239,880 has disclosed polyamides, polyamide-imide and polyimide polymers prepared from exotic poly nuclear aromatic diamines. Although the polymers manufactured using Darm's diamines show improved solubility, these polymers do not form rigid rod polymers.

It, thus, appears desirable to produce polyamides, polyamide-imides and polyimides based on poly-phenylated diamines according to the present invention which would form soluble rigid-rod polymers having outstanding modulus, tensile and compression strength, excellent energy absorption characteristics, very low coefficient of expansion, excellent thermal stability, non-conduction, excellent film processability and relatively less expense that existing polymers. These properties will make the polymers ideally suited for use in the production of antiballistics, reinforced molecular composites, microelectronic coatings and gas, liquid and solid membrane separation applications.

DESCRIPTION OF THE INVENTION

It is an object of this invention to provide novel rigid-rod, high modulus, soluble poly-phenylated polyamides, polyamide-imides, polyimides, polyamide-amide-acids, polyamide-acids based on poly-phenylated 1,4-diphenylbenzene diamines where the polyphenylation is at the center benzene ring.

A further aspect of this invention is to provide novel copolymers comprising repeat units based on the new poly-phenylated diamines and one or more polycarboxylic acids, acid chlorides, esters, and other carboxylic acid derivatives and repeat units based on one or more other organic diamine(s) and one or more polycarboxylic acids, acid chlorides, esters, and other carboxylic acid derivatives where the polycarboxylic is intended to connote organic species with two or more carboxylic functions, preferentially where the polycarboxylic or-
When At1, At2, and At3 are all three phenyl and R1 is a hydrogen atom, then the above generic synthetic scheme describes the synthesis of 1,4-bis-[4-amino-phenyl]-2,3,5-triphenylbenzene, a diamine of formula (I), where 2,5-bis-[4-nitrophenyl]-3,4-diphenylcyclopentadiene is the corresponding compound of formula (II) and phenylacetylene is the corresponding compound of formula (III). Thus for At1, At2, and At3 are all three phenyl and R1 is a hydrogen atom, 2,5-bis-[4-nitrophenyl]-3,4-diphenylcyclopentadiene is reacted with phenylacetylene in an appropriate solvent to form 1,4-bis-[4-nitrophenyl]-2,3,5-triphenylbenzene, the corresponding intermediate of formula (I). Dinitro intermediate (I) is converted into the desired diamine 1,4-bis-[4-amino-phenyl]-2,3,5-triphenylbenzene, a diamine of formula (I), by chemical reduction.

The compounds of the formulas (II) and (III) are preferably employed in stoichiometric amounts. However, the reaction can also be carried out with a slight excess of one or the other reactant.

Chemical hydrogenation of intermediate (I) can be performed using well known hydrogenation agent such as iron or tin in an acid medium including ferrous sulphate, CaCl2 or sodium hydrogen sulphate, titanium dichloride and/or tetrachloride in the presence of HCl, zinc in an acid or neutral medium, optionally with the addition of neutral salts, including CaCl2 and NH4Cl, lithium aluminium hydride, hydrazines, including hydrazine hydrate and phenylhydrazine, if necessary with the addition of Raney nickel catalysts, and sodium dithionate (Na2S2O4). All the above reduction system are well known in the literature.

Catalytic hydrogenation can also be used employing suitable commercial catalysts in the presence of hydrogen such as palladium, palladium-on-charcoal, platinum, platinum black, platinum oxide and, above all, Raney nickel. The catalytic reduction is appropriately carried out in a suitable inert organic solvent, such as dioxane or methylcellulose.

The compounds of the formulas (II) and (III) are known or can be manufactured easily in a manner which is in itself known.

The diamines of this invention can be used as polycondensation components for the manufacturing of homopolymers and copolymers of novel polyphenylated polyamides, polyamide-amide-acids or polyamide-acids by reacting the diamines of formula (I) with one or more polycarboxylate of formula (IV)

where Z is an organic radical selected from the group consisting of an aliphatic radical, a cyclo-aliphatic radical, a carbocyclic-aromatic radical, or an hetero-cyclic aromatic radical, where Y is a halogen atom, a hydroxy group, an unsubstituted or substituted phenoxy group, an alkxy group preferably having from about 1 to 12 carbon atoms, and particularly preferred having from about 1 to 12 carbon atoms, where k is a whole number having a numeric value of 2, 3 or 4 and where each COY is bonded to a different atom of Z and when k is equal to 3 and Z is a cyclic organic radical, then two of the COY groups are in an ortho orientation relative to each other and when k is equal to 4 and Z is a cyclic organic radical, then each pair of COY groups are arranged in an ortho orientation relative to one another and one or more diamine of formula (V) and from about 99 to 0 mole percent of a repeat unit of formula (VI):

where Q is a divalent organic radical selected from the representative and illustrative group consisting of an aliphatic radical having at least 2 carbon atoms, a carbocyclic aliphatic radical, a carbocyclic aromatic radical, or a heterocyclic radical. The resulting polymers comprise from about 1 to 100 mole percent of a repeat unit of formula (VII):

and from about 99 to 0 mole percent of a repeat unit of formula (VIII):

where At1, At2, At3 and R1 are as previously defined, n and m are whole numbers separately and independently having the numeric value of either 0 or 1 and the sum of n and m must be equal to m minus 2 where m represents the number of sites involved in reaction with a diamine of formula (I) or a diamine of formula (V), Z' and Z'' is separately and independently an organic radical selected from the group defining Z of formula (IV) above and Q is one or more divalent organic radical as previously defined for diamines of formula (V). When the k of formula (IV) is equal to 3, then either m or n of formulas (VI) and (VII) is equal to 1 (the other is equal to 0), and the resulting polymers and copolymers are called polyamide-amide-acids. When k of formula (IV) is equal to 4, then both m or n of formulas (VI) and (VII) are equal to 1 and the corresponding polymer and copolymers are called polyamide-acids.

Polyamide-amide-acids comprising 1 to 100 mole percent of repeat units of formula (VI) and 99 to 0 mole percent of repeat units of formula (VII) can be cyclized into polyamide-imides which have 1 to 100 mole percent of repeat units of formula (VIII) shown below:
The polymers and copolymers of the present invention are intended to include all manners of copolymers incorporating at least 1 mole percent of repeat units of formula (VI). Since the repeat units of formula (VI) can include one or more polycarboxy compounds of formula (IV), the resulting polymers can include portions that are polyamide, polycarboxy-dicarboxylic acids and polycarboxy-amide-acids or any combination thereof. Thus, cyclization of copolymers that contain various amounts of polycarboxy-amide-acids and polycarboxy-acids can be achieved by standard chemical cyclization procedure well known in the literature.

In preparing the above polymers, a mixture of diamines of formulas (I) and (V) are reacted with a mixture of one or more polycarboxy compounds of formula (IV). The amount of diamine and polycarboxy compound should be close to one to one molar mixture. However, 10% excess of either component is acceptable. Once the polymerization has completed, the last component to react will determine the polymer end group. The polymer end groups can, thus, be an amino group, a carboxy group or a mixture thereof. Alternatively, a chain termination reagent can be added to the polymerization to force termination of the growing polymer. Such chain termination reagents are used to limit the molecular weight of the polymer and are well known in the art. Amine termination reagents commonly employed include aniline or substituted anilines. Common carboxy terminating reagents include benzoic acid or phthalic acid or their acid derivatives such as esters, acid chloride or similar acid derivatives.

The above polymerization reaction which yields polyanides and copolyamides of the present invention can be optionally carried out in the presence of a promoter such as lithium chloride or lithium carbonate. These lithium reagents are dissolved in the polymerization solvent in amounts approximately equal to the amount of amide linkages to be formed during the polymerization or in amounts approximately equal to twice the amount of diamine added.

When $k$ is equal to 2, polycarboxy compounds of formula (IV) are selected from the representative and illustrative group consisting of: malonic acid, dimethylmalonic acid, succinic acid, glutaric acid, adipic acid, suberic acid, sebacic acid and dodecanedioic carboxylic acid, 1,3-cyclohexane-dicarboxylic acid, hexahydro-1,2-naphthalene-2,6-dicarboxylic acid, thiophene-2,5-dicarboxylic acid and pyridine-2,3-dicarboxylic acid as well as the corresponding dichlorides and diesters according to the definition.

When $k$ is equal to 3, polycarboxy compounds of formula (IV) are selected from the representative and illustrative group consisting of: trimellitic acid 1,2-anhydride-chloride, (1,3-dioxo-benzoxalane-5-carboxylic acid chloride), trimellitic acid anhydride and trimellitic acid as well as esters and anhydrides according to the definition.

When $k$ is equal to 4, polycarboxy compounds of formula (IV) are selected from the representative and illustrative group consisting of: pyromellitic acid dianhydride, 3,6-diphenylpyromellitic dianhydride, 3,3',4,4'-benzenophenonetetracarboxylic acid dianhydride, 2,3,3',4,4'-benzenophenonetetra carboxylic acid dianhydride, 2,2',3,3'-benzenophenonetetra carboxylic acid dianhydride, 3,3',4,4'-diphenyl-tetracarboxylic acid dianhydride, bis-(2,3-dicarboxyphenyl)-methane dianhydride, (bis(3,4-dicarboxyphenyl)-sulphone dianhydride, N,N-(3,4-dicarboxyphenyl)-N-methylamine dianhydride, bis(3,4-dicarboxyphenyl)-diethyilsilane dianhydride, 2,3,6,7- and 1,2,5,6-naphthalene-tetracarboxylic acid dianhydride, 2,6-dichloronaphthalene-1,4,5,8-tetracarboxylic acid dianhydride, thiophene-2,3,4,5-tetracarboxylic acid dianhydride, pyrazine-2,3,5,6-tetracarboxylic acid dianhydride and pyridine-2,3,5,6-tetracarboxylic acid dianhydride as well as esters or anhydrides of the above listed compounds.

When $Y$ of formula (IV) represents a substituted phenylene group, then the substituents are selected from the group consisting of nitro, chlorine, fluorine, alky having 1 to 2 carbon atoms or alkoxy having 1 to 2 carbon and preferably selected from the group consisting of 2-, 3- or 4-nitrophenoxymethyl, 2,4- or 3,5-dinitro-
phenoxyc group or 3,5-dichlorophenoxyc group or the pentachlorophenoxyc, 2-methylphenoxyc or 2-methoxyphenoxyc group. When Y represents an alkoy group, then the group is selected from the group consisting of methoxy, ethoxy, N-propoxy, propoxy, n-butoxy, tert-butoxy, hexyloxy, octyloxy, decyloxy, dodecyloxy, tetradecyloxy and octadecyloxy group and preferably selected from the group consisting of methoxy, ethoxy, N-propoxy, isopropoxy, n-butoxy, and tert-butoxy.

Diamines of formula (V) are selected from the representative and illustrative groups consisting of: aliphatic diamines such as, but not limited to, di-methylene diamine, tri-methylene diamine, tetra-methylene diamine, hexa-methylene diamine, hepta-methylene diamine, octa-methylene diamine and deca-methylene diamine, 2,2'-dimethylpropylene diamine, 2,5-dimethyl hexamethylene diamine, 4,4'-dimethylthiopropylene diamine, 3-methoxyhexamethylene diamine, 1,2-bis-(3-amino propoxy) ethane, N,N,N'-dimethylethylene diamine and N,N'-dimethyl-1,6-dimino hexane as well as the diamines of the formula H2N(CH2)30(CH2)nNH2 and N2N(CH2)5S(CH2)5S(CH2)5NH2 and similar polyethylene oxide or polydithioethylene diamines; carbocyclic aliphatic diamines such as, but not limited to, 1,2-diaminocyclohexane, 1,3-diaminocyclohexane, 1,4-diaminocyclohexane, 1,3-diamine cyclopentane, 1,4-diaminocyclopentane, and other similar cyclic aliphatic diamines; carbocyclic aromatic diamines such as, but not limited to, o-phenylenediamine, m-phenylenediamine, p-phenylenediamine, dianinotoluuenes, 2,4-diaminotoluene, 1,4-diamino-2-methoxybenzene, 2,5-diaminoxylen, 1,3-diamino-4-chlorobenzene, 4,4'-diaminodiphenyl methane, 4,4'-diaminodiphenyl ether, 4,4'-diaminodiphenyl triethenier, 4,4'-diaminophenyl sulphone, 2,2'-diaminobenzophenone, 4,4'-diaminodiphenylurea, 1,8-diaminonaphthalene, 1,5-diamononaphthalene, benzidine, 3,3'-dimethoxybenzidine, 2,2'-bis-(trifluoromethyl) benzidine, 2,2'-dimethyl benzidine, 2,2'-dichlorobenzidine, and other similarly substituted benzidines, 1,4-bis-(2-methyl-4 TMaminopentyl)-benzene and 1,4-bis-(aminomethyl)-benzene; heterocyclic diamines such as, but not limited to, 2,6-dianino pyridine, 2,4-dianinopyrimidine, 2,4-diamo-s-triazine and other similar heterocyclic diamines.

The polycondensation reaction of the compounds of the formula (I) with one or more compounds of the formula (IV) and, optionally, one or more compounds of the formula (V) is carried out in a manner which is in itself known, appropriately at temperatures of about -50 °C to +300 °C. The reaction can be carried out in the melt or, preferably, in an inert organic solvent or a solvent mixture. Temperatures of -20°C to +50°C are preferred for the polycondensation reaction in solution.

Examples of suitable organic solvents are: chlorinated aromatic hydrocarbons, such as chlorobenzene and dichlorobenzenes, chlorinated aliphatic hydrocarbons, such as methylene chloride, chloroform, tetra chloroethane and tetrachloroethene, aliphatic and cycloaliphatic ketones, such as acetone, methyl ethyl ketone, cyclopentanone and cyclohexanone, cyclic ethers, such as tetrahydrofuran, tetrahydropryan and dioxane, cyclic amines, such as N-methyl-2-pyrrolidone, N-propyl-2-pyrrolidone and N-methyl-a-caprolactam, N,N-dialkylides of aliphatic monocarbonylic acids with 1-3 carbon atoms in the acid part, such as N,N-dimethylformamide, N,N-dimethylacetamide, N,N-dimethylacetamide and N,N-dimethylthioacetamide, hexamethylene phosphoric acid triamide (hexametapal), N,N,N,N'-tetramethyleurea, tetrahydrophthene dioxide (sulphane) and dialkylsulphoxides, such as dimethylsulphoxide and diethylsulphoxide.

Preferred solvents are N,N-dialkylamides of aliphatic monocarboxylic acids with 1-3 carbon atoms in the acid part, especially N,N-dimethylacetamide, as well as cyclic amides, such as N-methyl-2-pyrrolidone.

The hydrochloric acid obtained during the condensation reaction with compounds of the formula (IV) in which Y represents chlorine can be removed by neutralization with basic substances, such as calcium hydroxide or triethylamine, or by reaction with an epoxide compound, such as ethylene oxide or propylene oxide, and by washing out with suitable solvents.

The condensation reactions are appropriately carried out with the exclusion of moisture, for example in an inert gas atmosphere, such as nitrogen.

As already mentioned initially, the reaction of the di- or tetra-carboxylic acid derivatives of the formula (IV) with the diamines of the formula (I) and/or (V) can also be carried out stepwise in order to manufacture polymers which, at least in part, have a block-type distribution of repeat units of the formula (VI) and repeat units of formula (VII). It is also possible to link polylamides, polylamide-amide-acids or polylamide-acids which have 1 to 100 mole percent of repeat units of formula (VI) and 99 to 0 mole percent of repeat units of formula (V) and have been manufactured separately, with one another and so-called block copolymers are formed by this means. In all of these cases, the reaction is carried out in a manner which is in itself known using a slight excess of one or the other reactant in order to obtain precopolymers which have end groups suitable for the further reaction, for example amino end groups, acid chloride groups and/or anhydride groups.

The optional cyclization of the polylamide-acids or polylamide-amide-acids has been carried out in a manner which is in itself known by chemical means or by means of heat.

The chemical cyclization is appropriately carried out by treatment with a dehydrating agent on its own or as a mixture with a tertiary amine. Reagents which can be used are, for example, acetic anhydride, propionic anhydride and diethylene carbodiimide or mixtures of acetic anhydride and triethylamine.

The cyclization by means of heat is carried out by heating to temperatures of about 50°C to 300°C and preferably of about 150°C to 250°C and optionally with the addition of an inert organic solvent.

The polylamides, polylamide-amide-acids and polylamide-acids according to the invention, as well as the corresponding cyclized derivatives, are suitable for the manufacture of shaped articles of very diverse types, such as fibers, films, sheets, coating compositions, foams, laminating resins, composite materials, molding powders, pressed articles and the like, in a manner which is in itself known, if desired with the use of customary additives, such as pigments, fillers and the like. The polymers according to the invention can also be processed easily from the melt and are distinguished by good mechanical, electrical and thermal properties as well as, in general, by good solubility in organic solvents, such as N,N-dimethyl acetamide, N,N-dimethylformamide and N-methyl-2-pyrrolidone.
BEST MODE FOR CARRYING OUT THE INVENTION

Applicants have found that the polyamides, polyamide-amide-acids and polyamide-acids, as well as the corresponding cyclized derivatives of polyamide-amide-acids, polyamide-imides, and polyamide-acids, polyimides, derived from diamines of formula (I) or copolymer having at least 1 mole percent polymer derived from diamines of formula (I) represent a new and novel class of soluble high performance polymers. When rigid, aromatic polycarboxy compounds of formula (IV) are used in the manufacture of polyamides, polyamide-imides and polyimides, these preferred polymers of the present invention represent a novel class of soluble and rigid-rod high performance polymers which maintain standard polyamides, polyamide-imides and polyimides properties such as outstanding modulus, tensile and compression strength, energy absorption, coefficient of expansion and electrical properties. The fact that these polymers have good solubility and can be manufactured with rigid structures makes them ideally suited for use in the production of antiballistics, reinforced molecular composites, microelectronic coatings and membranes because the polymers can be solution cast or deposited.

In preparing the homopolyamide, a near one to one mixture of a diamines of formula (I) such as 1,4-bis[4-aminophenyl]-2,3,5-triphenylbenzene (Ia) and a polycarboxylated compound of formula (IV) such as dichloro-teraphthalic acid (XII) are allowed to react under the above stated reaction conditions to from a polyamide of the following repeat formula (XIII)

\[
\begin{align*}
\text{(XIII)}
\end{align*}
\]

In preparing the homopolyamide-amide-acids, a near one to one mixture of a diamine of formula (I) such as 1,4-bis[4-aminophenyl]-2,3,5-triphenylbenzene (Ia) and a polycarboxylated compound of formula (IV) such as trimellitic acid (XV)

\[
\begin{align*}
\text{(XV)}
\end{align*}
\]

are allowed to react under the above stated reaction conditions to from a polyamide-amide-acid of the following repeat formula (XVI)

\[
\begin{align*}
\text{(XVI)}
\end{align*}
\]

The polymers of repeat unit of formula (XVI) can be cyclized to polyamide-imides by standard techniques well known in the literature.

In preparing the homopolyamide-acid, a near one to one mixture of a diamine of formula (I) such as 1,4-bis[4-aminophenyl]-2,3,5-triphenylbenzene (Ia) and a polycarboxylated compound of formula (XI) such as pyromellitic acid (XVII)

\[
\begin{align*}
\text{(XVII)}
\end{align*}
\]

are allowed to react under the above stated reaction conditions to from a polyamide-acid of the following repeat formula (XVIII)

\[
\begin{align*}
\text{(XVIII)}
\end{align*}
\]

The polymers of repeat unit of formula (XVIII) can be cyclized to polyamide-imides by standard techniques well known in the literature.

In preparing the copolymer consisting of 1 to 100 mole percent of repeat units of formula (IV) and 99 to 0 mole percent of repeat units of formula (V) it is understood that given molar amount of a combination of one or more diamines of formula (I) and one or more diamines of formula (XI) is mixed with a nearly equal molar amount of one or more polycarboxylated compound of formula (XI). This type of reaction mixture can be used to make copolyamides, copolyamide-amide-acids, copolyamide-imides, the cyclized version of cyclizable copolyamide-amide-acids, copolyamide-acids, and copolymides, the cyclized version of cyclizable copolyamide-acids. The cyclization reaction can be done thermally or chemically with the preferred method utilizing the chemical additive isoquinoline in amount between about 0.1% and 5% by weight of solvent.

The diamines of the formula (I) in which both amino groups (NH₂) are in the same position of the outer benzene rings, and in particular the diamines of the formula (I) in which both amino groups are in the para-position of the outer benzene rings is preferred.

Co-polyamides, co-polyamide-amide-acids and co-polyamide-acids having 10–90 mole percent of repeat units of formula (VII) and 90–10 mole percent of repeat units of formula (VII) are particularly preferred. These copolymers are preferred because they allow the expression of properties of the diamines of formula (I) and minimize the increased cost of using diamines of formula (I) in copolymers of the present invention.
In all the above polymers indicated by 1 to 100 mole percent of repeat units of formula (IV) and 99 to 0 mole percent of repeat units of formula (V), the end groups of the polymers can be either an acid, an amine, anhydride or a combination of any of these groups. The reaction can also be carried out in the presence of a chain limiting reagent such as a mono functional carboxylic acid or a mono functional amine containing organic reagent. These chain limiting reagents actually stop polymerization by introducing a non-reactive group at the end of a polymer chain. In the case of carboxylic acid chain limiting reagents, the reagent can be selected from the representative and illustrative group consisting of benzoic acid, naphthoic acid, or alkyl substituted benzoic acids. In the case of organic amine chain limiting reagents, the reagent can be selected from the group consisting of aniline, alkyl substituted anilines, or naphthyl amines.

The invention will be better understood by reference to the following examples which are included for purposes of illustration and not limitation.

PREPARATION OF DINITRO INTERMEDIATE OF FORMULA (I)

EXAMPLE 1

This example illustrates the preparation of 1,4-bis(4-nitrophenyl)-2,3,5,6-tetraphenylbenzene, a dinitro intermediate of formula (I) where Ar1, Ar2, and Ar3 are phenyl and R1 is hydrogen.

1,4-bis(4-nitrophenyl)-2,3,5,6-tetraphenylbenzene was prepared according to the procedure described in L. F., "Organic Experiments", 2nd Ed., 297 (1968). 19.3 grams (0.0362 moles) of 4-nitrophenyl)-2,3,5-terephthoyl chloride (TPC) was added to the solution of benzophenone in 130 mL of hydrochloric acid. When the temperature raised to 80°C, the reaction mixture was poured into 500 mL of hexane and filtered. The reaction yielded 22.9 grams of 1,4-bis(4-nitrophenyl)-2,3,5,6-tetraphenylbenzene, a 66% yield.

EXAMPLE 2

This example illustrates the preparation of 1,4-bis(4-nitrophenyl)-2,3,5,6-tetraphenylbenzene, a dinitro intermediate of formula (I), where Ar1, Ar2, and Ar3 are phenyl and R1 is hydrogen.

1,4-bis(4-nitrophenyl)-2,3,5,6-tetraphenylbenzene was prepared according to the procedure described in Example 1 with 1,4-bis(4-nitrophenyl)-2,3,5,6-tetraphenylbenzene being used at the same molar amount as 1,4-bis(4-nitrophenyl)-2,3,5-triphenylbenzene in Example 3, but the yield was only 45%. The crude product was recrystallized from pyridine.

The properties of 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene were: mp=453°C (DSC); IR (KBr) 3380 and 3450 cm⁻¹ (NH₂); 1H-NMR (CDC1₃) δ=3.3 ppm (s, 4H, NH₂); 13C-NMR (CDCl₃) δ= 150.2 ppm.

PREPARATION OF A HOMOPOLYAMIDE

EXAMPLE 5

This example illustrates the preparation of a polyamide consisting of 100 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5-triphenylbenzene, and the polycarboxylic acid compound of formula (IV) is terephthyl chloride.

0.4 grams (0.0008186 mole) of 1,4-bis(4-aminophenyl)-2,3,5-triphenylbenzene, synthesized according to example 2, was dissolved in 2.9 grams of NMP, cooled in an ice bath to below 5°C and kept under an inert atmosphere such as nitrogen during the course of the polymerization. 0.1662 grams (0.0008186 mole) of terephthyl chloride (TPC) was added to the solution and stirred for 30 minutes. This represented a 6% concentrated solution of starting material in solvent. The ice bath was removed shortly after TPC addition. The mixture became viscous and turned brown-yellow transparent color. After approximately 15 hours, the mixture was poured into 500 mL of methanol to affect coagulation. The polymer was filtered, dried and the yield was 97%.

The properties of 1,4-bis(4-aminophenyl)-2,3,5-triphenylbenzene were: [n]D₂₅ = 2.87 dL/g in NMP at 30°C; Tₘ = 50°C in nitrogen; Tₘ = 493°C in air; soluble in NMP at room temperature; tough transparent film.

It should be appreciated that any other polycarboxylic compound of formula (IV) can be substituted into the above synthesis to yield an analogous polymer.

EXAMPLE 6

This example illustrates the preparation of a polyamide consisting of 100 mole percent of repeat unit of
This polyamide was prepared according to Example 5 except the starting material concentration was 14%. The polymer solution was transparent.

Polymer properties: \[\eta_{inh} = 2.99 \text{ dL/g in NMP at } 30^\circ C; T_{-5\%} = 508^\circ C \text{ in nitrogen}; T_{-5\%} = 493^\circ C \text{ in air; soluble in NMP at room temperature; tough transparent film.}\]

EXAMPLE 7

This example illustrates the preparation of a polyamide consisting of 100 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene, and the polycarboxy compound of formula (IV) is terephthalamide. This polyamide was prepared according to Example 5 except isophthaloyl chloride was used instead of its isomeric analog terephthalamide and the starting material concentration was 14%. The polymer solution was transparent.

Polymer properties: \[\eta_{inh} = 0.88 \text{ dL/g in NMP at } 30^\circ C; T_{-5\%} = 509^\circ C \text{ in nitrogen}; T_{-5\%} = 520^\circ C \text{ in air; soluble in NMP at room temperature; tough transparent film.}\]

EXAMPLE 8

This example illustrates the preparation of a polyamide consisting of 100 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene, and the polycarboxy compound of formula (IV) is terephthalamide. This polyamide was prepared according to Example 5 except isophthaloyl chloride was used instead of 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene and the starting material concentration was 8%. The polymer precipitated and the yield was 78%.

Polymer properties: \[\eta_{inh} = 0.46 \text{ dL/g in sulfuric acid at } 30^\circ C; T_{-5\%} = 516^\circ C \text{ in nitrogen}; T_{-5\%} = 524^\circ C \text{ in air.}\]

EXAMPLE 9

This example illustrates the preparation of a polyamide consisting of 100 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene, and the polycarboxy compound of formula (IV) is terephthalamide. This polyamide was prepared according to Example 5 except 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene was used instead of 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenyldiamine and the starting material concentration was 8%. The polymer precipitated and the yield was 78%.

Polymer properties: \[\eta_{inh} = 0.46 \text{ dL/g in sulfuric acid at } 30^\circ C; T_{-5\%} = 516^\circ C \text{ in nitrogen}; T_{-5\%} = 524^\circ C \text{ in air.}\]

EXAMPLE 10

This example illustrates the preparation of a polyamide consisting of 100 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene, and the polycarboxy compound of formula (IV) is isophthaloyl chloride.

This polyamide was prepared according to Example 5 except 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene was used instead of 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene, isophthaloyl chloride was used instead of terephthalamide and the starting material concentration was 14%. The polymer precipitated and the yield was 91%.

Polymer properties: \[\eta_{inh} = 0.49 \text{ dL/g in sulfuric acid at } 30^\circ C; T_{-5\%} = 514^\circ C \text{ in nitrogen}; T_{-5\%} = 516^\circ C \text{ in air; a tough film was cast from sulfuric acid.}\]

EXAMPLE 11

This example illustrates the preparation of a polyamide consisting of 100 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene, and the polycarboxy compound of formula (IV) is isophthaloyl chloride.

This polyamide was prepared according to Example 5 except 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene was used instead of 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene, isophthaloyl chloride was used instead of terephthalamide, a two molar amount of lithium chloride based on the moles of diamine was added to the reaction after the monomer addition and the starting material concentration was 15%. The polymer solution was cloudy and the yield was 81%.

Polymer properties: \[\eta_{inh} = 0.46 \text{ dL/g in sulfuric acid at } 30^\circ C; T_{-5\%} = 514^\circ C \text{ in nitrogen}; T_{-5\%} = 516^\circ C \text{ in air; a tough film was cast from sulfuric acid.}\]

EXAMPLE 12

This example illustrates the preparation of a polyamide consisting of 100 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5,6-triphenylbenzene, and the polycarboxy compound of formula (IV) is pyromellitic dihydride (PMDA). 0.35 grams (0.0007163 mole) of 1,4-bis(4-aminophenyl)-2,3,5,6-triphenylbenzene, synthesized according to example 2, was dissolved in 9.0 grams of m-cresol under nitrogen. 0.1562 grams of pyromellitic dihydride (PMDA) (0.0007161 mole) was added to the solution along with 3 ml of toluene and 3 drops of isoquinoline. (Isoquinoline is an optional ingredient which promotes the cyclization reaction.) This solution represented a 5% concentration of starting material in solvent. The reaction was heated to 170° C. The toluene is added to azeotrope of the water formed during the reaction. When the distillation of toluene was over, 3 ml of fresh toluene was added. This operation was repeated three times to insure complete removal of the water of reaction. The temperature was then maintained at 190° C for 3 hours. After a couple of hours, an aliquot of the polymer mixture was cast into a film. After cooling, the reaction mixture was poured in to 500 mL of methanol to affect coagulation. The product...
obtained was washed with methanol and dried. The polymer was filtered and the yield was 94%.

Polymer properties: \([\eta]_{\text{inh}} = 3.16 \text{ dL/g in sulfuric acid at } 30^\circ\text{C}; T_{-5\%} = 583^\circ\text{C in nitrogen}; T_{-5\%} = 546^\circ\text{C.}\) in air; insoluble in CHCl₃ and NMP at room temperature.

It should be appreciated that any other polycarboxy compound of formula (IV) can be substituted into the above synthesis to yield an analogous polymer.

**EXAMPLE 13**

This example illustrates the preparation of a polyimide consisting of 100 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5-triphenylbenzene, and the polycarboxy compound of formula (IV) is pyromellitic dianhydride.

The polyimide was prepared according to Example 12 except the concentration of starting material in solvent was 15%. The solution was gelatinous and not homogeneous as in Example 12.

Polymer properties: \([\eta]_{\text{inh}} = 3.16 \text{ dL/g in sulfuric acid at } 30^\circ\text{C}; T_{-5\%} = 583^\circ\text{C in nitrogen}; T_{-5\%} = 546^\circ\text{C.}\) in air; insoluble in CHCl₃ and NMP at room temperature.

It should be appreciated that any other polycarboxy compound of formula (IV) can be substituted into the above synthesis to yield an analogous polymer.

**EXAMPLE 14**

This example illustrates the preparation of a polyimide consisting of 100 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5-triphenylbenzene, and the polycarboxy compound of formula (IV) is 3,3',4,4'-diphenyl-tetracarboxylic acid dianhydride.

The polyimide was prepared according to Example 12 except 3,3',4,4'-diphenyl-tetracarboxylic acid dianhydride was used instead of PMDA and the concentration of starting material in solvent was 7%. The yield of product was 92%.

Polymer properties: \([\eta]_{\text{inh}} = 1.89 \text{ dL/g in NMP at } 30^\circ\text{C}; T_{-5\%} = 601^\circ\text{C in nitrogen}; T_{-5\%} = 564^\circ\text{C.}\) in air; soluble in CHCl₃ and NMP at room temperature; a tough film was cast from NMP.

**EXAMPLE 15**

This example illustrates the preparation of a polyimide consisting of 100 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5-triphenylbenzene, and the polycarboxy compound of formula (IV) is bis(3,4-dicarboxyphenyl) ether dianhydride.

The polyimide was prepared according to Example 12 except bis(3,4-dicarboxyphenyl) ether dianhydride was used instead of PMDA and the concentration of starting material in solvent was 7%. The yield of product was 94%.

Polymer properties: \([\eta]_{\text{inh}} = 0.83 \text{ dL/g in sulfuric acid at } 30^\circ\text{C}; T_{-5\%} = 576^\circ\text{C in nitrogen}; T_{-5\%} = 539^\circ\text{C.}\) in air; soluble in CHCl₃ and NMP at room temperature; a tough film was cast from NMP.

**EXAMPLE 16**

This example illustrates the preparation of a polyimide consisting of 100 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5-triphenylbenzene, and the polycarboxy compound of formula (IV) is bis(3,4-dicarboxyphenyl)-sulphone dianhydride.

The polyimide was prepared according to Example 12 except bis(3,4-dicarboxyphenyl)-sulphone dianhydride was used instead of PMDA and the concentration of starting material in solvent was 7%. The yield of product was 90%.

Polymer properties: \([\eta]_{\text{inh}} = 0.66 \text{ dL/g in sulfuric acid at } 30^\circ\text{C}; T_{-5\%} = 504^\circ\text{C in nitrogen}; T_{-5\%} = 524^\circ\text{C.}\) in air; insoluble in CHCl₃ and NMP at room temperature; a tough film was cast from NMP.

**EXAMPLE 17**

This example illustrates the preparation of a polyimide consisting of 100 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene and the polycarboxy compound of formula (IV) is pyromellitic dianhydride.

The polyimide was prepared according to Example 12 except 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene was used instead of 1,4-bis(4-aminophenyl)-2,3,5-triphenylbenzene and the concentration of starting material in solvent was 7%. The yield was 97%.

Polymer properties: \([\eta]_{\text{inh}} = 1.14 \text{ dL/g in sulfuric acid at } 30^\circ\text{C}; T_{-5\%} = 572^\circ\text{C in nitrogen}; T_{-5\%} = 473^\circ\text{C.}\) in air; insoluble in CHCl₃ and NMP at room temperature.

It should be appreciated that any other polycarboxy compound of formula (IV) can be substituted into the above synthesis to yield an analogous polymer.

**EXAMPLE 18**

This example illustrates the preparation of a polyimide consisting of 100 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene, and the polycarboxy compound of formula (IV) is 3,3',4,4'-diphenyl-tetracarboxylic acid dianhydride.

The polyimide was prepared according to Example 12 except 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene was used instead of 1,4-bis(4-aminophenyl)-2,3,5-triphenylbenzene and the polycarboxy compound of formula (IV) is bis(3,4-dicarboxyphenyl)-tetracarboxylic acid dianhydride.

The polyimide was prepared according to Example 12 except 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene was used instead of 1,4-bis(4-aminophenyl)-2,3,5-triphenylbenzene, and the polycarboxy compound of formula (IV) is bis(3,4-dicarboxyphenyl) ether dianhydride.

**EXAMPLE 19**

This example illustrates the preparation of a polyimide consisting of 100 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene, and the polycarboxy compound of formula (IV) is bis(3,4-dicarboxyphenyl) ether dianhydride.
The polyimide was prepared according to Example 12 except 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene was used instead of 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene, bis(3,4-dicarboxyphenyl) ether dihydrid and the polycarboxy compound of formula (IV) was bis(3,4-dicarboxyphenyl)sulphone dihydrid.

The polycarboxy compound of formula (IV) is terephthoyl chloride. of formula (IV), pyromellitic dianhydride and bis-(3,4-dicarboxyphenyl)-2,3,5-triphenylbenzene, bis(3,4-dicarboxyphenyl)sulphone dihydrid was used instead of PMDA and the concentration of starting material in solvent was 9%. The yield of product was 95%.

Polymer properties: \[ \eta_{inh} = 0.61 \text{ Dl/g in sulfuric acid at } 30^\circ C; \ T_m = 571^\circ C \text{ in nitrogen; } T_m = 523^\circ C \text{ in air; partially soluble in NMP at room temperature; a tough film was cast from NMP.} \]

EXAMPLE 20

This example illustrates the preparation of a polycarboxy compound of formula (IV) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene, and the polycarboxy compound of formula (IV) is bis-(3,4-dicarboxyphenyl)sulphone dihydrid.

The polycarboxy compound of formula (IV) was dissolved in an ice bath and kept under an inert atmosphere such as nitrogen during the course of the polymerization. The yield of product was 94%.

Polymer properties: \[ \eta_{inh} = 0.67 \text{ Dl/g in sulfuric acid at } 30^\circ C; \ T_m = 527^\circ C \text{ in nitrogen; } T_m = 535^\circ C \text{ in air; soluble NMP at room temperature; a tough film was cast from NMP.} \]

PREPARATION OF A COPOLYMERS

EXAMPLE 21

This example illustrates the preparation of a copolyamide consisting of 50 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene, and the polycarboxy compound of formula (IV) is terephthohyl chloride and 50 mole percent of repeat unit of formula (VII) where the diamine of formula (V) is 4,4'-diaminodiphenyl ether and the polycarboxy compound of formula (IV) is terephthoyl chloride. 0.4 grams (0.0008186 mole) of 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene, synthesized according to example 2, and 0.1637 grams (0.0007163 moles) of terephthoyl chloride (TPC) and 0.1662 grams (0.0008186 moles) of isophthoyl chloride (IPC) is added to the solution along with 4 ml of NMP. The ice bath is removed shortly after TPC and IPC addition. The mixture becomes viscous and turned brown-yellow transparent color. After approximately 10 hours, the mixture is diluted in approximately 40 ml of NMP and poured into 1 liter of methanol to affect coagulation. The polymer is filtered and the yield was 97%.

It should be appreciated that any other diamine of formula (V) or any other polycarboxy compound of formula (IV) can be substituted into the above synthesis to yield an analogous polymer.

EXAMPLE 22

This example illustrates the preparation of a mixed copolyamide consisting of 50 mole percent of repeat unit of formula (VI) where the diamine of formula (I) represents the diamine prepared in example 2, 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene, and two polycarboxy compound of formula (IV), terephthohyl chloride and isophthohyl chloride, and 50 mole percent of repeat unit of formula (VII) where the diamine of formula (V) is 4,4'-diaminodiphenyl ether and two polycarboxy compound of formula (IV), terephthohyl chloride and isophthohyl chloride. 0.4 grams (0.0008186 mole) of 1,4-bis(4-aminophenyl)-2,3,5,6-tetraphenylbenzene, synthesized according to example 2, and 0.164 grams (0.0008186 moles) of 4,4'-diaminodiphenyl ether is dissolved in 12 ml NMP, cooled in an ice bath and kept under an inert atmosphere such as nitrogen during the course of the polymerization. The yield of product was 94%. The polymer is soluble in NMP and pourable into 1 liter of methanol to affect coagulation. The polymer is filtered and the yield was 97%.

It should be appreciated that any other diamine of formula (V) or any other polycarboxy compound of formula (IV) can be substituted into the above synthesis to yield an analogous polymer.

What is claimed is:
1. A polyamide, a polyamide-amide-acid or a polyamide-acid which comprises 1 to 100 mole percent of structural elements of the formula (VI)

\[
\begin{array}{c}
\text{O} \\
\text{(COOH)}_n \\
\text{Z} \\
\text{(HOOCC)m} \\
\text{C} \\
\text{Z} \\
\text{NH--Q--NH--} \\
\text{Z} \\
\text{C} \\
\text{(COOH)_m} \\
\end{array}
\]

and from 99 to 0 mole percent of a repeat unit of formula (VII)

\[
\begin{array}{c}
\text{O} \\
\text{(COOH)_n} \\
\text{Z} \\
\text{(HOOCC)m} \\
\text{C} \\
\text{Z} \\
\text{NH--Q--NH--} \\
\text{Z} \\
\end{array}
\]

where both imido (CONH) groups are independently in either the meta or para position with respect to the covalent bond to the center polysubstituted benzene ring, where \(A_1, A_2, A_3\) and \(A_4\) are separately and independently aryl groups selected from the group consisting of an aliphatic radical, a carbocyclic aromatic radical, a heterocyclic aromatic radical, or a heterocyclic radical.

2. A polyamide according to claim 1, wherein said polyamide comprises 100 mole percent of repeat unit formula (VI) where \(n\) and \(m\) are equal to the numeric value of either 0 or 1, \(Z'\) and \(Z''\) are separately and independently one or more divalent organic radical derived from a diamine of formula (V).

\[
\begin{array}{c}
\text{H}_2\text{N--Q--NH}_2 \\
\end{array}
\]

3. A polyamide according to claim 2, wherein said \(Q\) is one or more divalent aromatic radical derived from a diamine of formula (V)

\[
\begin{array}{c}
\text{H}_2\text{N--Q--NH}_2 \\
\end{array}
\]

4. A polyamide-amide-acid according to claim 1, wherein said polyamide-amide-acid comprises 100 mole percent of repeat unit formula (VI) where either \(m\) or \(n\) is equal to the numeric value of 1, and where \(Z'\) and \(Z''\) are separately and independently one or more trivalent radical derived from a polycarboxy compound of formula (IV)

\[
\begin{array}{c}
\text{Z(}\text{COY)k} \\
\end{array}
\]

where \(Z\) is an organic radical selected from the group consisting of an aliphatic radical, a cyclo-aliphatic radical, a carbocyclic-aromatic radical, or a heterocyclic radical.

5. A polyamide-amide-acid according to claim 4, wherein said polyamide-amide-acid is cyclized to the corresponding polyamide-imide.

6. A polyamide-amide-acid according to claim 6, wherein said polyamide-amide-acid is one or more divalent aromatic radical derived from a diamine of formula (V)

\[
\begin{array}{c}
\text{H}_2\text{N--Q--NH}_2 \\
\end{array}
\]

where \(Q\) is a divalent organic radical selected from the representative and illustrative group consisting of an aliphatic radical having at least 2 carbon atoms, a carbocyclic-aromatic radical, a carbocyclic-aromatic radical, or a heterocyclic radical.

7. A polyamide-amide-acid according to claim 7, wherein said polyamide-amide-acid is one or more divalent aromatic radical derived from a diamine of formula (V)

\[
\begin{array}{c}
\text{Z(}\text{COY)k} \\
\end{array}
\]

where \(Z\) is an organic radical selected from the group consisting of an aliphatic radical, a cyclo-aliphatic radical, a carbocyclic-aromatic radical, or a heterocyclic radical, where \(Y\) is a halogen atom, a hydroxy group, an unsubstituted or substituted phenoxy group or an alkoxy group preferably having from about 1 to 18 carbon atoms, and particularly preferred having from about 1 to 12 carbon atoms, where \(k\) is a whole number having a numeric value of 3 and where each \(COY\) is bonded to a different atom of \(Z\) and when \(k\) is equal to 3 and \(Z\) is a cyclic organic radical, then two of the \(COY\) groups are in an ortho orientation relative to each other.

8. A polyamide-amide-acid according to claim 7, wherein said polyamide-amide-acid is cyclized to the corresponding polyamide-imide.
about 1 to 12 carbon atoms, where \( k \) is a whole number having a numeric value of 4 and where each COY is bonded to a different atom of Z and when \( k \) is equal to 4 and \( Z \) is a cyclic organic radical, then each pair of COY groups are arranged in an ortho orientation relative to one another.

9. A polyamide-acid according to claim 8, wherein said polyamide-acid is cyclized to the corresponding polyimide.

10. A polyamide-acid according to claim 8, wherein said Q is one or more divalent aromatic radical derived from a diamine of formula (V)

\[
\text{H}_2\text{N}--\text{Q}--\text{NH}_2
\]

(V)

where \( Q \) is a divalent organic radical selected from the representative and illustrative group consisting of an aliphatic radical having at least 2 carbon atoms, a carbocyclic aliphatic radical, a carbocyclic aromatic radical, or a heterocyclic radical.

11. A polyamide-acid according to claim 10, wherein said polyamide-acid is cyclized to the corresponding polyimide.

12. A polyamide, a polyamide-amide-acid or a polyamide-acid according to claim 1, wherein said polyamide, polyamide-amide-acid or polyamide-acid consisting of 10 to 90 mole percent of repeat units of formula (VI) and 10 to 90 mole percent of repeat units of formula (VII).

13. A polyamide, polyamide-amide-acid, or polyamide-acid according to claim 12, wherein either \( n \) or \( m \) is equal to the numeric value of 1, and where \( Z' \) and \( Z'' \) are separately and independently one or more divalent radical derived from a polycarboxy compound of formula (IV)

\[
\text{H}_2\text{N}--\text{Q}--\text{NH}_3
\]

(V)

where \( Q \) is a divalent organic radical selected from the representative and illustrative group consisting of an aliphatic radical having at least 2 carbon atoms, a carbocyclic aliphatic radical, a carbocyclic aromatic radical, or a heterocyclic radical.

14. A polyamide-amide-acid or polyamide-acid according to claim 13, wherein said polyamide-amide-acid or polyamide-acid is cyclized to the corresponding polyamide-imide or polyimide.

15. The polyamide, a polyamide-acid polymer of claim 1 wherein the polymer is soluble, melt processable and has a heat stability in air as measured by \( T_{\text{50\%}} \) in air of 493°C. and higher and exists as a rod-shaped polymer. Darms is devoid of a teaching to a rod-shaped polymer.