

TELECHELIC N-ALKYLATED POLYAMIDE POLYMERS AND COPOLYMERSTECHNICAL FIELD

[0001] The present subject matter relates to telechelic polyamides that can be reacted into other polymeric materials to impart desirable properties.

BACKGROUND

[0002] Many polyamides, e.g. the various nylon polymers, are solids at temperatures of below about 80°C and thus would be difficult to homogenously react into other polymeric materials. N-alkylating the nitrogen atom of the polyamide or the nitrogen bearing precursor of the polyamide eliminates the hydrogen bonding making the polyamide of this disclosure lower melting and more soluble.

SUMMARY

[0003] The present subject matter relates to low molecular weight polyamide oligomers and telechelic polyamides (including copolymers) comprising N-alkylated amide groups in the backbone structure. These polymers are useful as soft segments in the preparation of thermoplastic, thermoset, or elastomer resins and water-borne dispersions of those resins. The unique feature of these polyamide polymers is their ability to be processed as liquids at temperatures from 20 to 80°C (such as 20 to 50°C), which makes them suitable for further reaction and polymerization forming various thermoplastic or thermoplastic elastomer compositions.

[0004] Thus, provided is a telechelic polyamide comprising: repeat units derived from polymerizing monomers connected by linkages between the repeat units and functional end groups selected from carboxyl or primary or secondary amine; a polyamide segment comprising at least two amide linkages characterized as being derived from reacting an amine with a carboxyl group, and the polyamide segment comprises repeat units derived from polymerizing dicarboxylic acid monomeric units with monomeric units of at least one other monomer type, wherein at least 50 percent of the dicarboxylic acid monomeric units each contain 34 to 36 carbon atoms, and wherein the at least one other monomer type comprises diamine monomeric units; wherein at least 10% of the total number of the heteroatom containing linkages connecting hydrocarbon type linkages are characterized as being amide; wherein less than 25% of the amide linkages are characterized as being tertiary amide linkages; and wherein at least

20 mole percent of the telechelic polyamide has exactly two functional end groups of the same functional type comprising amino end groups or carboxylic end groups.

[0005] The following embodiments of the present subject matter are contemplated:

[0006] 1. A telechelic polyamide comprising: (a) repeat units derived from polymerizing monomers connected by linkages between the repeat units and functional end groups selected from carboxyl or primary or secondary amine; (b) a polyamide segment comprising at least two amide linkages characterized as being derived from reacting an amine with a carboxyl group, and the polyamide segment comprises repeat units derived from polymerizing dicarboxylic acid monomeric units with monomeric units of at least one other monomer type, wherein at least 50 percent of the dicarboxylic acid monomeric units each contain 34 to 36 carbon atoms, and wherein the at least one other monomer type comprises diamine monomeric units; (c) wherein at least 10% of the total number of the heteroatom containing linkages connecting hydrocarbon type linkages are characterized as being amide; and (d) wherein less than 25% of the amide linkages are characterized as being tertiary amide linkages; wherein at least 20 mole percent of the telechelic polyamide has exactly two functional end groups of the same functional type comprising amino end groups or carboxylic end groups.

[0007] 2. The telechelic polyamide of embodiment 1, wherein at least 70 mole percent of the telechelic polyamide has exactly two functional end groups of the same functional type selected from the group consisting of amino or carboxylic end groups.

[0008] 3. The telechelic polyamide of either embodiment 1 or embodiment 2, wherein the at least one other monomer type further comprises at least one of lactams or aminocarboxylic acids.

[0009] 4. The telechelic polyamide of any one of embodiments 1 to 3, wherein the telechelic polyamide comprises repeat units derived from monomeric units of the dicarboxylic acid monomeric units and at least one of lactam monomeric units, aminocarboxylic acid monomeric units, or diamine monomeric units, such that the amide linkages in the telechelic polyamide comprise at least 50 wt.% of the telechelic polyamide.

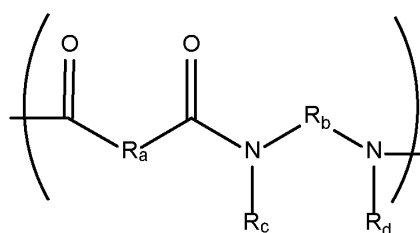
[0010] 5. The telechelic polyamide of any one of embodiments 1 to 4, wherein the telechelic polyamide comprises repeat units derived from monomeric units of the dicarboxylic acid monomeric units and at least one of lactam monomeric units, aminocarboxylic acid

monomeric units, or diamine monomeric units, such that the amide linkages in the telechelic polyamide comprise at least 80 wt.% of the telechelic polyamide.

[0011] 6. The telechelic polyamide of any one of embodiments 1 to 5, wherein the dicarboxylic acid monomeric units comprise at least 50 wt.% of the telechelic polyamide.

5 **[0012]** 7. The telechelic polyamide of any one of embodiments 1 to 6, wherein the dicarboxylic acid monomer units comprise at least 80 wt.% of the telechelic polyamide.

[0013] 8. The telechelic polyamide of any one of embodiments 1 to 7, wherein at least 50 wt.% of the polyamide segment comprises repeat units of the structure



10 wherein R_a is the alkylene portion of the dicarboxylic acid and is a branched alkylene of 34 to 36 carbon atoms, optionally including up to 1 heteroatom per 3 or 10 carbon atoms of the diacid; wherein R_b is a direct bond or a linear or branched (optionally being or including cyclic, heterocyclic, or aromatic portion(s)) alkylene group (optionally containing up to 1 or 3 heteroatoms per 10 carbon atoms) of 2 to 60 carbon atoms; and wherein R_c and R_d are individually hydrogen, a linear or branched alkyl group of 1 to 8 carbon atoms, or R_c and R_d connect together to form a single linear or branched alkylene group of 1 to 8 carbon atoms or optionally with one of R_c and R_d is connected to R_b at a carbon atom.

20 **[0014]** 9. The telechelic polyamide of any one of embodiments 1 to 8, wherein the polyamide segment has two carboxyl end groups and is further reacted with a polyether molecule having one terminal amine and one terminal hydroxyl group or two terminal hydroxyl groups to provide endblocks for the telechelic polyamide, the telechelic polyamide after adding the endblocks having at least 80 mole % terminal primary or secondary hydroxyl end groups.

[0015] 10. The telechelic polyamide of any one of embodiments 1 to 9, having functional end groups, wherein at least 80 mole % of the functional end groups are primary amine groups.

25 **[0016]** 11. The telechelic polyamide of any one of embodiments 1 to 10, wherein the telechelic polyamide has a weight average molecular weight from 200 to 10,000 g/mole, optionally from 400 to 10,000 g/mole.

[0017] 12. The telechelic polyamide of any one of embodiments 1 to 11, wherein the telechelic polyamide has a weight average molecular weight from 200 to 5,000 g/mole, optionally from 400 to 5,000 g/mole, further optionally from 500 to 5,000 g/mole.

5 [0018] 13. The telechelic polyamide of either embodiment 11 or embodiment 12, wherein the telechelic polyamide without solvents has a viscosity of less than 100,000 cps at 70°C as measured by a Brookfield circular disc viscometer with the circular disc spinning at 5 rpm.

[0019] 14. The telechelic polyamide of either embodiment 11 or embodiment 12, wherein the telechelic polyamide without solvents has a viscosity of less than 100,000 cps at 60°C as measured by a Brookfield circular disc viscometer with the circular disc spinning at 5 rpm.

10 [0020] 15. The telechelic polyamide of any one of embodiments 1 to 14, wherein the telechelic polyamide further comprises at least one oligomer segment selected from the group of a polyester segment, a polyether segment, and a polycarbonate segment.

[0021] 16. The telechelic polyamide of any one of embodiments 1 to 8, wherein the polyamide segment has two carboxyl end groups and is further reacted with an amino alcohol
15 having 3 to 16 carbon atoms and having one secondary amine group and one hydroxyl group to provide hydroxyl terminal groups for the telechelic polyamide.

DETAILED DESCRIPTION

[0022] As used herein, the indefinite article “a”/“an” is intended to mean one or more than one. As used herein, the phrase “at least one” means one or more than one of the following
20 terms. Thus, “a”/“an” and “at least one” may be used interchangeably. For example “at least one of A, B or C” means that just one of A, B or C may be included, and any mixture of two or more of A, B and C may be included, in alternative embodiments.

[0023] As used herein, the term “substantially” means that a value of a given quantity is within $\pm 10\%$ of the stated value. In other embodiments, the value is within $\pm 5\%$ of the stated
25 value. In other embodiments, the value is within $\pm 2.5\%$ of the stated value. In other embodiments, the value is within $\pm 1\%$ of the stated value.

[0024] As used herein, the term “substantially free of” means that a component does not include any intentional addition of the material which the component is “substantially free of”. For example, the component may include a material which the component is “substantially free
30 of” at no more than impurity levels, which may be the result of incomplete chemical reactions and/or unintended/undesired (but perhaps unavoidable) reaction products.

[0025] As used herein, the transitional term “comprising,” which is synonymous with “including,” “containing,” or “characterized by,” is inclusive or open-ended and does not exclude additional, un-recited elements or method steps. However, in each recitation of “comprising” herein, it is intended that the term also encompass, as alternative embodiments, the phrases “consisting essentially of” and “consisting of,” where “consisting of” excludes any element or step not specified and “consisting essentially of” permits the inclusion of additional un-recited elements or steps that do not materially affect the essential or basic and novel characteristics of the composition or method under consideration.

[0026] Parentheses are used herein to designate: (1) that the something is optionally present, such that “monomer(s)” means “monomer” and/or “monomers” or “(meth)acrylate” means “methacrylate” and/or “acrylate”; (2) to qualify or further define a previously mentioned term; or (3) to list narrower or different embodiments.

[0027] Telechelic polymers may be described as macromolecules that contain two reactive end groups, and may be used as cross-linkers, chain extenders, and important building blocks for various macromolecular structures, including block and graft copolymers, star, hyperbranched or dendritic polymers. Telechelic polymers of the polydiene, polyester, polyether, and polycarbonate type are known. These known telechelic polymers, with functional end groups selected from primary or secondary hydroxyl, primary or secondary amine, and carboxylic acid, have been reacted with complimentary reactants to form larger polymers with the properties of telechelic precursors. Easy-to-process polyamide telechelic polymers with low melting points were not available prior to work done by the inventor(s) hereof.

[0028] Polyester polyols render good mechanical properties and UV resistance, but they suffer from poor hydrolysis resistance. Polyether polyols have better hydrolytic stability than polyesters, but fall short in UV resistance. Polycarbonate polyols offer improved hydrolysis resistance over polyesters with some degree of increased hardness, but they are more expensive than other polyols. Polydiene polyols are useful but are too hydrophobic to interact well with polar substrates. Some polydiene polyols are hydrogenated to reduce degradation mechanisms relying on residual unsaturation from the diene monomer. Therefore, a new class of telechelic polyamide will help overcome these problems.

[0029] Amine terminated polyamide oligomers were made with low viscosity, low glass transition temperature, suppressed crystallinity, low acid number, with various nitrogen or amide:hydrocarbon weight ratios (or hydrophilic/hydrophobic balance), and with a controlled number of hydrogen bonding or non-hydrogen bonding amide groups.

5 **[0030]** A series of polyamide oligomers from conventional difunctional acids and amines were made. The initial oligomers contained amine terminations and in reaction with diisocyanates form polyamide-polyurea backbone. However, the presence of strong hydrogen bond in these structures makes them very hard (high glass transition) even at low molecular-weight and therefore not suitable for further structural modifications or preparation higher
10 molecular weight polymers or crosslinked networks. We discovered that substitution of N-alkyl groups on these polymers make them soft and easy to process.

[0031] The present subject matter relates to polyamide oligomers or telechelic polyamides resistant to chain scission, e.g. by hydrolysis or UV degradation, useful as macromonomers, prepolymers or polymer segments to make higher molecular weight polymers and/or
15 crosslinked polymer networks. The resulting polymers or networks have better thermal stability than similar polymers or networks from polyethers and/or polyesters due to the higher thermal stability of the amide bonds. Polymers built from moderate molecular weight polyamide oligomers and co-reactants that can form chemical bonds with co-reactive groups at the termini of the oligomers. These polymers have many of the properties of the polyamide oligomers from
20 which they are made as the oligomers form a substantial weight percent of the final polymer. Modifying the molecular weight and composition of the oligomers can be used to achieve the desired properties. They can also be used to make polyurea/urethanes. The term polyurea/urethane will be used to refer to polymers that have urea linkages, urethane linkages or blends of such linkages. The composition may contain small amounts of other polymers and
25 materials either as physical blends or where the other polymers or materials may be co-reacted into the polyamide.

[0032] The term polyamide oligomer will refer to an oligomer with two or more amide linkages, or sometimes the amount of amide linkages will be specified. A subset of polyamide oligomers will be telechelic polyamides. Telechelic polyamides will be polyamide oligomers
30 with high percentages, or specified percentages, of two functional groups of a single chemical type, e.g. two terminal amine groups (meaning either primary, secondary, or mixtures), two

terminal carboxyl groups, two terminal hydroxyl groups (again meaning primary, secondary, or mixtures), or two terminal isocyanate groups (meaning aliphatic, aromatic, or mixtures). Ranges for the percent difunctional that are preferred to meet the definition of telechelic are at least 70 or 80, more desirably at least 90 or 95 mole % of the oligomers being difunctional as opposed to higher or lower functionality. Reactive amine terminated telechelic polyamides will be telechelic polyamide oligomers where the terminal groups are both amine types, either primary or secondary and mixtures thereof, i.e. excluding tertiary amine groups.

[0033] A first portion of the present subject matter is the substitution of polyamide segments for polyester, polyether, or polycarbonate soft segments in a polymer made from telechelic oligomers. The replacement or substitution of polyamide segments for polyester, polyether, or polycarbonate segments can be partial or complete. Optimum environmental resistance, including thermal stability, would result from complete replacement of polyester and polyether segments, due to their potential for easier chain scission in polyethers and polyesters. In some embodiments some of the polyester and or polyether segments could be retained in the telechelic polyamide or polyamide oligomer for their ability to soften the elastomeric portion or modify the compatibility of the resulting polymer with other polymer surfaces. When polymer from polyesters or polyether are degraded by hydrolysis or UV activated chain scission the molecular weight of the polymer is decreased such that the polymer, or segment, soon loses its tensile strength, elongation to break, resistance to solvents, etc.

[0034] A second benefit of the first portion of the present subject matter, substituting soft polyamide segments for soft polyether or polyester segments, is that the polyamide segments tend to promote better wetting and adhesion to a variety of polar substrates, such as glass, nylon, and metals than polyester or polyether based polymers. The hydrophobic/hydrophilic nature of the polyamide can be adjusted by using different weight ratios of hydrocarbon to amide linkages, or nitrogen atoms, in the polyamide. Diacids, diamines, aminocarboxylic acids, and lactams with large aliphatic hydrocarbon portions relative to the amide linkage portion tend to be hydrophobic. When the hydrocarbon weight ratio to amide linkage, or nitrogen atoms, becomes smaller, the polyamide is more hydrophilic. Increasing the amount of polyamide in a polymer can increase adhesion to substrates that have similar or compatible surfaces to polyamides.

[0035] Polymers made from polyamide segments can have good solvent resistance. Solvents can cause deformation and swelling of a polymer thereby causing premature failure of the polymer. Solvents can cause a coating to swell and delaminate from a substrate at the interface between the two.

5 **[0036]** It should be noted that many conventional polyamides may be high melting point crystalline polyamides such as 6-nylon, 6,6-nylon, 6,10-nylon that melt at temperatures much too high, e.g. in excess of 100°C, to serve as soft segments if a blocky thermoplastic polymer is desired. In some of the prior art publications the polyamide, often a crystalline or high glass transition temperature (“Tg”) polyamide type, was added merely to increase the surface
10 interaction with a substrate that was compatible to polyamides. To create a lower Tg polymer, soft (low Tg) polyester, polyether or polycarbonates were added to the polyamide segment to provide a lower composite Tg elastomeric segment. In other prior art publications only a few polyamide linkages were inserted into a polymer to modify the polarity of the polymer, to increase solvent resistance, or to raise the softening temperature.

15 **[0037]** One objective of certain embodiments of the present subject matter is to use high percentages of amide linkages in a telechelic oligomer comprised of one or more polyamide segments to provide resistance to chain scission from hydrolysis and/or UV activated chain scission. Thus, many embodiments will describe soft segments with high percentages of total linkages between repeat units in the soft segment being amide linkages. Some embodiments
20 may allow for some linkages between repeat units to be other than amide linkages.

[0038] An important modification from conventional polyamides to get low Tg polyamide soft segments is the use of monomers with secondary amine terminal groups in forming the polyamide. The amide linkage formed from a secondary amine and a carboxylic acid type group is called a tertiary amide linkage. Primary amines react with carboxylic acid type groups
25 to form secondary amides. The nitrogen atom of a secondary amide has an attached hydrogen atom that often hydrogen bonds with a carbonyl group of a nearby amide. The intra-molecular H-bonds induce crystallinity with high melting point and can act as crosslinks reducing chain mobility. With tertiary amide groups the hydrogen on the nitrogen of the amide linkage is eliminated along with hydrogen bonding. A tertiary amide linkage that has one additional alkyl
30 group attached to it as compared to a secondary amide group, which has hydrogen attached to it, has reduced polar interactions with nearby amide groups when the polymer exists in a bulk

polymer sample. Reduced polar interactions mean that glassy or crystalline phases that include the amide linkage melt at lower temperatures than similar amide groups that are secondary amide groups. One way to source secondary amine reactant, a precursor to tertiary amide linkages, is to substitute one hydrogen atom on the nitrogen of the amine containing monomer with an alkyl group. Another way to source a secondary amine reactant is to use a heterocyclic molecule where the nitrogen of the amine is part of the ring structure. Piperazine is a common cyclic diamine where both nitrogen atoms are of the secondary type and part of the heterocyclic ring.

[0039] Another modification to reduce the Tg of the polyamide soft segments is to use at least one additional monomer beyond the minimum number of monomers to form the polyamide. Thus, for a polyamide formed from a lactam polymerization, such as from N-methyl-dodecyl lactam one would include an additional lactam, aminocarboxylic acid, diamine, or dicarboxylic acid in the monomers for the polymerization to change the spacing (among repeat units) between the amide linkages formed by the monomer so that the spacing between the amide linkages in the polyamide is irregular along the backbone, e.g. not the same physical dimension for some of the repeat units in each oligomer. For a polymerization of aminocarboxylic acid one would include additional lactam, aminocarboxylic acid, diamine, or dicarboxylic acid (with different physical length between the primary reactive groups of the monomer) in the monomer blend for the polymerization to change the spacing among repeat units between the amide linkages. Switching end groups on the monomers can also disrupt regularity in the spacing of the polar amide linkages and lower the effective Tg of the copolymer. Thus, co-polymerizing a C₆ amino carboxylic acid with a small portion of a C₆ diacid and C₆ diamine can disrupt regularity of the amide linkages as the diacid and diamine units would switch the orientation of the amide linkage from head to tail orientation to tail to head orientation, slightly disrupting uniformity of spacing of the amide linkages along the polyamide backbone. Typically, when following this procedure, one would try to add a disrupting monomer that increased or decreased the number of atoms between the amide forming end groups of the monomer(s) used as the primary monomer in the polyamide. One could also use a second disrupting monomer that had a cyclic structure, such as piperazine, a cyclic diamine monomer with where two methylene atoms form the top half of the ring and two methylene atoms form the bottom half of the ring, to disrupt the regularity of polyamide formed

from a diacid reacted with a diamine monomer with two methylene atoms between the nitrogen atoms of the diamine.

[0040] Generally, having nearly equal amounts of two or more different amide forming monomers results in different spacing between the amide linkages along the polyamide backbone and affords optimal reduction of the crystalline melting and glass transition temperatures. For example, a 50:50 mole blend of two different diamines would be desirable. A 50:50 mole blend of two different diacids would be desirable. A 33:33:33 mole blend of a lactam with a diacid and a diamine would be desirable.

[0041] We use the term low glass transition temperature ("Tg"), even though we realize most of the polyamide segments are initially low molecular weight and it would not be easily possible to measure the Tg of the low molecular weight oligomers, the measured value would be dramatically affected by molecular weight. High Tg polymers, e.g. having Tg values above 70, 80, or 90°C as measured by differential scanning calorimetry (DSC), form solids even at low molecular weights. Thus, the polyamide oligomers, telechelic polyamides, and even the oligomers from telechelic polyamides or polyamide oligomers are often described in this specification by their viscosity at specific temperatures. Low Tg polyamides oligomers will be defined as those compositions that would have Tg, if above 20,000 g/mole molecular weight, of below 50°C, more desirably below 25 or 0°C.

[0042] In one embodiment the telechelic oligomer or telechelic polyamide will have a viscosity measured by a Brookfield circular disc viscometer with the circular disc spinning at 5 rpm of less than 100,000 cps at a temperature of 70°C, more desirably less than 15,000 or 10,000 cps at 70°C, still more desirably less than 100,000 cps at 60 or 50°C, and more preferably less than 15,000 or 10,000 cps at 60°C; and still more preferable less than 15,000 or 10,000 cps at 50°C. Desirably, these viscosities are those of neat telechelic prepolymers or polyamide oligomers without solvent or plasticizers. These viscosity values will facilitate mixing the telechelic polyamide with co-reactants and or particulate materials under suitable conditions that desirable reactions occur at reasonable rates and undesirable reactions, e.g. side reactions, do not occur to any significant extent. In some embodiments the telechelic polyamide can be diluted with solvent to achieve viscosities in these ranges.

[0043] Many of the oligomers, telechelics, and polymers of this specification are made by condensation reactions of reactive groups on desired monomer(s). Lactam polymerization into

a polyamide results in similar amide linkages by a chain polymerization process and is well known in the art. These condensation reactions between carboxylic acid groups and amine or hydroxyl groups are well known and are driven by the removal of water and or catalysts. The formation of amides from the reaction of carboxylic acid groups and amine groups can be catalyzed by boric acid, boric acid esters, boranes, phosphorous acid, phosphates, phosphate esters, amines, acids, bases, silicates, and silsesquioxanes. Additional catalysts, conditions, etc. are available in textbooks such as "Comprehensive Organic Transformations" by Larock.

[0044] The condensation reaction of reactive groups will be defined as creating chemical linkages between the monomers. The portion of the monomer that is incorporated into the oligomer or polymer will be defined as the repeat unit from the particular monomer. Some monomers, such as aminocarboxylic acid, or one end of diacid reacting with one end of a diamine, lose one molecule of water as the monomer goes from a monomer to a repeat unit of a polymer. Other monomers, such as lactams, isocyanates, amines reacted with isocyanates, hydroxyl groups reacted with isocyanates, etc. do not release a portion of the molecule to the environment but rather retain all of the monomer in the resulting polymer.

[0045] We will define polyamide oligomer as a species below 20,000 g/mole molecular weight, e.g. often below 10,000; 5,000; 2,500; or 2000 g/mole, that has two or more amide linkages per oligomer. Later we will define preferred percentages of amide linkages or monomers that provide on average one amide linkage per repeat unit in various oligomeric species. A subset of polyamide oligomer will be telechelic oligomer. The telechelic polyamide has molecular weight preferences identical to the polyamide oligomer above. The term telechelic has been earlier defined. Multiple polyamide oligomers or telechelic polyamides can be linked with condensation reactions to form polymers, generally above 100,000 g/mole.

[0046] Generally, amide linkages are formed from the reaction of a carboxylic acid group with an amine group or the ring opening polymerization of a lactam, e.g. where an amide linkage in a ring structure is converted to an amide linkage in a polymer. In a preferred embodiment a large portion of the amine groups of the monomers are secondary amine groups or the nitrogen of the lactam is a tertiary amide group. Secondary amine groups form tertiary amide groups when the amine group reacts with carboxylic acid to form an amide. For the purposes of this disclosure, the carbonyl group of an amide, e.g. as in a lactam, will be considered as derived from a carboxylic acid group. The amide linkage of a lactam is formed

from the reaction of carboxylic group of an aminocarboxylic acid with the amine group of the same aminocarboxylic acid. In one embodiment we want less than 20, 10 or 5 mole percent of the monomers used in making the polyamide to have functionality in polymerization of amide linkages of 3 or more. This will reduce branching in the polyamide oligomer or telechelic polyamide.

[0047] The polyamide oligomers and telechelic polyamides of this disclosure can contain small amounts of ester linkages, ether linkages, urethane linkages, urea linkages, etc. if the additional monomers used to form these linkages are useful to the intended use of the polymers. This allows other monomers and oligomers to be included in the polyamide to provide specific properties, which might be necessary and not achievable with a 100% polyamide segment oligomer. Sometimes added polyether, polyester, or polycarbonate provides softer e.g. lower Tg, segments. Sometimes it is desirable to convert the carboxylic end groups or primary or secondary amine end groups of a polyamide to other functional end groups capable of condensation polymerizations. A telechelic polyamide with carboxylic end groups can be converted into an oligomer with hydroxyl end groups by reacting the telechelic polyamide with a polyether that has two hydroxyl end groups or a polyether that has one amino, primary or secondary, and one hydroxyl end group. Oligomers or polymers with polyether segments have susceptibility to chain breakage due to UV exposure. The effect of UV exposure on block copolymers of nylon 6-polyethylene glycol block copolymers is reported in Gauvin, Pascal; Lemaire, Jacques in Makromolekulare Chemie (1987), 188(5), 971-986. Sometimes an initiator for oligomer chain polymerization of a lactam is used that doesn't generate an amide linkage. Sometimes a polyether might be used as a segment or portion of a polyamide to reduce the Tg, or provide a soft segment, of the resulting polyamide oligomer. Sometimes a polyamide segment, e.g. possibly difunctional with carboxylic acid or amine terminal groups, can be functionalized with two polyether end segments (such as from Jeffamine™ D230) to further lower the Tg of, or provide a soft segment in, the polyamide oligomer and create a telechelic polyamide with amine or hydroxyl end groups. Sometimes a carboxylic acid terminated telechelic polyamide segment is functionalized by reacting with an amino alcohol, such as N-methylaminoethanol or $\text{HN}(\text{R}^a)(\text{R}^b)$ where R^a is a C_1 to C_4 alkyl group and R^b comprises an alcohol group and a C_2 to C_{12} alkylene group, alternatively R^a and R^b can be interconnected to form a C_3 to C_{16} alkylene group including a cyclic structure and pendant hydroxyl group (such

as in 2-hydroxymethyl piperidine), either of which can create a telechelic polyamide with terminal hydroxyl groups. The reaction of the secondary amine (as opposed to the hydroxyl group) with the carboxylic acid can be favored by using a 100% molar excess of the amino alcohol and conducting the reaction at 160 °C +/- 10 or 20 °. The excess amino alcohol can be removed by distillation after reaction. In one embodiment the functional primary or secondary amine groups of a telechelic polyamide are reacted with a lactone of 2 or 4 to 10 carbon atoms (e.g. a valero or caprolactone) and/or hydroxyl carboxylic acid of 3 to 30 carbon atoms to create one or two hydroxyl functional end groups derived from the lactone or the hydroxyl carboxylic acid on the telechelic polyamide. Optimally only one repeat unit from the lactone or hydroxyl carboxylic acid is added to each end of the telechelic polyamide.

[0048] As earlier indicated many amide forming monomers create on average one amide linkage per repeat unit. These include diacids and diamines when reacted with each other, aminocarboxylic acids, and lactams. When we discuss these monomers or repeat units from these monomers we generally mean these monomers, their repeat units and their reactive equivalents (meaning monomers that generate the same repeat unit as the named monomer). These reactive equivalents might include anhydride of diacids, esters of diacids, etc. These monomers, when reacted with other monomers in the same group, also create amide linkages at both ends of the repeat units formed. Thus we will use both percentages of amide linkages and mole percent and weight percentages of repeat units from amide forming monomers. Amide forming monomers will be used to refer to monomers that form on average one amide linkage per repeat unit in normal amide forming condensation linking reactions.

[0049] Provided are telechelic polyamides comprising: (a) repeat units derived from polymerizing monomers connected by linkages between the repeat units and functional end groups selected from carboxyl or primary or secondary amine; (b) a polyamide segment comprising at least two amide linkages characterized as being derived from reacting an amine with a carboxyl group, and the polyamide segment comprises repeat units derived from polymerizing dicarboxylic acid monomeric units with monomeric units of at least one other monomer type, wherein at least 50 (such as 55, 60, 65, 70, 75, 80, 85, 90, or 95) percent of the dicarboxylic acid monomeric units each contain 34 to 36 carbon atoms, and wherein the at least one other monomer type comprises diamine monomeric units; (c) wherein at least 10% of the total number of the heteroatom containing linkages connecting hydrocarbon type linkages are

characterized as being amide; and (d) wherein less than 25% (such as 20%, 15%, 10%, 9%, 8%, 7%, 6%, 5%, 4%, 3%, 2%, or 1%) of the amide linkages are characterized as being tertiary amide linkages; wherein at least 20 mole percent of the telechelic polyamide has exactly two functional end groups of the same functional type comprising amino end groups or carboxylic end groups.

[0050] In certain embodiments, substantially all or all of the dicarboxylic acid monomeric units contain an average across all units of 34 to 36 carbon atoms.

[0051] In certain embodiments, the telechelic polyamide is substantially free of, or free of amide linkages characterized as being tertiary amide linkages.

[0052] In certain embodiments, the dicarboxylic acid monomeric units which do not contain 34 to 36 carbon atoms may be any other suitable dicarboxylic acid monomeric units. For example, suitable dicarboxylic acids may be those where the alkylene portion of the dicarboxylic acid is a cyclic, linear, or branched (optionally including aromatic groups) alkylene of 2 to 36 carbon atoms, optionally including up to 1 heteroatom per 3 or up to 1 heteroatom per 10 carbon atoms of the diacid, such as from 4 to 36 carbon atoms (the diacid would include 2 more carbon atoms than the alkylene portion). These include dimer fatty acids, hydrogenated dimer acid, sebacic acid, etc. Generally suitable are diacids with larger alkylene groups, as this may provide polyamide repeat units with lower T_g value.

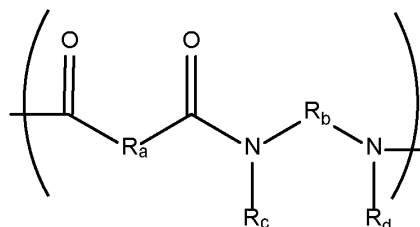
[0053] In certain embodiments, at least 70 (such as 75, 80, 85, 90, or 95) mole percent of the telechelic polyamide has exactly two functional end groups of the same functional type selected from the group consisting of amino or carboxylic end groups.

[0054] In certain embodiments, the at least one other monomer type further comprises at least one of lactams or aminocarboxylic acids.

[0055] In certain embodiments, the telechelic polyamide comprises repeat units derived from monomeric units of the dicarboxylic acid monomeric units and at least one of lactam monomeric units, aminocarboxylic acid monomeric units, or diamine monomeric units, such that the amide linkages in the telechelic polyamide comprise at least 50 (such as 55, 60, 65, 70, 75, or 80) wt.% of the telechelic polyamide.

[0056] In certain embodiments, the dicarboxylic acid monomeric units comprise at least 50 (such as 55, 60, 65, 70, 75, or 80) wt.% of the telechelic polyamide.

[0057] In certain embodiments, at least 50 (such as 55, 60, 65, 70, 75, 80, 85, 90, or 95) wt.% of the polyamide segment of the telechelic polyamide comprises repeat units of the structure



5 wherein R_a is the alkylene portion of the dicarboxylic acid and is a branched alkylene of 34 to 36 carbon atoms, optionally including up to 1 heteroatom per 3 or 10 carbon atoms of the diacid; wherein R_b is a direct bond or a linear or branched (optionally being or including cyclic, heterocyclic, or aromatic portion(s)) alkylene group (optionally containing up to 1 or 3 heteroatoms per 10 carbon atoms) of 2 to 60 carbon atoms; and wherein R_c and R_d are
 10 individually hydrogen, a linear or branched alkyl group of 1 to 8 carbon atoms, or R_c and R_d connect together to form a single linear or branched alkylene group of 1 to 8 carbon atoms or optionally with one of R_c and R_d is connected to R_b at a carbon atom.

[0058] In certain embodiments, the polyamide segment has two carboxyl end groups and is further reacted with a polyether molecule having one terminal amine and one terminal hydroxyl
 15 group or two terminal hydroxyl groups to provide endblocks for the telechelic polyamide, the telechelic polyamide after adding the endblocks having at least 80 mole % terminal primary or secondary hydroxyl end groups.

[0059] In certain embodiments, the telechelic polyamide has function end groups, wherein at least 80 mole % of the functional end groups are primary amine groups.

20 **[0060]** In certain embodiments, the telechelic polyamide has a weight average molecular weight from 200 to 10,000 g/mole, optionally from 400 to 10,000 g/mole.

[0061] In certain embodiments, the telechelic polyamide has a weight average molecular weight from 200 to 5,000 g/mole, optionally from 400 to 5,000 g/mole, further optionally from 500 to 5,000 g/mole.

25 **[0062]** In certain embodiments, the telechelic polyamide has a weight average molecular weight of from 200 (such as 220, 240, 260, 280, 300, 320, 340, 360, 380, 400, 420, 440, 460, 480, 500, 520, 540, 560, 580, 600, 620, 640, 660, 680, 700, 720, 740, 760, 780, 800, 820, 840,

860, 880, 900, 920, 940, 960, 980, 1,000, 1,500, 2,000, 2,500, 3,000, 3,500, 4,000, or 4,500) to 10,000 (such as 9,500, 9,000, 8,500, 8,000, 7,500, 7,000, 6,500, 6,000, 5,500, or 5,000) g/mole.

[0063] In certain embodiments, the telechelic polyamide without solvents has a viscosity of less than 100,000 cps at 70°C as measured by a Brookfield circular disc viscometer with the circular disc spinning at 5 rpm.

[0064] In certain embodiments, the telechelic polyamide without solvents has a viscosity of less than 100,000 cps at 60°C as measured by a Brookfield circular disc viscometer with the circular disc spinning at 5 rpm.

[0065] In certain embodiments, the telechelic polyamide further comprises at least one oligomer segment selected from the group of a polyester segment, a polyether segment, and a polycarbonate segment.

[0066] In certain embodiments, the polyamide segment has two carboxyl end groups and is further reacted with an amino alcohol having 3 to 16 carbon atoms and having one secondary amine group and one hydroxyl group to provide hydroxyl terminal groups for the telechelic polyamide.

[0067] In one embodiment desirably at least 10 mole %, such as at least 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 80, 85, 90, or 95 mole % of the total number of the heteroatom containing linkages connecting hydrocarbon type linkages are characterized as being amide linkages. Heteroatom linkages are linkages such as amide, ester, urethane, urea, ether linkages where a heteroatom connects two portions of an oligomer or polymer that are generally characterized as hydrocarbons (or having carbon to carbon bond, such as hydrocarbon linkages). As the amount of amide linkages in the polyamide increase the amount of repeat units from amide forming monomers in the polyamide increases.

[0068] In one embodiment desirably at least 25 wt. %, such as at least 30, 40, 50, 60, 70, 80, 90, or 95 wt.% of the polyamide oligomer or telechelic polyamide is repeat units from amide forming monomers, also identified as monomers that form amide linkages at both ends of the repeat unit. Such monomers include lactams, aminocarboxylic acids, dicarboxylic acid and diamines.

[0069] In one embodiment desirably less than 25, 20, 15, 10, or 5 mole percent of the amide linkages in the polyamide oligomer or telechelic polyamine are tertiary amide linkages. In certain embodiments, the polyamide oligomer or telechelic polyamine is substantially free of

or free of tertiary amide linkages. As used in this context, “substantially free of” means that every possible attempt was made to prevent tertiary amide linkages from forming during manufacture of the polyamide oligomer or telechelic polyamine. As earlier explained tertiary amide linkages result from ring opening polymerization of lactams with tertiary amides or
 5 reactions of secondary amines with carboxylic acid groups.

[0070] The % of tertiary amide linkages of the total number of amide linkages was calculated with the following equation:

$$\text{Tertiary amide linkage \%} = \frac{\sum_{i=1}^n (w_{tertN,i} \times n_i)}{\sum_{i=1}^n (w_{totalN,i} \times n_i)} \times 100$$

where n is the number of monomers, the index i refers to a certain monomer, w_{tertN} is the average
 10 number nitrogen atoms in a monomer that form or are part of tertiary amide linkages in the polymerizations, (note: end-group forming amines do not form amide groups during the polymerizations and their amounts are excluded from w_{tertN}), w_{totalN} is the average number nitrogen atoms in a monomer that form or are part of tertiary amide linkages in the polymerizations (note: the end-group forming amines do not form amide groups during the
 15 polymerizations and their amounts are excluded from w_{totalN}), and n_i is the number of moles of the monomer with the index i .

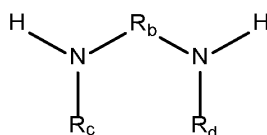
[0071] The % of amide linkages of the total number of all heteroatom containing linkages (connecting hydrocarbon linkages) was calculated by the following equation:

$$\text{Amide linkage \%} = \frac{\sum_{i=1}^n (w_{totalN,i} \times n_i)}{\sum_{i=1}^n (w_{totalS,i} \times n_i)} \times 100$$

20 where w_{totalS} is the sum of the average number of heteroatom containing linkages (connecting hydrocarbon linkages) in a monomer and the number of heteroatom containing linkages (connecting hydrocarbon linkages) forming from that monomer by the reaction with a carboxylic acid bearing monomer during the polyamide polymerizations. “Hydrocarbon linkages” are just the hydrocarbon portion of each repeat unit formed from continuous carbon
 25 to carbon bonds (i.e. without heteroatoms such as nitrogen or oxygen) in a repeat unit. This hydrocarbon portion would be the ethylene or propylene portion of ethylene oxide or propylene oxide; the undecyl group of dodecyl lactam, the ethylene group of ethylenediamine, and the $(CH_2)_4$ (or butylene) group of adipic acid.

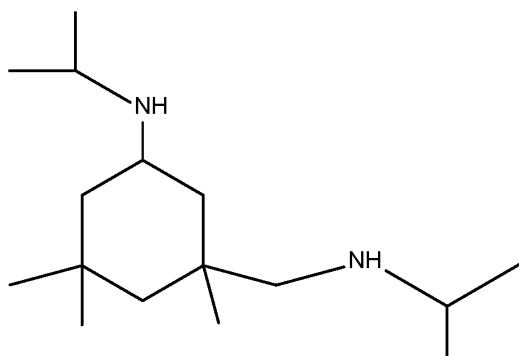
[0072] Preferred amide or tertiary amide forming monomers include dicarboxylic acids, diamines, aminocarboxylic acids and lactams. Preferred dicarboxylic acids are where the alkylene portion of the dicarboxylic acid is a cyclic, linear, or branched (optionally including aromatic groups) alkylene of 2 to 36 carbon atoms, optionally including up to 1 heteroatom per 3 or 10 carbon atoms of the diacid, more preferably from 4 to 36 carbon atoms (the diacid would include 2 more carbon atoms than the alkylene portion). These include dimer fatty acids, hydrogenated dimer acid, sebacic acid, etc. Generally we prefer diacids with larger alkylene groups as this generally provides polyamide repeat units with lower Tg value.

[0073] Preferred diamines include those with up to 60 carbon atoms, optionally including 1 heteroatom (besides the two nitrogen atoms) for each 3 or 10 carbon atoms of the diamine and optionally including a variety of cyclic, aromatic or heterocyclic groups providing that one or both of the amine groups are secondary amines, a preferred formula is:

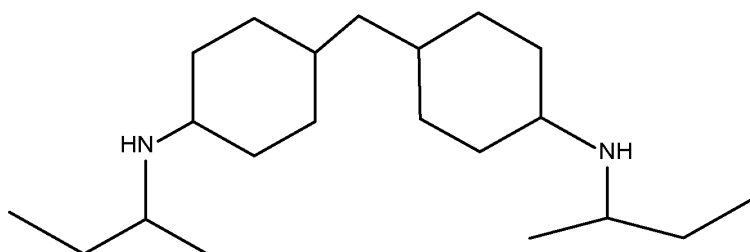


wherein R_b is a direct bond or a linear or branched (optionally being or including cyclic, heterocyclic, or aromatic portion(s)) alkylene group (optionally containing up to 1 or 3 heteroatoms per 10 carbon atoms of the diamine) of 2 to 36 carbon atoms and more preferably 2 or 4 to 12 carbon atoms and R_c and R_d are individually hydrogen, a linear or branched alkyl group of 1 to 8 carbon atoms, more preferably 1 or 2 to 4 carbon atoms, or R_c and R_d connect together to form a single linear or branched alkylene group of 1 to 8 carbon atoms or optionally with one of R_c and R_d is connected to R_b at a carbon atom, more desirably R_c and R_d being 1 or 2 to 4 carbon atoms.

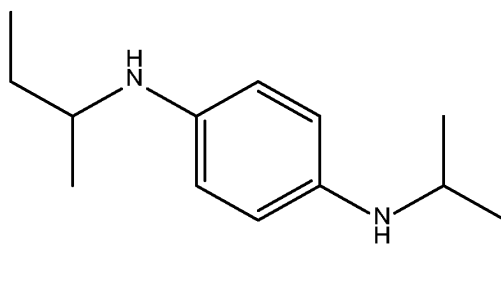
[0074] Such diamines include EthacureTM 90 from Albermarle (supposedly a N,N'-bis(1,2,2-trimethylpropyl)-1,6-hexanediamine); ClearlinkTM 1000 from Dorfketal, or JefflinkTM 754 from Huntsman; N-methylaminoethanol; dihydroxy terminated, hydroxyl and amine terminated or diamine terminated poly(alkyleneoxide) where the alkylene has from 2 to 4 carbon atoms and having molecular weights from about 40 or 100 to 2000; N,N'-diisopropyl-1,6-hexanediamine; N,N'-di(sec-butyl) phenylenediamine; piperazine; homopiperazine; and methyl-piperazine. JefflinkTM 754 has the structure



Clearlink™ 1000 has the structure



Another diamine with an aromatic group is: N,N'-di(sec-butyl) phenylenediamine, see structure
5 below:

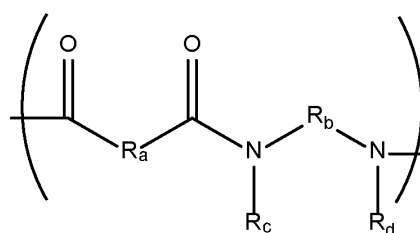


Preferred diamines are diamines wherein both amine groups are secondary amines.

- [0075] Preferred lactams include straight chain or branched alkylene segments therein of 4 to 12 carbon atoms such that the ring structure without substituents on the nitrogen of the lactam has 5 to 13 carbon atoms total (when one includes the carbonyl) and the substituent on the nitrogen of the lactam (if the lactam is a tertiary amide) is an alkyl of from 1 to 8 carbon atoms and more desirably an alkyl of 1 to 4 carbon atoms. Dodecyl lactam, alkyl substituted dodecyl lactam, caprolactam, alkyl substituted caprolactam, and other lactams with larger alkylene groups are preferred lactams as they provide repeat units with lower T_g values.
- 15 Aminocarboxylic acids have the same number of carbon atoms as the lactams. Desirably the

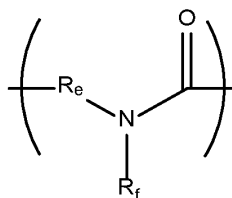
number of carbon atoms in the linear or branched alkylene group between the amine and carboxylic acid group of the aminocarboxylic acid is from 4 to 12 and the substituent on the nitrogen of the amine group (if it is a secondary amine group) is an alkyl group with from 1 to 8 carbon atoms, more preferably 1 or 2 to 4 carbon atoms. Aminocarboxylic acids with
 5 secondary amine groups are preferred.

[0076] In one embodiment desirably at least 50 wt.%, such as at least 60, 70, 80, or 90 wt.% of the polyamide oligomer or telechelic polyamide comprise repeat units from diacids and diamines of the structure of the repeat unit being:



10 wherein R_a is the alkylene portion of the dicarboxylic acid and is a branched alkylene of 34 to 36 carbon atoms, optionally including up to 1 heteroatom per 3 or 10 carbon atoms of the diacid and wherein R_b is a direct bond or a linear or branched (optionally being or including cyclic, heterocyclic, or aromatic portion(s)) alkylene group (optionally containing up to 1 or 3
 15 heteroatoms per 10 carbon atoms) of 2 to 36 or 60 carbon atoms and more preferably 2 or 4 to 12 carbon atoms and R_c and R_d are individually hydrogen, a linear or branched alkyl group of 1 to 8 carbon atoms, more preferably 1 or 2 to 4 carbon atoms, or R_c and R_d connect together to form a single linear or branched alkylene group of 1 to 8 carbon atoms or optionally with one of R_c and R_d is connected to R_b at a carbon atom, more desirably R_c and R_d being an alkyl group of 1 or 2 to 4 carbon atoms.

20 **[0077]** In one embodiment desirably at least 50 wt.%, more desirably at least 60, 70, 80 or 90 wt.% of the polyamide oligomer or telechelic polyamide comprise repeat units from lactams or amino carboxylic acids of the structure:



repeat units can be in a variety of orientations in the oligomer derived from lactams or amino carboxylic acid depending on initiator type, wherein each R_e independently is linear or branched alkylene of 4 to 12 carbon atoms and each R_f independently is hydrogen or a linear or branched alkyl of 1 to 8, more desirably 1 or 2 to 4, carbon atoms.

5 **[0078]** The above described polyamide oligomers and telechelic polyamide are useful to make polymers by reacting the polyamide oligomer or telechelic polyamide with co-reactants having two or more reactive groups that can form chemical bonds when reacted with the functional groups of the polyamide oligomers or telechelic polyamide (e.g. these functional groups of the polyamide include primary and secondary amine, primary or secondary hydroxyl, 10 or carboxylic acid group). The reactive groups on the co-reactants may be isocyanate, or with particular telechelic polyamides they could be hydroxyl, amine or carboxylic acid groups.

[0079] We made a series of polyamide oligomers from conventional difunctional acids and amines. These oligomers contained amine terminations and in reaction with diisocyanates form polyamide-polyurea backbone. The polyamide building blocks in our new dispersion polymers 15 provide excellent hydrolytic stability, superior heat and UV resistance, and better overall mechanical properties in comparison to polyester and polyether segments. In addition, the amine chain termination in these polyamide oligomers, when reacted with isocyanates, forms urea linkages, vs. urethane link from polyol reacted with isocyanates. These polyurea linkages are known to have stronger intermolecular attractions that act more like a true crosslinked 20 polymer, resulting in performance advantages over urethanes, including but not limited to better solvent resistance and elasticity.

[0080] The polyamide oligomer or telechelic polyamide described herein may be combined with compatible polymers and polymer dispersions by methods well known to those skilled in the art. Such polymers, polymer solutions, and dispersions include those described in A. S. 25 Teot. "Resins, Water-Soluble" in: Kirk-Othmer Encyclopedia of Chemical Technology. John Wiley & Sons. 3rd Ed., Vol. 20, H. F. Mark et al. Eds., pp. 207-230 (1982).

[0081] The oligomeric polyamides or telechelic polyamides described herein may be useful as components in polymer compositions used as coatings, films, fibers, adhesives, or molded or extruded goods.

30 **[0082]** Except where explicitly indicated or required by context, all numerical quantities in this description specifying amounts of materials, reaction conditions, molecular weights,

number of carbon atoms, and the like, are to be understood as modified by the word “about”. As used herein, the term “about” means that a value of a given quantity is within $\pm 20\%$ of the stated value. In other embodiments, the value is within $\pm 15\%$ of the stated value. In other embodiments, the value is within $\pm 10\%$ of the stated value. In other embodiments, the value is within $\pm 5\%$ of the stated value. In other embodiments, the value is within $\pm 2.5\%$ of the stated value. In other embodiments, the value is within $\pm 1\%$ of the stated value. In other embodiments, the value is within a range of the explicitly-described value which would be understood by those of ordinary skill, based on the disclosures provided herein, to perform substantially similarly to compositions including the literal amounts described herein.

10 **[0083]** It is to be understood that the upper and lower amount, range, and ratio limits set forth herein may be independently combined, and that any amount within a disclosed range is contemplated to provide a minimum or maximum of a narrower range in alternative embodiments (with the proviso, of course, that the minimum amount of a range must be lower than the maximum amount of the same range). Similarly, the ranges and amounts for each
15 element of the subject matter disclosed herein may be used together with ranges or amounts for any of the other elements.

[0084] While certain representative embodiments and details have been shown for the purpose of illustrating the subject matter disclosed herein, it will be apparent to those skilled in this art that various changes and modifications may be made therein without departing from
20 the scope of the subject matter. In this regard, the scope of the invention is to be limited only by the following claims.

What is claimed is:

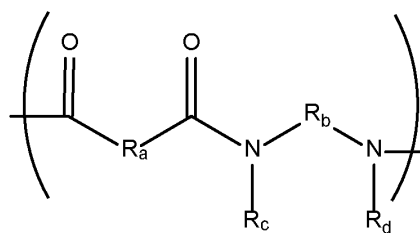
1. A telechelic polyamide comprising:
 - a) repeat units derived from polymerizing monomers connected by linkages between the repeat units and functional end groups selected from carboxyl or primary or secondary amine;
 - b) a polyamide segment comprising at least two amide linkages characterized as being derived from reacting an amine with a carboxyl group, and the polyamide segment comprises repeat units derived from polymerizing dicarboxylic acid monomeric units with monomeric units of at least one other monomer type, wherein at least 50 percent of the dicarboxylic acid monomeric units each contain 34 to 36 carbon atoms, and wherein the at least one other monomer type comprises diamine monomeric units;
 - c) wherein at least 10% of the total number of the heteroatom containing linkages connecting hydrocarbon type linkages are characterized as being amide; and
 - d) wherein less than 25% of the amide linkages are characterized as being tertiary amide linkages;wherein at least 20 mole percent of the telechelic polyamide has exactly two functional end groups of the same functional type comprising amino end groups or carboxylic end groups.
2. The telechelic polyamide of claim 1, wherein at least 70 mole percent of the telechelic polyamide has exactly two functional end groups of the same functional type selected from the group consisting of amino or carboxylic end groups.
3. The telechelic polyamide of either claim 1 or claim 2, wherein the at least one other monomer type further comprises at least one of lactams or aminocarboxylic acids.
4. The telechelic polyamide of any one of claims 1 to 3, wherein the telechelic polyamide comprises repeat units derived from monomeric units of the dicarboxylic acid monomeric units and at least one of lactam monomeric units, aminocarboxylic acid monomeric units, or diamine monomeric units, such that the amide linkages in the telechelic polyamide comprise at least 50 wt.% of the telechelic polyamide.

5. The telechelic polyamide of any one of claims 1 to 4, wherein the telechelic polyamide comprises repeat units derived from monomeric units of the dicarboxylic acid monomeric units and at least one of lactam monomeric units, aminocarboxylic acid monomeric units, or diamine monomeric units, such that the amide linkages in the telechelic polyamide comprise at least 80 wt.% of the telechelic polyamide.

6. The telechelic polyamide of any one of claims 1 to 5, wherein the dicarboxylic acid monomeric units comprise at least 50 wt.% of the telechelic polyamide.

7. The telechelic polyamide of any one of claims 1 to 6, wherein the dicarboxylic acid monomer units comprise at least 80 wt.% of the telechelic polyamide.

8. The telechelic polyamide of any one of claims 1 to 7, wherein at least 50 wt.% of the polyamide segment comprises repeat units of the structure



wherein R_a is the alkylene portion of the dicarboxylic acid and is a branched alkylene of 34 to 36 carbon atoms, optionally including up to 1 heteroatom per 3 or 10 carbon atoms of the diacid;

wherein R_b is a direct bond or a linear or branched (optionally being or including cyclic, heterocyclic, or aromatic portion(s)) alkylene group (optionally containing up to 1 or 3 heteroatoms per 10 carbon atoms) of 2 to 60 carbon atoms; and

wherein R_c and R_d are individually hydrogen, a linear or branched alkyl group of 1 to 8 carbon atoms, or R_c and R_d connect together to form a single linear or branched alkylene group of 1 to 8 carbon atoms or optionally with one of R_c and R_d is connected to R_b at a carbon atom.

9. The telechelic polyamide of any one of claims 1 to 8, wherein the polyamide segment has two carboxyl end groups and is further reacted with a polyether molecule having one

terminal amine and one terminal hydroxyl group or two terminal hydroxyl groups to provide endblocks for the telechelic polyamide, the telechelic polyamide after adding the endblocks having at least 80 mole % terminal primary or secondary hydroxyl end groups.

5 10. The telechelic polyamide of any one of claims 1 to 9, having functional end groups, wherein at least 80 mole % of the functional end groups are primary amine groups.

11. The telechelic polyamide of any one of claims 1 to 10, wherein the telechelic polyamide has a weight average molecular weight from 200 to 10,000 g/mole, optionally from 400 to
10 10,000 g/mole.

12. The telechelic polyamide of any one of claims 1 to 11, wherein the telechelic polyamide has a weight average molecular weight from 200 to 5,000 g/mole, optionally from 400 to 5,000 g/mole, further optionally from 500 to 5,000 g/mole.

15 13. The telechelic polyamide of either claim 11 or claim 12, wherein the telechelic polyamide without solvents has a viscosity of less than 100,000 cps at 70°C as measured by a Brookfield circular disc viscometer with the circular disc spinning at 5 rpm.

20 14. The telechelic polyamide of either claim 11 or claim 12, wherein the telechelic polyamide without solvents has a viscosity of less than 100,000 cps at 60°C as measured by a Brookfield circular disc viscometer with the circular disc spinning at 5 rpm.

25 15. The telechelic polyamide of any one of claims 1 to 14, wherein the telechelic polyamide further comprises at least one oligomer segment selected from the group of a polyester segment, a polyether segment, and a polycarbonate segment.

30 16. The telechelic polyamide of any one of claims 1 to 8, wherein the polyamide segment has two carboxyl end groups and is further reacted with an amino alcohol having 3 to 16 carbon atoms and having one secondary amine group and one hydroxyl group to provide hydroxyl terminal groups for the telechelic polyamide.