A polyimide precursor solid residuum is an admixture of an aromatic dianhydride or derivative thereof and an aromatic diamine or derivative thereof plus a complexing agent, which is complexed with the admixture by hydrogen bonding. The polyimide precursor solid residuum is effectively employed in the preparation of polyimide foam and the fabrication of polyimide foam structures.
Diamine + Solvent → Polyamic Acid

Solvent = Polar Aprotic Materials

Heat

Water By-Product

Polyimide

FIG. 1

Diamine + Dianhydride

Ball Mill

Monomer Mixture

Heat

Water By-Product

Polyimide Foam

FIG. 2
Dianhydride + Alcohol → Dialkylester-Diacid

Alkyl Alcohol

Polyimide Foam

Water & Alcohol

By-Product

Meta-Stable Mixture at Ambient Conditions

FIG 3

Dianhydride + Alcohol → Dialkylester-Diacid complexed with ether via hydrogen bonding

Alkyl Alcohol

Ether

Polyimide Foam

Ether, Water, & Alcohol By-Products

Stable Mixture at Ambient Conditions (Stabilized by the Ether)

FIG 5
R\_O\_R

ETHER

ETHER COMPLEX

FIG. 4
POLYIMIDE PRECURSOR SOLID RESIDUUM

CROSS REFERENCE TO RELATED APPLICATION

This application claims the benefit of U.S. provisional application No. 60/087,272, filed May 29, 1998.

ORIGIN OF THE INVENTION

The invention described herein was jointly made by employees of the United States Government and employees of Unitka Ltd. In the performance of work under NASA MOA SAA #385.

BACKGROUND OF THE INVENTION

1. Field of the Invention

The present invention relates generally to polyimides. It relates in particular to a polyimide precursor solid residuum, which is effectively employed in the preparation of polyimide foam and the fabrication of polyimide foam structures.

2. Description of the Related Art

High performance polyimides are presently used in the aerospace industry, for example, in joining metals to metals, or metals to composite structures. In addition, polyimides are rapidly finding new uses as foam insulation in cryogenic applications, and as structural foam having increased structural stiffness without large weight increases, in aerospace structures.

Polyimide foam materials have a number of beneficial attributes for next generation space vehicles, such as high temperature and solvent resistance, flame resistance, low smoke generation, high modulus and chemical and hot water resistance. Another area for polyimide foams is in the manufacture of low density insulation for thermal and acoustic applications, and reinforcement for the maritime industry.

U.S. Pat. Nos. 5,147,966 and 5,478,916 disclose polyimides that can be melt processed into various useful forms such as coatings, adhesives, composite matrix resins and films. These polyimides are prepared from various diamines and dianhydrides in various solvents. The use of monoanhydrides as endcapping agents is also disclosed in these patents to control the molecular weight of the polymers and, in turn, to make them easier to process in molten form. The use of ethers to make polyimide adhesives was disclosed in U.S. Pat. No. 4,065,345, which demonstrates another method to produce polyimide resin systems. FIG. 1 shows the method employed by these patents to produce polyimides.

U.S. Pat. No. 3,483,144 discloses a process for making polyimide foam by ball milling a mixture of monomers and heating the mixture to 300° C. In all cases, the foams produced by this patent are the result of dianhydrides or tetracids being dissolved by a diamine upon melting. The ensuing reaction produces water and thus foams the molten material. FIG. 2 illustrates the process to make foam by this patent.

The state-of-the-art technology for making polyimide foams as disclosed in U.S. Pat. Nos. 5,298,531, 5,122,546, 5,077,318, and 4,900,761 utilizes solutions of diamines and dianhydride derivatives in a low molecular weight alkyl alcohol solvent. Polyimide precursor solutions and powders therefrom are then processed into foams through the expulsion of water and alcohol (R-OH) during the thermal imidization process. In these cases the alcohol solvent reacts initially with the dianhydride to form a covalently bonded specie referred to as a dialkylester-diacid (DADA) before the aromatic diamine is added. The aforementioned patents also illustrate the use of blowing agents to aid in the foaming process. The blowing agents utilized by these patents serve as a separate entity and usually result in a foam that has residual blowing agent within its cell walls. FIG. 3 demonstrates the state-of-the-art in this foam technology.

Howsoever useful, these related art processes for preparing foams from polyimide precursors are all found wanting, in that none provide foam densities over a wide range. Indeed, many commercially-available polyimido foams are made by employing an added step of densifying a low density foam to produce a foam of desired higher density. Moreover, in many of these related art processes undesirably high foaming temperatures are required, because the foaming agents are volatile by-products generated in the polycondensation reaction. Furthermore, many of these related art processes do not provide for uniformity in the blowing of the foam, and they result in non-uniformity of the foamed product. In particular, many of these foaming processes start out slowly, build tip to a maximum, and then taper off toward the end. The foamed products produced often exhibit non-uniformity of density and cell size throughout the foamed structure, which results in low yields of acceptable product. Additionally, some of the polyimide precursors employed in these related art processes (especially those involving isocyanate chemistry) have poor hydrolytic stability and poor stability toward other types of decomposition.

SUMMARY OF THE INVENTION

It is accordingly a primary object of the present invention to provide what is lacking in the related art, viz., a polyimide precursor solid residuum which possesses hydrolytic stability and storage stability toward other types of decomposition, and which affords a uniform foaming action, especially in respect of the length and intensity of the blowing process. It is another primary object of the present invention to provide a polyimide precursor solid residuum which affords the preparation of polyimide foams at moderate temperatures to effect foam products having a wide range of densities and a high degree of product uniformity, especially in respect of uniformity of density and cell size throughout the foamed structure. It is another primary object of the present invention to provide a polyimide precursor solid residuum which affords an efficient and efficacious foaming process, and which provides a polyimide foam having high temperature and solvent resistance, flame resistance, low smoke generation, high modulus, low density, and chemical and hot water resistance.

These objects and their attending benefits are achieved, and the disadvantages of the related art are avoided, by the present invention, which is a polyimide precursor solid residuum. This residuum is an admixture of an aromatic compound (A), which is an aromatic dianhydride or derivative thereof and aromatic compound (B), which is an aromatic diamine or derivative thereof, plus a complexing agent (C), which is complexed with the admixture by hydrogen bonding, the complexing agent (C) being present in an amount sufficient to provide from about 1 to about 15% by weight, based on the total weight of the polyimide precursor solid residuum.
Especially beneficial results are obtained if the polyimide precursor solid residuum according to the present invention is an admixture of an aromatic compound (A) and an aromatic compound (B).

\[
\begin{align*}
R_1\text{OOCC}H_2\text{N}-&\text{R}_3\text{C}(\text{O})\text{COOH} + \text{R}_1\text{HNOC}(\text{O})\text{COOH} \\
\text{HOC}^\text{CONH}_2\text{COOH} &\text{HOC}^\text{CONH}_2\text{COOH}
\end{align*}
\]

wherein \( n \) is an integer from 0 to 3, and \( R_1 \) is hydrogen or an alkyl group, and \( R_3 \) is a tetravalent aromatic radical having 1 to 5 benzenoid-unsaturated rings of 6 carbon atoms, \( R_1 \) is a divalent aromatic radical having 1 to 5 benzenoid-unsaturated rings of 6 carbon atoms, along with a complexing agent (C) being present in an amount sufficient to provide from about 1 to about 15% by weight, based on the total weight of the polyimide precursor solid residuum.

Outstanding results are obtained if the aromatic compound (A) and the aromatic compound (B) as specified above are present in the admixture in approximately equimolar amounts.

Especially good results are obtained if the complexing agent is an ether, especially one of, or a suitable mixture of tetrahydrofuran, glyme, diglyme, dioxane, 2-methoxy ethanol and 2-ethoxy ethanol.

The polyimide precursor solid residuum of the present invention advantageously has an apparent density of from about 1 to about 40 pounds/ft\(^3\), and is beneficially in the form of a powder, or friable balloons (as hereinafter described), or flake (as hereinafter described).

Very beneficial results are obtained if the aromatic compound (A) is at least one derivative of a dihydridene selected from ODPA; BPDA; BTDA; DSDA; PMDA and BPADA; and the aromatic compound (B) is at least one diamin which is selected from 3,4'ODA; 4,4'ODA; APB; BPB; m-PDA; p-PDA; 3,3'DDS; 4,4' DDS; 4,4 BAPS; 4,3 BAPS; 3 BAPB; 4 BAPB and BAPP.

Equally beneficial results are obtained if the polyimide precursor solid residuum is a combination of two or more different admixtures of aromatic compound (A) and aromatic compound (B) along with complexing agent (C), the aromatic compound (A) being at least one derivative of a dihydridene selected from ODPA; BPDA; BTDA; DSDA; PMDA and BPADA; and the aromatic compound (B) being at least one diamin which is selected from the group consisting of 3,4'ODA; 4,4'ODA; APB; BPB; m-PDA; p-PDA; 3,3'DDS; 4,4' DDS; 4,4 BAPS; 4,3 BAPS; 3 BAPB; 4 BAPB and BAPP.

The present invention also comprehends a process for producing the instant polyimide precursor solid residuum. This process includes the sequential steps of:

(a) dissolving an admixture of an aromatic compound (A), which is an aromatic dihydridene or derivative thereof, and an aromatic compound (B), which is an aromatic diamine or derivative thereof, in a solvent containing a complexing agent (C), which is capable of complexing with the admixture by hydrogen bonding, to form a solution having a solid content of at least 20% by weight, based on the total weight of the solution;

(b) condensing the solution by heating to effect a condensed solution having a solid content of up to about 85% by weight, based on the total weight of the condensed solution;

(c) cooling the condensed solution to effect solidification thereof;

(d) crushing the solidified condensed solution; and

(e) heating the crushed solidified condensed solution at a temperature between about 70°C. and about 140°C. to form a polyimide precursor solid residuum of an apparent density from about 1 to about 40 pounds/ft\(^3\) and a residual content of complexing agent (C) between about 1 and 15% by weight, based on the total weight of the polyimide precursor solid residuum.

Especially beneficial results are obtained if aromatic compound (A) of the admixture is

\[
\begin{align*}
R_1\text{OOCC}H_2\text{N}-&\text{R}_3\text{C}(\text{O})\text{COOH} + \text{R}_1\text{HNOC}(\text{O})\text{COOH} \\
\text{HOC}^\text{CONH}_2\text{COOH} &\text{HOC}^\text{CONH}_2\text{COOH}
\end{align*}
\]

and an aromatic compound (B)

\[
\begin{align*}
H_2N-R_1\text{HNCO}R_2\text{CONHR} &\text{R}_3\text{NH}_2
\end{align*}
\]

wherein \( n \) is an integer from 0 to 3, and \( R_1 \) is hydrogen or an alkyl group, and \( R_3 \) is a tetravalent aromatic radical having 1 to 5 benzenoid-unsaturated rings of 6 carbon atoms, \( R_1 \) is a divalent aromatic radical having 1 to 5 benzenoid-unsaturated rings of 6 carbon atoms, along with a complexing agent (C) being present in an amount sufficient to provide from about 1 to about 15% by weight, based on the total weight of the polyimide precursor solid residuum.

Outstanding results are obtained if the aromatic compound (B) as specified above are present in the admixture in approximately equimolar amounts.

Especially good results are obtained if aromatic compound (A) is an ether, especially one of, or a suitable mixture of tetrahydrofuran, glyme, diglyme, dioxane, 2-methoxy ethanol and 2-ethoxy ethanol.

**BRIEF DESCRIPTION OF THE DRAWINGS**

For a more complete understanding of the present invention, including its primary objects and attending benefits, reference should be made to the Description of the Preferred Embodiments, which is set forth below. This Description should be read together with the accompanying Drawings, wherein:

FIG. 1 is a process schematic representing a standard process for preparing polyimides; FIG. 2 is a process schematic representing the preparation of polyimide foam by ball milling; FIG. 3 is a process schematic representing the state-of-the-art process for making polyimide foams; FIG. 4 is a representation of the formation of an ether complex, according to the present invention; and FIG. 5 is a process schematic representing a process for polyimide foam production using a polyimide precursor solid residuum according to the present invention.
Polyimide precursor solid residuum in any ratio to yield the flake.

The tern polyimide means a thermally-stable organic compound which becomes a polyimide by thermal imidization.

(3) Apparent density

Apparent density is estimated, for example, according to ASTM D-3574A.

(4) Compression strength

Compression strength is estimated, for example, according to ASTM D-3574C.

(5) Glass Transition temperature \( (T_g) \)

This thermal property is determined from the inflection point associated with a change in heat capacity at constant pressure employing Differential Scanning Calorimetry (DSC).

(6) An approximately equimolar mixture

A mixture of two components wherein the molar ratio is within the range of 0.95 to 1.05.

(7) The following abbreviations used herein are understood to mean:

ODPA=4,4'-oxydipthalic anhydride

ODP-TA=4,4'-oxydipthalic acid (a tetraacid)

BPDA=3,3',4,4'-biphenyl tetracarboxylic dianhydride

BTDA=3,3',4,4'-benzophenone tetracarboxylic dianhydride

DSDA=3,3',4,4'-diphenylsulfone tetracarboxylic dianhydride

PMDA=p-phenylenediamine

BPADA=2,2-bis(4-(3,4-carboxyphenyl)) propane dianhydride

3.4ODA=3,4'-oxydianiline

4.4ODA=4,4'-oxianiline

m-PDA=m-phenylenediamine

p-PDA=p-phenylenediamine

3.3DDS=3,3'-dianinodiphenylsulfone

4.4DDS=4,4'-dianinodiphenylsulfone

4,4 BAPS=4,4'-bis(3-aminoxyphenyl) sulfone

4,3 BAPS=4,3'-bis(3-aminoxyphenyl) sulfone

3 BAPS=4,4'-bis(3-aminoxyphenyl) biphenyl

4 BAPS=4,4'-bis(4-aminoxyphenyl) biphenyl

BAPP=2,2-bis(4-(4-aminoxy phenyl) propane

Friable balloons is understood to mean a particular form of the polyimide precursor solid residuum according to the present invention, which has an apparent density of less than about 10 pounds/ft\(^3\) and which preferably has a substantially spherical shape.

Flake means a blend of a polyimide precursor solid residuum according to the present invention with other polymers such as thermoplastic or thermosetting polymers. Thermoplastic polymers include thermoplastic polyimide and polyamide imide. Thermosetting polymers include epoxy, phenolic, and thermosetting polyimide. These other polymers are blended with the polyimide precursor solid residuum in any ratio to yield the flake.

Examples of preferred polyimides comprehend herein include polyimides from ODPA/3,4'ODA (Formula 2), BPADA/3,4'ODA (Formula 3), BTDA/4,4'ODA (Formula 4), BTDA/4,4'ODA (Formula 5), BTDA/3,4'ODA (Formula 6), DSDA/BPP (Formula 7), BTDA/BPP (Formula 8), and BPDA/BPP (Formula 9).
(11) The polyimide comprehended in the present invention may be a copolyimide.

Examples of preferred copolyimides include polyimides of ODPA/3,4′ODA/PDA (90/10 molar ratio) (Formula 10), BPDA/3,4′ODA/APB (85/15 molar ratio) (Formula 11); BTDA/BPB/4,4′ODA (1/1 molar ratio) (Formula 12); and BTDA/PMDA (8/2 molar ratio)/4,4′ODA (Formula 13).
The following Examples are illustrative of the present invention and are not intended to limit its scope.

1. 756 g (2.4 mol) of ODPA was dissolved in THF at room temperature. This solution was treated at 70°C for 6 hours in order to convert the ODPA into ODP-DADA complexed with THF by hydrogen bonding. 488 g (2.4 mol) of 3,4'ODA was added to the resulting solution of ODP-DADA and stirred for 2 hours to yield a homogeneous polyimide precursor solution. The resulting polyimide precursor solution had a solids content of 70 wt% and a viscosity of 20 poise at 20°C. The solution was then charged into a stainless-steel vat and treated at 70°C for 14 hours in order to evaporate off the solvent (THF and MeOH). The resulting material was allowed to cool and crushed into a fine powder (2 to 500 μm). (These powders are sieved using a mesh, if needed.) The polyimide precursor solid residuum was then treated for an additional amount of time (0 to 300 minutes) at 80°C to further reduce residual solvents to around 1-10 wt% depending on the final foam density desired. Residual amounts of THF were determined by measuring proton NMR spectra of the powders.

2. The procedure of Example 1 was repeated except that 480 g of Glyme was used in replacement of THF.

3. The procedure of Example 1 was repeated except that 334 g of EtOH was used in replacement of MeOH.

4. The procedure of Example 1 was repeated except that Dioxane was used in replacement of THF.

5. The procedure of Example 1 was repeated except that BTDA was used in replacement of ODPA.

6. The procedure of Example 1 was repeated except that BPDA was used in replacement of ODP-DADA.

7. The procedure of Example 1 was repeated except that DSDA was used in replacement of ODPA.

8. The procedure of Example 1 was repeated except that PMDA and 3-BAPB was used, in replacement of ODPA and 3,4'ODA, respectively.

9. The procedure of Example 1 was repeated except that BPDA and m-PDA was used, in replacement of ODP and 3,4'ODA, respectively.

10. The procedure of Example 1 was repeated except that a mixture of 3,4'ODA and p-PDA (mol ratio: 90/10) was used in replacement of 3,4'ODA.

11. The procedure of Example 5 was repeated except that 4,4'ODA was used in replacement of 3,4'ODA.

12. The procedure of Example 2 was repeated except that 4,4'DDS and BTDA were used in replacement of 3,4'ODA and ODP-DADA, respectively.

13. The procedure of Example 5 was repeated except that 4,3BAPS was used in replacement of 3,4'ODA.

14. The procedure of Example 6 was repeated except that a mixture of 3,4'ODA and APB (mol ratio: 85/15) was used in replacement of 3,4'ODA.

15. 227 g (1.1 mol) of 3,4'ODA was dissolved in a mixture of 1120 g of THF and 280 g of MeOH at room temperature. To the stirring 3,4'ODA solution, 176 g (0.57 mol) of ODPA was added gradually at 15°C over 40 min. The resulting mixture was stirred for 2 hr at 15°C to yield a homogeneous solution. To this solution, 197 g (0.57 mol) of ODP-DADA formed precipitate. This solution was added gradually and the mixture was stirred for 24 hr at 30°C. To yield a homogeneous precursor solution. Solid contents and viscosity of the resulting solution were 30 wt% and 0.2 poise, respectively. From this solution, polyimide precursor solid residuum was prepared in a manner similar to Example 1.

16. 336 g (0.78 mol) of 4,4 BAPS was dissolved in a mixture of 1120 g of THF and 280 g of MeOH at room temperature. To the stirring BAPS solution, 125 g (0.39 mol) of BTDA was added gradually at 15°C over 40 min. The resulting mixture was stirred for 2 hr at 15°C to yield a homogeneous solution. To this solution, 139 g (0.39 mol) of BTDA-TA (tetra-acid form of BTDA) was added gradually and the mixture was stirred for 24 hr at 30°C to yield a homogeneous precursor solution. Solid content and viscosity of the resulting solution were 30 wt% and 0.2 poise, respectively. From this solution, polyimide precursor solid residuum was prepared in a manner similar to Example 1.
31. Polyimide precursor solid residua obtained by the procedure of the Example 22 were further treated at 110° C. to expand them without thermal imidization so that their apparent densities were decreased without thermal imidization.

32. The procedure of Example 15 was repeated except that ODPA Dimethyl ester was used in replacement of ODPA-TA.

33. Polyimide precursor solid residua obtained by the procedure of the Example 32 were further treated at 100° C. to expand them without thermal imidization so that their apparent densities were decreased without thermal imidization.

34. The procedure of Example 1 was repeated, except that a mixture of BTDA and Na(5-norbornene-2,3-dicarboxylic anhydride) (mol ratio: 2.1/2.0) was used in replacement of ODPA (mol ratio of BTDA/3,4ODA is 0.68).

35. Polyimide precursor solid residua prepared by the process of the above examples were expanded to form foam as follows: 0.5 g of polyimide precursor solid residuum was charged into a standard glass tube (diameter 16 mm/length 180 mm). The glass tube was placed in a circulating air oven set to 140° C., and the material was held up to 15 minutes at this temperature. The hold at 140° C. allowed foaming to occur. Once the hold was completed the test tube was returned to ambient conditions and the height of the resulting foam was measured.

In Table 1, values of the foam height are summarized, as well as properties of the polyimide precursor powder of these examples.

Comparative Examples

The following examples, which do not exemplify the polyimide precursor solid residuum of the present invention, are set forth for comparative purposes, as they are characteristic of the related art.

### Comparative Examples

#### Properties of Polyimide Precursor Solid Residuum

<table>
<thead>
<tr>
<th>Example</th>
<th>PI</th>
<th>Precursor Powder No.</th>
<th>Ether contents (wt %)</th>
<th>Average Particle size (μm)</th>
<th>Apparent Density (pcf)</th>
<th>Expandability (mm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>ODPA/3,4ODA</td>
<td>(A)</td>
<td>2.7</td>
<td>155</td>
<td>43</td>
<td>32</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(B)</td>
<td>3.5</td>
<td>400</td>
<td>37</td>
<td>60</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(C)</td>
<td>3.2</td>
<td>300</td>
<td>37</td>
<td>55</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(D)</td>
<td>2.8</td>
<td>200</td>
<td>42</td>
<td>45</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(E)</td>
<td>1.9</td>
<td>100</td>
<td>36</td>
<td>17</td>
</tr>
<tr>
<td>2</td>
<td>ODPA/3,4ODA</td>
<td>(F)</td>
<td>6.1</td>
<td>140</td>
<td>41</td>
<td>83</td>
</tr>
<tr>
<td>3</td>
<td>ODPA/3,4ODA</td>
<td>(G)</td>
<td>4.1</td>
<td>130</td>
<td>38</td>
<td>85</td>
</tr>
<tr>
<td>4</td>
<td>ODPA/3,4ODA</td>
<td>(H)</td>
<td>5.8</td>
<td>140</td>
<td>40</td>
<td>82</td>
</tr>
<tr>
<td>5</td>
<td>BTDA/3,4ODA</td>
<td>(I)</td>
<td>4.2</td>
<td>150</td>
<td>37</td>
<td>82</td>
</tr>
<tr>
<td>6</td>
<td>BPDA/3,4ODA</td>
<td>(J)</td>
<td>3.9</td>
<td>140</td>
<td>39</td>
<td>80</td>
</tr>
<tr>
<td>7</td>
<td>DSDA/3,4ODA</td>
<td>(K)</td>
<td>4.0</td>
<td>160</td>
<td>40</td>
<td>85</td>
</tr>
<tr>
<td>8</td>
<td>PMDA/3APB</td>
<td>(L)</td>
<td>3.8</td>
<td>150</td>
<td>40</td>
<td>82</td>
</tr>
<tr>
<td>9</td>
<td>BPDA/m-PDA</td>
<td>(M)</td>
<td>4.0</td>
<td>150</td>
<td>41</td>
<td>84</td>
</tr>
<tr>
<td>10</td>
<td>ODPA/3,4ODA</td>
<td>(N)</td>
<td>3.3</td>
<td>120</td>
<td>36</td>
<td>81</td>
</tr>
<tr>
<td>11</td>
<td>BTDA/4,4ODA</td>
<td>(O)</td>
<td>4.0</td>
<td>80</td>
<td>26</td>
<td>95</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(P)</td>
<td>4.3</td>
<td>300</td>
<td>30</td>
<td>105</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(Q)</td>
<td>4.1</td>
<td>200</td>
<td>31</td>
<td>91</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(R)</td>
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<td>100</td>
<td>27</td>
<td>80</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(S)</td>
<td>3.7</td>
<td>50</td>
<td>26</td>
<td>40</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(T)</td>
<td>3.2</td>
<td>50</td>
<td>25</td>
<td>20</td>
</tr>
<tr>
<td>12</td>
<td>BTDA/4,4DOS</td>
<td>(U)</td>
<td>6.4</td>
<td>150</td>
<td>48</td>
<td>84</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(V)</td>
<td>6.7</td>
<td>150</td>
<td>49</td>
<td>100</td>
</tr>
<tr>
<td>13</td>
<td>BTDA/4,3APS</td>
<td>(W)</td>
<td>4.6</td>
<td>150</td>
<td>41</td>
<td>81</td>
</tr>
</tbody>
</table>
We claim:

1. A polyimide precursor solid residuum comprising an admixture of an aromatic compound (A), which is an aromatic dianhydride or a derivative of an aromatic dianhydride, and an aromatic compound (B), which is an aromatic diamine or a derivative of an aromatic diamine, plus a complexing agent (C), which is an ether, the complexing agent (C) being present in an amount sufficient to provide from about 1 to about 15% by weight, based on the total weight of the polyimide precursor solid residuum.

2. A polyimide precursor solid residuum comprising an admixture of an aromatic compound (A)

\[
R_1\text{OOC} - R_2\text{CONH} - R_3\text{HNOC} - R_4\text{COOR}_5
\]

and an aromatic compound (B)

\[
H_2N - R_1\text{HNOC} - R_2\text{CONH} - R_3\text{NH}_2
\]

wherein \( n \) is an integer from 0 to 3, and \( R_1 \) is hydrogen or an alkyl group, and \( R_2 \) is a tetravalent aromatic radical having 1 to 5 benzenoid-unsaturated rings of 6 carbon atoms, \( R_3 \) is a divalent aromatic radical having 1 to 5 benzenoid-unsaturated rings of 6 carbon atoms; plus a complexing agent (C), which is an ether, the complexing agent being present in an amount sufficient to provide from about 1 to about 15% by weight, based on the total weight of the polyimide precursor solid residuum.

3. A polyimide precursor solid residuum comprising an approximately equimolar admixture of an aromatic compound (A)

\[
R_1\text{OOC} - R_2\text{CONH} - R_3\text{HNOC} - R_4\text{COOR}_5
\]

and an aromatic compound (B)

\[
H_2N - R_1\text{HNOC} - R_2\text{CONH} - R_3\text{NH}_2
\]

wherein \( n \) is an integer from 0 to 3, and \( R_1 \) is hydrogen or an alkyl group, and \( R_2 \) is a tetravalent aromatic radical having 1 to 5 benzenoid-unsaturated rings of 6 carbon atoms.
carbon atoms, \( R_i \) is a divalent aromatic radical having 1 to 5 benzenoid-unsaturated rings of 6 carbon atoms; plus a complexing agent (C), which is ether, the complexing agent being present in an amount sufficient to provide from about 1 to about 15\% by weight, based on the total weight of the polyimide precursor solid residuum.

4. A polyimide precursor solid residuum according to claim 3, wherein the ether is selected from the group consisting of tetrahydrofuran, glyme, diglyme, dioxane, 2-methoxy ethanol and 2-ethoxy ethanol.

5. A polyimide precursor solid residuum according to claim 3, wherein the aromatic compound (A) is at least one derivative of a diamine or derivative thereof, in a solvent containing a complexing agent being present in an amount sufficient to provide from about 1 to about 15\% by weight, based on the total weight of the solution; and the aromatic compound (B) is at least one derivative of a diamine which is a member selected from the group consisting of 3,4' ODA; 4,4' ODA; APB; p-PDA; 4,4 BAPS; 4,3 BAPS; 3 BAPB; and the aromatic compound (B) is at least one derivative of a dianhydride selected from the group consisting of ODPA; BPDA; BTDA; DSDA; PMDA and BPADA; and the aromatic compound (B) is at least one derivative of an aromatic compound which is a member selected from the group consisting of 3,4' ODA; 4,4' ODA; APB; p-PDA; 3,3' DDS; 4,4' DDS; 4,4 BAPS; 4,3 BAPS; 3 BAPB; 4 BAPB and BAPP.

6. A polyimide precursor solid residuum according to claim 3, which is a combination of two or more different admixtures of aromatic compound (A) and aromatic compound (B) along with complexing agent (C), the aromatic compound (A) being at least one derivative of a diamine or derivative thereof, in a solvent containing a complexing agent (C), which is complexed with the admixture by hydrogen bonding, to form a solution having a solid content of at least 20\% by weight, based on the total weight of the solution; and an aromatic compound (B) is at least one derivative of a diamine which is a member selected from the group consisting of 3,4' ODA; 4,4' ODA; APB; BPDA; p-PDA; 3,3' DDS; 4,4' DDS; 4,4 BAPS; 4,3 BAPS; 3 BAPB; 4 BAPB and BAPP.

7. A polyimide precursor solid residuum according to claim 3, which has a density of from about 1 to about 10 pounds/ft\(^3\).

8. A process for producing the polyimide precursor solid residuum of claim 1, which process comprises:

(a) dissolving an admixture of an aromatic compound (A), which is an aromatic diamine or derivative thereof, and an aromatic compound (B), which is an aromatic diamine or derivative thereof, in a solvent containing a complexing agent (C), which is complexed with the admixture by hydrogen bonding, to form a solution having a solid content of at least 20\% by weight, based on the total weight of the solution;

(b) cooling the condensed solution to effect solidification thereof;

(c) dissolving an approximately equimolar admixture of an aromatic compound (A)

R\(_1\)OOC--R\(_2\)--CONH--R\(_3\)--HNOC--R\(_4\)--COOR\(_5\)

and an aromatic compound (B)

H\(_2\)N--R\(_6\)--CO--R\(_7\)--CONH--R\(_8\)--NH--R\(_9\),

wherein \( n \) is an integer from 0 to 3, and \( R_i \) is hydrogen or an alkyl group, and \( R_{1-9} \) is a tetravalent aromatic radical having 1 to 5 benzenoid-unsaturated rings of 6 carbon atoms, \( R_i \) is a divalent aromatic radical having 1 to 5 benzenoid-unsaturated rings of 6 carbon atoms, \( n \) is a divalent aromatic radical having 1 to 5 benzenoid-unsaturated rings of 6 carbon atoms, and the aromatic compound (B) is at least one derivative of a diamine or derivative thereof, in a solvent containing a complexing agent (C), which is complexed with the admixture by hydrogen bonding, to form a solution having a solid content of at least 20\% by weight, based on the total weight of the solution; and (e) heating the crushed solidified condensed solution at a temperature between about 70\°C and about 140\°C.

9. A process for producing a polyimide precursor solid residuum of claim 3, which process comprises:

(a) dissolving an approximately equimolar admixture of an aromatic compound (A)

R\(_1\)OOC--R\(_2\)--CONH--R\(_3\)--HNOC--R\(_4\)--COOR\(_5\)

and an aromatic compound (B)

H\(_2\)N--R\(_6\)--CO--R\(_7\)--CONH--R\(_8\)--NH--R\(_9\),

wherein \( n \) is an integer from 0 to 3, and \( R_i \) is hydrogen or an alkyl group, and \( R_{1-9} \) is a tetravalent aromatic radical having 1 to 5 benzenoid-unsaturated rings of 6 carbon atoms, \( R_i \) is a divalent aromatic radical having 1 to 5 benzenoid-unsaturated rings of 6 carbon atoms, and the aromatic compound (B) is at least one derivative of a diamine or derivative thereof, in a solvent containing a complexing agent (C), which is complexed with the admixture by hydrogen bonding, to form a solution having a solid content of at least 20\% by weight, based on the total weight of the solution; and (e) heating the crushed solidified condensed solution at a temperature between about 70\°C and about 140\°C.

10. A process for producing a polyimide precursor solid residuum of claim 3, which process comprises:

(a) dissolving an approximately equimolar admixture of an aromatic compound (A)

R\(_1\)OOC--R\(_2\)--CONH--R\(_3\)--HNOC--R\(_4\)--COOR\(_5\)

and an aromatic compound (B)

H\(_2\)N--R\(_6\)--CO--R\(_7\)--CONH--R\(_8\)--NH--R\(_9\),

wherein \( n \) is an integer from 0 to 3, and \( R_i \) is hydrogen or an alkyl group, and \( R_{1-9} \) is a tetravalent aromatic radical having 1 to 5 benzenoid-unsaturated rings of 6 carbon atoms, \( R_i \) is a divalent aromatic radical having 1 to 5 benzenoid-unsaturated rings of 6 carbon atoms, and the aromatic compound (B) is at least one derivative of a diamine or derivative thereof, in a solvent containing a complexing agent (C), which is complexed with the admixture by hydrogen bonding, to form a solution having a solid content of at least 20\% by weight, based on the total weight of the solution; and (e) heating the crushed solidified condensed solution at a temperature between about 70\°C and about 140\°C.

11. The process of claim 10, wherein the complexing agent (C) is ether.

12. The process of claim 11, wherein the ether is selected from the group consisting of tetrahydrofuran, glyme, diglyme, dioxane, 2-methoxy ethanol and 2-ethoxy ethanol.

13. A polyimide precursor solid residuum according to claim 3, which is in the form of a member selected from the group consisting of powder, friable balloons, and flake.

14. A polyimide precursor solid residuum according to claim 13, which is in the form of friable balloons and has a density of from about 1 to about 10 pounds/ft\(^3\).