TO: NIT-44/Scientific and Technical Information Division
   Attn: Shirley Peigare

FROM: GP-4/Office of Assistant General Counsel
      for Patent Matters

SUBJECT: Announcement of NASA-Owned U.S. Patents in STAR

In accordance with the procedures agreed upon by Code GP-4 and
Code NST-44, the attached NASA-owned U.S. Patent is being
forwarded for abstracting and announcement in NASA STAR.

The following information is provided:

U.S. Patent No. : 4,450,268 Issued: 5/22/84

Government or Contractor Employee:

Pursuant to Section 305(a) of the National Aeronautics and Space
Act, the name of the Administrator of NASA appears on the first
page of the patent; however, the name of the actual inventor
(author) appears at the heading of Column No. 1 of the
specification, following the words "...with respect to an
invention of...."
The invention relates to metal, primarily divalent metal such as divalent copper, cobalt and nickel, 4,4',4'', 4'' phthalocyanine tetracarboxylic acids and to polymers prepared by heating the same, and to a method of preparing the tetracarboxylic acids.

A method is provided for preparing the aforesaid tetracarboxylic acids in purer form than heretofore. Upon heating these acids decarboxylate and produce polymers.

These polymers have diphenyl (conjugated) types of linkage; they are thermally and oxidatively stable and useful as sheet polymers for various purposes such as heat shielding.

The monomeric tetracarboxylic acid is shown in FIG. 1, the polymer in FIG. 2 and thermal resistance in FIG. 3.

The monomeric tetracarboxylic acids are prepared in purer form than heretofore, the resulting polymers are more stable thermally and oxidatively than metal phthalocyanine polymers and monomers heretofore made.

12 Claims, 3 Drawing Figures
$M = \text{Cu, Co, Ni, ...}$

FIG. 2
FIG. 3

TGA & DTG CURVES OF THE SHEET POLYMERS OF COPPER PHTHALOCYANINE COBALT PHTHALOCYANINE NICKEL PHTHALOCYANINE

RATE OF WEIGHT LOSS (mg/min

0 0.9 0.8 0.7 0.6 0.5 0.4 0.3 0.2 0.1

0 1000 900 800 700 600 500 400 300 200

0 10 20 30 40 50 60 70 80 90

WEIGHT LOSS %

TEMPERATURE, °C
METAL PHTHALOCYANINE POLYMERS

DESCRIPTION

ORIGIN OF THE INVENTION

The invention described herein was made in the performance of work under a NASA contract and is subject to the provisions of Section 305 of the National Aeronautics and Space Act of 1958, Public Law 85-568 (72 Stat. 435; 42 U.S.C. 2457).

FIELD OF THE INVENTION

This invention relates to metal phthalocyanine tetracarboxylic acids, to a novel method of synthesizing the same and to polymers of the same.

BACKGROUND OF THE INVENTION

In recent years there has been an increasing demand in industry and defense for polymeric substances which would either remain intact or continue to serve in a more or less degraded state under conditions where temperatures far above the ordinary are encountered. In many cases highly cross-linked organic polymers have been found promising. The phthalocyanine structure is one of the most thermally stable organic structures known. Attractive properties like resistance to chemical attack, electrical properties, catalytic activity and moderate cost of manufacture with good coloring properties have made phthalocyanines the object of intensive world-wide investigations. Many attempts to synthesize polymers based on phthalocyanines have failed to provide the expected thermal stability. Phthalocyanine polymers so far produced have shown less thermal stability than the phthalocyanine monomer itself because of the presence of impurities, weak chemical bonds and low degree of polymerization with structural inhomogeneity. Impurities have the considerable effect of decreasing thermal stability of the phthalocyanine monomers as well as the polymers produced from them. It is well known that the way in which the repeating units are linked is reflected in the properties of the polymers. If the repeating phthalocyanine mer units are linked in the way phenyl groups are linked in biphenyl, the conjugation extends throughout the macromolecule thereby increasing the extend of delocalization of the \( \pi \)-electrons. This is expected to increase the conductivity as well as the thermal stability of the phthalocyanine polymers.

OBJECTS OF THE INVENTION

Objects of the invention include methods of preparing metal phthalocyanine monomers in a pure state and the preparation from such monomers of thermally stable sheet polymers which have biphenyl type linkages between the mer units, high thermal and thermal oxidative stability and electrical conductivity, also good flame and fire resistance.

BRIEF DESCRIPTION OF THE INVENTION

A metal phthalocyanine tetracarboxylic acid (MPTC) is prepared in pure form by reaction of trimellitic anhydride with urea and a soluble metal salt in the presence of a suitable catalyst and in a suitable solvent. The metal is that which it is desired to introduce into the intended metal complex. The resulting MPTC, after separation and purification, is polymerized by heat in a vacuum or in an atmosphere of an inert gas such as nitrogen. The MPTC prepared by this method is purer than MPTC's prepared heretofore and the polymer derived from it is free of impurities which detract from its useful properties.

DESCRIPTION OF THE DRAWINGS

FIG. 1 is a structural formula of the metal 4,4',4",4"'-tetracarboxy phthalocyanine which is to be polymerized.

FIG. 2 is a structural formula of the resulting polymer.

FIG. 3 is a graph showing TGA and DTG curves for these polymers, including the polymer of FIG. 2.

DETAILED DESCRIPTION OF THE INVENTION

EXAMPLE 1

Preparation of Copper (II) phthalocyanine tetracarboxylic acid 1

This compound has the formula

\[
\text{HOOC} \quad \text{N} \quad \text{M} \quad \text{N} \quad \text{COOH}
\]

wherein M is copper (II).

It was prepared as follows: 12.0 g Copper (II) sulfate pentahydrate, 33.8 g trimellitic anhydride, 4.5 g ammonium chloride, 0.5 g ammonium molybdate and 60 g urea were finely ground together and placed in a 500 ml three-necked flask containing 25 ml of nitrobenzene. The flask was provided with a thermometer, condenser and a mechanical stirrer. The temperature of the flask was slowly increased to 180° C. and maintained at 185° C. for four hours. The color of the reaction mixture gradually deepened and finally a deep colored solid was obtained. The product was ground well and washed with methanol until it was free from nitrobenzene. The solid product was added to 500 ml of 1.0N hydrochloric acid saturated with sodium chloride, boiled briefly, cooled to room temperature and filtered. The resulting solid material was treated with 500 ml of 2.0N sodium hydroxide containing 200 g sodium chloride and heated at 90° C. until the evolution of ammonia ceased. The solution after filtration was treated with 500 ml 2.0N hydrochloric acid and the product was separated by centrifugation. The residue was redissolved in 0.1N sodium hydroxide and filtered to separate the insoluble materials. The compound was re-precipitated with 1.0N...
hydrochloric acid and centrifuged to obtain the solid material. Dissolution and precipitation steps were repeated twice. Then the compound was washed until chloride free and finally washed with methanol. The blue product was dried at 100° in vacuum.

EXAMPLES 2 AND 3

The cobalt and nickel analogues of 1 were prepared by the same method using in Example 2 cobalt sulfate and in Example 3 nickel sulfate. The empirical formulae and analytical results obtained were as follows:

EXAMPLE 2 (Cobalt)

Compound C₃₂H₁₆N₈O₇(Cu), (CuPTC): calcd: C, 57.5; H, 2.14; N, 14.9; Cu, 8.45; Eq. wt. 188.0, Found: C, 57.1; H, 2.3; N, 15.0; Cu, 8.5; Eq. wt. 187.6. IR absorption bands (cm⁻¹): 3500-2500 (broad), 1691 (broad), 1614m, 1578s, 1536s, 1508m, 1329m, 1279m, 1246m (broad), 1188m, 1149m, 1089s, 1050w, 968s, 940w, 851w, 785w, 774w, 736s.

EXAMPLE 3 (Nickel)

Compound C₃₂H₁₆N₈O₇Ni, (NiPTC): calcd: C, 57.8; H, 2.15; N, 15.0; Ni, 7.88; Eq. wt. 186.9, found: C, 57.5; H, 2.2; N, 15.2; Co, 7.90; Eq. wt. 186.7. IR absorption bands (cm⁻¹): 3500-2500 (broad), 1696 (broad), 1613m, 1586m, 1521m, 1330m, 1281m, 1246m (broad), 1189m, 1149m, 1090s, 1050w, 9730w, 944w, 848w, 782w, 773w, 742s.

Polymersization of MPTC—Method 1

About 1-2 g of metal phthalocyanine tetracarboxylic acid was finely ground in a small vibrating ball mill and placed in a polymerization tube. The tube was connected to an apparatus provided with a tube to condense volatile products, stopcocks to connect gas collection tube and IR cell for gaseous analysis. The apparatus was connected to vacuum system. Reaction tube was carefully evacuated to 10⁻⁵-10⁻⁶ torr pressure and heated to 450°C. The gaseous and volatile products were condensed in separate traps using liquid nitrogen. The reaction was found to be completed after one hour of heating at 450°C. in vacuum.

EXAMPLE 5

Polymersization of MPTC—Method 2

Finely ground metal phthalocyanine tetracarboxylic acid was taken in a reaction tube fitted with a glass enclosed iron-constantan thermocouple and inlet and outlet tubes. The polymerization tube was carefully purged with nitrogen by repeated evacuation and refilling. Then it was gradually heated to 450°C. in a current of nitrogen and maintained temperature of 450°-500°C. in a current of nitrogen for one hour.

Methods 1 and 2 were applied to produce the divalent copper, cobalt and nickel compounds. Analytical results were the same for products of the two methods and were as follows:

(1) Compound C₂₂H₁₅₂₅NaCu: calcd: C, 67.18; H, 2.11; N, 19.58; Cu, 11.2, found: C, 66.9; H, 2.21; N, 19.8; Cu, 11.5. IR absorption bands (cm⁻¹): 1610w, 1503w, 1407w, 1328w, 1271m, 1164m, 1118w, 1089m, 1065m, 898w, 779w, 754w, 738s.

(2) Compound C₂₂H₁₅₂₅Co: calcd: C, 67.73; H, 2.13; N, 19.75; Co, 10.39, found: C, 67.6; H, 2.3; N, 19.95; Co, 10.46. IR absorption bands (cm⁻¹): 1599w, 1502w,
TABLE I

<table>
<thead>
<tr>
<th>POLYMER</th>
<th>PDTo</th>
<th>CHAR YIELD (800°C)</th>
<th>CONDUCTIVITY (22–300°C)</th>
</tr>
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<tbody>
<tr>
<td>CuPC-Sheet</td>
<td>760</td>
<td>90.5%</td>
<td>3.2 × 10⁻⁴–3.8 × 10⁻⁵</td>
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<td>Polymer</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>CoPC-Sheet</td>
<td>860</td>
<td>89.0%</td>
<td>1.8 × 10⁻⁵–2.3 × 10⁻⁵</td>
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<td></td>
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</tr>
<tr>
<td>NiPC-Sheet</td>
<td>890</td>
<td>93.0%</td>
<td>1.4 × 10⁻⁵–3.9 × 10⁻⁵</td>
</tr>
<tr>
<td>Polymer</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

(3) Compound C₂₂H₁₂Ni₄: calcd: C, 67.76; H, 2.13; N, 19.75; Ni, 10.35, found: C, 67.8; H, 2.18; N, 19.92; Ni, 10.43. IR absorption bands (cm⁻¹): 1602w, 1519w, 1514w, 1334m, 1287m, 1157m, 1115w, 1086m, 1053w, 944w, 914w, 782w, 745w, 743m.

The polymerization reaction proceeds by way of the elimination of carboxyl groups and the formation of a sheet polymer in which the mer units are linked by biphenyl linkage. The formula of the polymer may be expressed as follows:

In FIG. 3 of the drawing the results plotted are for Cu, Co and Ni sheet polymers heated in an atmosphere of nitrogen. The ordinate scale on the left represents percent decomposition and the ordinate scale on the right represents rate of weight loss. As will be seen the polymers did not undergo substantial decomposition below 750°C to 800°C. (See curves at top.) The rate of decomposition (lower set of curves) did not become substantial until about 750° to 900°C.

It will be apparent that purer metal phthalocyanine tetracarboxylic acids are provided, that a new and useful method of preparing them has been provided and that new and useful polymers and methods of producing them have been provided.

It is claimed:
1. A method of preparing a heat stable and oxidation resistant sheet polymer of a metal phthalocyanine which comprises heating a metal 4,4',4''-tetracarboxylic phthalocyanine to a temperature sufficient to result in the formation of gaseous products and to form a sheet polymer having the repeating structure...
wherein M is a metal.

2. The method of claim 1 wherein the heating is carried out under vacuum.

3. The method of claim 1 wherein the heating is carried out in an atmosphere of an inert gas.

4. The method of claim 1 wherein the metal M is a divalent metal having an atomic radius of approximately 1.35 Å.

5. The method of claim 4 wherein M is Cu.

6. The method of claim 4 wherein M is Co.

7. The method of claim 4 wherein M is Ni.

8. A heat stable oxidation resistant sheet polymer having the repeating structure

9. The polymer of claim 8 wherein the metal M is a divalent metal having an atomic radius of approximately 1.35 Å.

10. The polymer of claim 9 wherein M is Cu.

11. The polymer of claim 9 wherein M is Co.

12. The polymer of claim 9 wherein M is Ni.

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