

United States Patent [19]

Gibson et al.

[11] Patent Number: 5,021,583

[45] Date of Patent: Jun. 4, 1991

[54] FORMATION OF REISSERT COMPOUND OF BISBENZIMIDAZOLE

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[21] Appl. No.: 586,463

[22] Filed: Sep. 21, 1990

Related U.S. Application Data

[62] Division of Ser. No. 418,319, Oct. 6, 1989, Pat. No. 4,977,271.

[51] Int. Cl.⁵ C07D 403/06

[52] U.S. Cl. 548/328

[58] Field of Search 548/328

[56] References Cited PUBLICATIONS

Popp, J. Heterocyclic Chem., vol. 24, pp. 1349 to 1351, (1987).

Uff et al.; J. Chem. Soc., (London), Chem. Commun., vol. of 1984, pp. 1245 and 1246.

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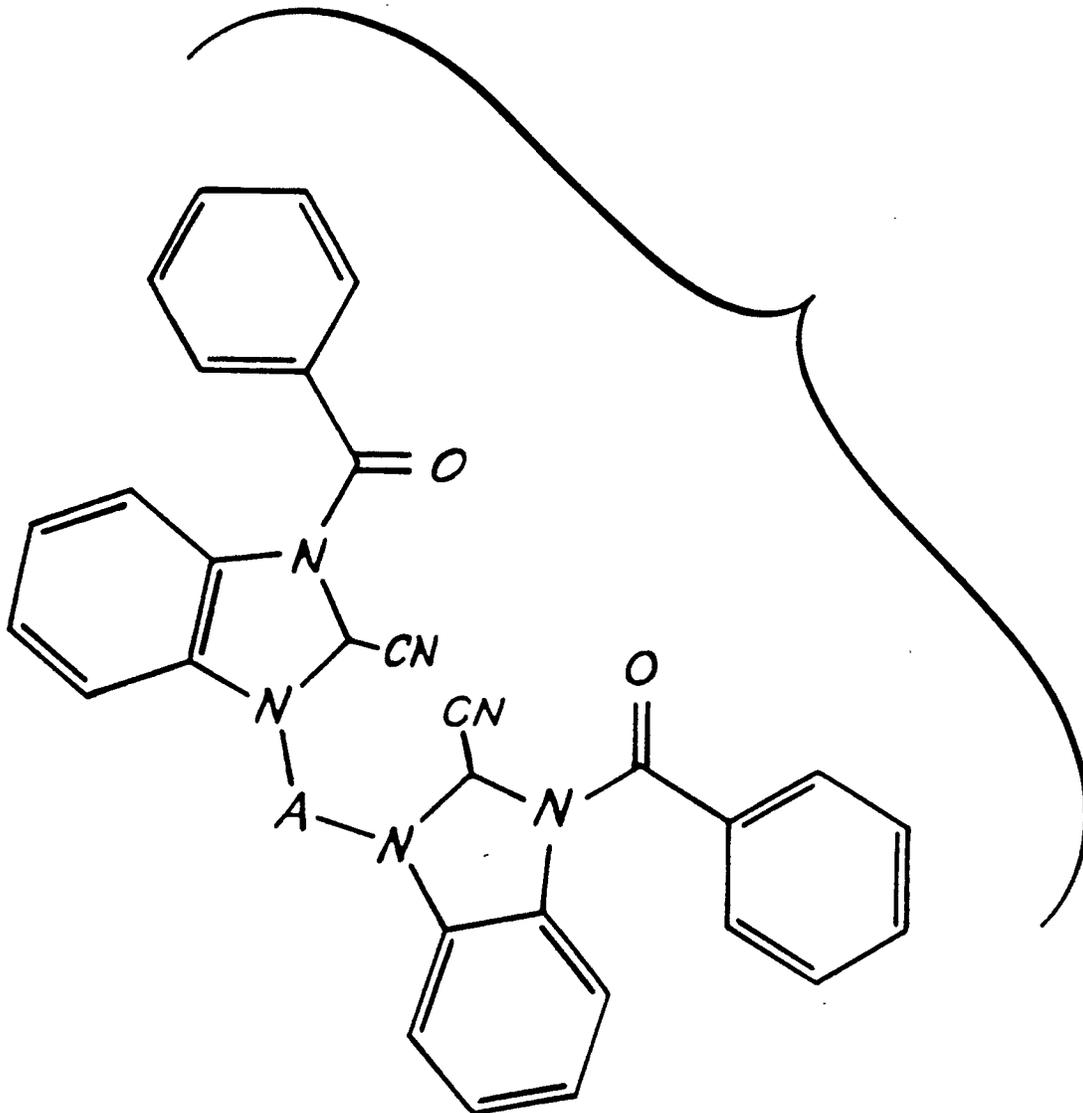
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[57] ABSTRACT

A Reissert compound of bisbenzimidazole can be formed by first reacting benzimidazole with an aliphatic diacid chloride to form bisbenzimidazole and then reacting the bisbenzimidazole with an aliphatic acid chloride and cyanide to form the Reissert compound thereof.

5 Claims, 1 Drawing Sheet



FORMATION OF REISSERT COMPOUND OF BISBENZIMIDAZOLE

This is a division of application Ser. No. 418,319 filed Oct. 6, 1989, now U.S. Pat. No. 4,977,271.

BACKGROUND OF THE INVENTION

Because of the excellent mechanical and thermal properties imbued to polymers of benzimidazole (see *Encycl. of Polymer. Sci. and Engineering*, Second Edition, Vol. 7, pp. 639-665, 1987 and *Encycl. of Polymer Sci. and Engineering*, Second Edition, Vol. 11, pp. 572-601, 1988), it is desirable to incorporate this nucleus into these "Reisert compound polymers". However, only two examples of a Reisert-type compound from benzimidazole have been reported, and they were formed from chloroformates and not from acid chlorides (see *J. Heterocyclic Chem.* 1987, 24, 1349-1351). Attempts to use acid chlorides led to destruction of the benzimidazole nucleus (see *J. Chem. Soc., Chem. Comm.*, 1984, 1245-1246). Thus, for the production of benzimidazole polymers by Reisert chemistry difunctionality was required as well as a means of producing Reisert compound moieties from acid chlorides. This invention relates to the discovery of methods for producing difunctional benzimidazole Reisert compounds.

DESCRIPTION OF THE INVENTION

The foregoing problem in regard to undesired bisbenzimidazole formation can be overcome by use of a two step process which yields a poly Reisert. The first step involves the reaction of benzimidazole to form bisbenzimidazole which is then reacted with acid chloride and cyanide to form the Reisert compound thereof.

The first step in the instant process involves reaction of benzimidazole with an aliphatic diacid chloride of the general formula ClC(O)RC(O)Cl , where R is alkylene (e.g., C_4 - C_6 alkylene) to yield a bisbenzimidazole. This reaction can be conveniently performed under the following general conditions: Organic solvent (e.g., dimethyl formamide), base (e.g., triethylamine), temperatures of about 5°C . to 20°C ., with mechanical stirring.

The bisbenzimidazole compound is then reacted with an aliphatic acid chloride and cyanide (e.g., trimethylsilyl cyanide) to yield the Reisert compound thereof having the structure shown in the Figure. This reaction can be conducted in the same type of solvent used in the preceding step, at temperatures from 10°C . to 20°C . using amounts of the reagents which can range from 1 to 1.2 on a molar ratio basis.

The following Examples further illustrate the present invention.

EXAMPLE 1

This shows synthesis of the type of compound shown as (I) in the Figure, specifically, bis(benzimidazoyl)-1,6-dicarbonyl hexane.

To a well stirred solution of benzimidazole (0.022 mole, 2.6 gm) and triethylamine (0.011 mole, 0.11 gm) in dimethylformamide (10 ml), adipoyl chloride (0.01

mole, 1.83 gm) in dichloromethane (10 ml) was added. The resulting reaction media was stirred at room temperature for twenty-four hours. The reaction mixture was poured into water and was stirred for three hours. An essentially quantitative yield of the product was obtained by filtering the solid from the liquid mixture and was washed with water, ethanol (2 and 10 ml portions), ether (2 and 25 ml portions) and was dried. The crude product was taken up in hot dimethylformamide (35 ml) and was precipitated by pouring into water (250 ml), was filtered, was crystallized from tetrahydrofuran, and was dried. The yield was 3.2 gm (93%).

Melting point: 250° - 252°C . (dec)

IR (KBr): 3150-2900 (C-H), 1721 (N-CO), 1680, 1677, 1610, 1507 (aromatic), 1477, 1451, 1419, 1387, 1348, 1335, 1309, 1284, 1235, 1204, and 1165 cm^{-1} .

^1H NMR (DMSO-d): δ 8.96 (s, 2H, C_{21} -H), 8.25-8.15 (m, 2H, Ar-H), 7.78-7.74 (m, 2H, Ar-H), 7.45-7.35 (m, 4H, Ar-H), 3.3-3.2 (s, 4H OCH_2) and 1.95-1.80 (s, 4H CH_2CH_2).

EXAMPLE 2

Benzoyl chloride (0.0022 mole, 0.31 gm) was added to a well stirring suspension of bis(benzimidazoyl)-1,6-dicarbonyl hexane from Example 1 (0.001 mole, 0.346 gm) in N-methylpyrrolidone (5 ml). Addition of trimethylsilyl cyanide (0.0022 mole, 0.3 ml) followed. The reaction mixture was stirred at room temperature for thirty-two hours and was quenched by pouring into 150 ml of water. The mixture was extracted with dichloromethane (3×50 ml). The organic layers were combined, washed with 8% hydrochloric acid (2×50 ml), aqueous saturated bicarbonate (2×50 ml), water (2×50 ml) and were dried. The crude product was treated with activated carbon (NORIT brand) and recrystallized from ethylacetate hexane. The yield was 350 gm (58%). The melting point was 145° - 150°C .

IR (KBr): 2940, 2920 (C-H), 1660 (N-CO), 1595, 1485, 1370, 1360, 1345, 1275, and 1230 cm^{-1} .

^1H NMR (CDCl_3): 8.3-6.1 (m, 20H, Ar-H), 3.1-2.4 (m, 4H, COCH_2), 2.2-1.4 (m, 4H, CH_2CH_2).

We claim:

1. A process for the formation of a Reisert compound of bisbenzimidazole which comprises:

- reaction of benzimidazole with a diacid chloride to form bisbenzimidazole; and
- reaction of the bisbenzimidazole with acid chloride and cyanide to form the Reisert compound thereof.

2. A process as claimed in claim 1 wherein the diacid chloride is adipoyl chloride.

3. A process as claimed in claim 1 wherein the cyanide is trimethylsilyl cyanide.

4. A process as claimed in claim 1 wherein the diacid chloride is of the formula ClC(O)RC(O)Cl where R is C_4 to C_6 alkylene.

5. A process as claimed in claim 1 wherein the diacid chloride is of the formula ClC(O)RC(O)Cl where R is C_4 to C_6 alkylene and the cyanide is trimethylsilyl cyanide.

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