

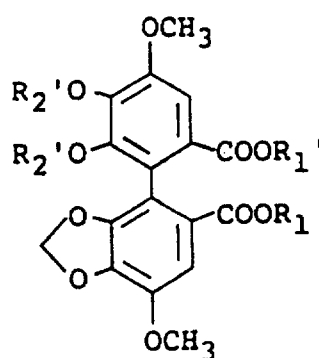
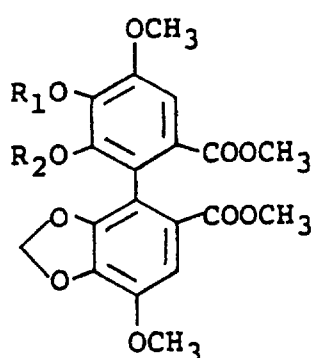
Description

This invention relates to novel biphenyl derivatives useful for prophylaxis and treatment of hepatic diseases, processes for preparing the same, and novel intermediates therefor.

The liver is an important organ which has various functions, for example, participates in various metabolisms, reserve of nutriments, but it is injured by various factors such as virus, drugs, alcohol, and thereby induced into various hepatic diseases such as acute hepatitis, chronic hepatitis, fatty liver, and finally hepatocirrhosis. It is known that these chronic hepatitis and hepatocirrhosis associated with cellular infiltration and hepatic fibrosis are one of intractable diseases.

As a drug for treating these hepatic diseases, there have been known symmetric octa-substituted biphenyl compounds such as dimethyl 4,4'-dimethoxy-5,6,5',6'-bis(methylenedioxy)-2,2'-biphenyldicarboxylate (DDB) [cf. Japanese Patent First Publication (Kokai) No. 209582/1985].

Japanese Patent First Publication (Kokai) Nos. 294677/1987 and 88176/1988 disclose that the following octa-substituted biphenyl derivatives are effective for the treatment of hepatic diseases such as chronic or acute hepatitis.

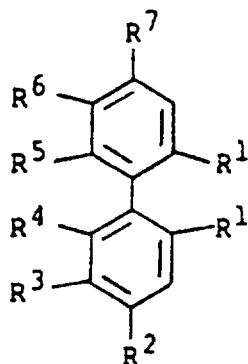


wherein R_1 and R_2 are methyl or both combine to form $O=C$ and R_1' is hydrogen atom or a lower alkyl group and R_2 is a lower alkyl group.

The present inventors have intensively studied as to various substituted biphenyl derivatives and have found that novel hexa- or hepta-substituted biphenyl derivatives have excellent therapeutic and prophylactic effects on hepatic diseases.

An object of the invention is to provide novel hexa- or hepta-substituted biphenyl derivatives useful for the prophylaxis and treatment of various hepatic diseases. Another object of the invention is to provide processes for preparing the novel biphenyl derivatives. A further object of the invention is to provide a pharmaceutical composition for the prophylaxis and treatment of hepatic diseases containing as an active ingredient the novel biphenyl derivative. Still further object of the invention is to provide novel intermediate compounds useful for preparing the novel hexa- or hepta-substituted biphenyl derivatives. These and other objects and the advantages of the invention will be apparent to those skilled in the art from the following description.

The novel hexa- or hepta-substituted biphenyl derivatives of the invention have the following formula:



(I)

wherein R¹ is a substituted or unsubstituted aminocarbonyl group, aminothiocarbonyl group, a substituted or unsubstituted C₁₋₆ alkoxy-carbonyl group, cyano group, or a group of the formula:



one or two of R² to R⁷ are hydrogen atom, and the remaining groups are the same or different and are each a C₁₋₆ alkoxy group, a phenyl-C₁₋₆ alkoxy group or hydroxy group, or the adjacent two groups thereof combine to form a C₁₋₆ alkylenedioxy group, and Alk¹ is a C₁₋₆ alkylene group.

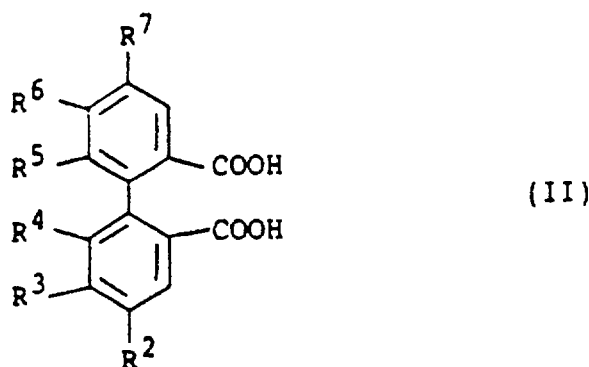
In an embodiment of the invention, the biphenyl derivatives of the invention are the compounds of the formula (I) wherein R¹ is an aminocarbonyl group having optionally one or two substituents selected from a hydroxy-C₁₋₆ alkyl group and a C₁₋₆ alkyl group; aminothio-carbonyl group; a C₁₋₆ alkoxy-carbonyl group being optionally substituted by a C₁₋₆ alkoxy group; cyano group; or a 4,4-di(C₁₋₆ alkyl)-4,5-dihydro-2-oxazolyl group.

Preferred compounds are the compounds of the formula (I) wherein among R² to R⁴, either one of R² and R⁴ is hydrogen atom or a C₁₋₆ alkoxy group, and the remaining two groups are a C₁₋₆ alkoxy group, a phenyl-C₁₋₆ alkoxy group or hydroxy group, or both combine to form a C₁₋₆ alkylenedioxy group; R⁵ and R⁶ combine to form a C₁₋₆ alkylenedioxy group; and R⁷ is hydrogen atom.

More preferred compounds are the compounds of the formula (I) wherein

- (i) R² and R³ are each a C₁₋₆ alkoxy or both combine to form a C₁₋₆ alkylenedioxy group, and R⁴ is hydrogen atom;
 - (ii) R² is hydrogen atom, and R³ and R⁴ combine to form a C₁₋₆ alkylenedioxy group;
 - (iii) R² to R⁴ are all a C₁₋₆ alkoxy group;
 - (iv) R² is a lower alkoxy group; R³ and R⁴ are combine to form a C₁₋₆ alkylenedioxy group, or R³ is a phenyl-C₁₋₆ alkoxy group and R⁴ is hydrogen atom, or
 - (v) R² is a C₁₋₆ alkoxy group, R³ is hydroxy group, and R⁴ is hydrogen atom;
- and in any case of the above (i) to (v), R⁵ and R⁶ combine to form a C₁₋₆ alkylenedioxy group, and R⁷ is hydrogen atom.

Among the compounds of this invention, the compounds of the formula (I) wherein R¹ is a substituted or unsubstituted aminocarbonyl group or a substituted or unsubstituted C₁₋₆ alkoxy-carbonyl group can be prepared by reacting a biphenyldicarboxylic acid of the formula:

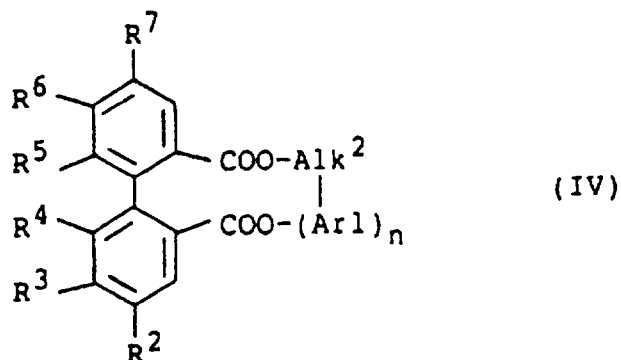


wherein R² to R⁷ are as defined above, or a reactive derivative of the carboxyl groups thereof or a salt thereof with a compound of the formula:

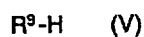


wherein R⁸ is a substituted or unsubstituted amino group or a substituted or unsubstituted C₁₋₆ alkoxy group, or a salt thereof.

The compounds of the formula (I) wherein R¹ is a substituted or unsubstituted C₁₋₆ alkoxy-carbonyl group can also be prepared by reacting a cyclic diester compound of the formula:

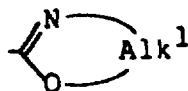


15 wherein Alk² is a C₁₋₆ alkylene, Ar₁ is an arylene group, n is 0 or 1, and R² to R⁷ are as defined above, or a salt thereof with an alcohol compound of the formula:

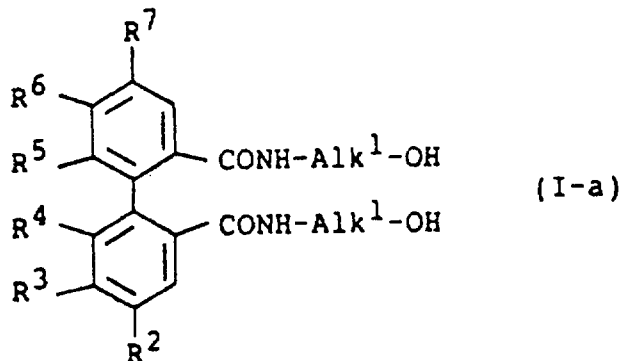


20 wherein R⁹ is a substituted or unsubstituted C₁₋₆ alkoxy group.

The compounds of the formula (I) wherein R¹ is a group of the formula:



(Alk¹ is as defined above) can be prepared by subjecting a bis(hydroxyalkylamide) compound of the formula:

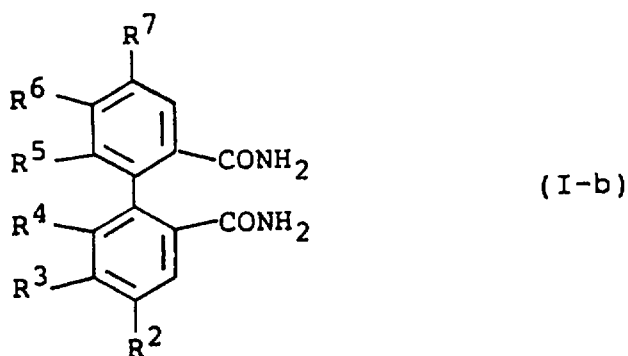


45 wherein R² to R⁷ and Alk¹ are as defined above, or a reactive derivative thereof or a salt thereof to an intramolecular ring closure reaction.

The compounds of the formula (I) wherein R¹ is cyano group can be prepared by dehydrating a bis-(carboxamide) compound of the formula:

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wherein R^2 to R^7 are as defined above, or a salt thereof.

Moreover, the compounds of the formula (I) wherein R^1 is aminothiocarbonyl group can be prepared by thiocarbonylation of the compound of the formula (I) wherein R^1 is cyano group, or a salt thereof.

The condensation reaction of a reactive derivative of the biphenyldicarboxylic acid compound (II) with the compound (III) can be carried out in an appropriate solvent in the presence or absence of an acid scavenger. The reactive derivative includes any conventional reactive derivative, for example, the corresponding acid halide, mixed acid anhydride, active ester. The acid scavenger includes, for example, pyridine, N,N-dialkylanilines, N-alkylmorpholins, alkali metal hydroxides, alkali metal carbonates, alkali metal hydrogen carbonates; trialkylamines in the presence or absence of N,N-dialkylaminopyridines. Besides, the condensation reaction of a free biphenyldicarboxylic acid compound (II) with the compound (III) can be carried out, for example, in the presence or absence of a condensation agent. The condensation agent includes any conventional condensation agent, for example, N,N'-dicyclohexylcarbodiimide, N-cyclohexyl-N'-morpholinocarbodiimide, N-ethyl-N'-(3-dimethylaminopropyl)carbodiimide. The above both condensation reactions are carried out in a conventional solvent such as dimethylformamide, dimethylacetamide, tetrahydrofuran, dichloromethane, ethyl acetate, acetonitrile, at a temperature of from cooling temperature to room temperature, for example, 0° to 20°C .

The reaction of the cyclic diester compound (IV) with the alcohol compound (V) can be carried out by a conventional ester exchange reaction, for example in the presence of an acid or a base in an appropriate solvent or without any solvent. The acid includes inorganic or organic acids such as sulfuric acid, hydrochloric acid, formic acid, acetic acid, p-toluenesulfonic acid, methanesulfonic acid, and the base includes organic or inorganic bases such as alkali metal hydroxides, alkaline earth metal hydroxides, alkali metal hydrides. The solvent includes lower alcohols, dioxane, tetrahydrofuran, diethyl ether, dimethylformamide, dimethylsulfoxide, acetonitrile. The reaction can be carried out at a warm or heat temperature, preferably at 30° to 80°C .

The intramolecular ring closure reaction of the bis(hydroxyalkylamide) compound (I-a) or a reactive derivative thereof can be carried out in the presence or absence of a halogenating agent in an appropriate solvent or without any solvent in a usual manner. The halogenating agent includes any conventional agent, for example, thionyl chloride, phosphorus pentachloride, phosphorus oxychloride. The solvent includes dichloromethane, dichloroethane, chloroform. The reaction preferably proceeds at a temperature of from cooling to room temperature, for example -20°C to 20°C .

The dehydrating reaction of the bis(carboxamide) compound (I-b) can be carried out in the presence or absence of a dehydrating agent. The dehydrating agent includes any conventional agent such as phosphorus oxychloride, thionyl chloride, phosphorus pentachloride, benzenesulfonic acid, aluminum chloride. The reaction can be carried out without using any solvent or in an appropriate solvent such as benzene, tetrahydrofuran, dimethylformamide, pyridine, and the like and preferably proceeds under heating, for example 60° to 100°C .

The thiocarbonylation of the compound of the formula (I) wherein R^1 is cyano group, can be carried out by treating the compound with a thiocarbonylation agent in an appropriate solvent in the presence or absence of an acid scavenger. The thiocarbonylation agent includes any conventional agents, for example hydrogen sulfide gas, phosphorus pentachloride, a dimer of p-methoxyphenylthionophosphinesulfide. The acid scavenger includes any substance as mentioned hereinbefore. The solvent includes preferably dioxane, pyridine, and the like. The reaction preferably proceeds at a temperature of from room temperature to heat temperature, for example at 20° to 100°C .

Besides, the compounds of the formula (I) wherein one of R^2 to R^7 is a phenyl- C_{1-6} alkoxy group (e.g. benzyl group) may be converted into the corresponding reduction products of the formula (I) wherein said

group is hydroxy group. The reduction can be carried out by a conventional method, for example, by catalytic hydrogenation in hydrogen gas stream in the presence of an appropriate catalyst (e.g. palladium-carbon). The solvent includes any conventional solvent such as lower alkanols, dioxane, tetrahydrofuran. The reaction preferably proceeds under atmospheric pressure at a temperature of from room temperature to heat temperature, for example at 10° to 50° C.

In the above reactions, the starting compounds may be used as they stand or in the form of a salt thereof. For instance, the starting compounds of the formulae (II), (V) and (I-a) and of the formulae (III), (IV) and (I-b) which have a hydroxy group may be used in the form of an alkali metal salt, alkaline earth metal salt or ammonium salt, and the compounds of the formula (III) which have a substituted or unsubstituted amino group may be used in the form of an inorganic or organic acid addition salt.

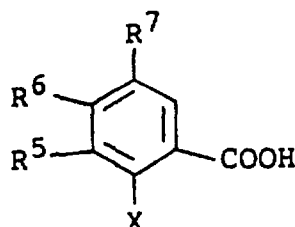
The desired compounds (I) of this invention are novel and have excellent properties for the prophylaxis and treatment of hepatic diseases and can be used as a medicament. Specifically, the compounds (I) of this invention have excellent therapeutic, ameliorating and protecting effects on various hepatic damages such as denature and necrosis of hepatic cells, hepatic fibrosis, fatty retention, and are useful as a therapeutic or prophylactic drug for chronic hepatitis, acute hepatitis, fatty liver, hepatic congestion, and hepatocirrhosis. For instance, a test compound was orally administered to mice and thereafter carbon tetrachloride was orally administered, and then increase of the blood plasma glutamic pyruvic transaminase (GPT), i.e., an index of hepatic disease, was measured. As a result, the object compounds (I) of this invention, dimethyl 5,6-methylenedioxy-4'-methoxy-5',6'-methylenedioxy-2,2'-bisphenyldicarboxylate and N,N,N',N'-tetraethyl-5,6-methylenedioxy-4',5'-dimethoxy-2,2'-biphenylbiscarboxamide showed 90 % or more inhibitory ratio of increase of the blood plasma GPT activity in a dose of 100 mg/kg.

The compounds (I) of this invention can be used as a drug either in a free form or in a form of a pharmaceutically acceptable salt thereof. The pharmaceutically acceptable salt includes inorganic or organic base salts, for example, alkali metal salts (e.g. sodium salt, potassium salt,), alkaline earth metal salts (e.g. calcium salt,), ammonium salts (e.g. tetramethylammonium salt, tetraethylammonium salt,).

The compounds (I) of this invention can be administered orally or parenterally, but preferably orally. The compounds can be used in the form of a conventional pharmaceutical preparation, for example tablets, capsules, powders, and injections.

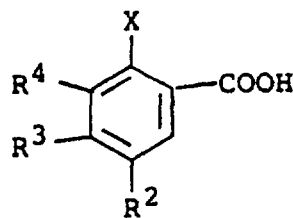
The dose of the compounds (I) may vary depending on the age, body weight or state of the patients, severity of the diseases and the like, but is usually in the range of 0.1 to 500 mg/kg/day, preferably 1 to 300 mg/kg/day.

The starting compounds of the formulae (II) and (IV) used in the process for preparing the desired compounds (I) of the invention can be prepared by condensing a 2-halogenobenzoic acid of the formulae:



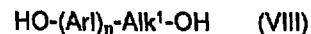
(VI)

and

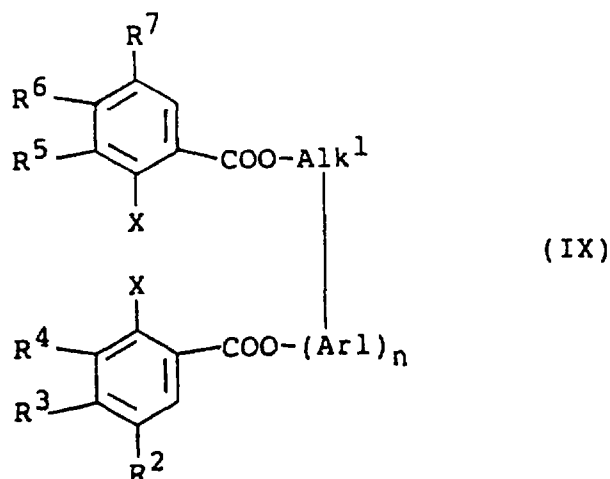


(VII)

wherein X is a halogen atom, and R² to R⁷ are as defined above, or a reactive derivative thereof with a compound of the formula:



wherein Ar, n and Alk¹ are as defined above, to give a dihalogeno compound of the formula:



wherein R^2 to R^7 , Alk^1 , Arl , and n are as defined above, and subjecting said compound (IX) to an intramolecular ring closure reaction, for example, in the presence of copper powder or copper(I) iodide, optionally followed by hydrolysis.

The compounds and preparation thereof of this invention are illustrated by the following Examples but should not be construed to be limited thereto.

Example 1

(1-a) Piperonal (500 g) and cyclohexylamine (330 g) are dissolved in benzene, and the mixture is refluxed for about 3 hours while dehydrating with Diene Schtark device. The solvent is distilled off from the reaction mixture, and the residue is recrystallized from a mixture of isopropyl ether and hexane to give N-(3,4-methylenedioxybenzylidene)-N-cyclohexylamine (597 g).

The product (158 g) is dissolved in tetrahydrofuran. To the solution is added 2.5M n-butyllithium (30 ml) below -50°C , followed by dropwise addition of a solution of iodine (172 g) in tetrahydrofuran. The reaction mixture is warmed to room temperature and is poured into water. The precipitated crystals are separated by filtration, washed with water and dissolved in a mixture of dioxane, acetic acid and water. The mixture is allowed to stand at room temperature overnight. The solvent is distilled off from the mixture, and the residue is diluted with water. The precipitated crystals are collected by filtration, washed and dried to give 2-iodo-3,4-methylenedioxybenzaldehyde (118 g). M.p. $124 - 126^\circ\text{C}$.

(1-b) 2-Bromo-4,5-dimethoxybenzaldehyde dimethylacetal (59 g) is dissolved in tetrahydrofuran and thereto is added 2.5M n-butyllithium (89 ml) under cooling at -70°C . The reaction mixture is kept below -50°C for 30 minutes and thereto is added dropwise a solution of iodine (51.4 g) in tetrahydrofuran below -50°C . The mixture is kept at the same temperature for 15 minutes and thereto are added aqueous acetic acid to adjust the pH of the solution at 2. The mixture is allowed to stand at room temperature overnight, and the solvent is distilled off from the reaction mixture. The residue is dissolved in ethyl acetate. The solution is washed and dried, and the solvent is distilled off. The residue is crystallized from ethyl acetate to give 2-iodo-4,5-dimethoxybenzaldehyde (43 g). M.p. $139 - 141^\circ\text{C}$.

(2-a) A solution of the product obtained in (1-a) (15 g) in dioxane (400 ml) and resorcinol (7.8 g) are added to 0.2 N acetate buffer (200 ml), and thereto is gradually added sodium chlorite (5.9 g) with stirring. The mixture is stirred for 2 hours and then concentrated. The pH of the mixture is adjusted at 1 with conc. hydrochloric acid, and the precipitated crystals are separated by filtration, washed and dried to give 2-iodo-3,4-methylenedioxybenzoic acid (14 g).

The above product (13 g), thionyl chloride (16 ml) and pyridine (0.1 ml) are added to dioxane, and the mixture is refluxed for 30 minutes. The reaction mixture is distilled to remove the solvent and further subjected to azeotropic distillation with toluene to give 2-iodo-3,4-methylenedioxybenzoic chloride (14.5 g). M.p. $69 - 70^\circ\text{C}$.

(2-b) The product obtained in (1-b) is treated in the same manner as described in (2-a) to give 2-iodo-4,5-dimethoxybenzoic chloride. M.p. $151 - 154^\circ\text{C}$.

(3) Salicyl alcohol (22.4 g) is dissolved in dimethylacetamide (150 ml) and thereto are added triethylamine (64.1 ml) and 4-dimethylaminopyridine (100 ml) under nitrogen gas stream, and the mixture is cooled to -30°C . To the mixture is added dropwise with stirring a suspension of the product

obtained in (2-a) (62.1 g) in dimethylacetamide (100 ml) below -25 °C. The mixture is stirred at the same temperature for 30 minutes and the reaction temperature is gradually raised up to room temperature. The mixture is further stirred for 5 hours at room temperature. After the reaction is over, the reaction mixture is cooled to -30 °C and thereto is added dropwise triethylamine (30.7 ml) followed by addition of a solution of the product obtained in (2-b) (52.8 g) in dimethylacetamide (100 ml). The mixture is stirred at room temperature overnight and is poured into a mixture of water-acetic acid. The organic layer is separated, washed, dried and distilled under reduced pressure to remove the solvent. The residue is recrystallized from toluene to give 2-iodo-3,4-methylenedioxybenzoic acid 2-[(2-iodo-4,5-dimethoxyphenyl)-carbonyloxy]benzyl ester (89.4 g). M.p. 128 - 130 °C.

(4) A suspension of copper powder (413 g) in dimethylformamide (1.5 liter) is refluxed with stirring. To the suspension is added dropwise a solution of the product obtained in (3) (89.5 g) in dimethylformamide (1000 ml) over a period of 3 hours, and the mixture is refluxed for 2 hours. Undissolved substances are filtered off from the reaction mixture, and the filtrate is concentrated under reduced pressure. To the residue is added ethyl acetate, and the mixture is washed with water, dried and distilled under reduced pressure to remove the solvent. The residue is recrystallized from toluene to give 5,6-methylenedioxy-4',5'-dimethoxy-2'-(2-hydroxymethylphenyloxycarbonyl)-2-biphenylcarboxylic acid lactone (53.7 g). M.p. 176 - 179 °C.

(5) The product obtained in (4) (1.5 g), ethyleneglycol monomethyl ether (30 ml) and sulfuric acid (1 ml) are heated at 110 °C for 60 hours. The reaction mixture is poured into a mixture of ethyl acetate (200 ml) and water (200 ml), and the organic layer is separated. The organic layer is washed, dried and distilled under reduced pressure to remove the solvent. The residue is purified with silica gel column chromatography [solvent, ethyl acetate-hexane (1 : 2)], and the eluate is distilled under reduced pressure to remove the solvent, and the residue is recrystallized from a mixture of diethyl ether-diisopropyl ether to give 5,6-methylenedioxy-4',5'-dimethoxy-2,2'-biphenyldicarboxylic acid bis(methoxyethyl) ester (900 mg) as colorless crystal. M.p. 59 - 61 °C.

Example 2

(1) 2-Iodo-3,4-methylenedioxybenzoyl chloride (4.66 g), 1,3-propanediol (9.0 g) and 4-dimethylaminopyridine (50 ml) are dissolved in a mixture of tetrahydrofuran (70 ml) and triethylamine (4.21 ml), and the mixture is stirred at room temperature overnight. The reaction mixture is concentrated under reduced pressure, and the residue is extracted with methylene chloride. The solution is washed, dried and evaporated to dryness in vacuo to give colorless crystals (3.84 g). The crystals are dissolved in a mixture of tetrahydrofuran (50 ml) and triethylamine (2.95 ml). To the solution is added 4-dimethylaminopyridine (50 mg), followed by dropwise addition of a solution of 2-iodo-4,5-dimethoxybenzoyl chloride (3.5 g) in dimethylacetamide (50 ml) at -30 °C. The mixture is stirred at room temperature overnight and poured into a mixture of water and ethyl acetate, and the organic layer is separated. The organic layer is washed, dried and concentrated under reduced pressure. The residue is recrystallized from ether to give 2-iodo-3,4-methylenedioxybenzoic acid 2-[(2-iodo-4,5-dimethoxyphenyl)-carbonyloxy]propyl ester (4.82 g). M.p. 131 - 133 °C.

(2) The product obtained in (1) is treated in the same manner as in Example 1-(4) to give 5,6-methylenedioxy-4',5'-dimethoxy-2,2'-biphenyldicarboxylic acid trimethylene ester (1.8 g). M.p. 197 - 199 °C.

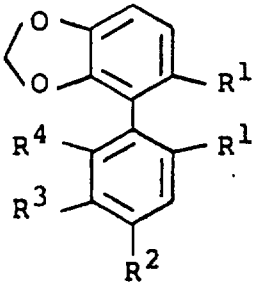
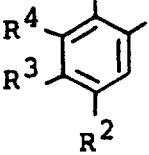
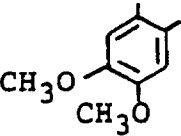
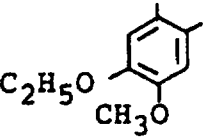
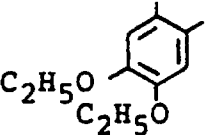
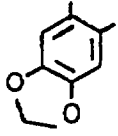
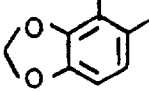
(3) The product obtained in (2) is treated in the same manner as in Example 1-(5) to give 5,6-methylenedioxy-4',5'-dimethoxy-2,2'-biphenyldicarboxylic acid bis(methoxyethyl) ester as colorless crystals.

This product has the same physicochemical properties as those of the product of Example 1.

Examples 3 to 11

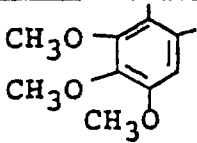
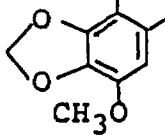
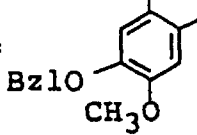
The corresponding starting compounds are treated in the same manner as in Example 1 or Example 2 to give the compounds as shown in Table 1.

Table 1

Ex. No.			
		R ¹	Physical properties, etc.
3		-COOCH ₃	M.p. 121°C
4	The same as above	-COOC ₂ H ₅	M.p. 82°C
5		-COOCH ₃	M.p. 148 - 150°C
6		The same as above	M.p. 122 - 123°C
7		The same as above	M.p. 139 - 141°C
8		The same as above	M.p. 186°C

- to be continued -

Table 1 (Continued)

9		-COOCH ₃	M.p. 142 - 143°C
10		The same as above	M.p. 131°C
11	* 	The same as above	IR ν Nujol (cm ⁻¹): 1700 Max Mass (M ⁺): 450 (m/e)

*) Bzl means a benzyl group.

Example 12

(1-a) To 5,6-methylenedioxy-4',5'-dimethoxy-2'-(2-hydroxymethylphenyloxycarbonyl)-2-biphenylcarboxylic acid lactone (4 g) are added methanol (100 ml) and an aqueous solution (50 ml) of potassium hydroxide (4 g). The mixture is refluxed for 30 minutes, and concentrated under reduced pressure to remove methanol. The pH of the resulting aqueous solution is adjusted at 1 with 10 % hydrochloric acid. The products are extracted with ethyl acetate. The extract is washed, dried and concentrated to dryness under reduced pressure. The residue is triturated with diisopropyl ether to give 5,6-methylenedioxy-4',5'-dimethoxy-2,2'-biphenyldicarboxylic acid (2.84 g). M.p. 207 - 210 °C.

(1-b) 5,6-Methylenedioxy-4',5'-dimethoxy-2,2'-biphenyldicarboxylic acid trimethylene ester is treated in the same manner as in the above (1-a) to give 5,6-methylenedioxy-4',5'-dimethoxy-2,2'-biphenyldicarboxylic acid (1.4 g).

This product has the same physicochemical properties as those of the product in (1-a).

(2) The product obtained in (1-a) or (1-b) (1.3 g) is dissolved in dioxane (10 ml). To the solution are added thionyl chloride (1.76 ml) and dimethylformamide (0.2 ml), and the mixture is refluxed for one hour. The solvent is distilled off from the reaction mixture, and the residue is dissolved in tetrahydrofuran (100 ml). The solution is added with stirring to a mixture of diethylamine hydrochloride (2.06 g), triethylamine (2.8 ml) and tetrahydrofuran (50 ml). The mixture is stirred at room temperature overnight and concentrated to dryness under reduced pressure. The residue is poured into a mixture of ethyl acetate and water, and the organic layer is separated. The organic layer is washed, dried and distilled to remove the solvent. The residue is recrystallized from a mixture of diethyl ether and hexane to give N,N,N',N'-tetraethyl-5,6-methylenedioxy-4',5'-dimethoxy-2,2'-biphenyldicarboxamide (1.23 g) as colorless crystal. m.p. 148 - 149 °C.

Example 13

(1) 2-Iodo-3,4-methylenedioxybenzoic chloride is treated in the same manner as in Example 1-(3) and (4) to give 5,6-methylenedioxy-5',6'-methylenedioxy-2'-(2-hydroxymethylphenyloxycarbonyl)-2-biphenylcarboxylic acid lactone. M.p. 249 - 252 °C.

(2) The product obtained in (1) (4 g) is treated in the same manner as in Example 12-(1-a) to give 5,6-methylenedioxy-5',6'-methylenedioxy-2,2'-biphenyldicarboxylic acid (2.1 g). M.p. 253 - 255 °C.

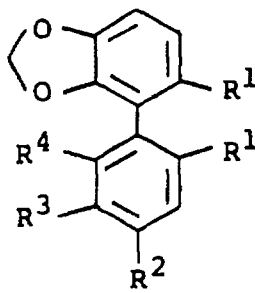
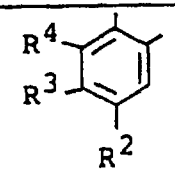
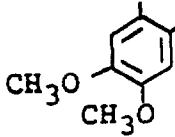
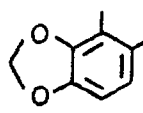
(3) The product obtained in (2) (20 g) is suspended in dioxane. There to is added thionyl chloride (20 ml), and the mixture is refluxed. The solvent is distilled off under reduced pressure from the reaction mixture, and the residue is subjected to azeotropic distillation with toluene and then dissolved in dioxane (200 ml). To the solution is added dropwise conc. aqueous ammonia (200 ml) at -20 °C, and the mixture is stirred at room temperature overnight. The solvent is distilled off from the reaction mixture, and the

residue is adjusted to pH 2 with conc. hydrochloric acid. The precipitated crystals are separated by filtration, washed with water, and then recrystallized from methanol to give 5,6-methylenedioxy-5',6'-methylenedioxy-2,2'-biphenyldicarboxamide (11 g) as colorless crystal. M.p. 249 - 250 °C.

5 Examples 14 to 16

The corresponding starting compounds are treated in the same manner as in Example 12 or 13-(2) and -(3) to give the compounds as shown in Table 2.

Table 2

Ex. No.			
		R ¹	Physical properties etc.
14		both -CONHC(CH ₃) ₂ CH ₂ OH	M.p. 182 - 184 °C
15	The same as above	both -CONHC ₂ H ₅	M.p. 196 - 197 °C
16		both -CONHC(CH ₃) ₂ CH ₂ OH	M.p. 173 - 175 °C

Example 17

N,N'-Bis(3-hydroxy-2-methylpropan-2-yl)-5,6-methylenedioxy-5',6'-methylenedioxy-2,2'-biphenyldicarboxamide (3 g) is added to thionyl chloride (20 ml), and the mixture is stirred at room temperature for 2 hours. The reaction mixture is distilled under reduced pressure to remove thionyl chloride, and the residue is dissolved in ethyl acetate. The solution is washed, dried and distilled to remove the solvent, and the residue is crystallized from diisopropyl ether to give 5,6-methylenedioxy-5',6'-methylenedioxy-2,2'-bis(4,4-dimethyl-4,5-dihydro-2-oxazolyl)biphenyl (2 g) as colorless crystal. M.p. 152 - 154 °C.

Example 18

5,6-Methylenedioxy-5',6'-methylenedioxy-2,2'-biphenyldicarboxamide (8 g) is refluxed in phosphorus oxychloride (70 ml) for 4 hours. The reaction mixture is cooled and then poured into ice water. The precipitated crystals are separated by filtration and dissolved in chloroform. The solution is washed with water, dried and distilled to remove the solvent to give 5,6-methylenedioxy-5',6'-methylenedioxy-2,2'-biphenyldicarbonitrile (6.5 g) as colorless crystal. M.p. 213 - 215 °C.

Example 19

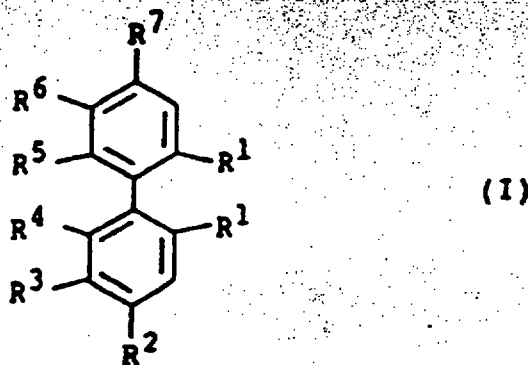
5,6-Methylenedioxy-5',6'-methylenedioxy-2,2'-biphenyldicarbonitrile (5 g) is suspended in a mixture of pyridine (100 ml) and dioxane (100 ml), and thereto is added triethylamine (4.7 ml) and further hydrogen sulfide gas is passed through the mixture over a period of 5 hours. The reaction vessel is sealed and allowed to stand at room temperature for 2 hours. The reaction mixture is distilled under reduced pressure to remove the solvent. The residue is extracted with chloroform, and water is added to the extract. The precipitated crystals are separated by filtration to give 5,6-methylenedioxy-5',6'-methylenedioxy-2,2'-biphenylbis(thiocarboxamide) (5.5 g). M.p. 237 - 238 °C (decomp.)

Example 20

Dimethyl 5,6-methylenedioxy-4'-methoxy-5'-benzyloxy-2,2'-biphenyldicarboxylate (1.15 g) is dissolved in tetrahydrofuran (50 ml) and methanol (50 ml), and the solution is stirred under 40 psi hydrogen gas atmosphere in the presence of palladium-carbon (0.2 g) for 16 hours. After the reaction, the catalyst is filtered off, and the filtrate is evaporated into dryness under reduced pressure. The residue is treated with methanol to give dimethyl 5,6-methylenedioxy-4'-methoxy-5'-hydroxy-2,2'-biphenyldicarboxylate (0.86 g) as colorless crystal. M.p. 172 - 173 °C.

Claims

1. A biphenyl derivative of the formula:



wherein R¹ is a substituted or unsubstituted aminocarbonyl group, aminothiocarbonyl group, a substituted or unsubstituted C₁₋₆ alkoxy-carbonyl group, cyano group, or a group of the formula:



among R² to R⁴, either one of R² and R⁴ is hydrogen atom or a C₁₋₆ alkoxy group and the remaining two groups are a C₁₋₆ alkoxy group, a phenyl-C₁₋₆ alkoxy group or hydroxy group, or both combine to form a C₁₋₆ alkylenedioxy group; R⁵ and R⁶ combine to form a C₁₋₆ alkylenedioxy group; R⁷ is hydrogen atom; and Alk¹ is a C₁₋₆ alkylene group, provided that, when R¹ is a C₁₋₆ alkoxy-carbonyl group, and when R³ and R⁴ combine to form a C₁₋₆ alkylenedioxy group, R² is not hydrogen atom; or



Example 18

5,6-Methylenedioxy-5',6'-methylenedioxy-2,2'-biphenyldicarboxamide (8 g) is refluxed in phosphorus oxychloride (70 ml) for 4 hours. The reaction mixture is cooled and then poured into ice water. The precipitated crystals are separated by filtration and dissolved in chloroform. The solution is washed with water, dried and distilled to remove the solvent to give 5,6-methylenedioxy-5',6'-methylenedioxy-2,2'-biphenyldicarbonitrile (6.5 g) as colorless crystal. M.p. 213 - 215 °C.

Example 19

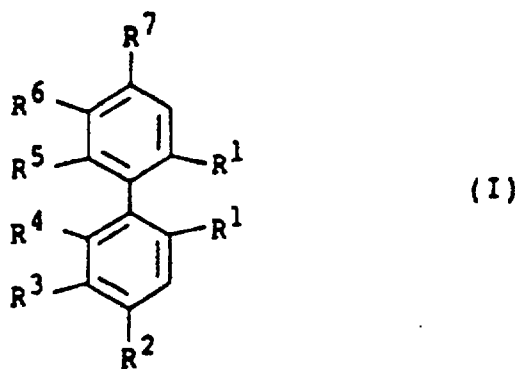
5,6-Methylenedioxy-5',6'-methylenedioxy-2,2'-biphenyldicarbonitrile (5 g) is suspended in a mixture of pyridine (100 ml) and dioxane (100 ml), and thereto is added triethylamine (4.7 ml) and further hydrogen sulfide gas is passed through the mixture over a period of 5 hours. The reaction vessel is sealed and allowed to stand at room temperature for 2 hours. The reaction mixture is distilled under reduced pressure to remove the solvent. The residue is extracted with chloroform, and water is added to the extract. The precipitated crystals are separated by filtration to give 5,6-methylenedioxy-5',6'-methylenedioxy-2,2'-biphenylbis(thiocarboxamide) (5.5 g). M.p. 237 - 238 °C (decomp.)

Example 20

Dimethyl 5,6-methylenedioxy-4'-methoxy-5'-benzyloxy-2,2'-biphenyldicarboxylate (1.15 g) is dissolved in tetrahydrofuran (50 ml) and methanol (50 ml), and the solution is stirred under 40 psi hydrogen gas atmosphere in the presence of palladium-carbon (0.2 g) for 16 hours. After the reaction, the catalyst is filtered off, and the filtrate is evaporated into dryness under reduced pressure. The residue is treated with methanol to give dimethyl 5,6-methylenedioxy-4'-methoxy-5'-hydroxy-2,2'-biphenyldicarboxylate (0.86 g) as colorless crystal. M.p. 172 - 173 °C.

Claims

1. A biphenyl derivative of the formula:



wherein R¹ is a substituted or unsubstituted aminocarbonyl group, aminothiocabonyl group, a substituted or unsubstituted C₁₋₆ alkoxy-carbonyl group, cyano group, or a group of the formula:

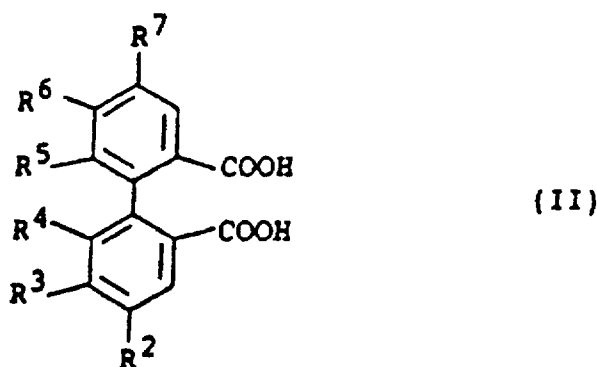


among R² to R⁴, either one of R² and R⁴ is hydrogen atom or a C₁₋₆ alkoxy group and the remaining two groups are a C₁₋₆ alkoxy group, a phenyl-C₁₋₆ alkoxy group or hydroxy group, or both combine to form a C₁₋₆ alkylendioxy group; R⁵ and R⁶ combine to form a C₁₋₆ alkylendioxy group; R⁷ is hydrogen atom; and Alk¹ is a C₁₋₆ alkylene group, provided that, when R¹ is a C₁₋₆ alkoxy-carbonyl group, and when R³ and R⁴ combine to form a C₁₋₆ alkylendioxy group, R² is not hydrogen atom; or



a pharmaceutically acceptable salt thereof.

2. The compound according to claim 1, wherein R¹ is an aminocarbonyl group having optionally one or two substituents selected from a hydroxy-C₁₋₆ alkyl group and a C₁₋₆ alkyl group; aminothiocarbonyl group; a C₁₋₆ alkoxy-carbonyl group being optionally substituted by a C₁₋₆ alkoxy group; cyano group; or a 4,4-di(C₁₋₆ alkyl)-4,5-dihydro-2-oxazolyl group, provided that, when R¹ is a C₁₋₆ alkoxy-carbonyl group, and when R³ and R⁴ combine to form a C₁₋₆ alkylenedioxy group, R² is not hydrogen atom.
3. The compound according to claim 2, wherein R² and R³ are each a C₁₋₆ alkoxy or both combine to form a C₁₋₆ alkylenedioxy group, R⁴ is hydrogen atom, R⁵ and R⁶ combine to form a C₁₋₆ alkylenedioxy group, and R⁷ is hydrogen atom.
4. The compound according to claim 2, wherein R¹ is an aminocarbonyl group having optionally one or two substituents selected from a hydroxy-C₁₋₆ alkyl group and a C₁₋₆ alkyl group; aminothiocarbonyl group; a C₁₋₆ alkoxy-C₁₋₆ alkoxy-carbonyl group; cyano group; or a 4,4-di(C₁₋₆ alkyl)-4,5-dihydro-2-oxazolyl group, R² is hydrogen atom, R³ and R⁴ combine to form a C₁₋₆ alkylenedioxy group, R⁵ and R⁶ combine to form a C₁₋₆ alkylenedioxy group, and R⁷ is hydrogen atom.
5. The compound according to claim 2, wherein R² to R⁴ are all a C₁₋₆ alkoxy group, R⁵ and R⁶ combine to form a C₁₋₆ alkylenedioxy group, and R⁷ is hydrogen atom.
6. The compound according to claim 2, wherein R² is a C₁₋₆ alkoxy group, R³ and R⁴ combine to form a C₁₋₆ alkylenedioxy group, or R³ is a phenyl-C₁₋₆ alkoxy group and R⁴ is hydrogen atom, R⁵ and R⁶ combine to form a C₁₋₆ alkylenedioxy group, and R⁷ is hydrogen atom.
7. The compound according to claim 2, wherein R² is a C₁₋₆ alkoxy group, R³ is hydroxy group, R⁴ is hydrogen atom, R⁵ and R⁶ combine to form a C₁₋₆ alkylenedioxy group, and R⁷ is hydrogen atom.
8. A process for preparing a compound of the formula (I) as set forth in claim 1 wherein R¹ is a substituted or unsubstituted aminocarbonyl group or a substituted or unsubstituted C₁₋₆ alkoxy-carbonyl group, provided that, when R¹ is a C₁₋₆ alkoxy-carbonyl group, and when R³ and R⁴ combine to form a C₁₋₆ alkylenedioxy group, R² is not hydrogen atom, or a pharmaceutically acceptable salt thereof, which comprises reacting a biphenyldicarboxylic acid of the formula:



wherein R² to R⁷ are as defined in claim 1, or a reactive derivative of the carboxyl groups thereof or a salt thereof with a compound of the formula:

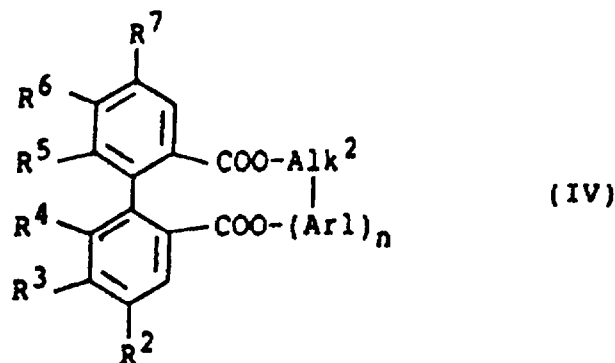


wherein R⁸ is a substituted or unsubstituted amino group or a substituted or unsubstituted C₁₋₆ alkoxy group, or a salt thereof, optionally followed by converting the product into a pharmaceutically acceptable salt thereof.

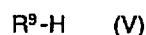


9. The process according to claim 8, wherein R^8 is an amino group having optionally one or two substituents selected from a hydroxy- C_{1-6} alkyl group and a C_{1-6} alkyl group, or a C_{1-6} alkoxy being optionally substituted by a C_{1-6} alkoxy group.

10. A process for preparing a compound of the formula (I) as set forth in claim 1 wherein R^1 is a substituted or unsubstituted C_{1-6} alkoxy-carbonyl group, provided that, when R^1 is a C_{1-6} alkoxy-carbonyl group, and when R^3 and R^4 combine to form a C_{1-6} alkylendioxy group, R^2 is not hydrogen atom, or a pharmaceutically acceptable salt thereof, which comprises reacting a cyclic diester compound of the formula:



wherein Alk^2 is a C_{1-6} alkylene, $Ar1$ is an arylene group, n is 0 or 1, and R^2 to R^7 are as defined in claim 1, or a salt thereof with an alcohol compound of the formula:



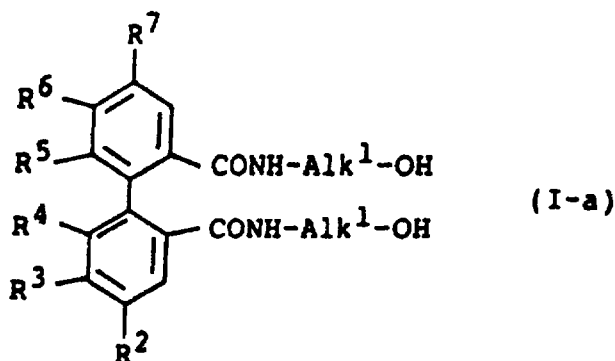
wherein R^9 is a substituted or unsubstituted C_{1-6} alkoxy group, optionally followed by converting the product into a pharmaceutically acceptable salt thereof.

11. The process according to claim 10, wherein R^9 is a C_{1-6} alkoxy group being optionally substituted by a C_{1-6} alkoxy group, and $Ar1$ is 1,2-phenylene group.

12. A process for preparing a compound of the formula (I) as set forth in claim 1 wherein R^1 is a group of the formula:



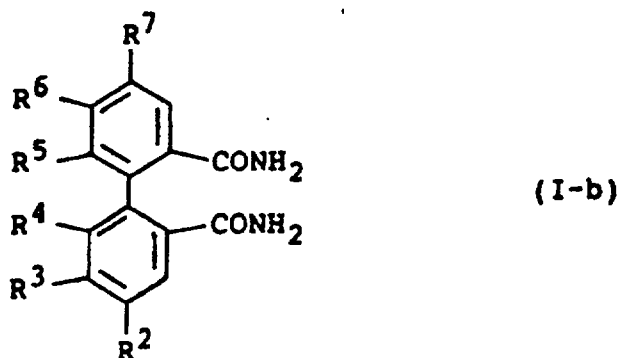
(Alk^1 is as defined in claim 1); or a pharmaceutically acceptable salt thereof, which comprises subjecting a bis(hydroxyalkylamide) compound of the formula:





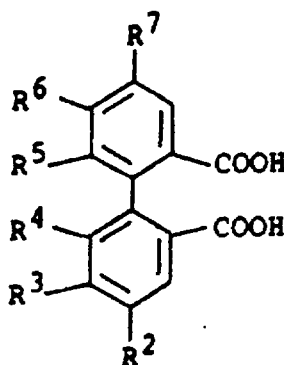
wherein R^2 to R^7 and Alk^1 are as defined in claim 1, or a reactive derivative thereof or a salt thereof to an intramolecular ring closure, optionally followed by converting the product into a pharmaceutically acceptable salt thereof.

13. A process for preparing a compound of the formula (I) as set forth in claim 1 wherein R^1 is cyano group or aminothiocarbonyl group, or a pharmaceutically acceptable salt thereof, which comprises dehydrating a biphenylbis(carboxamide) compound of the formula:



wherein R^2 to R^7 are as defined in claim 1, or a salt thereof, optionally followed by thiocarbonylation, and further optionally followed by converting the product into a pharmaceutically acceptable salt thereof.

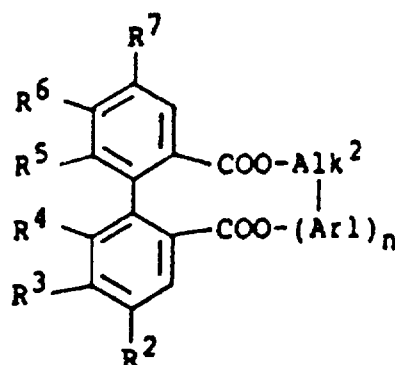
14. A biphenyldicarboxylic acid compound of the formula:



wherein among R^2 to R^4 , either one of R^2 and R^4 is hydrogen atom or a C_{1-6} alkoxy group and the remaining two groups are a C_{1-6} alkoxy group, a phenyl- C_{1-6} alkoxy group or hydroxy group, or both combine to form a C_{1-6} alkylenedioxy group; R^5 and R^6 combine to form a C_{1-6} alkylenedioxy group; and R^7 is hydrogen atom, provided that, when R^3 and R^4 combine to form a C_{1-6} alkylenedioxy group, R^2 is not hydrogen atom, or a salt thereof.



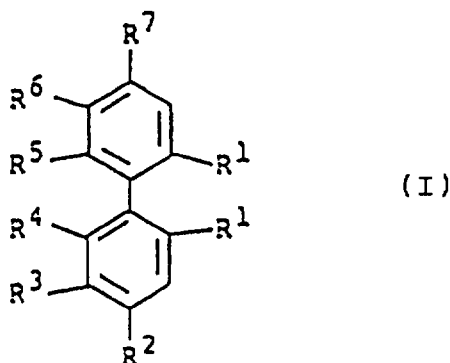
15. A cyclic diester compound of the formula:



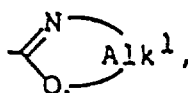
wherein Alk^2 is a C_{1-6} alkylene group; Ar^1 is an arylene group; among R^2 to R^4 , either one of R^2 and R^4 is hydrogen atom or a C_{1-6} alkoxy group and the remaining two groups are a C_{1-6} alkoxy group, a phenyl- C_{1-6} alkoxy group or hydroxy group, or both combine to form a C_{1-6} alkylendioxy group; R^5 and R^6 combine to form a C_{1-6} alkylendioxy group; R^7 is hydrogen atom, and n is 0 or 1, or salt thereof.

Patentansprüche

1. Biphenylderivat der Formel:



worin R^1 eine substituierte oder unsubstituierte Aminocarbonylgruppe, eine Aminothiocarbonylgruppe, eine substituierte oder unsubstituierte C_{1-6} Alkoxycarbonylgruppe, eine Cyanogruppe oder eine Gruppe der Formel

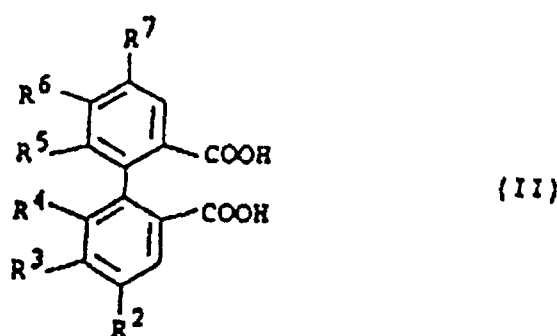


ist;

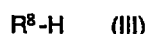
von R^2 bis R^4 ist entweder eines von R^2 und R^4 ein Wasserstoffatom oder eine C_{1-6} -Alkoxygruppe, und die übrigen beiden Gruppen sind eine C_{1-6} -Alkoxygruppe, eine Phenyl- C_{1-6} -alkoxygruppe oder eine Hydroxygruppe, oder beide verbinden sich unter Bildung einer C_{1-6} -Alkylendioxygruppe; R^5 und R^6 verbinden sich unter Bildung einer C_{1-6} -Alkylendioxygruppe; R^7 ist ein Wasserstoffatom; und Alk^1 ist eine C_{1-6} -Alkylengruppe, unter der Voraussetzung, daß wenn R^1 eine C_{1-6} -Alkoxycarbonylgruppe ist, und wenn R^3 und R^4 sich unter Bildung einer C_{1-6} -Alkylendioxygruppe verbinden, R^2 kein Wasserstoffatom ist; oder ein pharmazeutisch annehmbares Salz davon.



2. Verbindung nach Anspruch 1, worin R^1 eine Aminocarbonylgruppe mit wahlweise einem oder zwei Substituenten ausgewählt aus einer Hydroxy- C_{1-6} -Alkylgruppe und einer C_{1-6} -Alkylgruppe; eine Aminothiocarbonylgruppe; eine C_{1-6} -Alkoxy-carbonylgruppe, die wahlweise mit einer C_{1-6} -Alkoxygruppe substituiert ist; eine Cyanogruppe oder eine 4,4-Di(C_{1-6} -alkyl)-4,5-dihydro-2-oxazolylgruppe ist, unter der Voraussetzung, daß wenn R^1 eine C_{1-6} -Alkoxy-carbonylgruppe ist, und wenn R^3 und R^4 sich unter Bildung einer C_{1-6} -Alkylendioxygruppe verbinden, R^2 ein Wasserstoffatom ist.
3. Verbindung nach Anspruch 2, worin R^2 und R^3 jeweils eine C_{1-6} -Alkoxygruppe sind, oder sich beide unter Bildung einer C_{1-6} -Alkylendioxygruppe verbinden, R^4 ist ein Wasserstoffatom, R^5 und R^6 verbinden sich unter Bildung einer C_{1-6} -Alkylendioxygruppe und R^7 ist ein Wasserstoffatom.
4. Verbindung nach Anspruch 2, worin R^1 eine Aminocarbonylgruppe mit wahlweise einem oder zwei Substituenten, ausgewählt aus einer Hydroxy- C_{1-6} -alkylgruppe und einer C_{1-6} -Alkylgruppe; eine Aminothiocarbonylgruppe, eine C_{1-6} -Alkoxy- C_{1-6} -alkoxy-carbonylgruppe; eine Cyanogruppe; oder eine 4,4-Di(C_{1-6} -alkyl)-4,5-dihydro-2-oxazolylgruppe ist, R^2 ist ein Wasserstoffatom, R^3 und R^4 verbinden sich unter Bildung einer C_{1-6} -Alkylendioxygruppe, R^5 und R^6 verbinden sich unter Bildung einer C_{1-6} -Alkylendioxygruppe und R^7 ist ein Wasserstoffatom.
5. Verbindung nach Anspruch 2, worin R^2 bis R^4 jeweils eine C_{1-6} -Alkoxygruppe ist, R^5 und R^6 verbinden sich unter Bildung einer C_{1-6} -Alkylendioxygruppe und R^7 ist ein Wasserstoffatom.
6. Verbindung nach Anspruch 2, worin R^2 eine C_{1-6} -Alkoxygruppe ist, R^3 und R^4 verbinden sich unter Bildung einer C_{1-6} -Alkylendioxygruppe oder R^3 ist eine Phenyl- C_{1-6} -alkoxygruppe und R^4 ist ein Wasserstoffatom, R^5 und R^6 verbinden sich unter Bildung einer C_{1-6} -Alkylendioxygruppe und R^7 ist ein Wasserstoffatom.
7. Verbindung nach Anspruch 2, worin R^2 eine C_{1-6} -Alkoxygruppe ist, R^3 ist eine Hydroxygruppe, R^4 ist ein Wasserstoffatom, R^5 und R^6 verbinden sich unter Bildung einer C_{1-6} -Alkylendioxygruppe und R^7 ist ein Wasserstoffatom.
8. Verfahren zur Herstellung einer Verbindung der Formel (I) gemäß Anspruch 1, worin R^1 eine substituierte oder unsubstituierte Aminocarbonylgruppe oder eine substituierte oder unsubstituierte C_{1-6} -Alkoxy-carbonylgruppe ist, unter der Voraussetzung daß, wenn R^1 eine C_{1-6} -Alkoxy-carbonylgruppe ist, und wenn R^3 und R^4 sich unter Bildung einer C_{1-6} -Alkylendioxygruppe verbinden, R^2 kein Wasserstoffatom ist, oder ein pharmazeutisch annehmbares Salz davon, welches das Umsetzen einer Biphenyldicarbonsäure der Formel:



worin R^2 bis R^7 wie in Anspruch 1 definiert sind, oder eines reaktiven Derivats der Carbonylgruppen davon, oder eines Salzes davon, mit einer Verbindung der Formel:

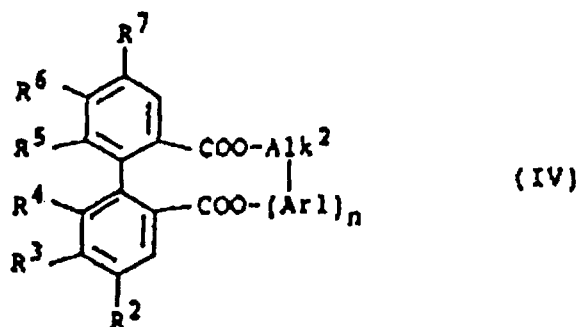


worin R^8 eine substituierte oder unsubstituierte Aminogruppe oder eine substituierte oder unsubstituierte C_{1-6} -Alkoxygruppe, oder ein Salz davon ist, umfaßt, worauf sich gegebenenfalls die Umwandlung des Produktes in ein pharmazeutisch annehmbares Salz davon anschließt.

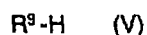


9. Verfahren nach Anspruch 8, worin R^8 eine Aminogruppe mit wahlweise einem oder zwei Substituenten, ausgewählt aus einer Hydroxy- C_{1-6} -alkylgruppe und einer C_{1-6} -Alkylgruppe, oder eine C_{1-6} -Alkoxygruppe ist, welche wahlweise mit einer C_{1-6} -Alkoxygruppe substituiert ist.

10. Verfahren zur Herstellung einer Verbindung der Formel (I) gemäß Anspruch 1, worin R^1 eine substituierte oder unsubstituierte C_{1-6} -Alkoxy-carbonylgruppe ist, unter der Voraussetzung, daß, wenn R^1 eine C_{1-6} -Alkoxy-carbonylgruppe ist, und wenn R^3 und R^4 sich unter Bildung einer C_{1-6} -Alkylendioxygruppe verbinden, R^2 kein Wasserstoffatom ist, oder ein pharmazeutisch annehmbares Salz davon, welches das Umsetzen einer zyklischen Diester-Verbindung der Formel:



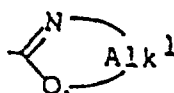
worin Alk^2 eine C_{1-6} -Alkyl-, $Ar1$ eine Arylgruppe ist, n ist 0 oder 1 und R^2 bis R^7 sind wie in Anspruch 1 definiert, oder ein Salz davon, mit einer Alkohol-Verbindung der Formel



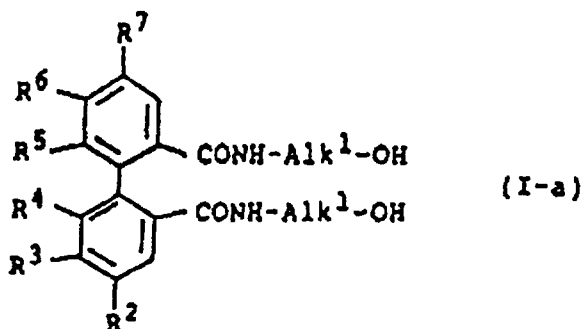
umfaßt, worin R^9 eine substituierte oder unsubstituierte C_{1-6} -Alkoxygruppe ist, woran sich gegebenenfalls die Umwandlung des Produkts in ein pharmazeutisch annehmbares Salz davon anschließt.

11. Verfahren nach Anspruch 10, worin R^9 eine C_{1-6} -Alkoxygruppe ist, die wahlweise durch eine C_{1-6} -Alkoxygruppe substituiert ist und $Ar1$ ist eine 1,2-Phenylengruppe.

12. Verfahren zur Herstellung einer Verbindung der Formel (I) nach Anspruch 1, worin R^1 eine Gruppe der Formel



ist (Alk^1 ist wie in Anspruch 1 definiert); oder ein pharmazeutisch annehmbares Salz davon, welches die Bildung eines intermolekularen Ringschlusses einer Bis(hydroxyalkylamid)-Verbindung der Formel

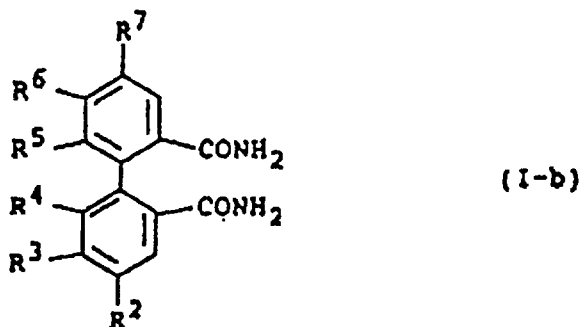


worin R^2 bis R^7 und Alk^1 wie in Anspruch 1 definiert sind, oder eines reaktiven Derivats davon oder



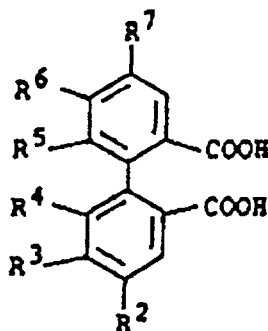
eines Salzes davon, umfaßt, woran sich gegebenenfalls die Umwandlung des Produktes in ein pharmazeutisch annehmbares Salz davon anschließt.

13. Verfahren zur Herstellung einer Verbindung der Formel (I) gemäß Anspruch 1, worin R¹ eine Cyano-
gruppe oder eine Aminothiocarbonylgruppe, oder ein pharmazeutisch annehmbares Salz davon ist,
welches die Dehydratisierung der Biphenylbis(carboxamid)-Verbindung der Formel:



umfaßt, worin R² bis R⁷ wie in Anspruch 1 definiert sind, oder eines Salzes davon, woran sich
gegebenenfalls eine Thiocarbonylierung, und gegebenenfalls zusätzlich eine Umwandlung des Produk-
tes in ein pharmazeutisch annehmbares Salz davon anschließt.

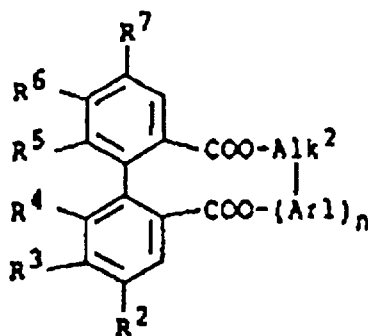
14. Biphenyldicarbonsäureverbindung der Formel



worin von R² bis R⁴ entweder eines von R² und R⁴ ein Wasserstoffatom oder eine C₁₋₆-Alkoxygruppe
ist und die beiden übrigen Gruppen eine C₁₋₆-Alkoxygruppe, eine Phenyl-C₁₋₆-alkoxygruppe oder eine
Hydroxygruppe sind, oder beide verbinden sich unter Bildung einer C₁₋₆-Alkylendioxygruppe; R⁵ und
R⁶ verbinden sich unter Bildung einer C₁₋₆-Alkylendioxygruppe und R⁷ ist ein Wasserstoffatom, unter
der Voraussetzung, daß wenn R³ und R⁴ sich unter Bildung einer C₁₋₆-Alkylendioxygruppe verbinden,
R² kein Wasserstoffatom ist, oder ein Salz davon.



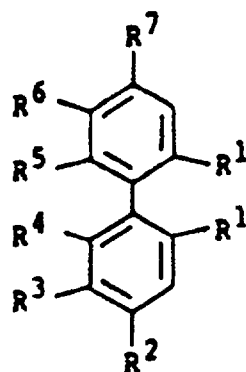
15. Zyklische Diesterverbindung der Formel:



worin Alk^2 eine $\text{C}_1\text{--}_6$ -Alkylengruppe ist, Ar^1 ist eine Arylengruppe, von R^2 bis R^4 ist entweder eines von R^2 und R^4 ein Wasserstoffatom oder eine $\text{C}_1\text{--}_6$ -Alkoxygruppe und die übrigen beiden Gruppen sind eine $\text{C}_1\text{--}_6$ -Alkoxygruppe, eine Phenyl- $\text{C}_1\text{--}_6$ -alkoxygruppe oder eine Hydroxygruppe, oder beide verbinden sich unter Bildung einer $\text{C}_1\text{--}_6$ -Alkylendioxygruppe; R^5 und R^6 verbinden sich unter Bildung einer $\text{C}_1\text{--}_6$ -Alkylendioxygruppe; R^7 ist ein Wasserstoffatom und n ist 0 oder 1, oder ein Salz davon.

Revendications

1. Dérivé biphényle de formule :



(I)

dans laquelle R^1 est un groupe aminocarbonyle substitué ou non substitué, un groupe aminothiocabonyle, un groupe alcoxy-carbonyle en $\text{C}_1\text{--}_6$ substitué ou non substitué, un groupe cyano, ou un groupe de formule



parmi R^2 à R^4 , soit l'un de R^2 et de R^4 est un atome d'hydrogène ou un groupe alcoxy en $\text{C}_1\text{--}_6$ et les deux groupes restant sont un groupe alcoxy en $\text{C}_1\text{--}_6$, un groupe phényl-alcoxy en $\text{C}_1\text{--}_6$ ou un groupe hydroxy, soit les deux se combinent pour former un groupe alkylendioxy en $\text{C}_1\text{--}_6$; R^5 et R^6 se combinent pour former un groupe alkylendioxy en $\text{C}_1\text{--}_6$; R^7 est un atome d'hydrogène; et Alk^1 est un groupe alkylène en $\text{C}_1\text{--}_6$ étant entendu que quand R^1 est un groupe carbonyl-alcoxy en $\text{C}_1\text{--}_6$, et quand R^3 et R^4 se combinent pour former un groupe alkylendioxy en $\text{C}_1\text{--}_6$, R^2 n'est pas un atome d'hydrogène; ou un de ses sels pharmaceutiquement acceptables.

2. Composé selon la revendication 1, dans lequel R^1 est un groupe aminocarbonyle ayant de façon facultative un ou deux substituants sélectionnés parmi un groupe hydroxyalkyle en $\text{C}_1\text{--}_6$ et un groupe



alkyle en C₁₋₆ ; un groupe aminothiocabonyl ; un groupe carbonyl-alcoxy en C₁₋₆ étant de façon facultative substitué par un groupe alcoxy en C₁₋₆ ; un groupe cyano ; ou un groupe 4,4-di(alkyle en C₁₋₆)-4,5-dihydro-2-oxazolyle ; étant entendu que quand R¹ est un groupe carbonyl-alcoxy en C₁₋₆, et quand R³ et R⁴ se combinent pour former un groupe alkylènedioxy en C₁₋₆, R² n'est pas un atome d'hydrogène.

3. Composé selon la revendication 2, dans lequel R² et R³ sont chacun un groupe alcoxy en C₁₋₆ ou se combinent ensemble pour former un groupe alkylènedioxy en C₁₋₆, R⁴ est un atome d'hydrogène, R⁵ et R⁶ se combinent pour former un groupe alkylènedioxy en C₁₋₆, et R⁷ est un atome d'hydrogène.

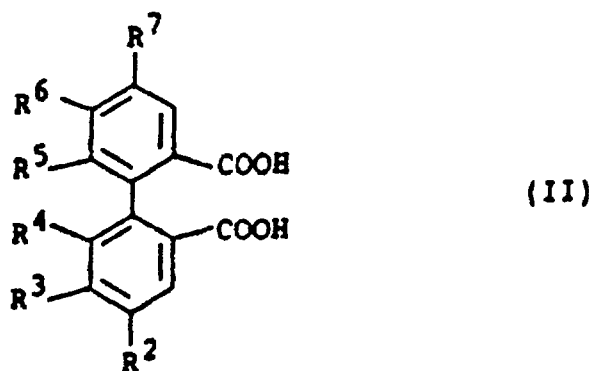
4. Composé selon la revendication 2, dans lequel R¹ est un groupe aminocarbonyl ayant de façon facultative un ou deux substituants sélectionnés parmi un groupe hydroxyalkyle en C₁₋₆ et un groupe alkyle en C₁₋₆ ; un groupe aminothiocabonyl ; un groupe carbonyl-alcoxy en C₁₋₆ ; un groupe cyano ; ou un groupe 4,4-di(alkyle en C₁₋₆)-4,5-dihydro-2-oxazolyle, R² est un atome d'hydrogène, R³ et R⁴ se combinent pour former un groupe alkylènedioxy en C₁₋₆, R⁵ et R⁶ se combinent pour former un groupe alkylènedioxy en C₁₋₆ et R⁷ est un atome d'hydrogène.

5. Composé selon la revendication 2, dans lequel R² à R⁴ sont chacun un groupe alcoxy en C₁₋₆, R⁵ et R⁶ se combinent pour former un groupe alkylènedioxy en C₁₋₆ et R⁷ est un atome d'hydrogène.

6. Composé selon la revendication 2, dans lequel R² est un groupe alcoxy en C₁₋₆, R³ et R⁴ se combinent pour former un groupe alkylènedioxy en C₁₋₆ ou R³ est un groupe phényl-alcoxy en C₁₋₆ et R⁴ est un atome d'hydrogène, R⁵ et R⁶ se combinent pour former un groupe alkylènedioxy en C₁₋₆, et R⁷ est un atome d'hydrogène.

7. Composé selon la revendication 2, dans lequel R² est un groupe alcoxy en C₁₋₆, R³ est un groupe hydroxy, R⁴ est un atome d'hydrogène, R⁵ et R⁶ se combinent pour former un groupe alkylènedioxy en C₁₋₆, et R⁷ est un atome d'hydrogène.

8. Procédé pour la préparation d'un composé de formule (I) comme exposé dans la revendication 1, dans laquelle R¹ est un groupe aminocarbonyl substitué ou non substitué ou un groupe carbonyl-alcoxy en C₁₋₆ substitué ou non substitué, étant entendu que quand R¹ est un groupe carbonyl-alcoxy en C₁₋₆ et quand R³ et R⁴ se combinent pour former un groupe alkylènedioxy en C₁₋₆, R² n'est pas un atome d'hydrogène, ou d'un de ses sels pharmaceutiquement acceptables, qui comprend la réaction d'un acide biphenyle dicarboxylique de formule :



dans laquelle R² à R⁷ sont tels que définis dans la revendication 1, ou d'un dérivé réactif de ses groupes carboxyles ou d'un de ses sels avec un composé de formule :

R⁸-H (III)

dans laquelle R⁸ est un groupe amino substitué ou non substitué ou un groupe alcoxy en C₁₋₆ substitué ou non substitué, ou l'un de ses sels, suivi de façon facultative par la conversion du produit en l'un de ses sels pharmaceutiquement acceptables.



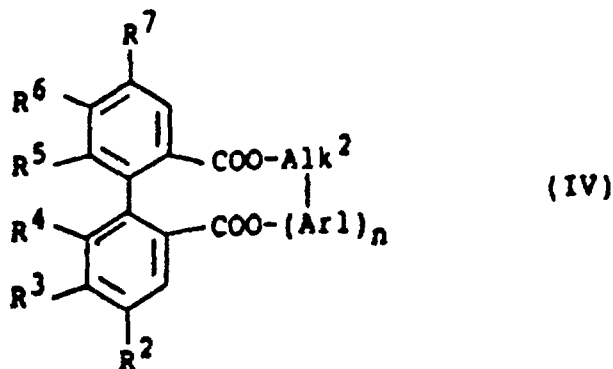
9. Procédé selon la revendication 8, dans lequel R^8 est un groupe amino ayant de façon facultative un ou deux substituants sélectionnés parmi un groupe hydroxy-alkyl en C_{1-6} et un groupe alkyl en C_{1-6} , ou un groupe alcoxy en C_{1-6} substitué de façon facultative par un groupe alcoxy en C_{1-6} .

5 10. Procédé pour la préparation d'un composé de formule (I) tel qu'exposé dans la revendication 1 dans laquelle R^1 est un groupe carbonyl-alcoxy en C_{1-6} substitué ou non substitué, étant entendu que quand R^1 est un groupe carbonyl-alcoxy en C_{1-6} , et quand R^3 et R^4 se combinent pour former un groupe alkylènedioxy en C_{1-6} , R^2 n'est pas un atome d'hydrogène, ou d'un de ses sels pharmaceutiquement acceptables, qui comprend la réaction d'un composé diester cyclique de formule :

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25 dans laquelle Alk^2 est un alkylène en C_{1-6} , $Ar1$ est un groupe arylène, n est 0 ou 1, et R^2 à R^7 sont tels que définis dans la revendication 1, ou d'un de ses sels avec un composé alcool de formule :



30 dans laquelle R^9 est un groupe alcoxy en C_{1-6} substitué ou non substitué, suivi de façon facultative par la conversion du produit en l'un de ses sels pharmaceutiquement acceptables.

11. Procédé selon la revendication 8, dans lequel R^9 est un groupe alcoxy en C_{1-6} substitué de façon facultative par un groupe alcoxy en C_{1-6} , et $Ar1$ est un groupe 1,2-phénylène.

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12. Procédé pour la préparation d'un composé de formule (I) tel qu'exposé dans la revendication 1 dans laquelle R^1 est un groupe de formule :

40



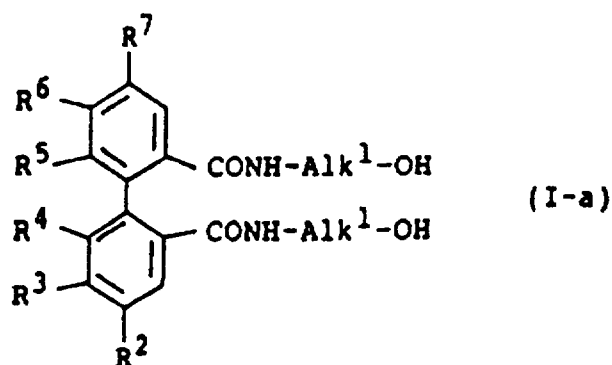
(Alk^1 est tel que défini dans la revendication 1) ; ou d'un de ses sels pharmaceutiquement acceptables,

45

qui consiste à soumettre un composé bis(hydroxyalkylamide) de formule :

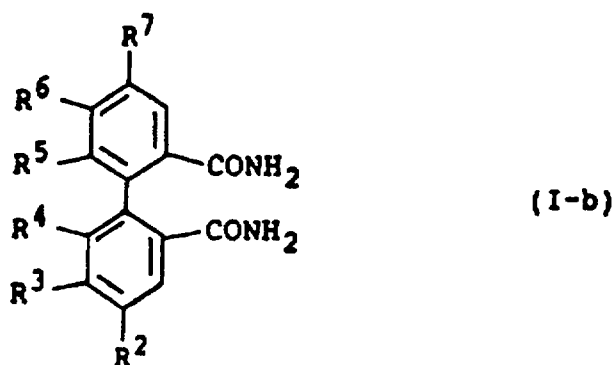
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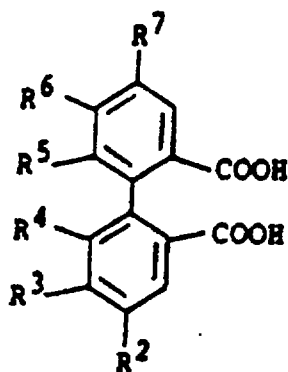
15 dans laquelle R^2 à R^7 et Alk^1 sont tels que définis dans la revendication 1, ou un de ses dérivés réactifs ou l'un de ses sels, à une cyclisation intramoléculaire, suivie de façon facultative par la conversion du produit en l'un de ses sels pharmaceutiquement acceptables.

20 13. Procédé pour la préparation d'un composé de formule (I) tel que défini dans la revendication 1 dans laquelle R^1 est un groupe cyano ou un groupe aminothiocarbonyle, ou d'un de ses sels pharmaceutiquement acceptables, qui comprend la déshydratation d'un composé biphenylebis(carboxamide) de formule :



40 dans laquelle R^2 à R^7 sont tels que définis dans la revendication 1, ou d'un de ses sels, suivie de façon facultative par une thiocarbonylation, et ensuite suivie de façon facultative par la conversion du produit en l'un de ses sels pharmaceutiquement acceptables.

45 14. Composé d'acide biphenyldicarboxylique de formule :

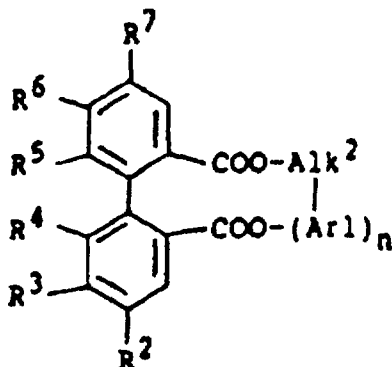


dans laquelle parmi R^2 à R^4 , soit l'un de R^2 et R^4 est un atome d'hydrogène ou un groupe alcoxy en



C_{1-6} et les deux groupes restant sont un groupe alcoxy en C_{1-6} , un groupe phényl-alcoxy en C_{1-6} ou un groupe hydroxy, soit les deux se combinent pour former un groupe alkylènedioxy en C_{1-6} , R^5 et R^6 se combinent pour former un groupe alkylènedioxy en C_{1-6} ; et R^7 est un atome d'hydrogène, étant entendu que, quand R^3 et R^4 se combinent pour former un groupe alkylènedioxy en C_{1-6} , R^2 n'est pas un atome d'hydrogène, ou l'un de ses sels.

15. Composé diester cyclique de formule :



dans laquelle Alk^2 est un groupe alkylène en C_{1-6} ; $Ar1$ est un groupe arylène; parmi R^2 à R^4 , soit l'un de R^2 et R^4 est un atome d'hydrogène ou un groupe alcoxy en C_{1-6} et les deux groupes restant sont un groupe alcoxy en C_{1-6} , un groupe phényl-alcoxy en C_{1-6} ou un groupe hydroxy, soit les deux se combinent pour former un groupe alkylènedioxy en C_{1-6} ; R^5 et R^6 se combinent pour former un groupe alkylènedioxy en C_{1-6} ; R^7 est un atome d'hydrogène, et n est 0 ou 1, ou l'un de ses sels.