



159159

PROCESS FOR PREPARING NITRILE COMPOUNDS FROM
ETHYLENICALLY UNSATURATED COMPOUNDS

The present invention concerns a process for
5 hydrocyanating ethylenically unsaturated organic
compounds to compounds containing at least one nitrile
function.

It relates more particularly to the hydrocyanation of
10 diolefins such as butadiene or of substituted olefins
such as alkenenitriles, for instance pentenenitriles.
The hydrocyanation of butadiene to pentenenitriles is
an important reaction which has been implemented
industrially for a number of years, particularly in the
15 process for synthesizing adiponitrile, a major chemical
intermediate which allows access in particular to the
monomers of numerous polymers, foremost among which are
the polyamides.

20 French patent 1 599 761 describes a process for
preparing nitriles by addition of hydrocyanic acid with
organic compounds having at least one ethylenic double
bond, in the presence of a nickel catalyst and a
triaryl phosphite. This reaction may be conducted in
25 the presence or absence of a solvent.

When a solvent is used in this prior art process it is
preferably a hydrocarbon, such as benzene or xylenes,
or a nitrile such as acetonitrile.

30 The catalyst employed is an organometallic complex of
nickel, containing ligands such as phosphines, arsines,
stibines, antimonites, arsenites, phosphites, phos-
phinites or phosphonites.

35 The processes for hydrocyanating dienes generally
comprise two steps: a first hydrocyanation, leading to
branched and linear unsaturated mononitriles, and a
second step which allows the dinitriles to be obtained.



G00002

Often only the linear nitriles are of interest for the synthesis of new products such as, for example, adiponitrile. These processes therefore also include an intermediate step, which is referred to as the isomerization step and consists in treating the branched unsaturated mononitriles to convert them into linear unsaturated mononitriles.

The presence of a promoter to activate the catalyst, such as a boron compound or a metal salt, generally a Lewis acid, is also recommended for carrying out the second step.

Patent FR-A-2 338 253 proposed carrying out the hydrocyanation of compounds having at least one ethylenic unsaturation in the presence of an aqueous solution of a compound of a transition metal, particularly nickel, palladium or iron, and of a sulphonated phosphine.

The sulphonated phosphines described in said patent are sulphonated triarylphosphines and more particularly sulphonated triphenylphosphines.

This process permits proper hydrocyanation, particularly of the butadiene and the pentenenitriles, easy separation of the catalytic solution, by simple decanting, and, consequently, goes as far as possible to avoid the discharge of effluent or of waste containing the metals used as catalyst.

Research has nevertheless been conducted in order to find new catalytic systems which give greater performance in terms of catalytic activity and of selectivity and stability.

One of the aims of the present invention is to propose a new class of ligands which with the transition metals make it possible to obtain catalytic systems exhibiting

The present invention accordingly provides a process
5 for hydrocyanating a hydrocarbon compound containing at
least one ethylenic unsaturation by reacting it in a
liquid medium with hydrogen cyanide in the presence of
a catalyst comprising a metallic element selected from
transition metals and an organic ligand, characterized
10 in that the organic ligand corresponds to the general
formula I below:



15

R₁, R₂, R₃ and R₄, which are identical or different, represent a substituted or unsubstituted, aromatic, aliphatic or cycloaliphatic radical comprising one or more rings, which are in fused form or not and which may contain one or more heteroatoms, where the radicals R₁ and R₂ on the one hand and R₃ and R₄ on the other hand may be interconnected by a covalent bond, a hydrocarbon chain or a heteroatom.

U₁, U₂, U₃ and U₄, which are identical or different, represent an oxygen atom or a radical of formula NR in which R denotes a monovalent alkyl, aryl, cycloalkyl, 30 sulphonyl or carbonyl radical,

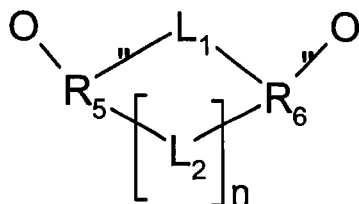
R_5 and R_6 , which are identical or different, represent an aryl or cycloaliphatic group which may comprise heteroatoms and/or one or more rings, in fused form or not, and which are substituted or unsubstituted,

n is an integer equal to 0 or 1,

L_1 , when n is 0, represents a divalent radical selected from the group consisting of the groups NR_7 , PR_8 , SiR_9R_{10} , BR_{11} , S , POR_{12} , SO_2 and CO , in which R_7 is as defined for R above, R_8 and R_{12} may represent the radical OR_{13} , and R_8 , R_9 , R_{10} , R_{11} , R_{12} and R_{13} represent alkyl, aryl or cycloalkyl radicals,

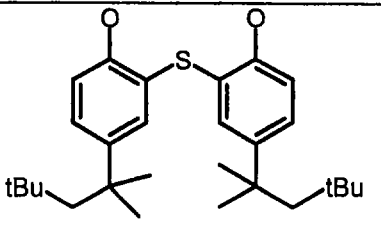
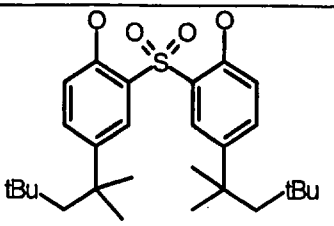
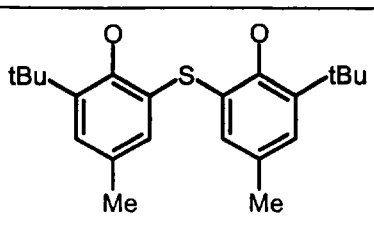
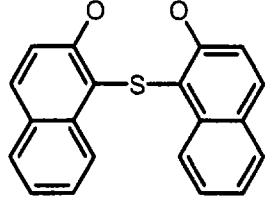
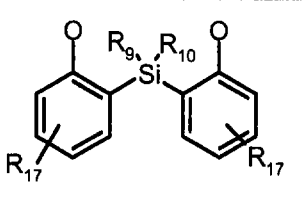
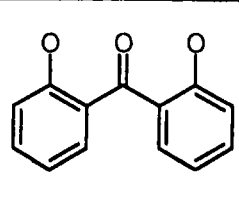
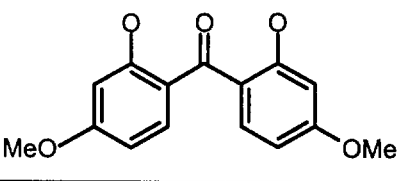
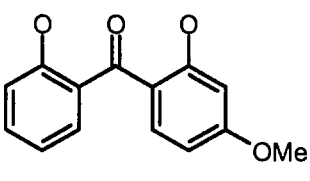
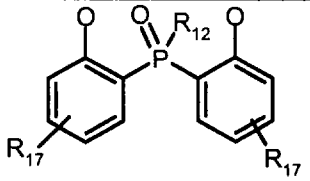
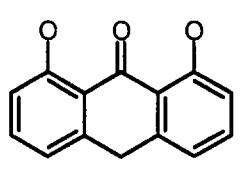
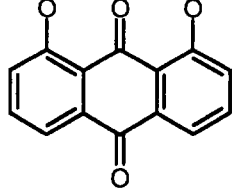
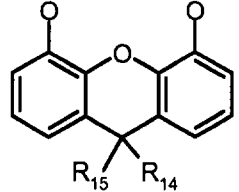
L_1 and L_2 , when n is 1, are identical or different and represent a covalent bond or a radical selected from the group consisting of the groups O , NR_7 , PR_8 , SiR_9R_{10} , BR_{11} , S , POR_{12} , SO_2 , CO and $-CR_{14}R_{15}-$, in which R_7 is as defined for R above, R_8 and R_{12} may represent the radical OR_{13} , and R_8 , R_9 , R_{10} , R_{11} , R_{12} , R_{13} , R_{14} and R_{15} represent alkyl, aryl or cycloalkyl radicals, it being possible also for R_{14} and R_{15} to represent a hydrogen atom.

Examples of structures



include the following formulae listed in Table I below:

Table I

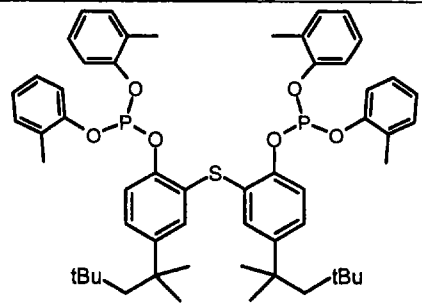
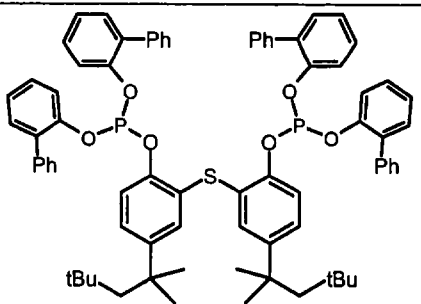
in which R₁₇ represents an alkyl, aryl, halogen, alkoxy, thiol, cyano, nitro, aryloxy, alkoxycarbonyl, acyl or formyl radical.

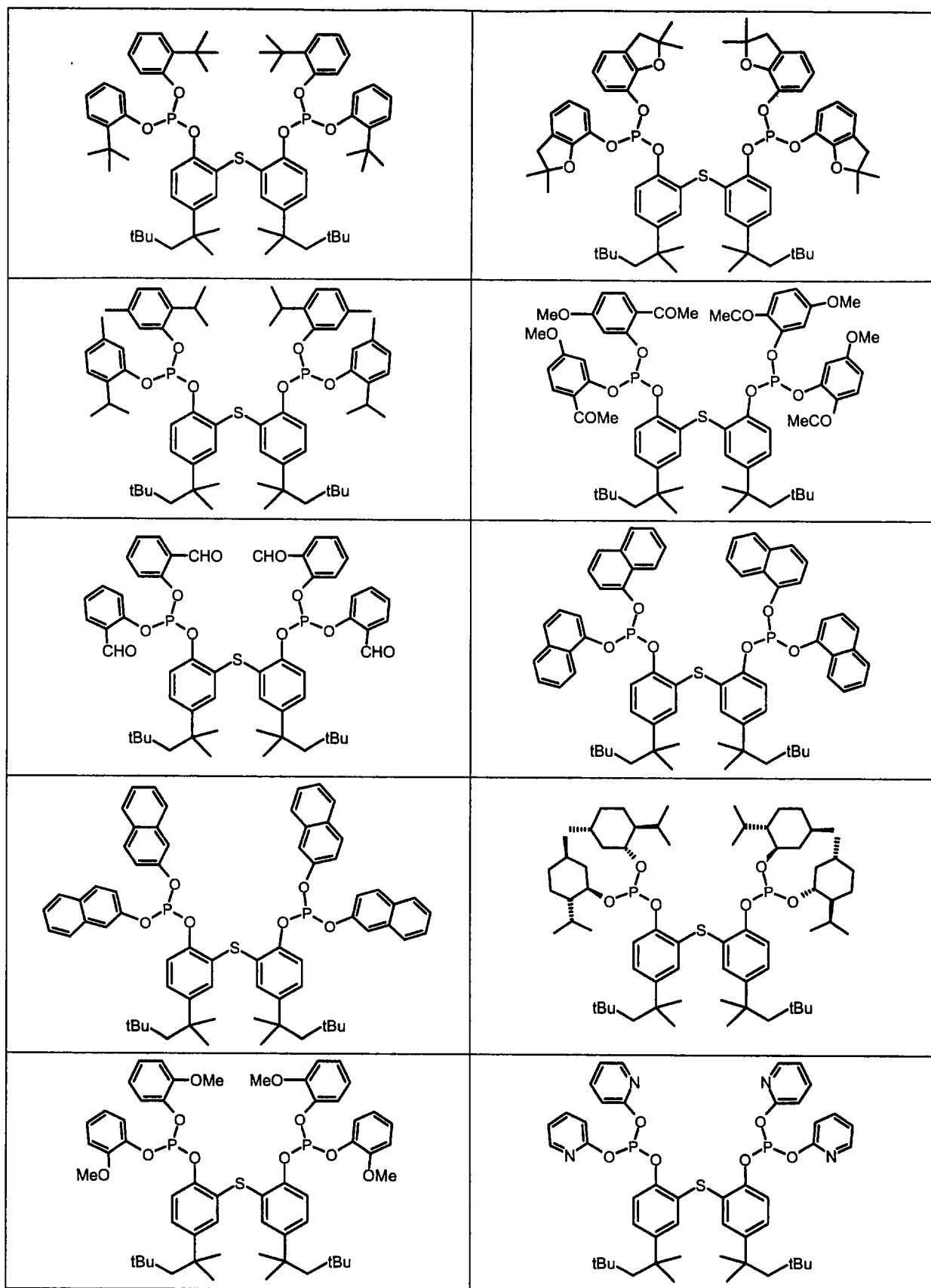
5

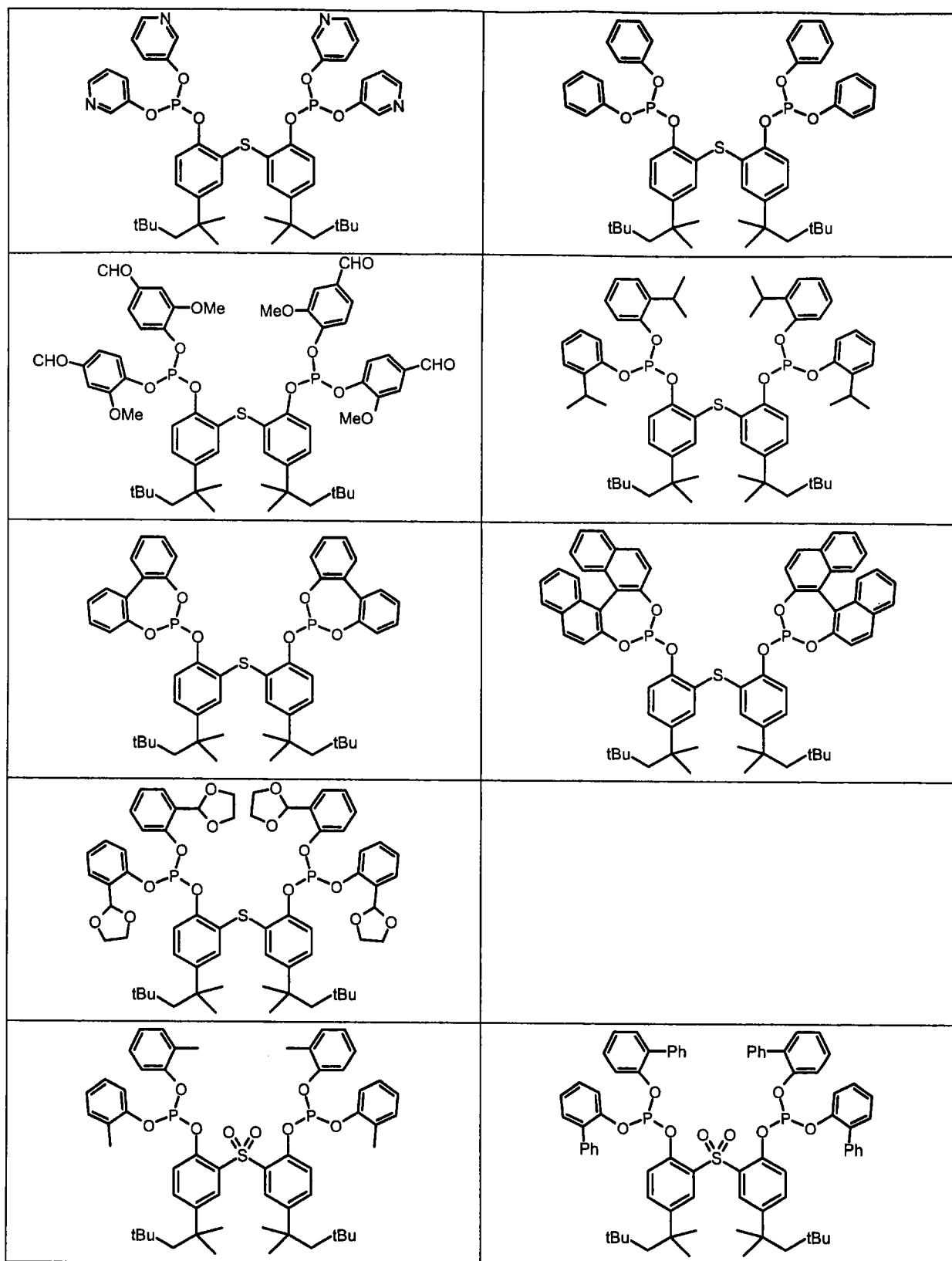
Ligands in accordance with the invention include the following compounds listed in Table II below:

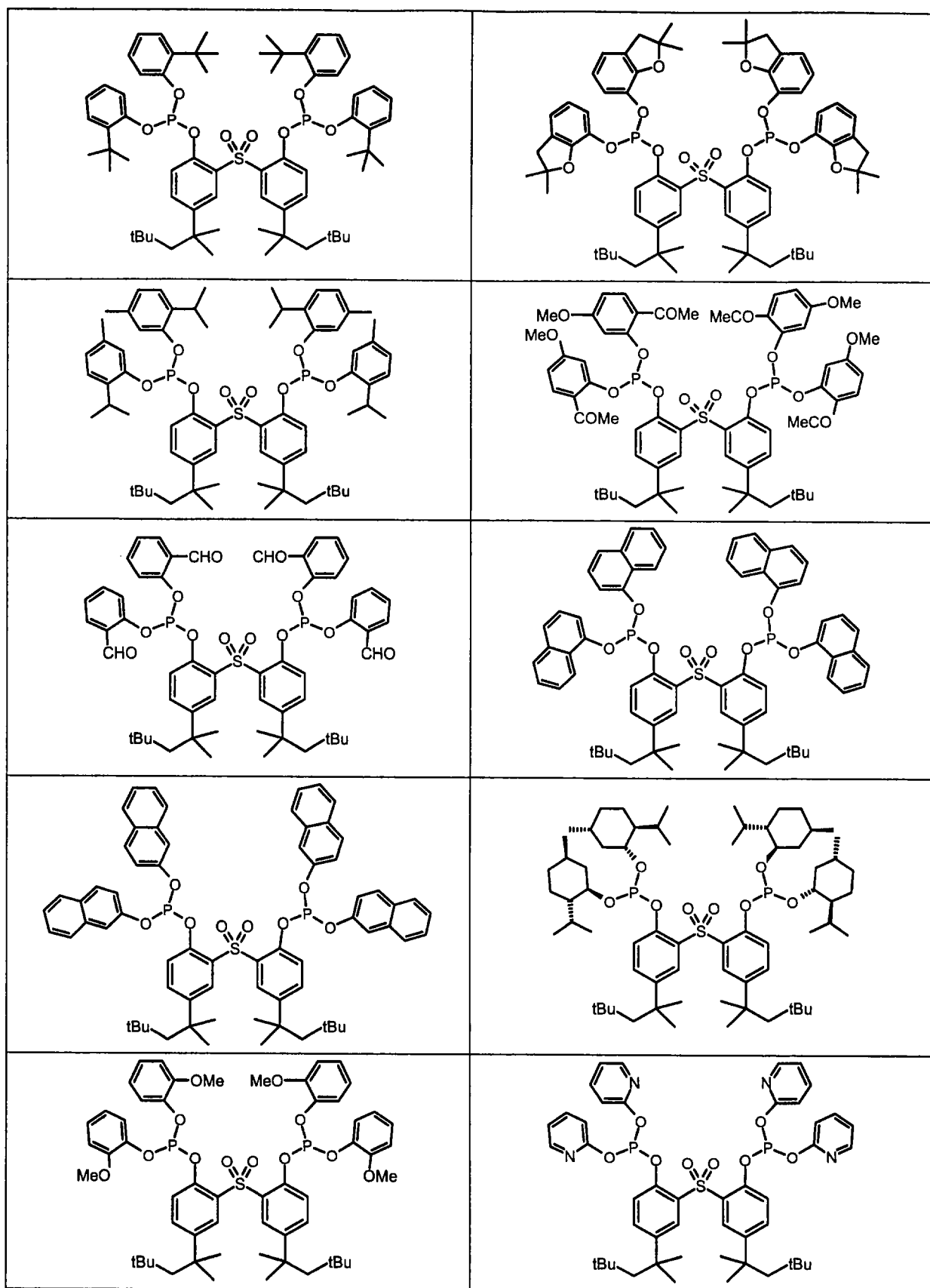
Table II

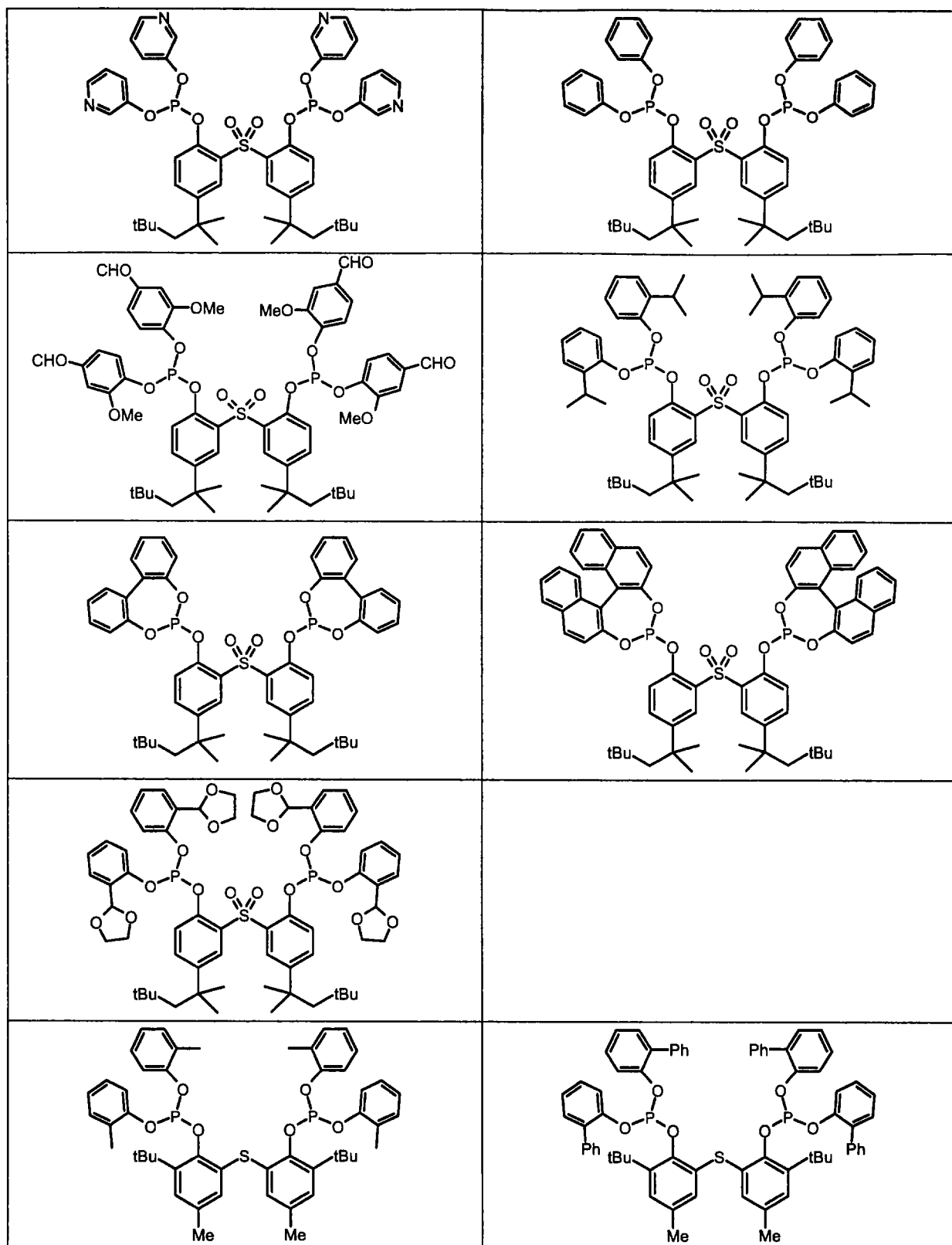
10

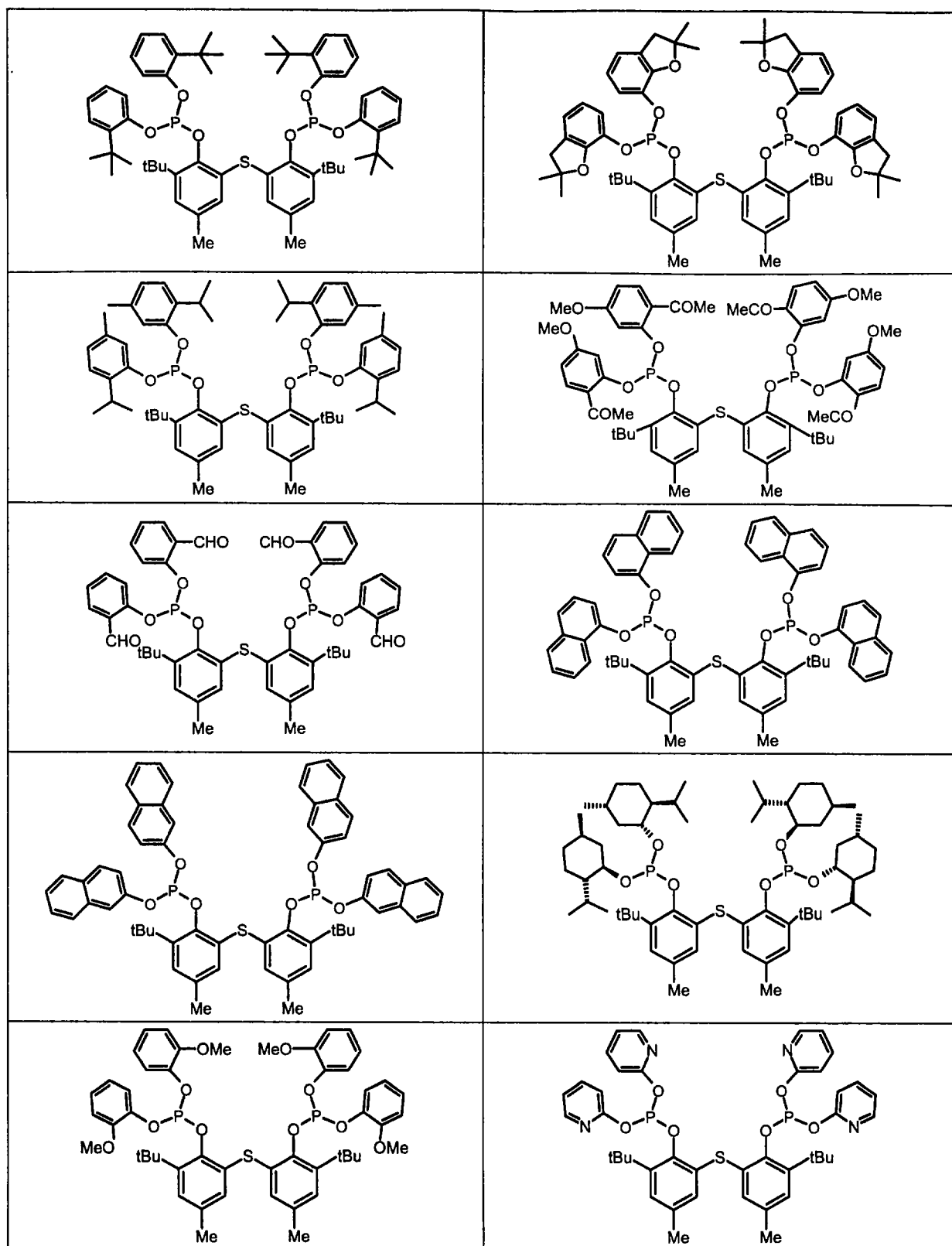
	
---	---

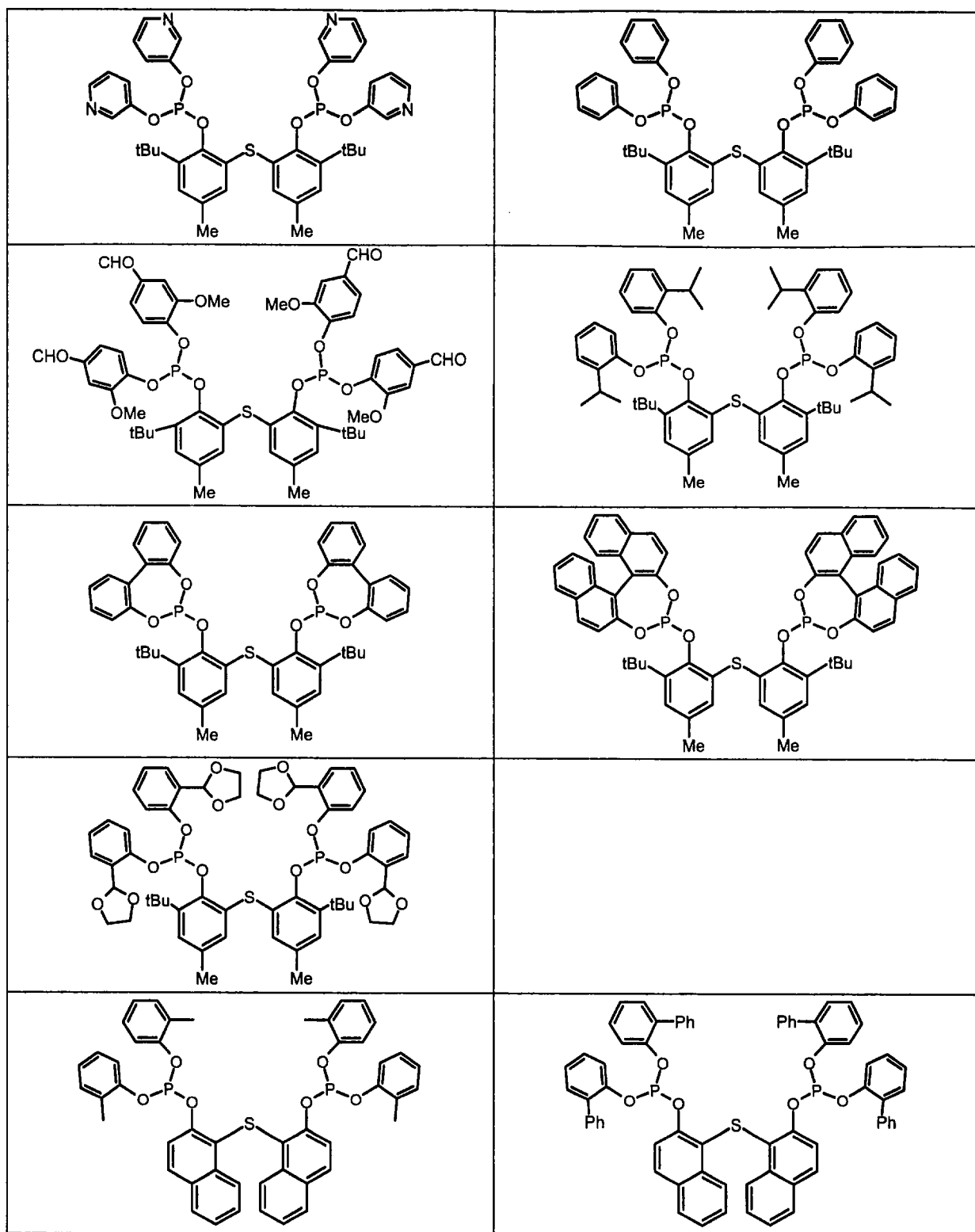


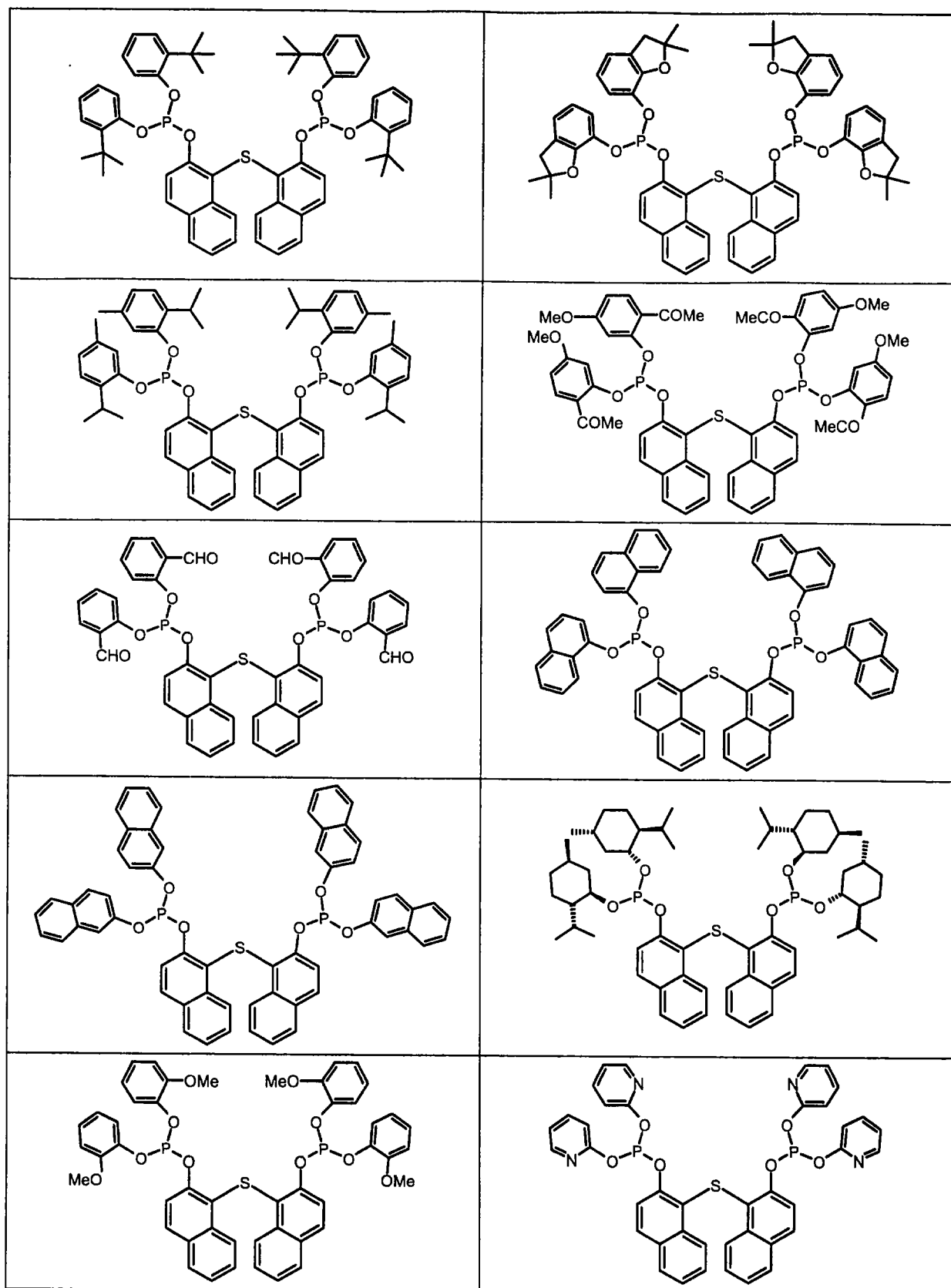


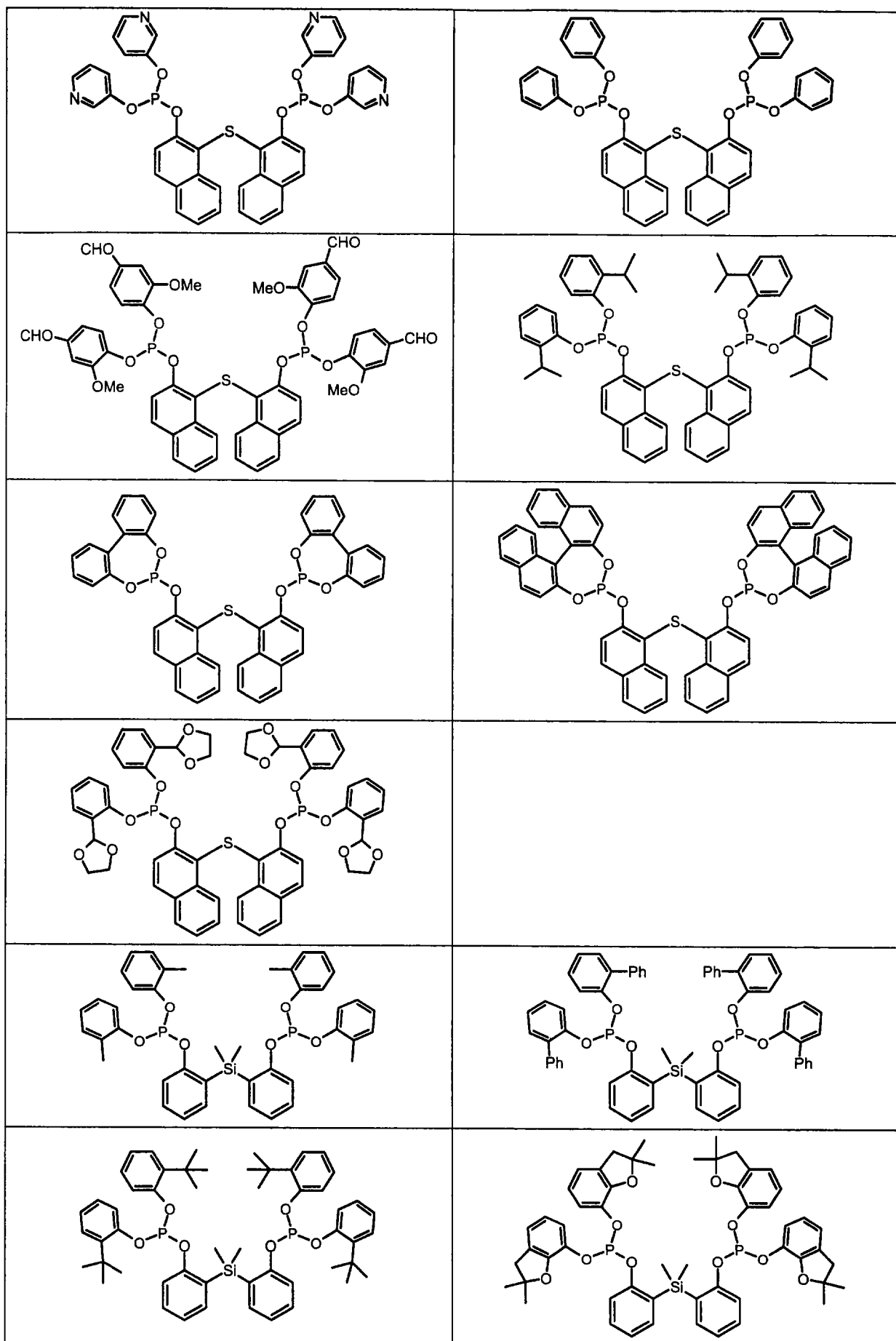


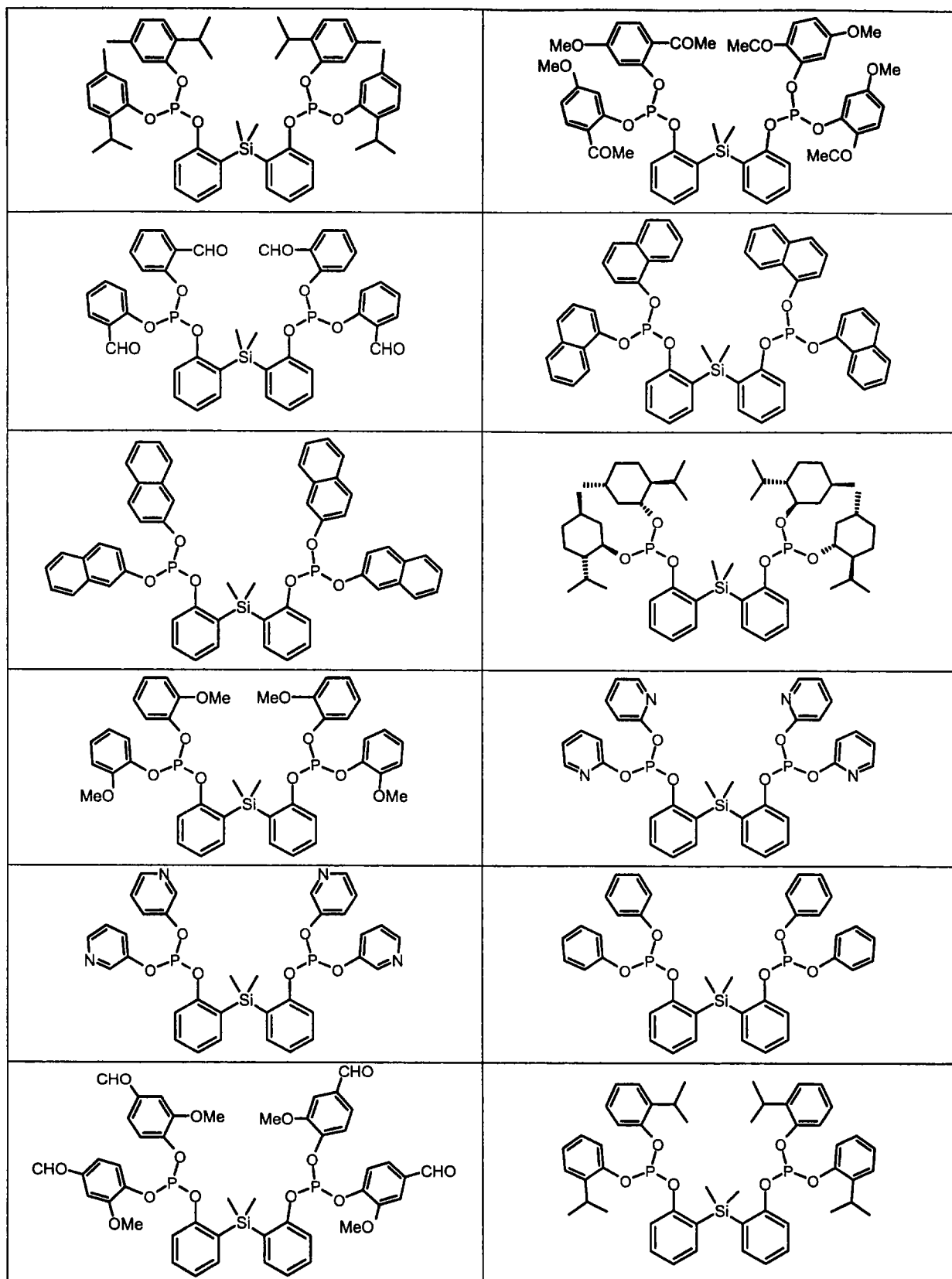


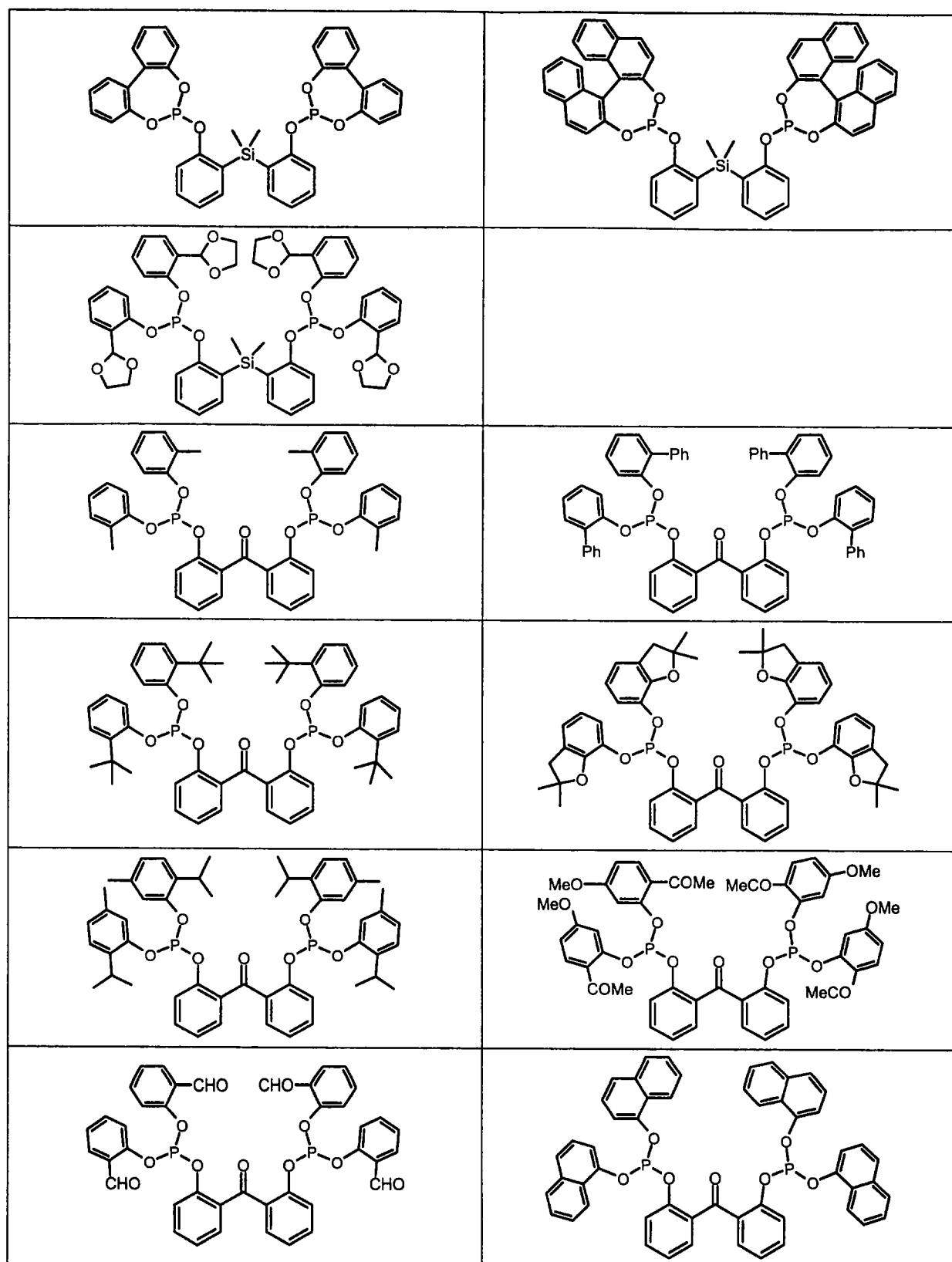


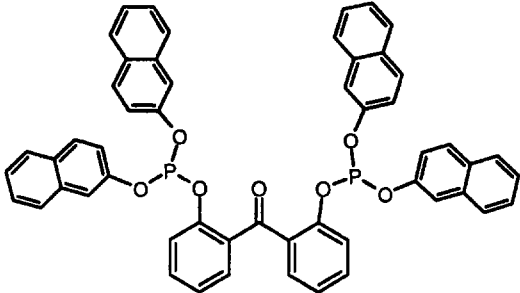
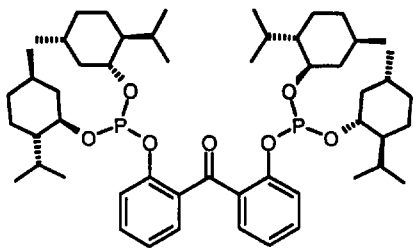
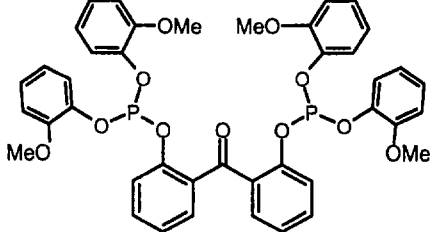
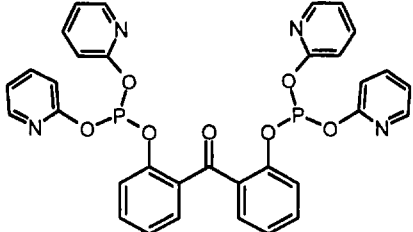
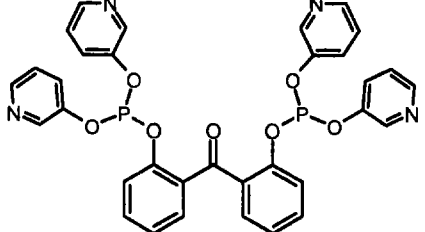
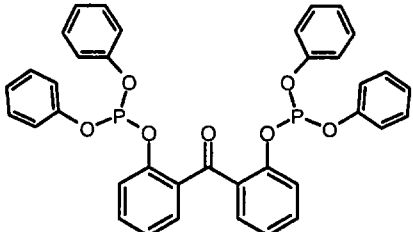
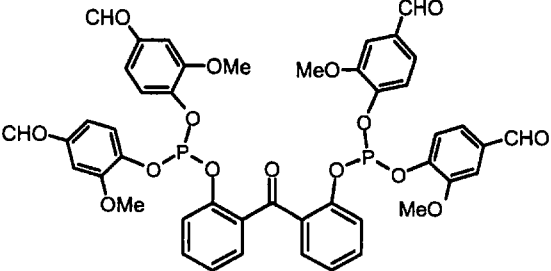
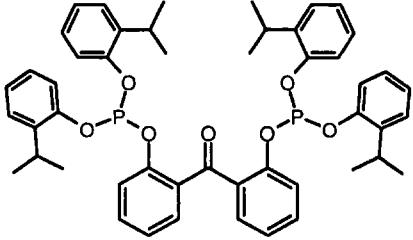
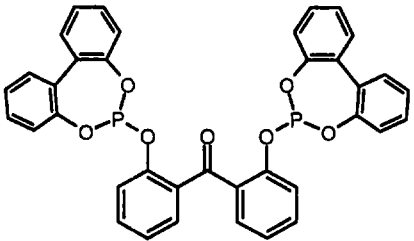
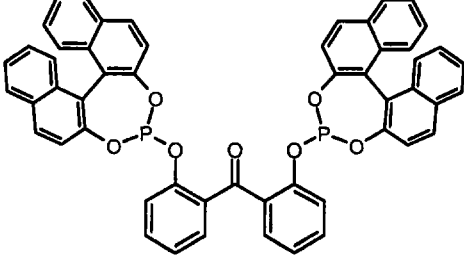
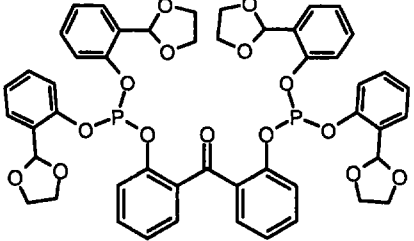


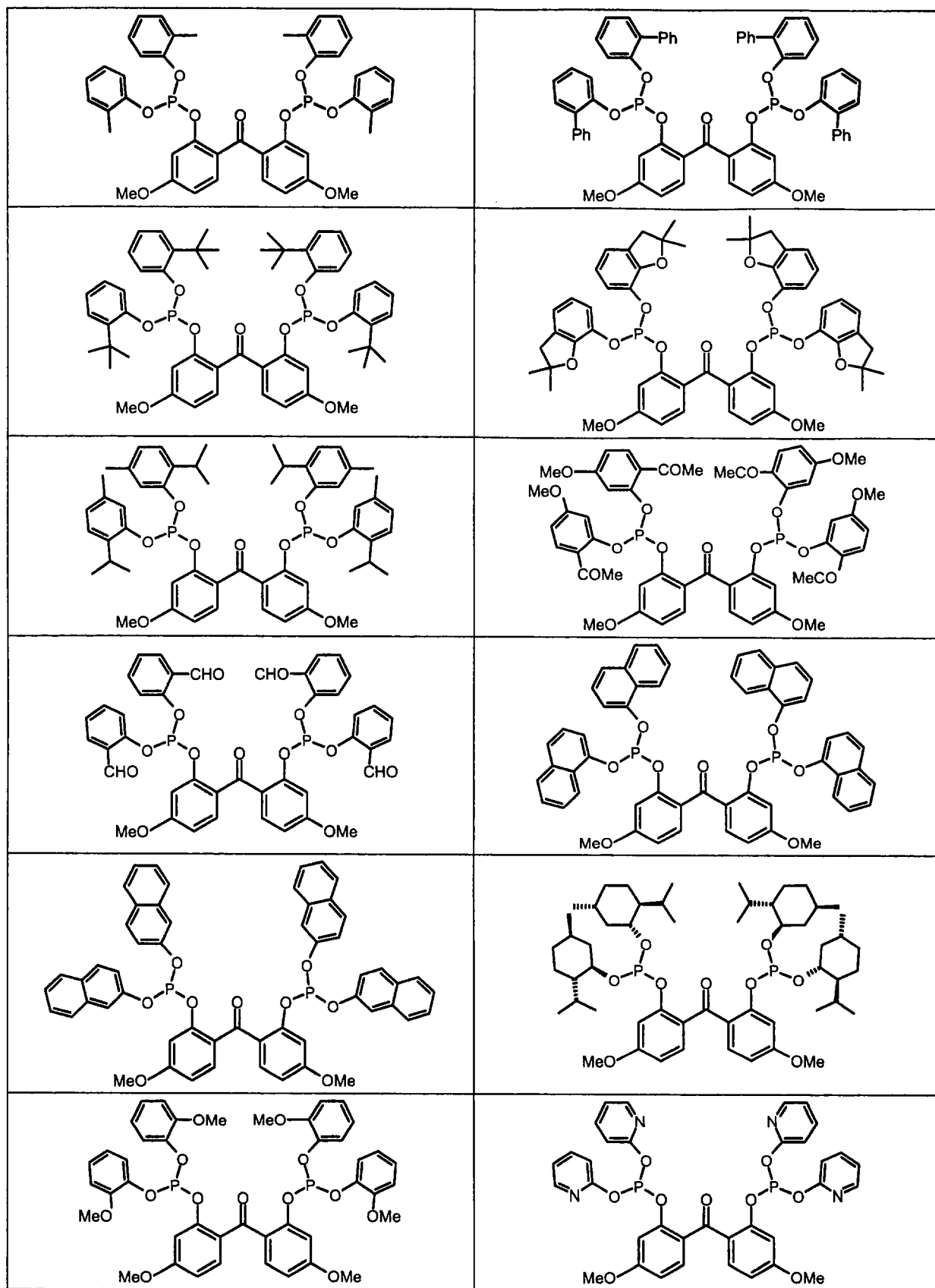


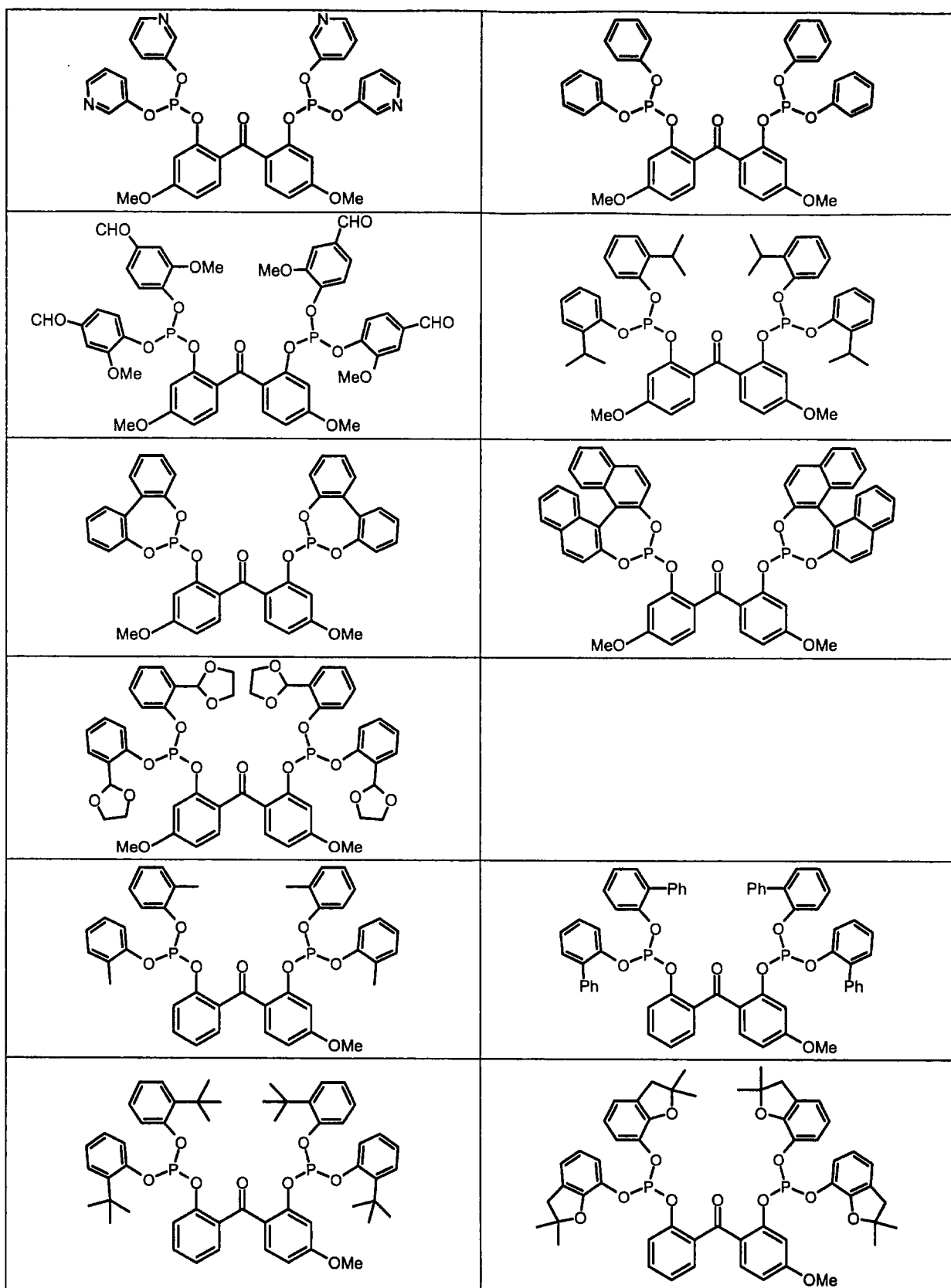


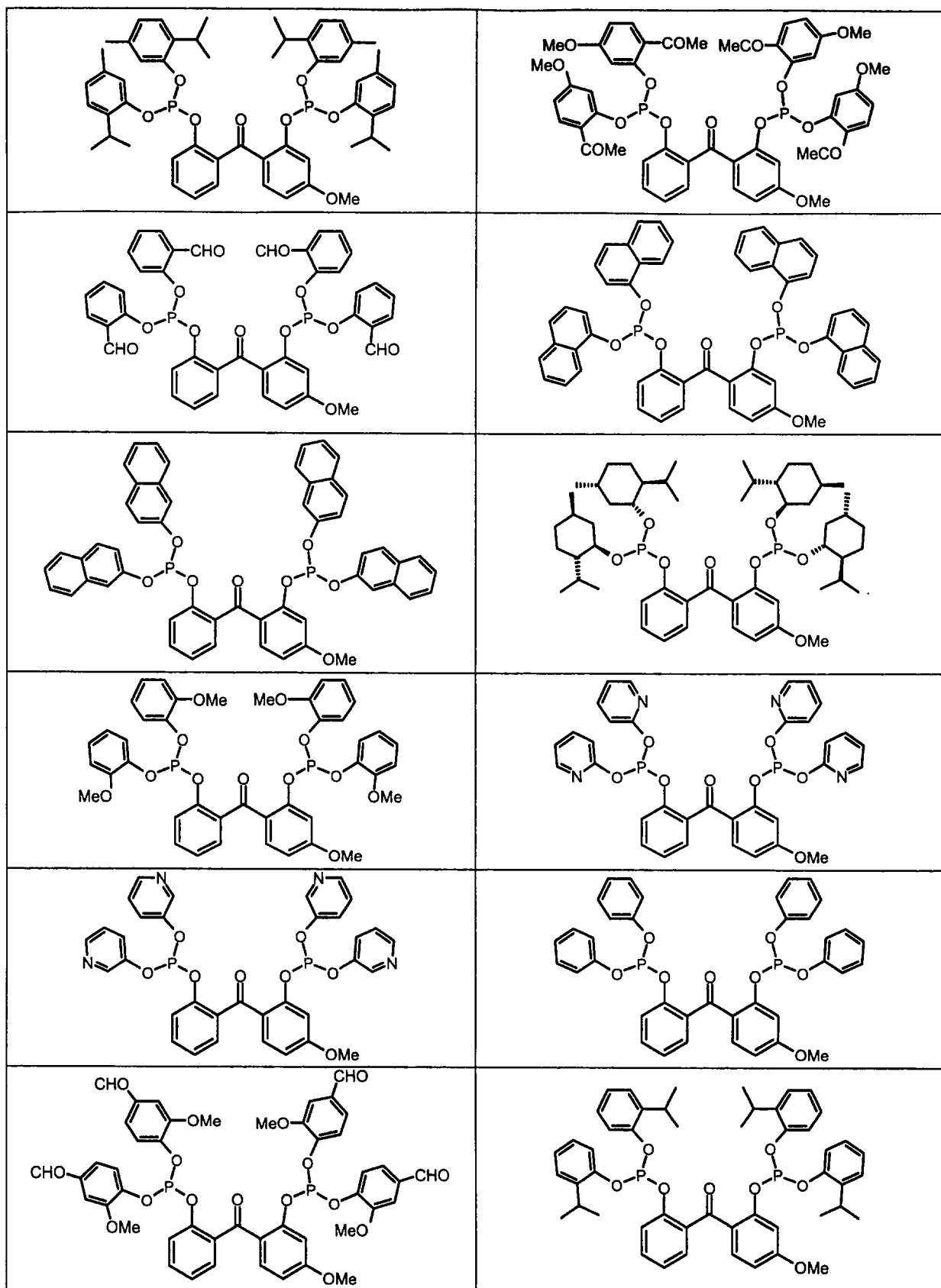


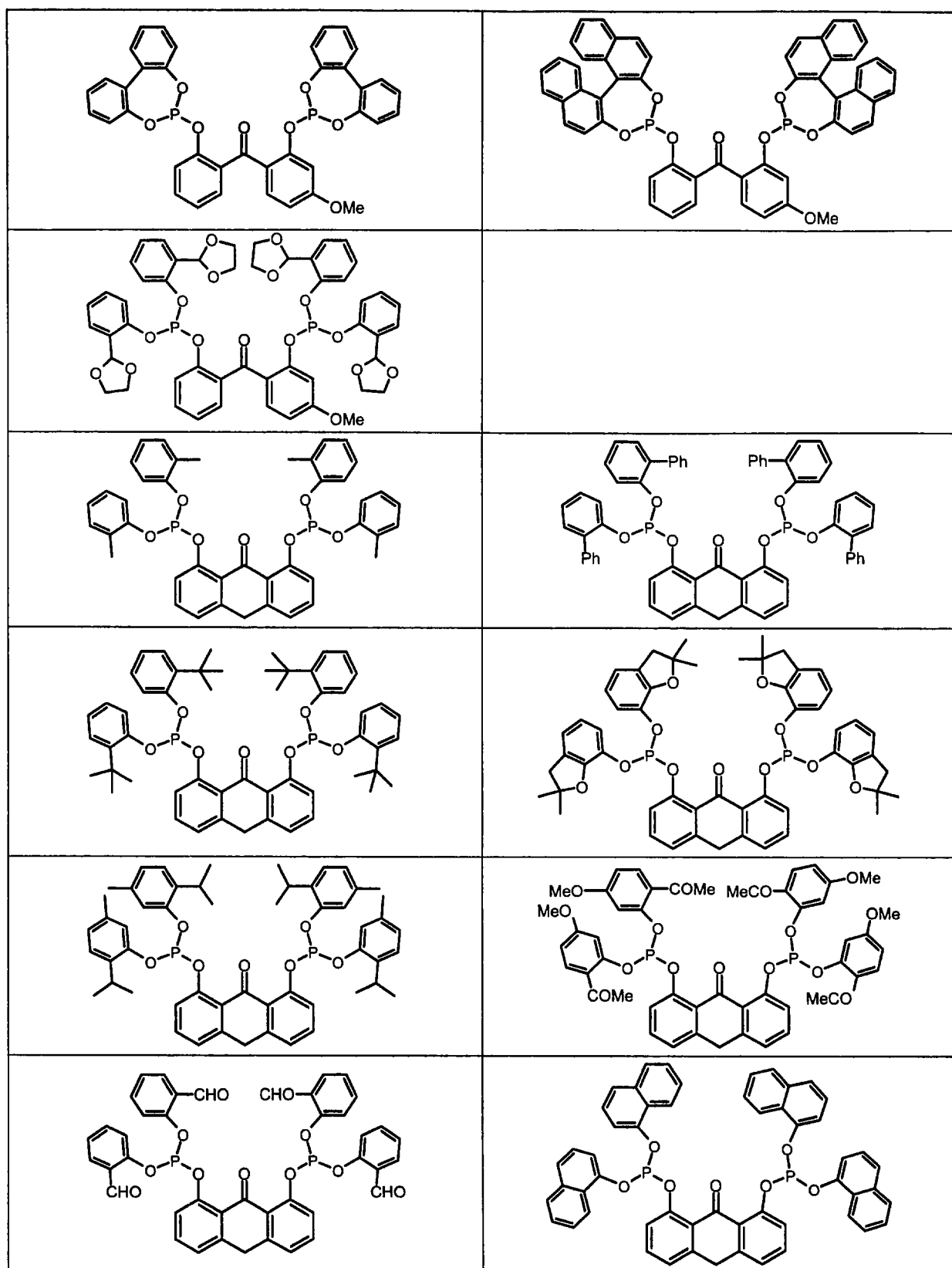


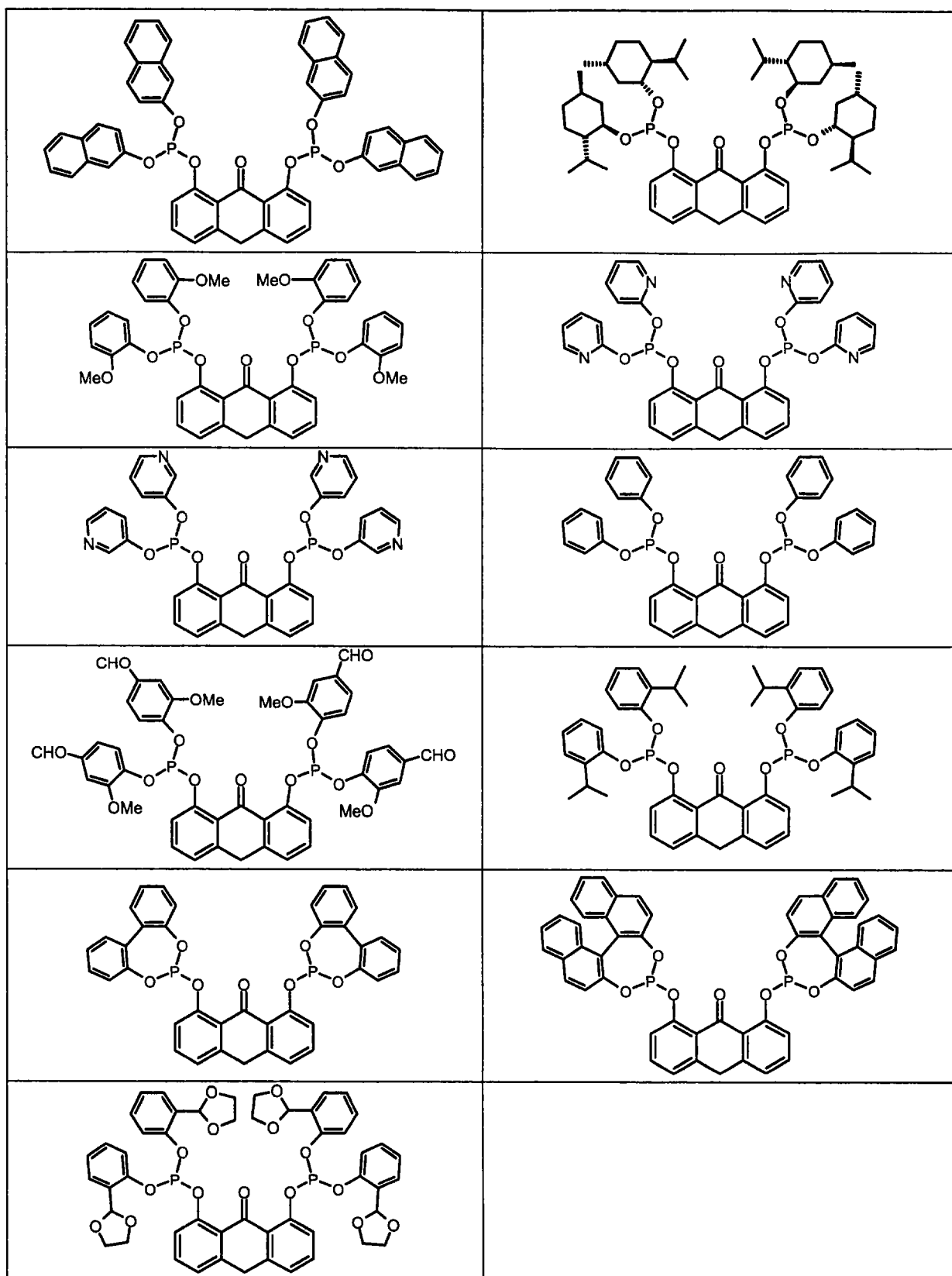
	
	
	
	
	
	

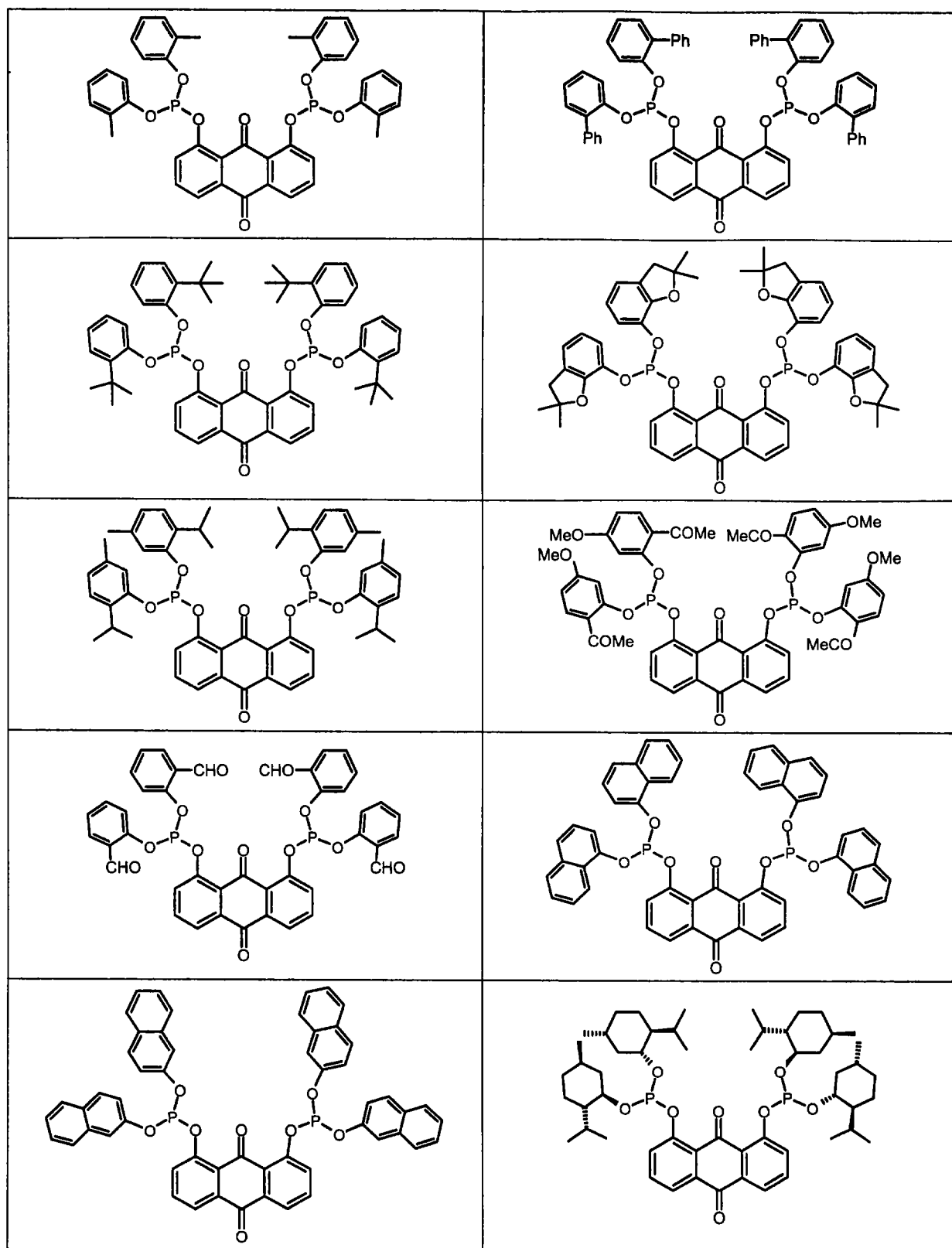


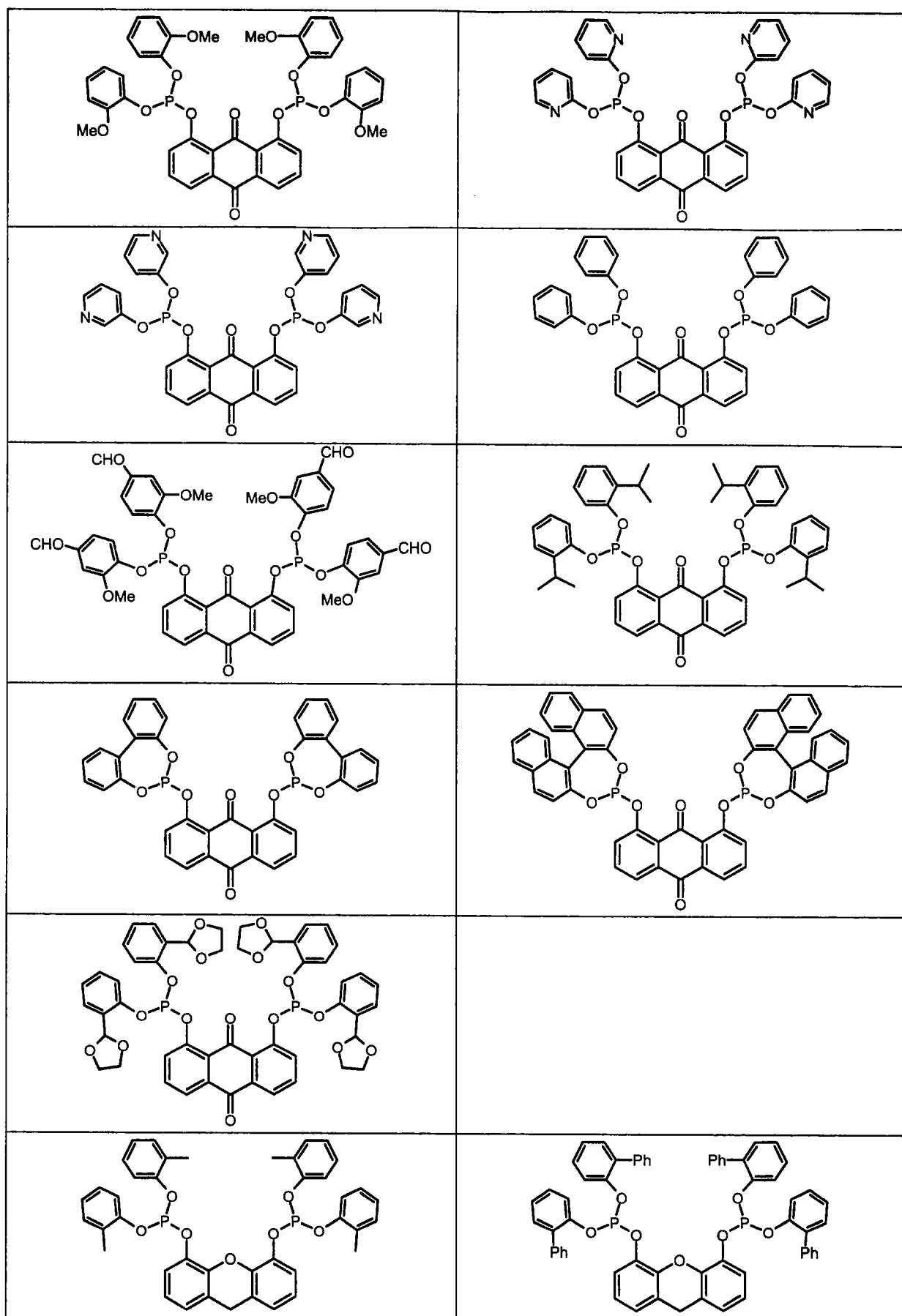


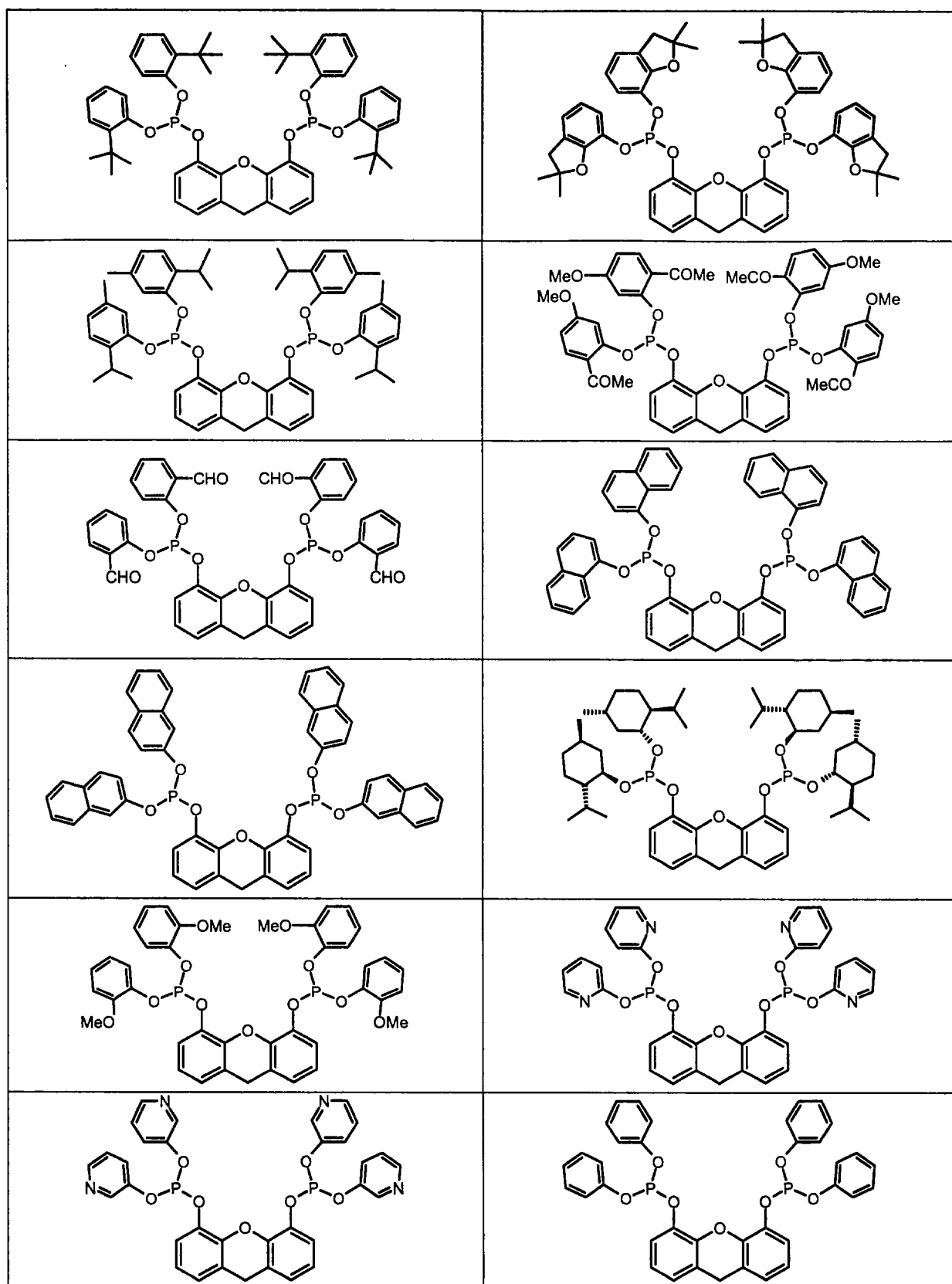


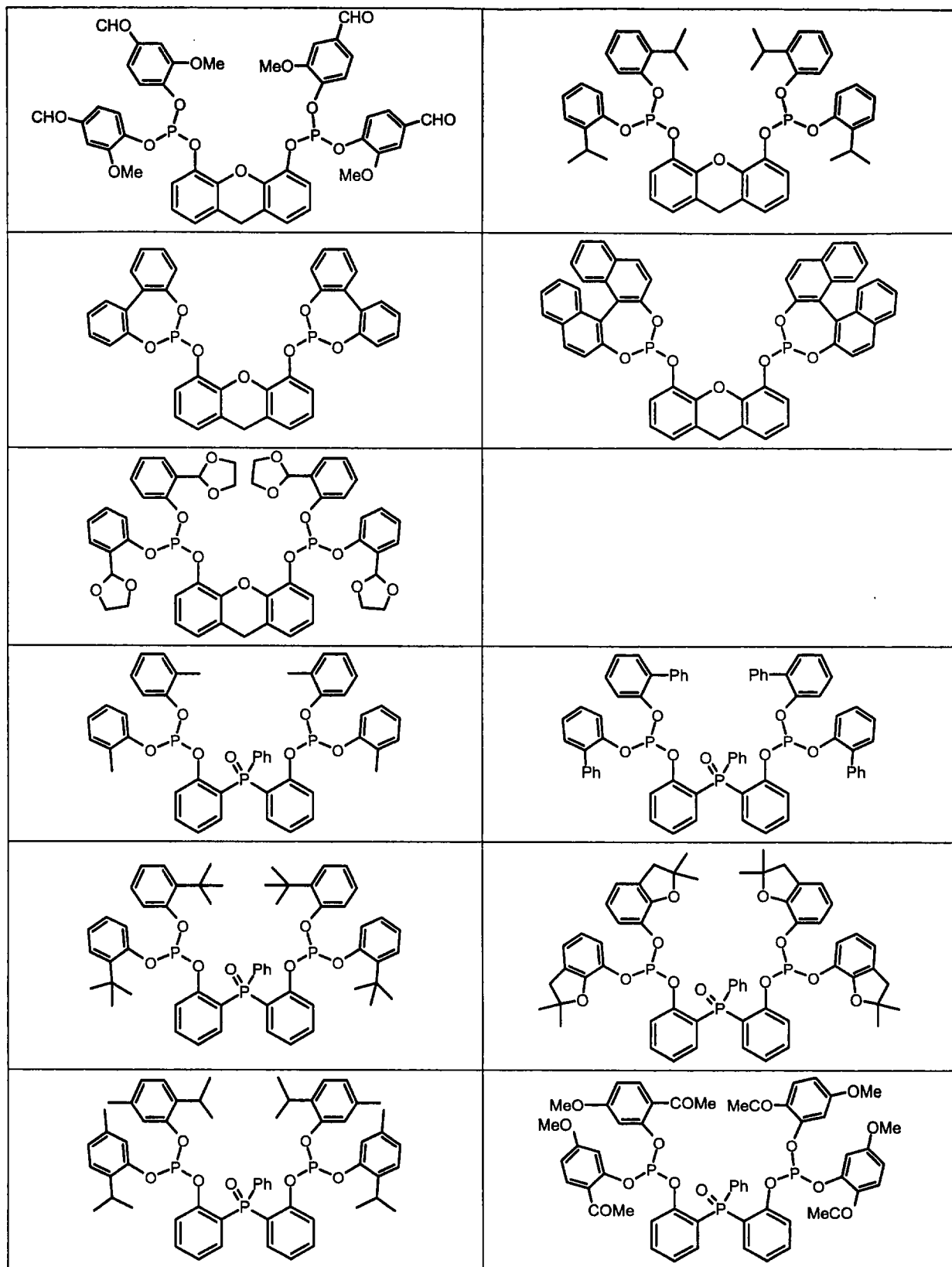


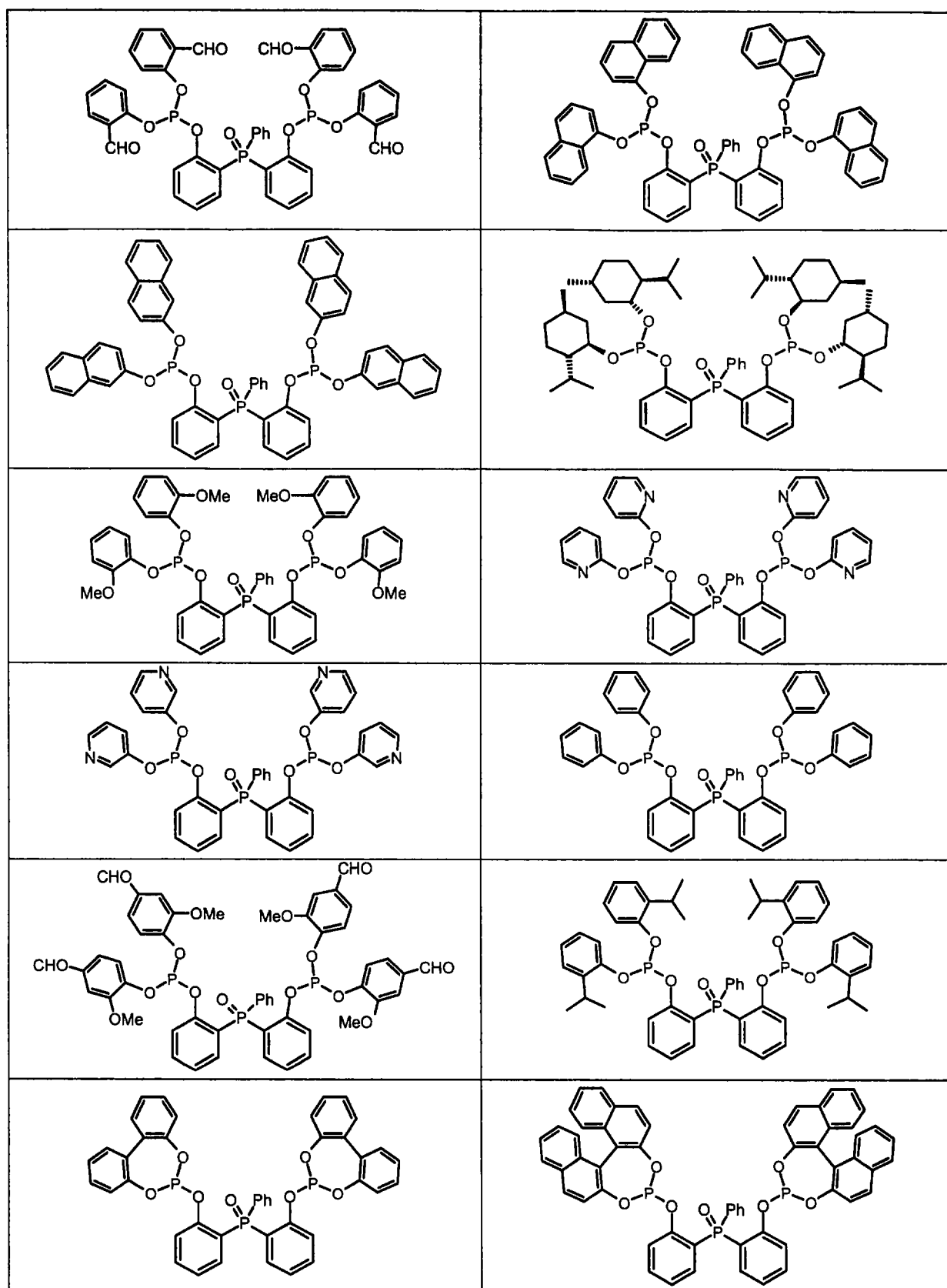


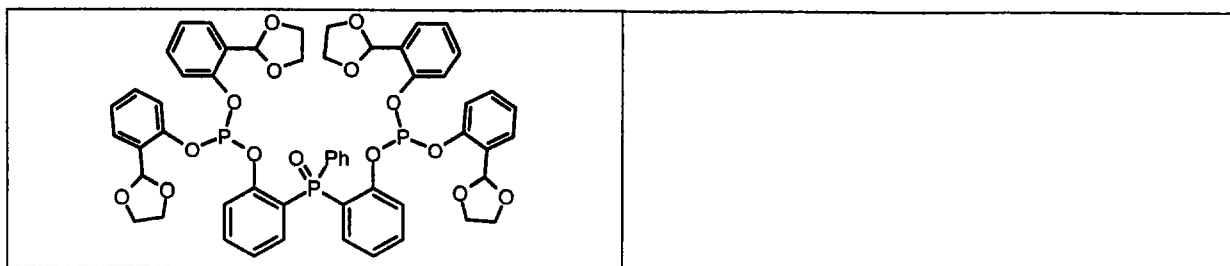












According to the invention the catalyst corresponds advantageously to the general formula (II):



5 in which

M is a transition metal,

L_f represents the organic ligand of formula (I) and

t represents a number between 1 and 4 (inclusive).

10 Metals which may be complexed by the organic ligands of the invention are generally all transition metals of groups 1b, 2b, 3b, 4b, 5b, 6b, 7b and 8 of the Periodic Table of the Elements, as published in the "Handbook of Chemistry and Physics", 51st Edition (1970-1971) from
15 The Chemical Rubber Company.

Among these metals mention may be made more particularly of the metals mention may be made as non-limiting examples, of nickel, cobalt, iron, ruthenium,
20 rhodium, palladium, osmium, iridium, platinum, copper, silver, gold, zinc, cadmium and mercury.

Organometallic complexes comprising the organic ligands of the invention may be prepared by contacting a
25 solution of a compound of the selected metal with a solution of the organic ligand of the invention.

The metal compound may be dissolved in a solvent.

30 The metal in the compound employed may be in the same oxidation state which it will have in the organometallic complex or in a higher oxidation state.

By way of example it is possible to indicate that in the organometallic complexes of the invention rhodium is in oxidation state (I), ruthenium in oxidation state (II), platinum in oxidation state (0), palladium in oxidation state (0), osmium in oxidation state (II), iridium in oxidation state (I), cobalt in oxidation stage (I) and nickel in oxidation state (0).

If in the course of the preparation of the organometallic complex the metal is employed in a higher oxidation state it will be possible for it to be reduced in situ.

The organometallic complexes comprising the organic ligands of the invention may be used as catalysts in olefin hydrocyanation reactions.

As transition metal the transition metal compounds, more particularly the compounds of nickel, of palladium, of cobalt, of iron or of copper, are preferably used.

Among the aforementioned compounds the most preferred compounds are those of nickel.

Non-limiting examples include:

- compounds in which nickel is in oxidation state zero, such as potassium tetracyanonickelate $K_4[Ni(CN)_4]$, bis(acrylonitrile)nickel zero, bis(cycloocta-1,5-diene)nickel zero (also called $Ni(cod)_2$) and derivatives containing ligands, such as tetrakis(triphenylphosphine)nickel zero;
- compounds of nickel such as carboxylates (especially the acetate), carbonate, bicarbonate, borate, bromide, chloride, citrate, thiocyanate, cyanide, formate, hydroxide, hydrophosphite, phosphite, phosphate and derivatives, iodide,

nitrate, sulphate, sulphite, aryl- and alkylsulphonates.

When the nickel compound used corresponds to a nickel
5 oxidation state greater than zero a nickel reducing
agent which reacts preferentially with the nickel under
the reaction conditions is added to the reaction
mixture. This reducing agent may be organic or
inorganic. Non-limiting examples include borohydrides
10 such as BH_4Na and BH_4K , Zn powder, magnesium or
hydrogen.

When the nickel compound used corresponds to the zero
oxidation state of nickel it is also possible to add a
15 reducing agent of the aforementioned type, but such
addition is not mandatory.

When an iron compound is used the same reducing agents
are suitable.

20

In the case of palladium the reducing agents may be,
furthermore, elements of the reaction mixture
(phosphine, solvent, olefin).

25 The organic compounds containing at least one ethylenic
double bond that are more particularly employed in the
present process are diolefins such as butadiene,
isoprene, hexa-1,5-diene, cycloocta-1,5-diene,
ethylenically unsaturated aliphatic nitriles,
30 especially linear pentenenitriles such as pent-3-ene-
nitrile and pent-4-enenitrile, monoolefins such as
styrene, methylstyrene, vinylnaphthalene, cyclohexene,
methylcyclohexene and mixtures of two or more of these
compounds.

35

The pentenenitriles in particular may contain amounts,
generally minor amounts, of other compounds, such as
2-methylbut-3-enenitrile, 2-methylbut-2-enenitrile,
pent-2-enenitrile, valeronitrile, adiponitrile,

2-methylglutaronitrile, 2-ethylsuccinonitrile or butadiene, originating for example from the prior hydrocyanation reaction of the butadiene to unsaturated nitriles.

5

The reason for this is that the hydrocyanation of the butadiene is accompanied by the formation, along with the linear pentenenitriles, of not inconsiderable amounts of 2-methylbut-3-enenitrile and 2-methylbut-2-enenitrile.

10

The catalyst system used for the hydrocyanation according to the process of the invention may be prepared before it is introduced into the reaction zone; for example by adding the appropriate amount of selected transition metal compound and where appropriate of reducing agent to the ligand in accordance with the invention, alone or in solution in a solvent. It is also possible to prepare the catalyst system "in situ" by simply adding the ligand and the transition metal compound to the hydrocyanation reaction mixture before or after adding the compound to be hydrocyanated.

15

20

The amount of compound of nickel or of another transition metal used is selected so as to give a concentration, in moles of transition metal per mole of organic compounds to be hydrocyanated or isomerized, of between 10^{-4} and 1, and preferably between 0.005 and 0.5 mol of nickel or of the other transition metal employed.

25

30

The amount of organic ligands of the invention used to form the catalyst is selected such that the number of moles of this compound relative to 1 mol of transition metal is from 0.5 to 50 and preferably from 1 to 10.

35

Although the reaction is generally conducted without solvent it may be advantageous to add an inert organic

solvent. The solvent may be a solvent for the catalyst which is miscible with the phase comprising the compound to be hydrocyanated at the hydrocyanation temperature. Examples of such solvents include
5 aromatic, aliphatic or cycloaliphatic hydrocarbons.

This solvent may also be partially miscible with the compounds to be hydrocyanated, particularly when the reaction mixture is at a temperature lower than the
10 reaction temperature. Hence at such temperatures a two-phase system may be obtained. Where the catalyst system is soluble in said solvent its extraction from the reaction mixture is thereby facilitated. Partially or immiscible solvents of this kind may be water or
15 organic salts in melt form with an ionic character. Such solvents are used particularly when the organic ligand contains anionic radicals which render it soluble in ionic media. These radicals are, for example, sulphonate, carbonate, carboxylate, phosphate,
20 ammonium, guanidinium and imidazolium groups which are substituents on the aromatic radicals of the ligand.

The hydrocyanation reaction is generally carried out at a temperature from 10°C to 200°C and preferably from
25 30°C to 120°C. Advantageously it is carried out in a single-phase medium at the reaction temperature.

The process of the invention may be implemented continuously or batchwise.
30

The hydrogen cyanide employed may be prepared from metal cyanides, especially sodium cyanide, or from cyanohydrins, such as acetone cyanohydrin, or by any other known synthesis process.
35

The hydrogen cyanide is introduced into the reactor in gaseous form, as a gas mixture or in liquid form. It may also be dissolved beforehand in an organic solvent.

In the context of a batchwise implementation it is possible in practice to charge a reactor, purged beforehand by means of an inert gas (such as nitrogen or argon) either with a solution containing the entirety or a part of the various constituents, such as the organic ligand of the invention, the transition metal compound, the reducing agents and solvents where appropriate, or with said constituents separately. Generally the reactor is then taken to the selected temperature and then the compound to be hydrocyanated is introduced. The hydrogen cyanide is then itself introduced, preferably continuously and regularly.

When the reaction (whose development can be monitored by assaying samples) is at an end the reaction mixture is withdrawn, after cooling, and the reaction products are isolated, by distillation for example.

An enhancement of the process for hydrocyanating ethylenically unsaturated compounds according to the present invention relates in particular to the hydrocyanation of said ethylenically unsaturated nitrile compounds by reaction with hydrogen cyanide and consists in using a catalyst system in accordance with the present invention with a cocatalyst composed of at least one Lewis acid.

The ethylenically unsaturated compounds which may be employed in this improvement are in general those which were mentioned for the basic process. However, it is more particularly advantageous to apply it to the hydrocyanation reaction to dinitriles of the ethylenically unsaturated aliphatic mononitriles, particularly to apply it to linear pentenenitriles such as pent-3-enenitrile, pent-4-enenitrile and mixtures thereof.

These pentenenitriles may contain amounts, generally minor amounts, of other compounds, such as 2-methylbut-

3-enenitrile, 2-methylbut-2-enenitrile, pent-2-enenitrile, valeronitrile, adiponitrile, 2-methylglutaronitrile, 2-ethylsuccinonitrile or butadiene, originating from the prior hydrocyanation reaction of
5 the butadiene and/or from the isomerization of 2-methylbut-3-enenitrile to pentenenitriles.

The Lewis acid used as cocatalyst makes it possible in particular, in the case where ethylenically unsaturated
10 aliphatic nitriles are hydrocyanated, to improve the linearity of the dinitriles obtained, in other words the percentage of linear dinitriles relative to the entirety of the dinitriles formed, and/or to enhance the activity and service life of the catalyst.

15 A Lewis acid in the present text, according to the usual definition, refers to compounds which accept electron pairs.

20 Use may be made in particular of the Lewis acids cited in the work edited by G.A. Olah, "Friedel-Crafts and related Reactions", volume I, pages 191 to 197 (1963).

The Lewis acids which may be employed as cocatalysts in
25 the present process are selected from compounds of the elements of groups Ib, IIb, IIIa, IIIb, IVa, IVb, Va, Vb, VIb, VIIb and VIII of the Periodic Table of the Elements. These compounds are most often salts, especially halides, such as chlorides or bromides,
30 sulphates, sulphonates, haloalkylsulphonates, perhaloalkylsulphonates, especially fluoroalkylsulphonates or perfluoroalkylsulphonates, haloalkylacetates, perhaloalkylacetates, especially fluoroalkylacetates or perfluoroalkylacetates, carboxylates and phosphates.

35 Non-limiting examples of such Lewis acids include zinc chloride, zinc bromide, zinc iodide, manganese chloride, manganese bromide, cadmium chloride, cadmium bromide, stannous chloride, stannous bromide, stannous

sulphate, stannous tartrate, indium trifluoromethylsulphonate, indium trifluoroacetate, zinc trifluoroacetate, the chlorides or bromides of rare earth elements such as lanthanum, cerium, praseodymium, 5 neodymium, samarium, europium, gadolinium, terbium, dysprosium, hafnium, erbium, thallium, ytterbium and lutetium, and cobalt chloride, ferrous chloride and yttrium chloride.

10 As a Lewis acid it is also possible to use organometallic compounds such as triphenylborane or titanium isopropoxide.

It will be appreciated that mixtures of two or more Lewis acids may be employed.

15 Among Lewis acids particular preference is given to zinc chloride, zinc bromide, stannous chloride, stannous bromide and triphenylborane and to zinc chloride/stannous chloride mixtures, indium trifluoromethylsulphonate, indium trifluoroacetate and 20 zinc trifluoroacetate.

The Lewis acid cocatalyst employed represents generally from 0.01 to 50 mol per mole of transition metal 25 compound, more particularly of nickel compound, and preferably from 0.5 to 10 mol per mole.

As for the implementation of the basic process of the invention, the catalyst solution used for the 30 hydrocyanation in the presence of Lewis acid may be prepared prior to its introduction into the reaction zone; for example, by adding the ligand of formula (I), the appropriate amount of selected transition metal compound, Lewis acid and, where appropriate, reducing 35 agent to the reaction mixture. It is also possible to prepare the catalyst solution "in situ" by simply mixing these various constituents.

It is also possible, under the conditions of the

hydrocyanation process of the present invention, and especially when working in the presence of the above-described catalyst comprising at least one ligand of formula (I) and at least one transition metal compound,
5 to carry out the isomerization of 2-methylbut-3-enenitrile to pentenenitriles, and more generally of branched unsaturated nitriles to linear unsaturated nitriles, in the absence of hydrogen cyanide.

10 The 2-methylbut-3-enenitrile subjected to the isomerization according to the invention may be employed alone or in a mixture with other compounds.

Thus 2-methylbut-3-enenitrile may be deployed as a
15 mixture with 2-methylbut-2-enenitrile, pent-4-enenitrile, pent-3-enenitrile, pent-2-enenitrile, butadiene, adiponitrile, 2-methylglutaronitrile, 2-ethylsuccinonitrile or valeronitrile.

20 It is particularly advantageous to treat the reaction mixture originating from the hydrocyanation of butadiene by HCN in the presence of at least one ligand of formula (I) and at least one transition metal compound, more preferably a compound of nickel in
25 oxidation state 0, as defined above.

In the context of this preferred version the catalyst system is already present for the butadiene hydrocyanation reaction and it is therefore sufficient
30 to stop any introduction of hydrogen cyanide in order to allow the isomerization reaction to take place.

It is possible, where appropriate, in this version to effect a gentle purging of the reactor by means of an
35 inert gas, such as nitrogen or argon for example, in order to expel the hydrocyanic acid which might still be present.

The isomerization reaction is generally carried out at

a temperature from 10°C to 200°C and preferably from 60°C to 160°C.

5 In the preferred case of an isomerization immediately following the butadiene hydrocyanation reaction it will be advantageous to operate at the temperature at which the hydrocyanation was conducted.

10 As for the process of hydrocyanating ethylenically unsaturated compounds, the catalyst system used for the isomerization may be prepared prior to its introduction into the reaction zone; for example, by adding the ligand of formula (I), the appropriate amount of selected transition metal compound and, where
15 appropriate, reducing agent to a solvent. It is also possible to prepare the catalyst system "in situ" by simply adding these various constituents to the reaction mixture. The amount of transition metal compound, and more particularly of nickel compound,
20 that is used, and the amount of ligand of formula (I), are the same as for the hydrocyanation reaction.

Although the isomerization reaction is conducted generally without solvent it may be advantageous to add
25 an inert organic solvent, which may be that of the subsequent extraction. This is the case particularly when such a solvent has been employed in the butadiene hydrocyanation reaction that was used to prepare the mixture which is subjected to the isomerization
30 reaction. Such solvents may be selected from those which were identified above for the hydrocyanation.

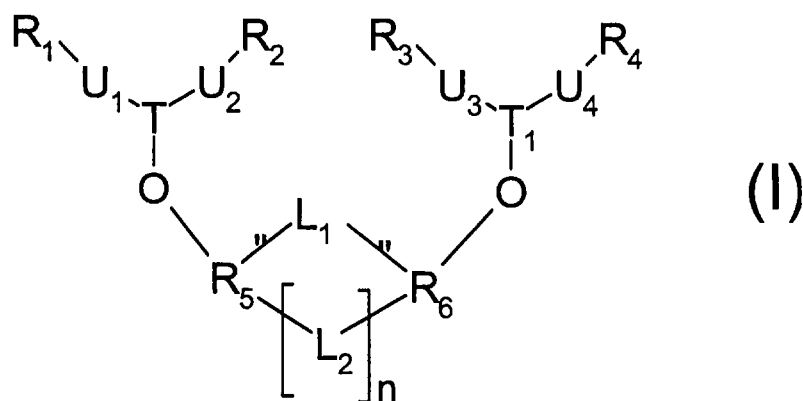
The preparation of dinitrile compounds by hydrocyanation of an olefin such as butadiene may be
35 carried out using a catalyst system in accordance with the invention for the above steps of formation of unsaturated mononitriles and the isomerization step; the hydrocyanation reaction of unsaturated mononitriles to dinitriles can be implemented with a catalyst system

in accordance with the invention or any other catalyst system already known for this reaction.

Similarly the hydrocyanation reaction of the olefin to
 5 unsaturated mononitriles and the isomerization thereof may be carried out with a catalyst system other than that of the invention; the hydrocyanation step of the unsaturated mononitriles to dinitriles is employed with a catalyst system in accordance with the invention.

10

The invention likewise provides organophosphorus compounds corresponding to the general formula I below:



15

in which:

T and T₁, which are identical or different, represent a phosphorus, arsenic or antimony atom,

20

R₁, R₂, R₃ and R₄, which are identical or different, represent a substituted or unsubstituted, aromatic, aliphatic or cycloaliphatic radical comprising one or more rings, which are in fused form or not and which
 25 may contain one or more heteroatoms, where the radicals R₁ and R₂ on the one hand and R₃ and R₄ on the other hand may be interconnected by a covalent bond, a hydrocarbon chain or a heteroatom,

30

U₁, U₂, U₃ and U₄, which are identical or different, represent an oxygen atom or a radical of formula NR in

which R denotes a monovalent alkyl, aryl, cycloalkyl, sulphonyl or carbonyl radical,

5 R₅ and R₆, which are identical or different, represent an aryl or cycloaliphatic group which may comprise heteroatoms and/or one or more rings, in fused form or not, and which are substituted or unsubstituted,

n is an integer equal to 0 or 1,

10

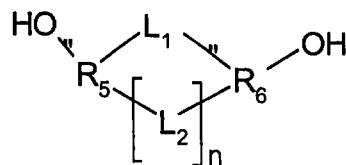
L₁, when n is 0, represents a divalent radical selected from the group consisting of the groups NR₇, PR₈, SiR₉R₁₀, BR₁₁, S, POR₁₂, SO₂ and CO, in which R₇ is as defined for R above, R₈ and R₁₂ may represent the radical OR₁₃, and R₈, R₉, R₁₀, R₁₁, R₁₂ and R₁₃ represent alkyl, aryl or cycloalkyl radicals,

L₁ and L₂, when n is 1, are identical or different and represent a covalent bond or a radical selected from the group consisting of the groups O, NR₇, PR₈, SiR₉R₁₀, BR₁₁, S, POR₁₂, SO₂, CO and -CR₁₄R₁₅-, in which R₇ is as defined for R above, R₈ and R₁₂ may represent the radical OR₁₃, and R₈, R₉, R₁₀, R₁₁, R₁₂, R₁₃, R₁₄ and R₁₅ represent alkyl, aryl or cycloalkyl radicals, it being possible also for R₁₄ and R₁₅ to represent a hydrogen atom.

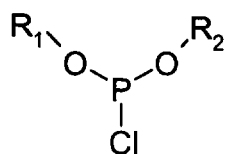
The invention relates in particular to the organophosphorus compounds listed in Table II above.

30

These compounds are obtained in particular by mixing a solution in toluene of a diphenol compound corresponding to the structure of the following fomula:

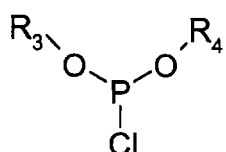


with a toluenic solution of a chlorophosphite of following formula(e):



5

and/or



10

In the above formulae the symbols have the same meanings as in formula I.

15 The mixture is stirred for a number of hours (of the order of 8 to 15 hours) at ambient temperature (between 15°C and 30°C). The organophosphorus compound is recovered, for example, by filtering the mixture.

20 This preparation procedure is given solely by way of example and indication. Other preparation processes may be used without departing from the scope of the invention.

25 Moreover, the phenols and chlorophosphites are generally available commercially or may be synthesized by conventional processes described for obtaining these compounds.

The examples which follow illustrate the invention.

30

In the examples the abbreviations used have the meanings indicated below.

- cod: 1,5-cyclooctadiene
eq: equivalent
3PN: 3-pentenitrile
4PN: 4-pentenitrile
5 3+4PN: 3PN + 4PN
DC(Y): degree of conversion of the product to be hydrocyanated, Y, corresponding to the ratio of the number of moles of Y converted to the initial number of moles of Y
10 Linearity (L): ratio of the number of moles of adiponitrile (AdN) formed to the number of moles of dinitriles formed (sum of the moles of AdN, ethylsuccinonitrile (ESN) and methylglutaronitrile (MGN))
15 GC: gas chromatography
ml: millilitre
mol: mole(s)
mmol: millimole(s).
20 The various products in accordance with the invention may be prepared by one of the two following procedures:

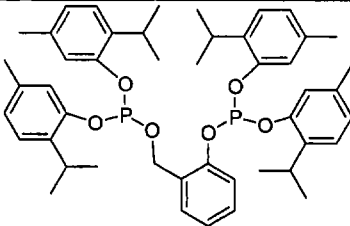
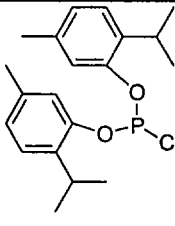
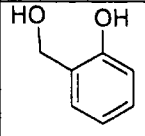
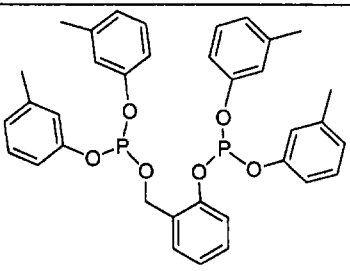
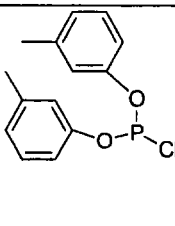
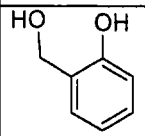
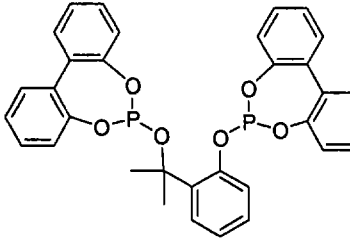
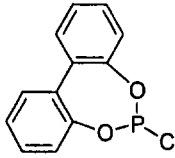
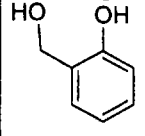
Procedure I:

- 25 A phosphochloridite is used which is prepared from a phenol or biphenol derivative and PCl_3 according to a conventional procedure (see, for example, the method described by G. Buisman et al. in Tetrahedron, Asymmetry, vol. 4, (7), pp. 1625-34 (1993)), and a diol
30 available commercially. The following general procedure is representative:
In a 100 ml reactor under argon 6 mmol of phosphochloridite are dissolved in 20 ml of anhydrous toluene. The solution is stirred at -10°C . A solution
35 of 3 mmol of diol and 10 mmol of triethylamine in 20 ml of anhydrous toluene is introduced dropwise into the reaction mixture, which is held at -10°C : a white precipitate forms. The suspension is stirred vigorously at 25°C for 18 h and then filtered under argon over a

bed of basic alumina. After rinsing with toluene, the filtrate is concentrated under reduced pressure to give the desired product, which is used without further purification.

5

The following ligands were prepared according to the procedure described above:

Example	Ligand	Structure	Phospho- chloridite	Diol
1	A			
2	B			
3	C			

10

Procedure II:

A phosphoramidite is used which is prepared from Cl_2PNEt_2 (available commercially) and a substituted or
15 unsubstituted phenol according to the following procedure:

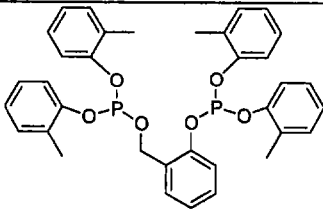
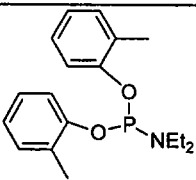
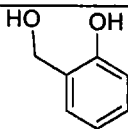
A mixture of Cl_2PNEt_2 (0.20 mol) and triethylamine (0.44 mol) in toluene (800 ml) at 0°C is admixed slowly, with vigorous stirring, with a solution of
20 phenol (0.40 mol) in toluene (100 ml). The formation of

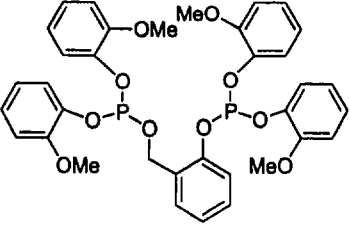
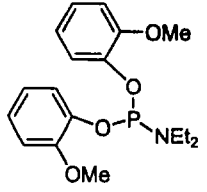
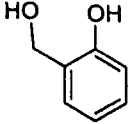
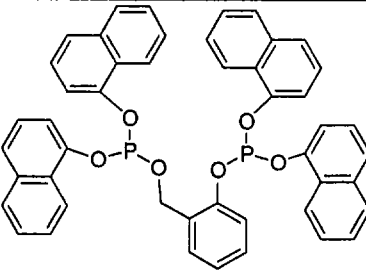
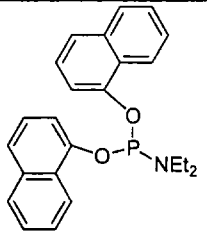
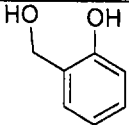
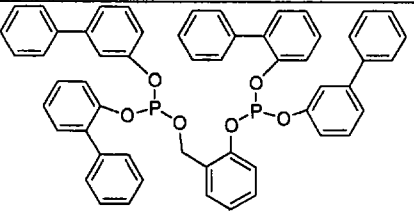
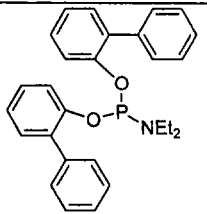
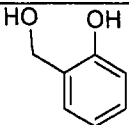
a white precipitate is observed. The mixture is allowed to climb to ambient temperature and is stirred, still vigorously, for 2 h. The mixture is subsequently filtered on a bed of silica and concentrated under vacuum to give the phosphoramidite $(\text{ArO})_2\text{PNEt}_2$ with a purity of more than 95%.

Additionally a commercially available diol is used. The following general procedure is representative:

In a 100 ml reactor under argon 6 mmol of phosphoramidite are introduced in 20 ml of anhydrous toluene. The solution is stirred at 0°C and 7.5 ml of a 2 M solution of hydrochloric acid in ether are added thereto over 30 minutes. The formation of a white precipitate is observed, and the mixture is stirred at ambient temperature for 1 hour. A solution of 3 mmol of diol and 10 mmol of triethylamine in 20 ml of anhydrous toluene is subsequently introduced dropwise into the reaction mixture, which is cooled at -10°C . The suspension is stirred vigorously at 25°C for 18 hours and then filtered under argon on a bed of basic alumina. After rinsing with toluene, the filtrate is concentrated under reduced pressure to give the desired product, which is used without further purification.

The following ligands were prepared according to the procedure described above:

Example	Ligand	Structure	Phosphoramidite	Diol
4	D			

5	E			
6	F			
7	G			

Examples of hydrocyanation of 3-pentenitrile (3PN) to adiponitrile (AdN)

5

The general procedure used is as follows:

Under an argon atmosphere a 60 ml Schott glass tube equipped with a septum stopper is charged in succession with

10

- the ligand (2.5 eq),
- 1.21 g (15 mmol; 30 eq) of anhydrous 3PN,
- 138 mg (0.5 mmol; 1 eq) of Ni(cod)₂ and
- the Lewis acid (0.5 mmol; 1 eq).

15

The mixture is taken to 70°C with stirring. Acetone cyanohydrin is injected into the reaction mixture by means of a syringe driver at a rate of 0.45 ml per hour. After 3 hours of injection the syringe driver is

20 stopped. The mixture is cooled to ambient temperature, diluted with acetone and analysed by gas chromatography.

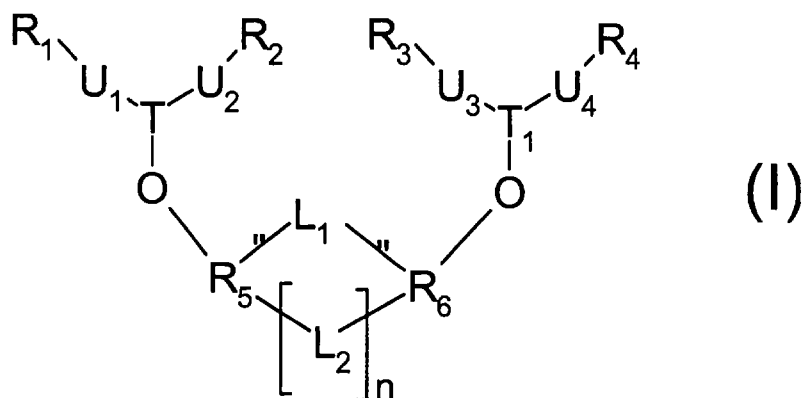
The results are compiled in the table below:

Example	Ligand	Lewis acid	DC (3PN)	Linearity
8	A	ZnCl ₂	64%	70%
9	B	ZnCl ₂	14%	71%
10	C	ZnCl ₂	20%	73%
11	D	ZnCl ₂	44%	76%
12	E	ZnCl ₂	11%	72%
13	E	CoCl ₂	32%	74%
14	F	ZnCl ₂	16%	67%
15	G	ZnCl ₂	48%	80%
16	G	YCl ₃	64%	88%

CLAIMS

1. Process for hydrocyanating a hydrocarbon compound containing at least one ethylenic unsaturation by reacting it in a liquid medium with hydrogen cyanide in the presence of a catalyst comprising a metallic element selected from transition metals and an organic ligand, characterized in that the organic ligand corresponds to the general formula I below:

10



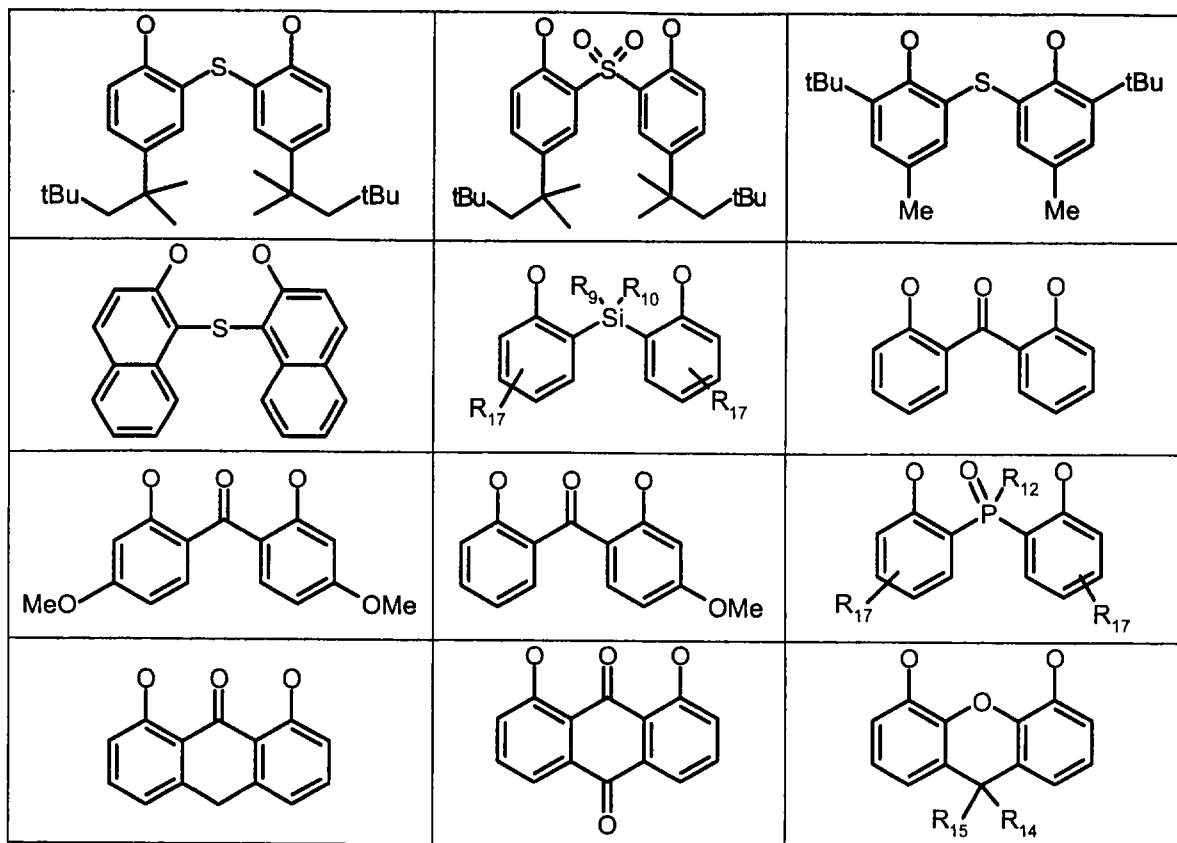
in which:

15

T and T₁, which are identical or different, represent a phosphorus, arsenic or antimony atom,

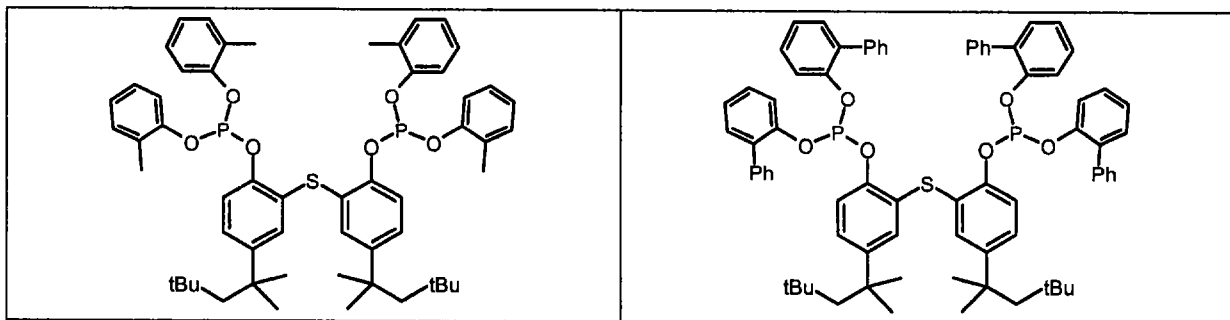
20 R₁, R₂, R₃ and R₄, which are identical or different, represent a substituted or unsubstituted, aromatic, aliphatic or cycloaliphatic radical comprising one or more rings, which are in fused form or not and which may contain one or more heteroatoms, where the radicals R₁ and R₂ on the one hand and R₃ and R₄ on the other hand
 25 may be interconnected by a covalent bond, a hydrocarbon chain or a heteroatom,

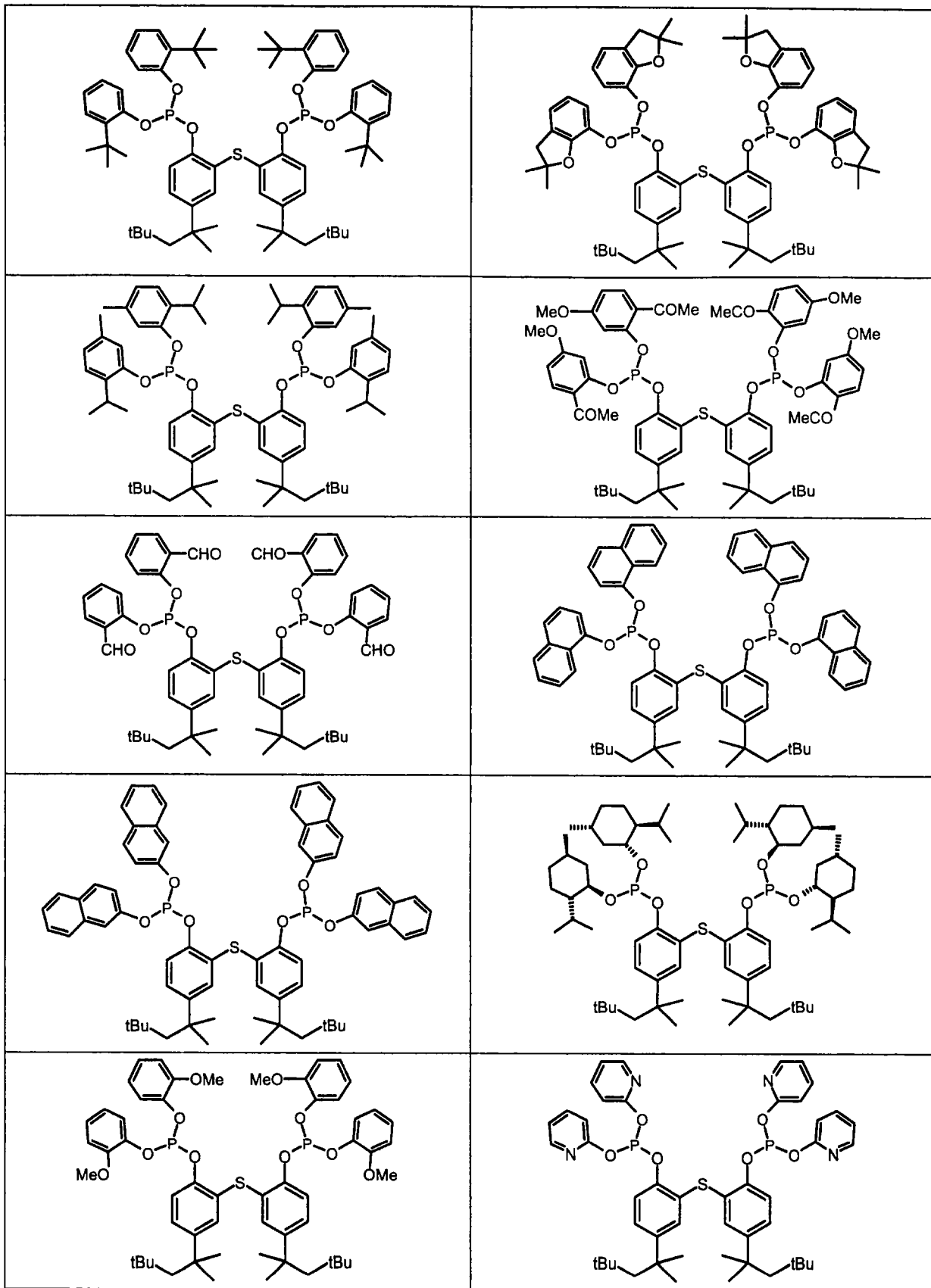
U₁, U₂, U₃ and U₄, which are identical or different, represent an oxygen atom or a radical of formula NR in
 30 which R denotes a monovalent alkyl, aryl, cycloalkyl, sulphonyl or carbonyl radical,

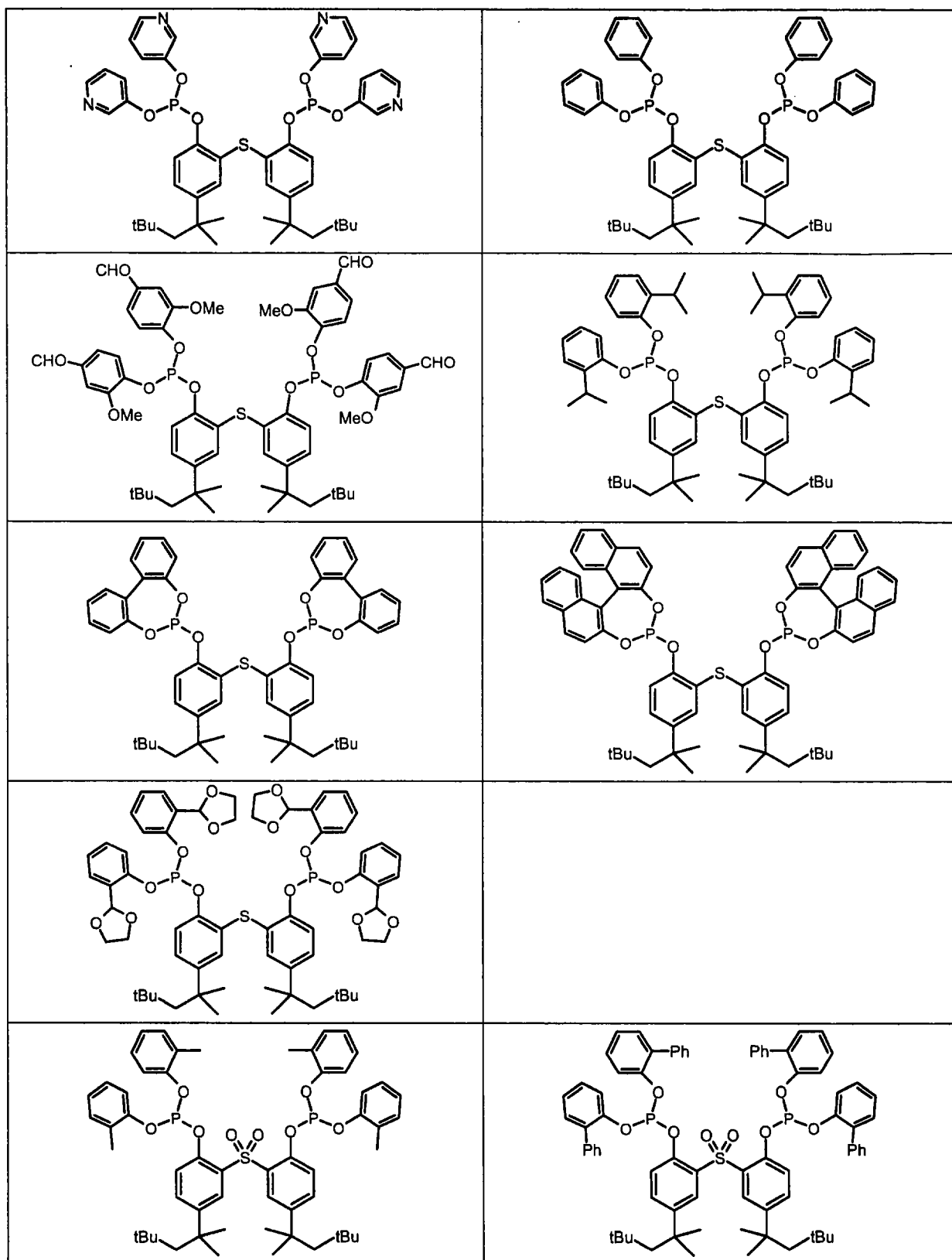


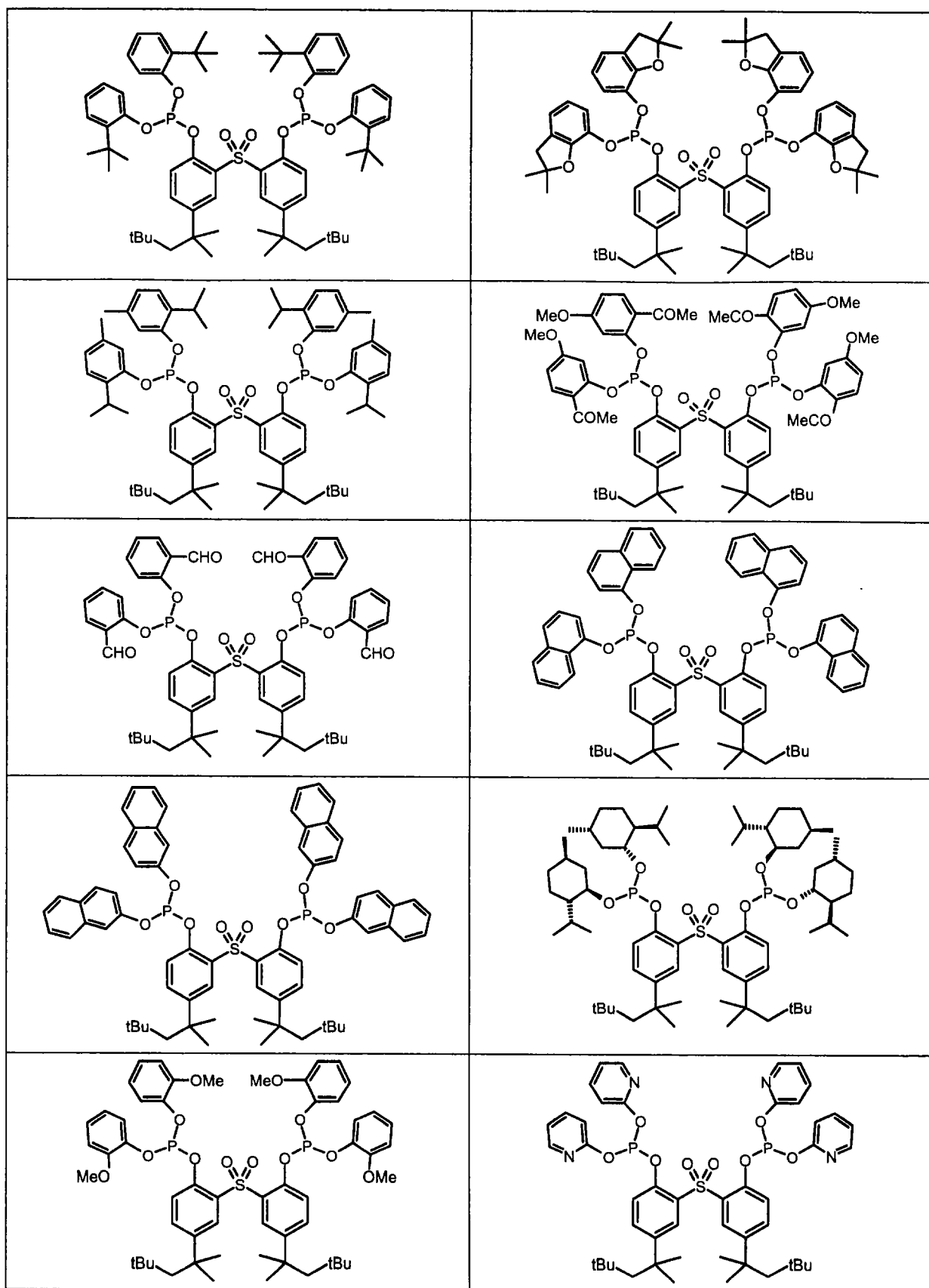
in which R_{17} represents an alkyl, aryl, halogen, alkoxy, thiol, cyano, nitro, aryloxy, alkoxy carbonyl, acyl or formyl radical.

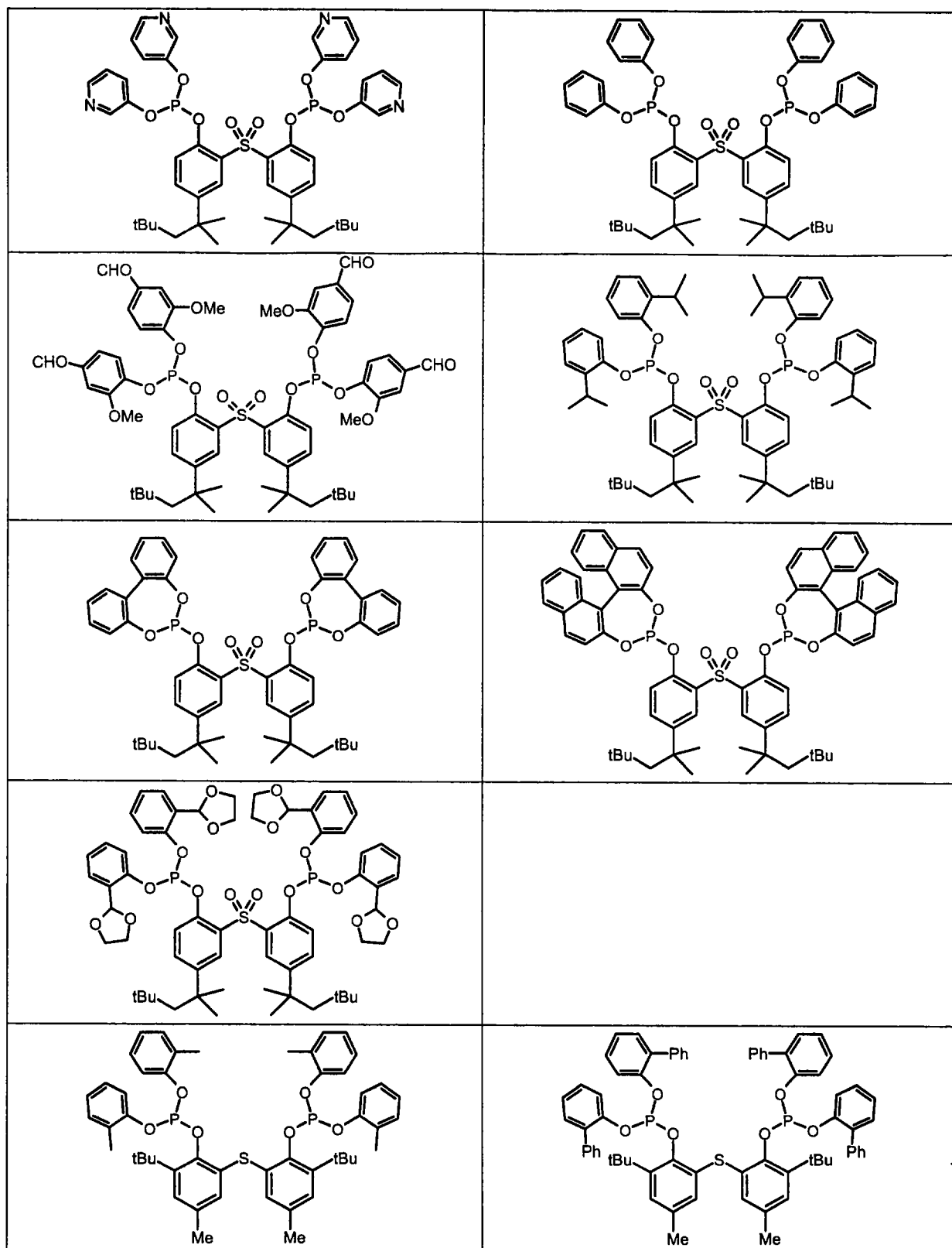
- 5 3. Process according to Claim 1 or 2, characterized in that the organic ligand of formula I is selected from the group consisting of:

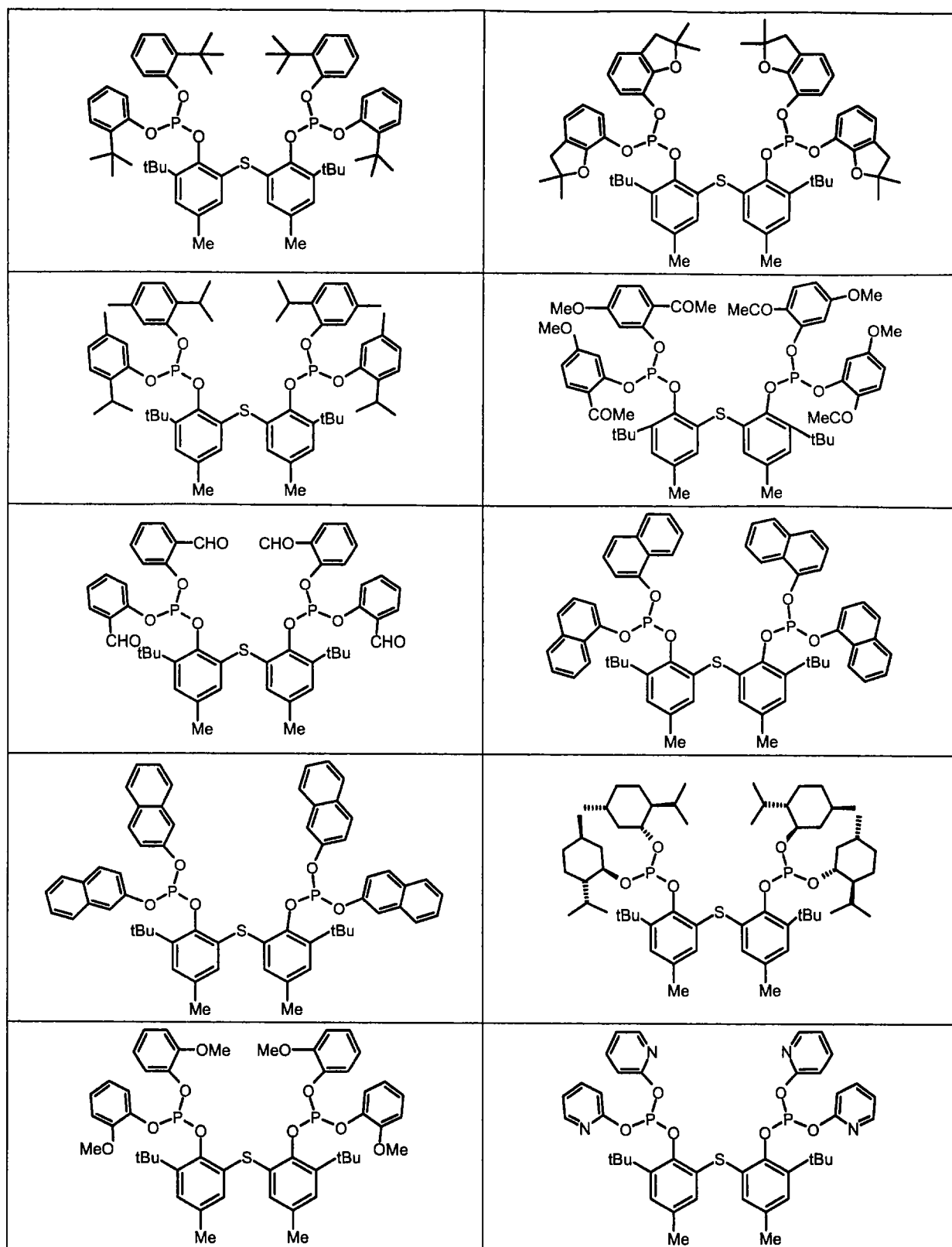


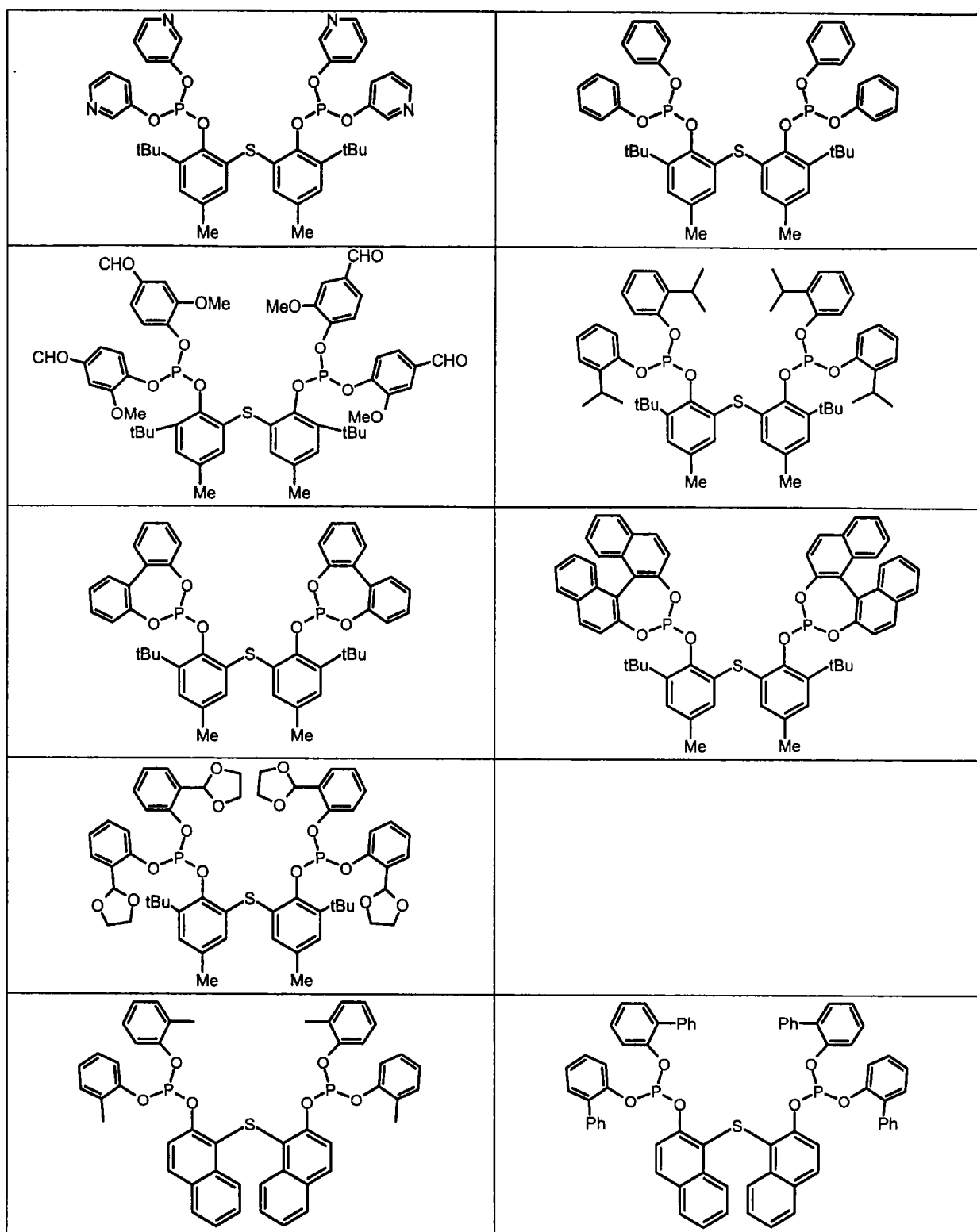


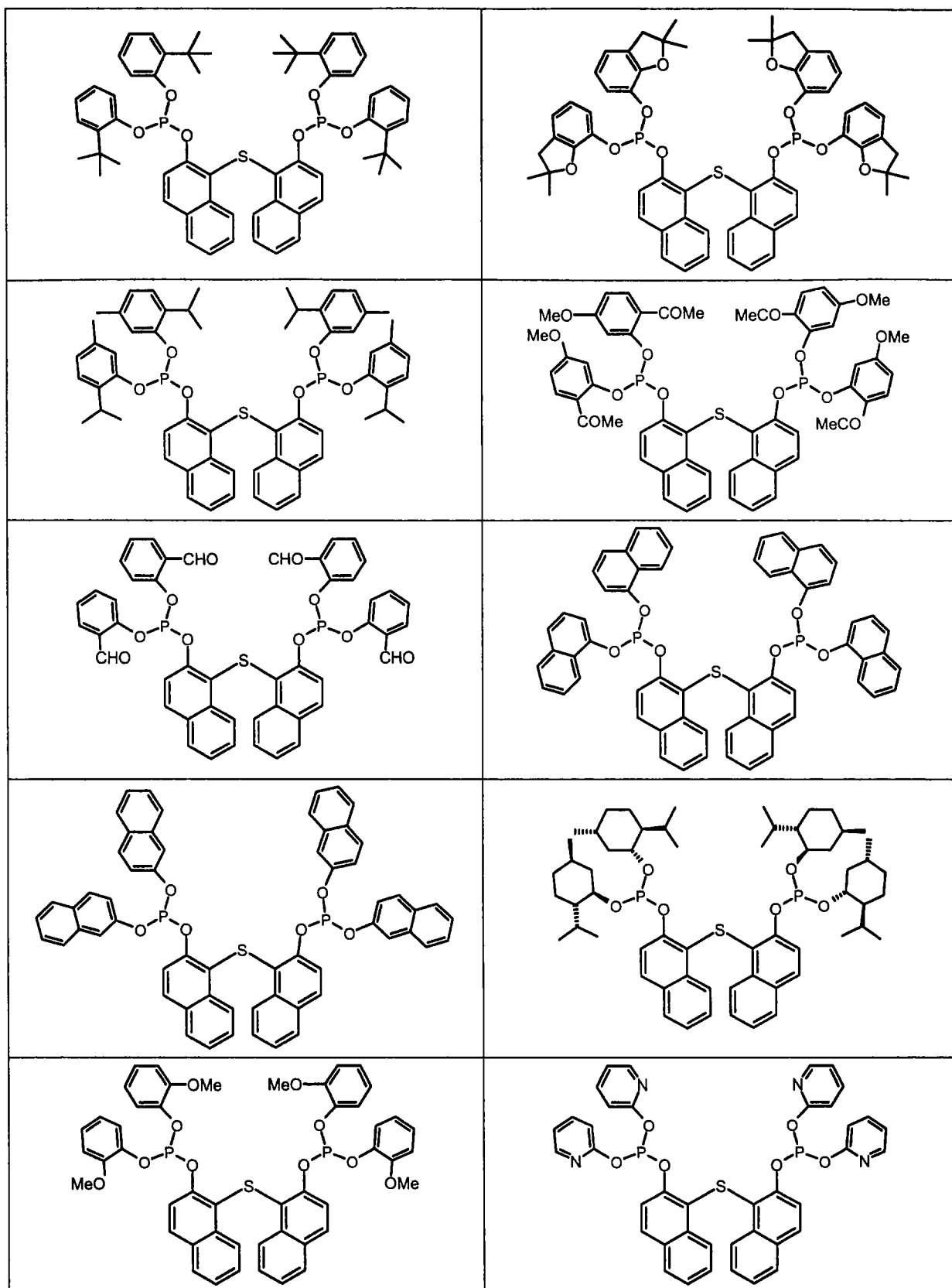


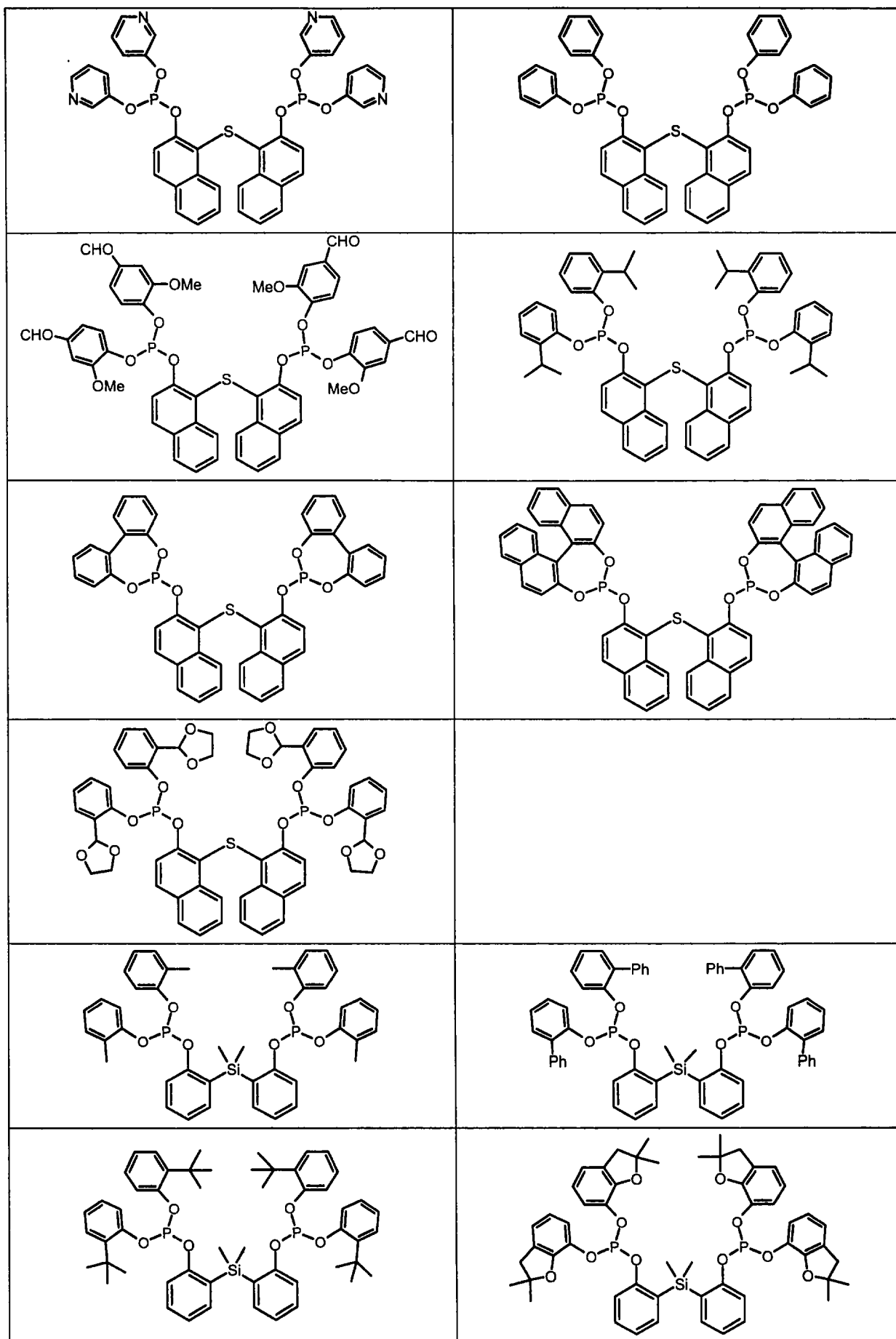


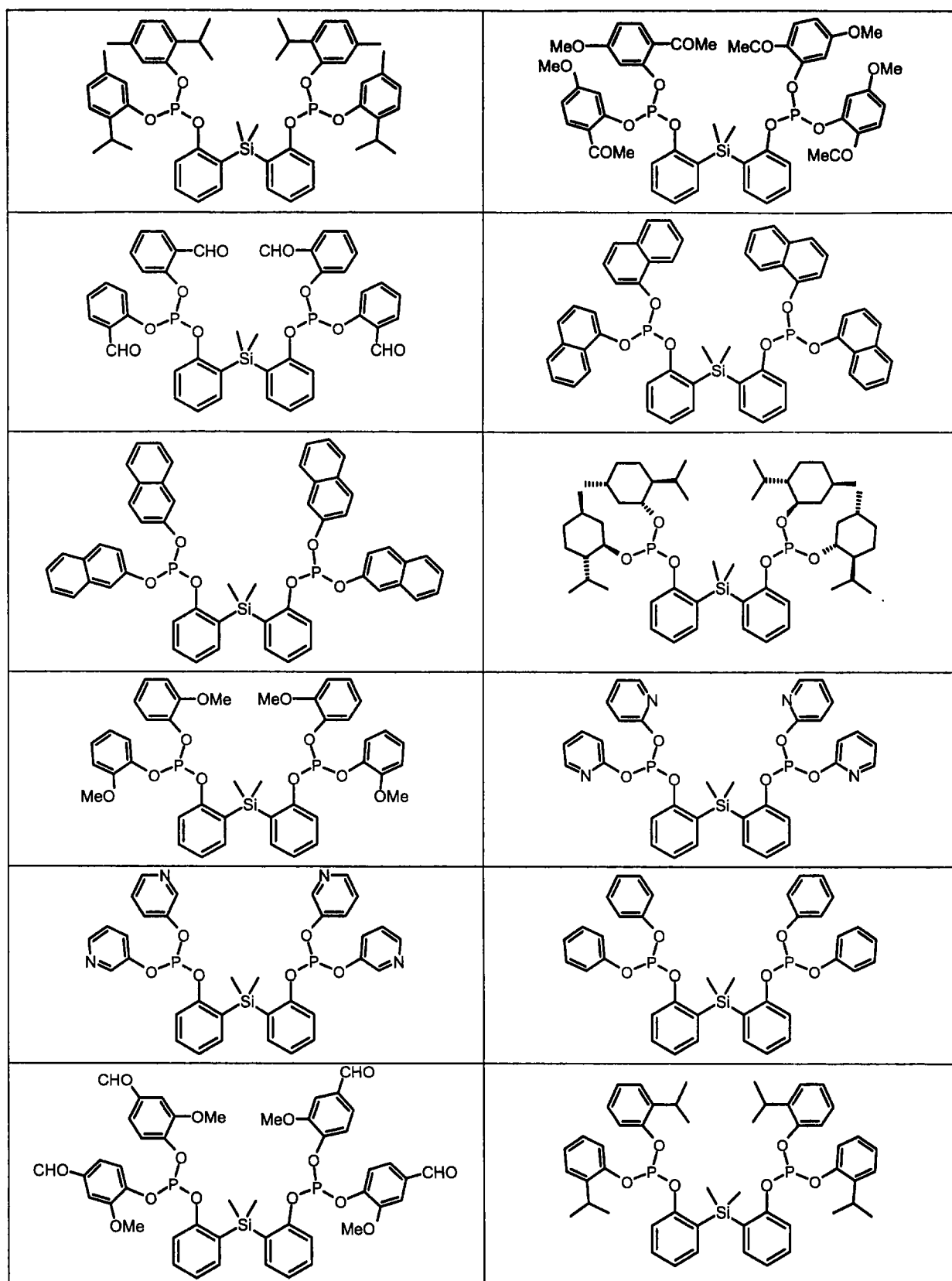


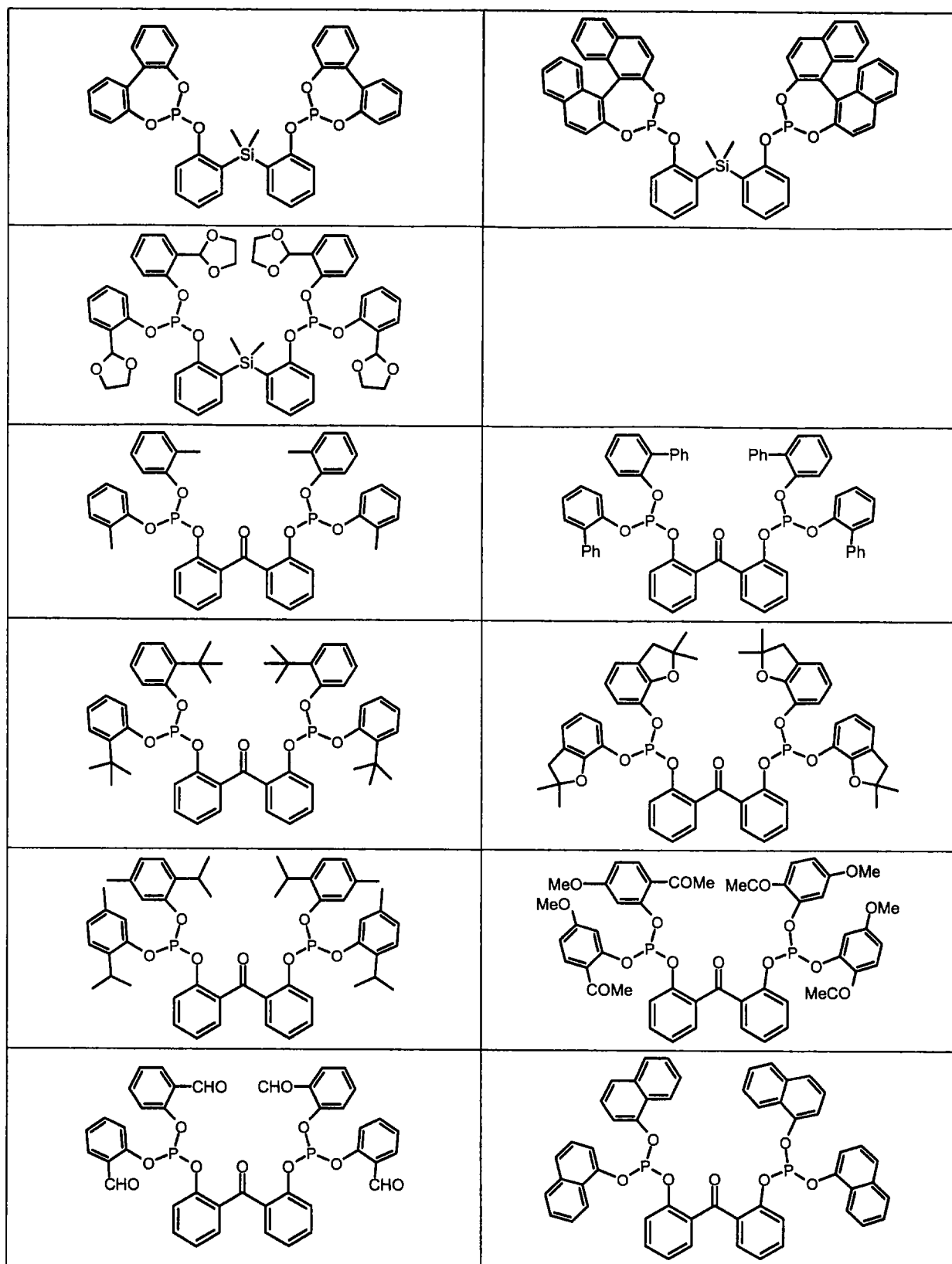


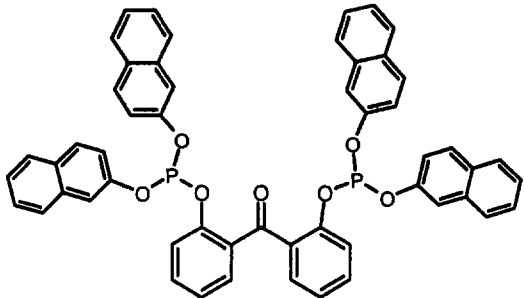
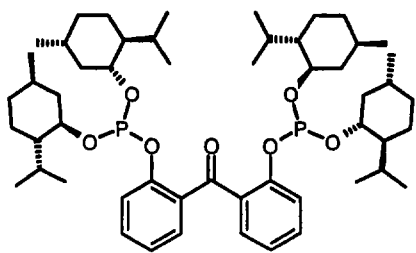
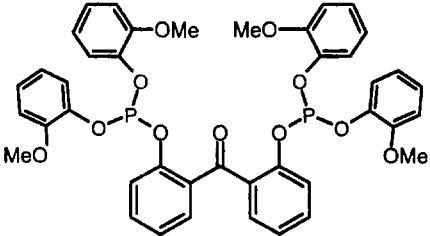
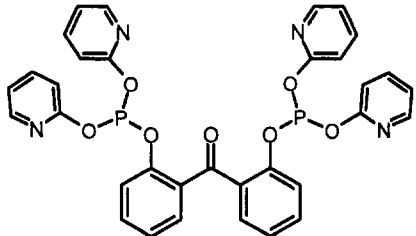
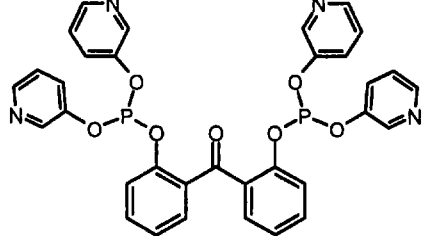
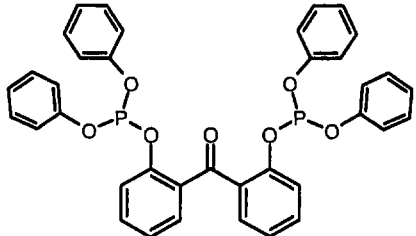
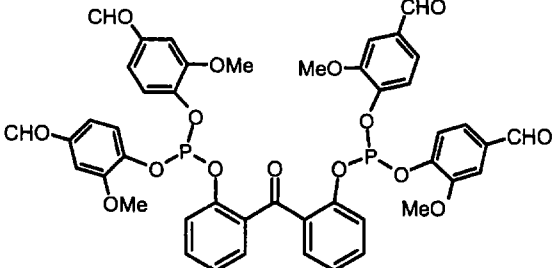
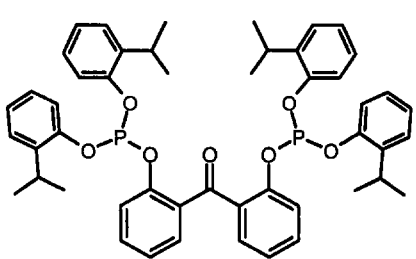
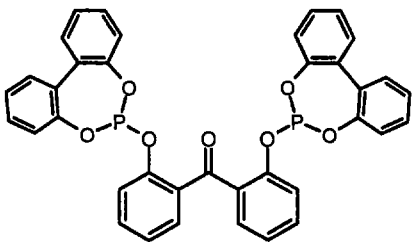
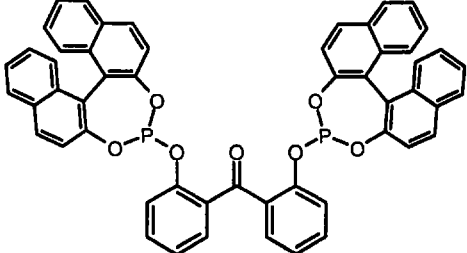
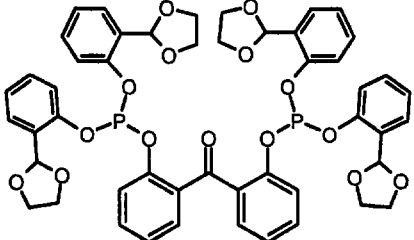


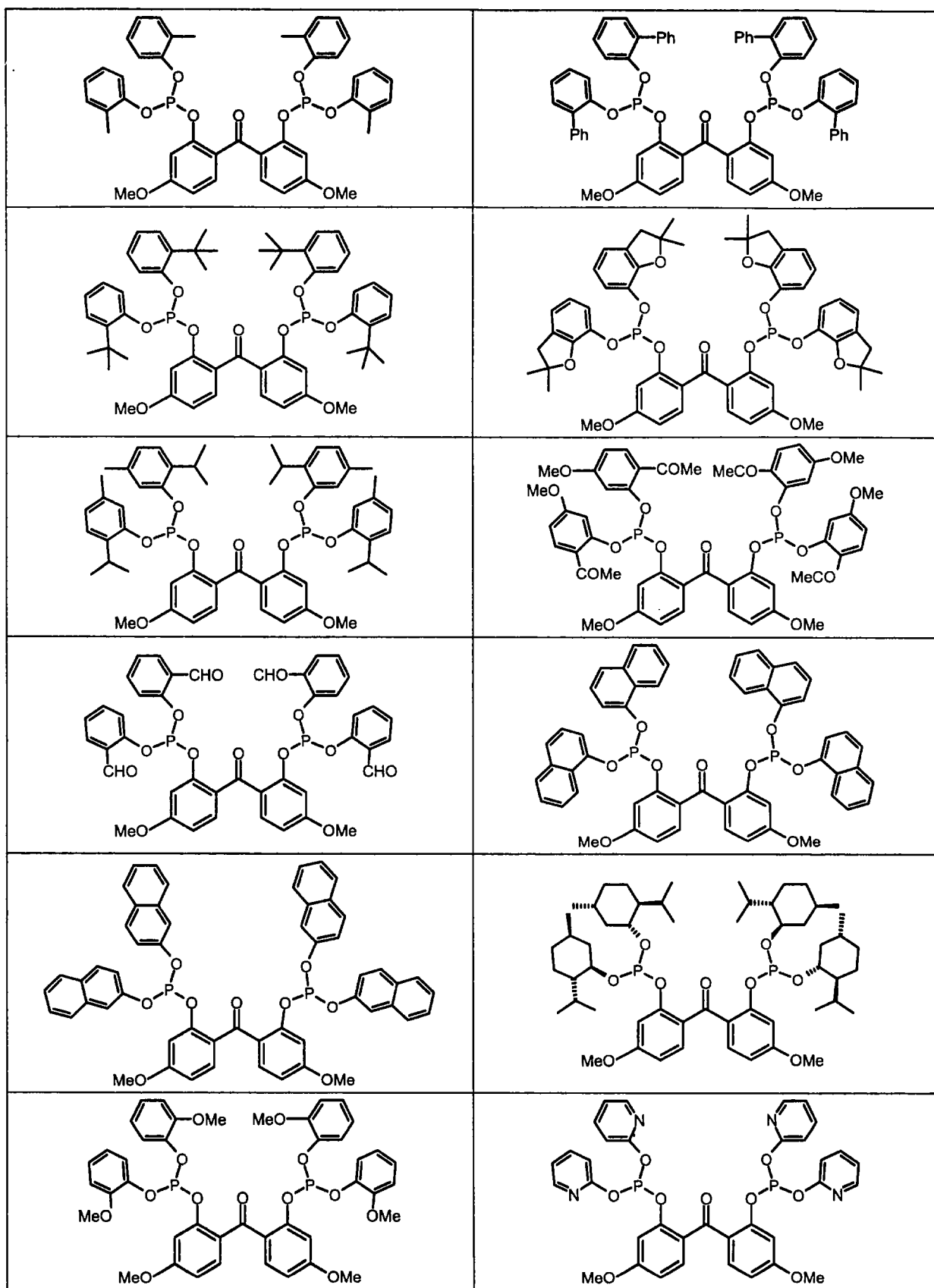


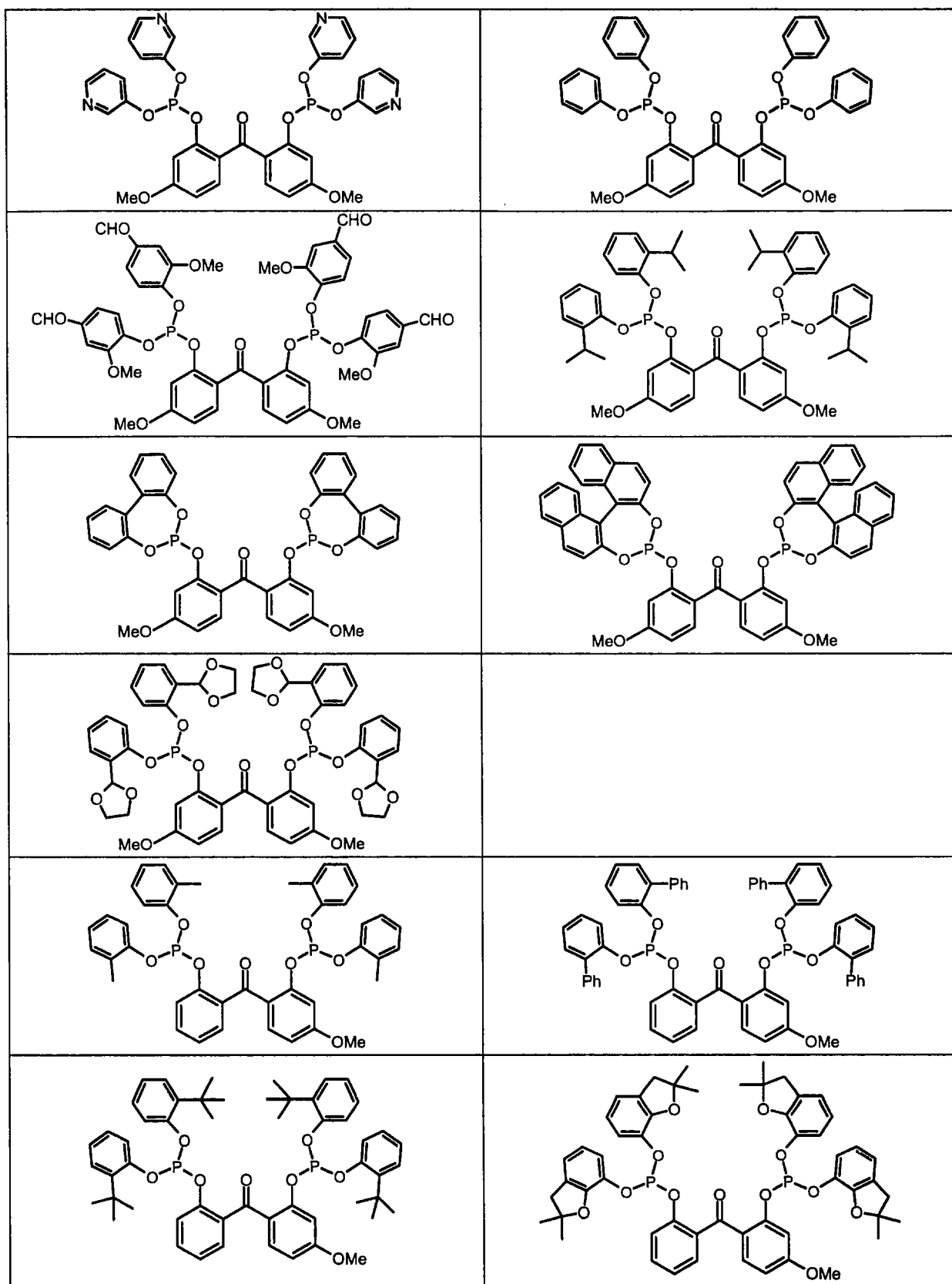


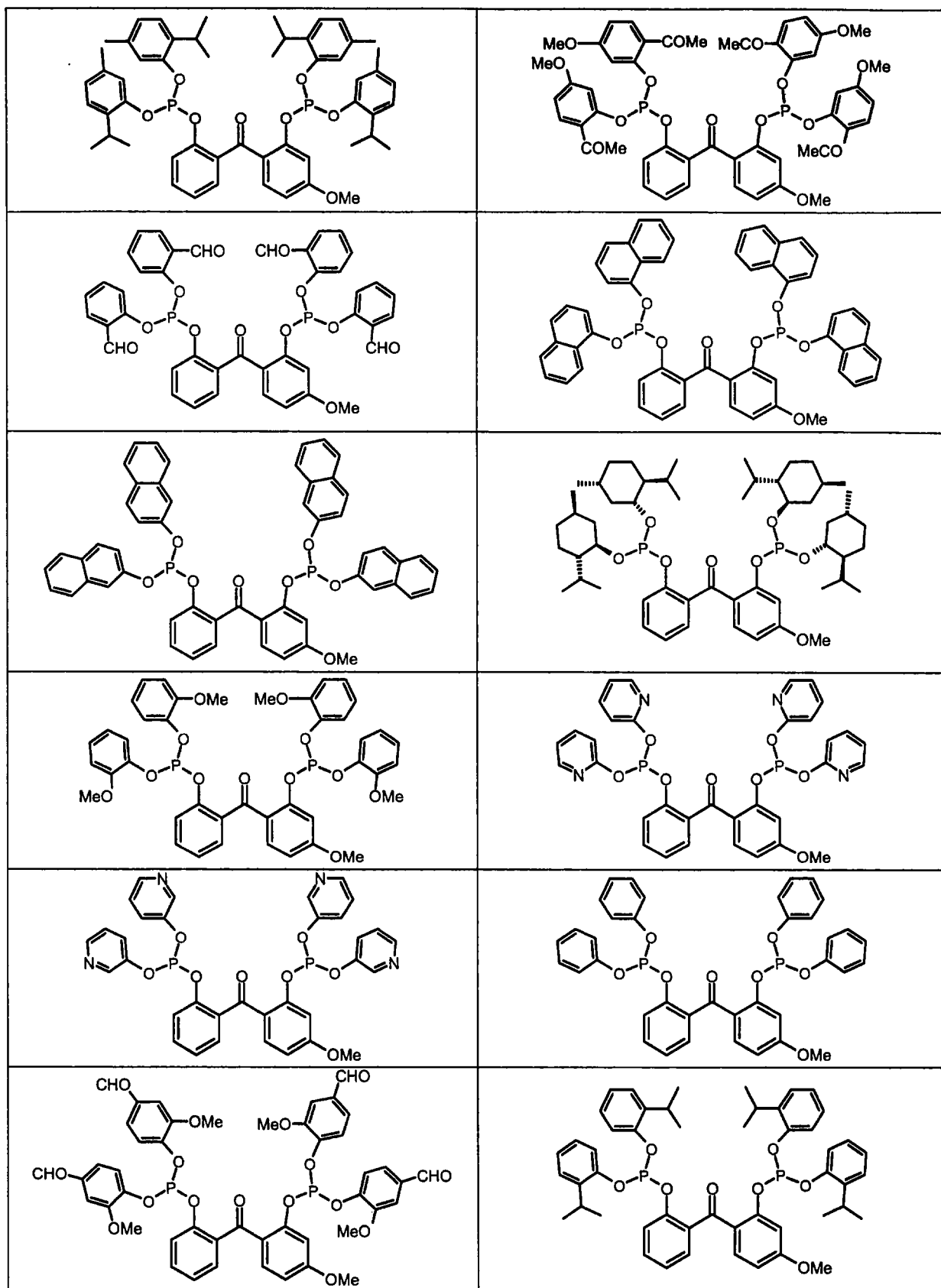


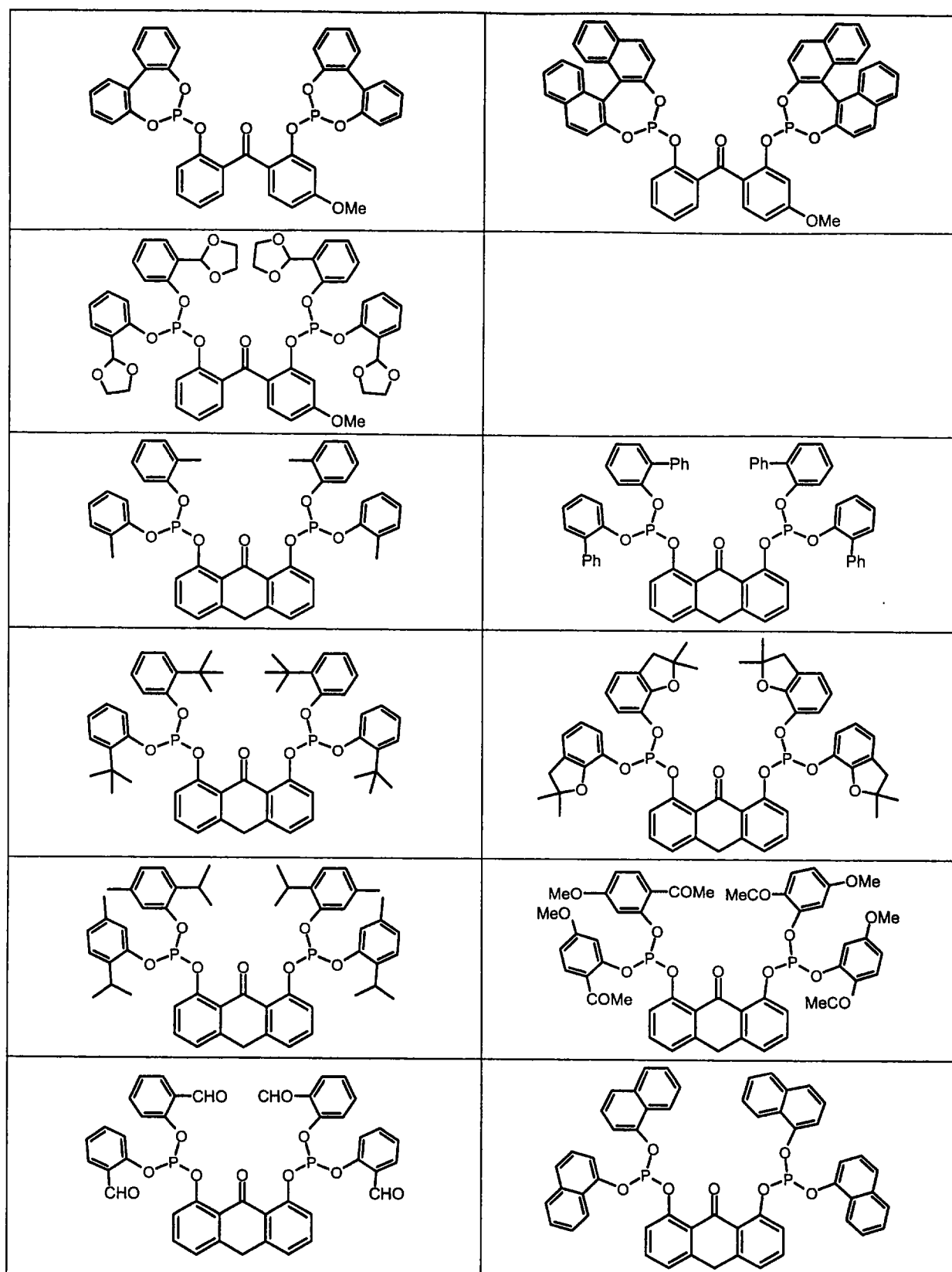


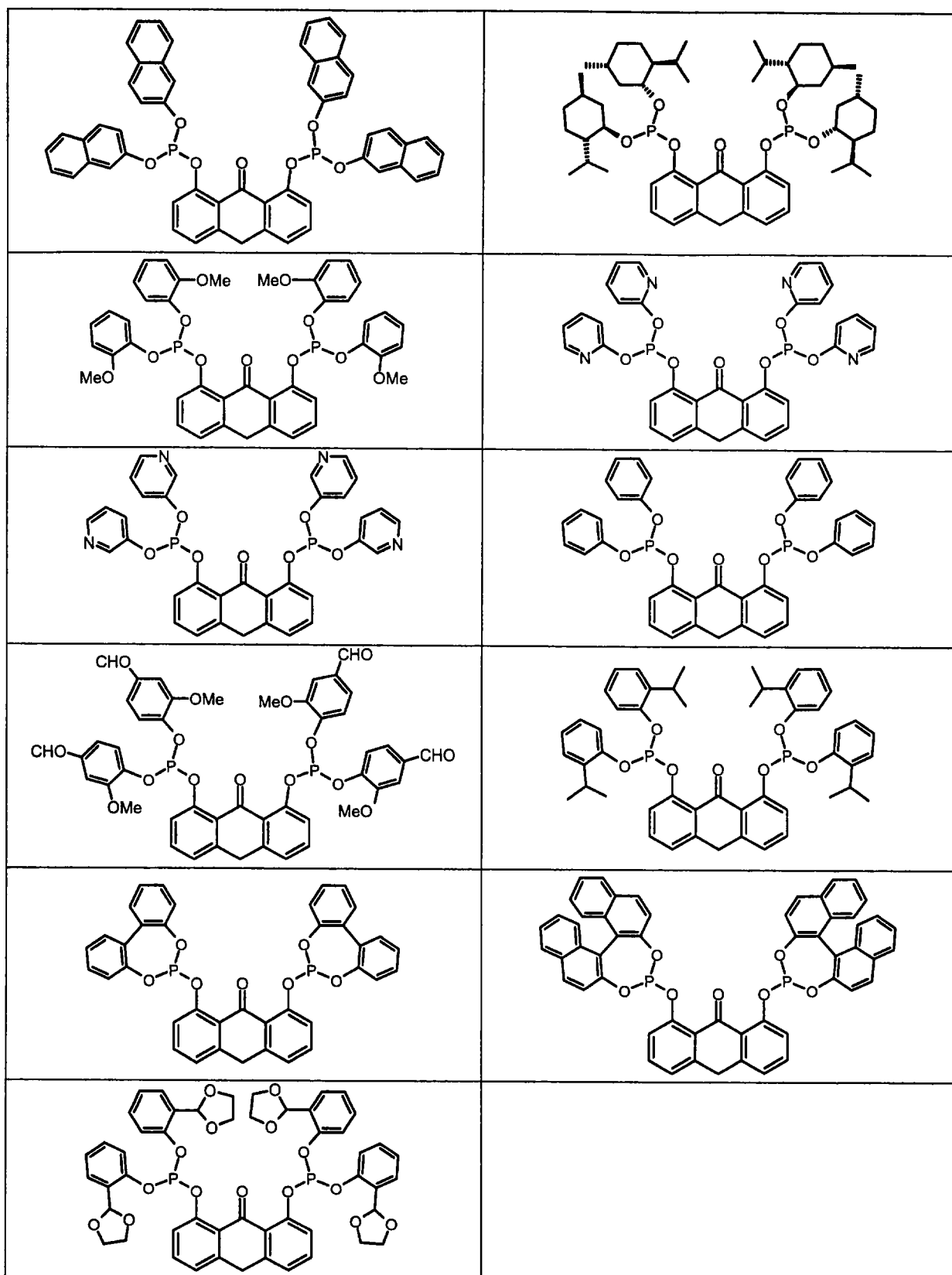
	
	
	
	
	
	

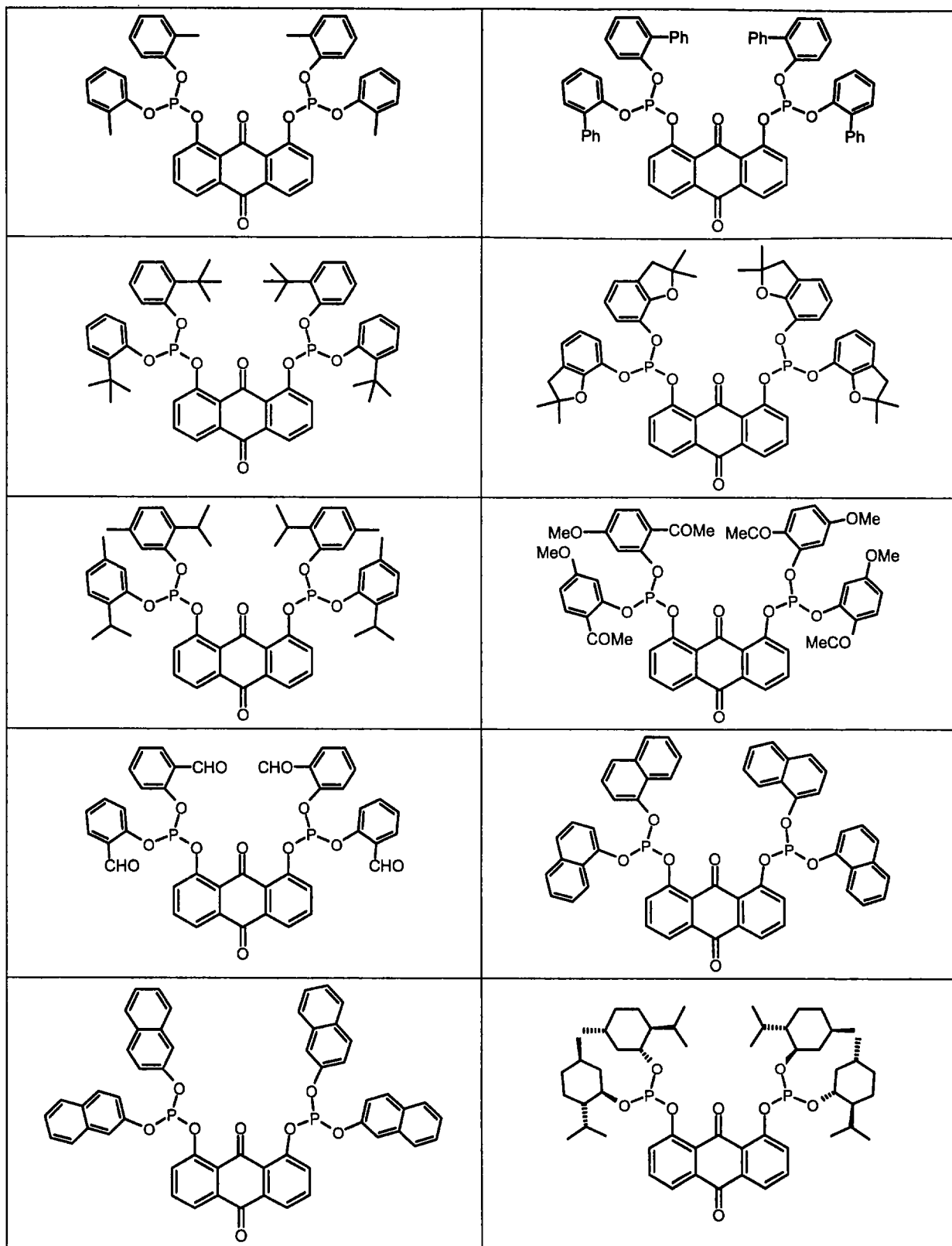


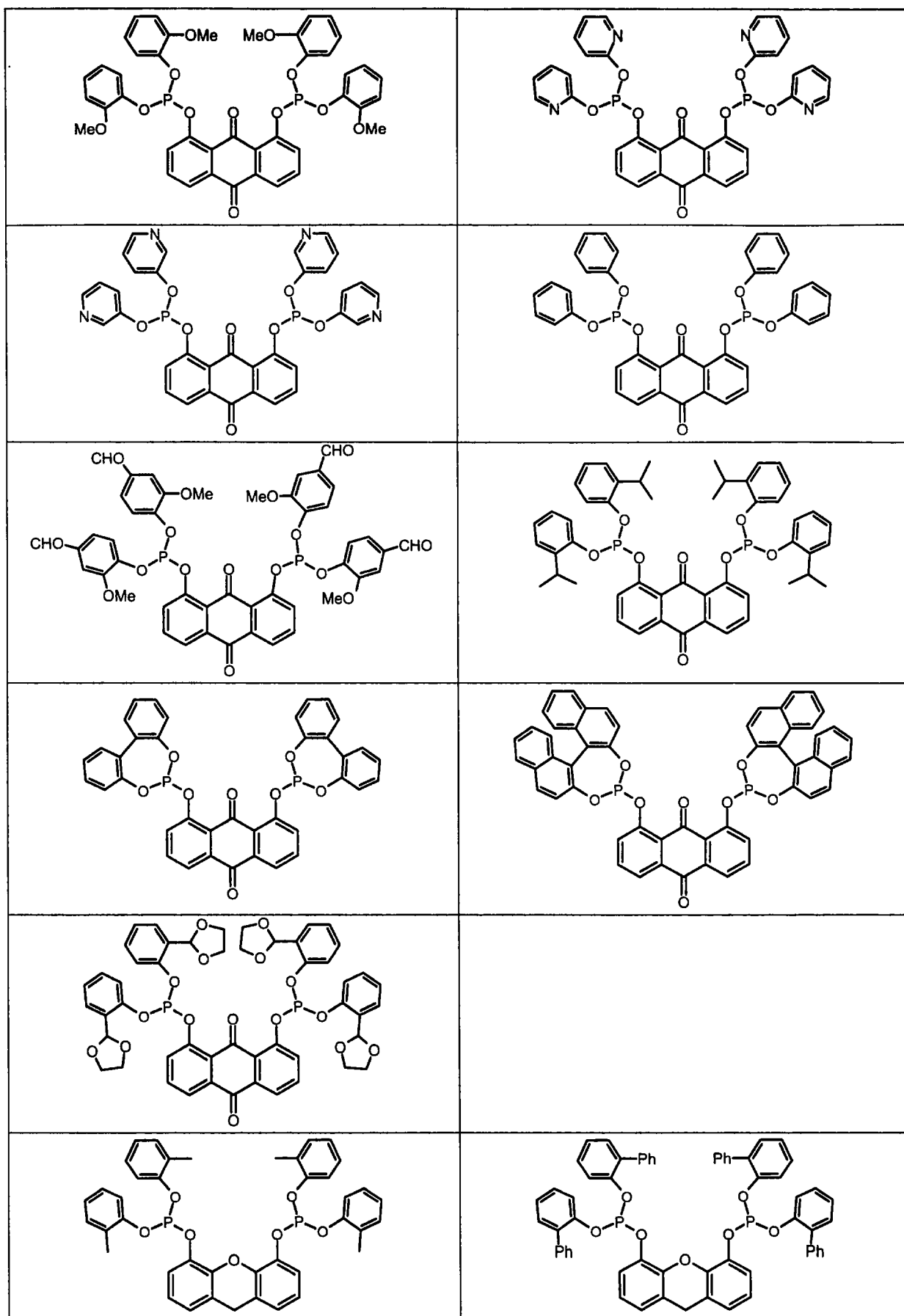


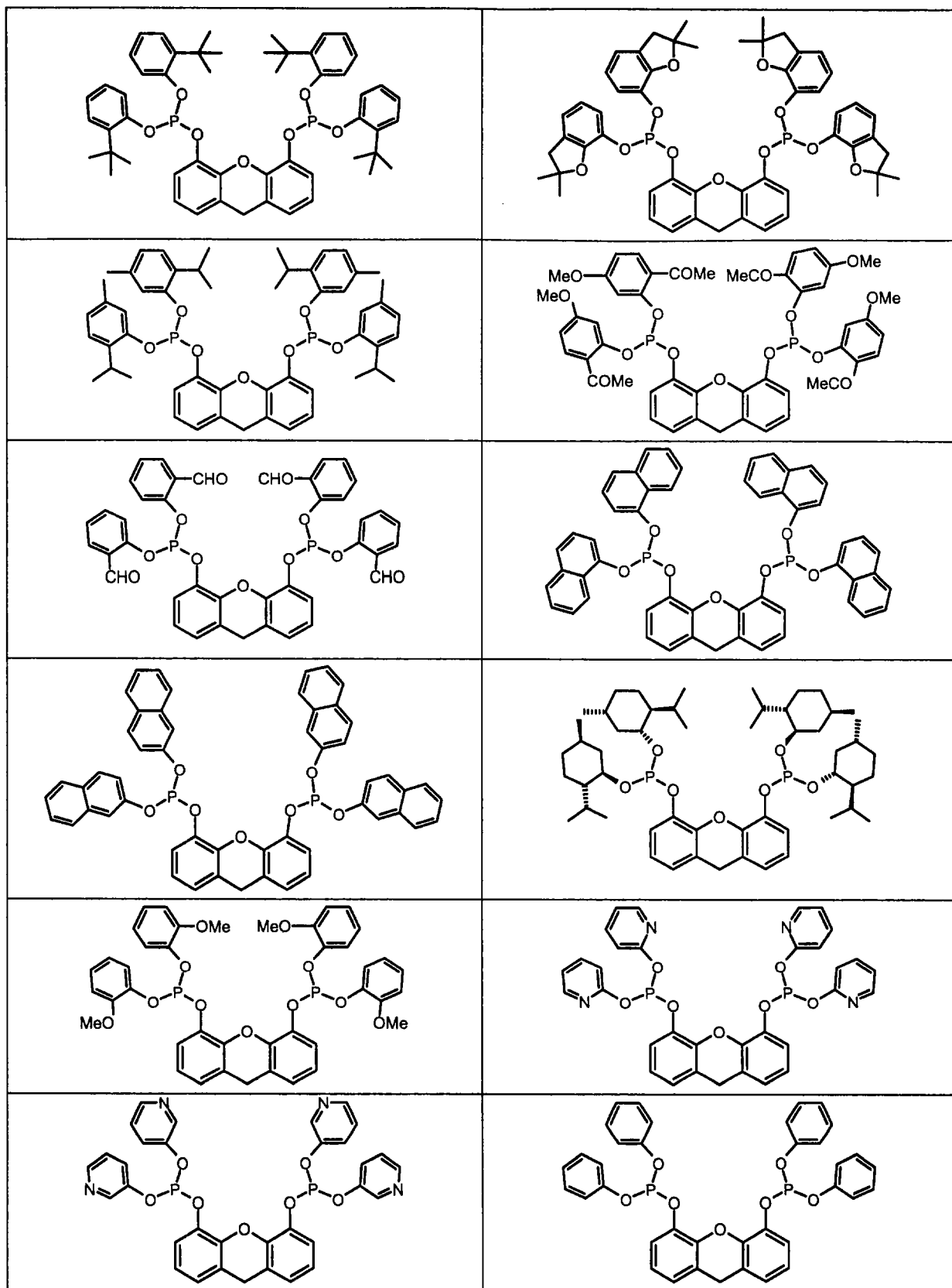


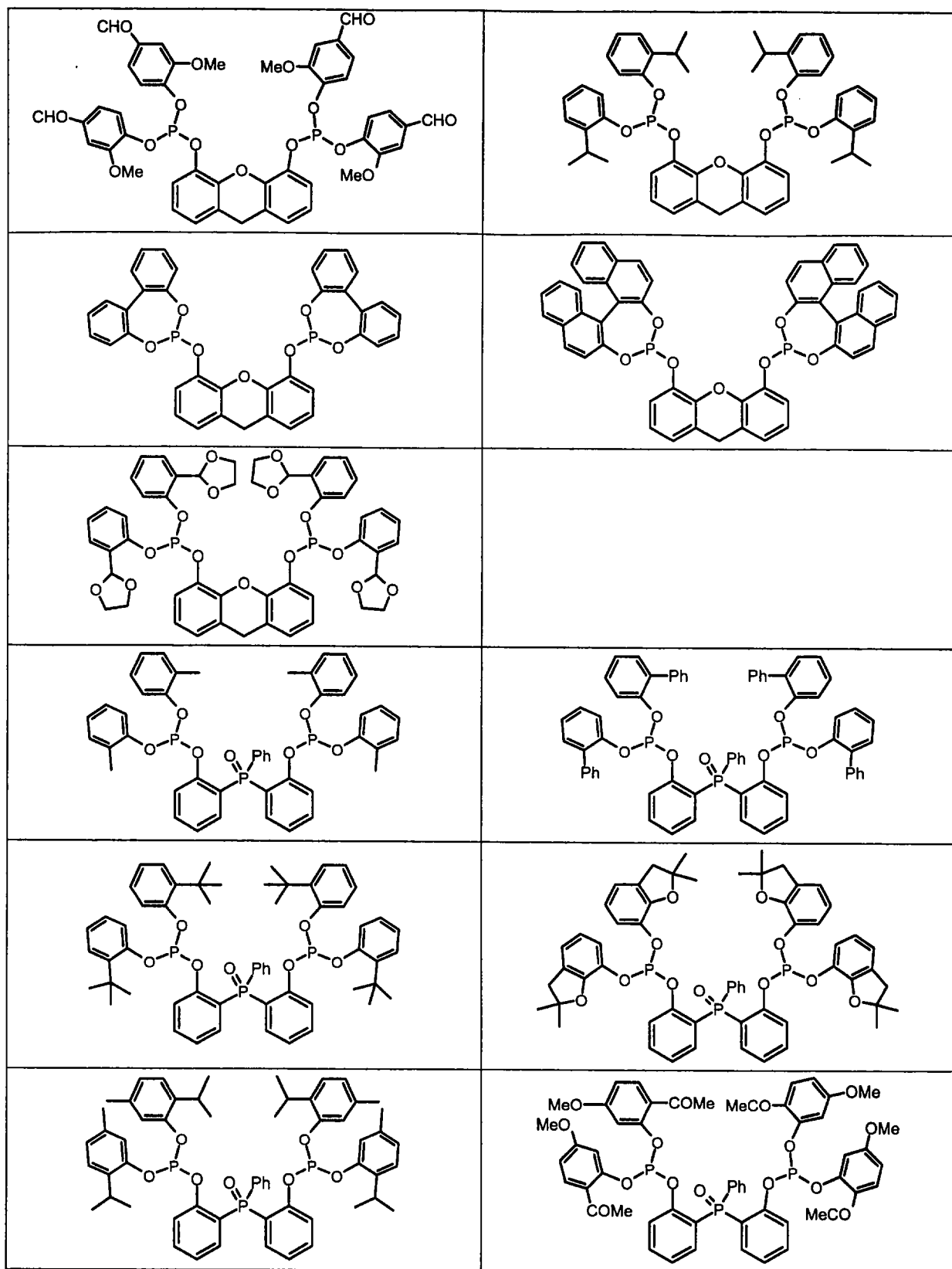


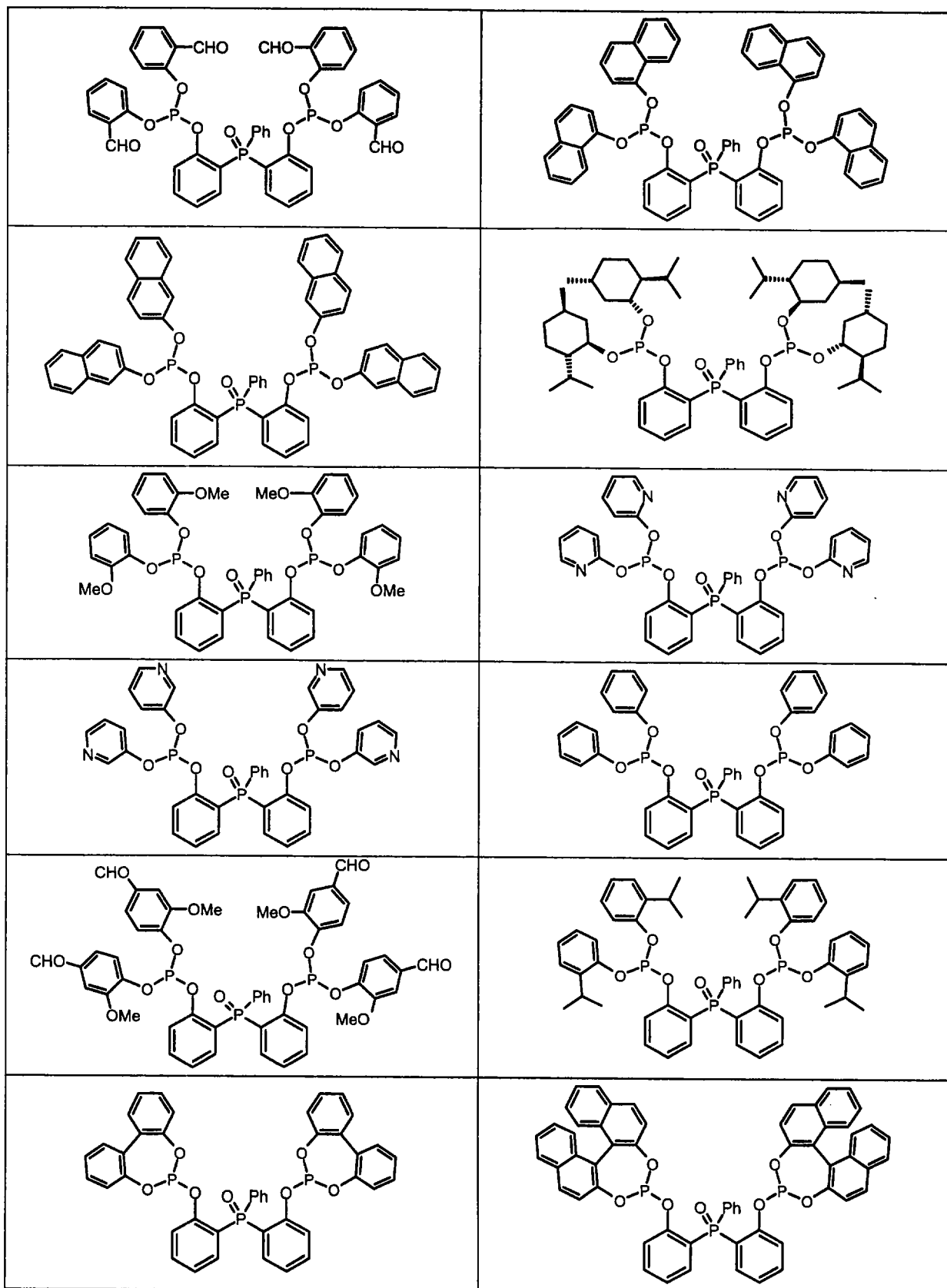


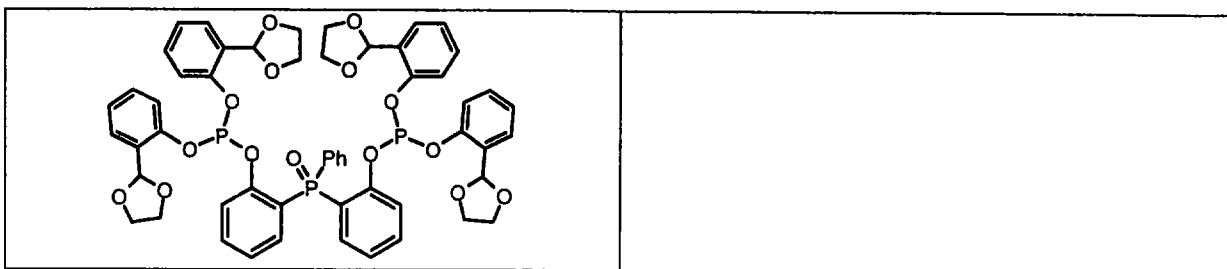












4. Process according to one of Claims 1 to 3,
characterized in that the metallic element is selected
from the group consisting of nickel, cobalt, iron,
5 ruthenium, rhodium, palladium, osmium, iridium,
platinum, copper, silver, gold, zinc, cadmium and
mercury.

5. Process according to one of the preceding claims,
10 characterized in that the reaction is carried out in a
single-phase medium.

6. Process according to one of the preceding claims,
characterized in that the catalyst corresponds to the
15 general formula (II):



in which

M is a transition metal,

L_f represents the organic ligand of formula (I) and

20 t represents a number between 1 and 4 (inclusive).

7. Process according to one of the preceding claims,
characterized in that the reaction mixture comprises a
solvent for the catalyst which is miscible with the
25 phase comprising the compound to be hydrocyanated at
the hydrocyanation temperature.

8. Process according to one of the preceding claims,
characterized in that the transition metal compounds
30 are nickel compounds selected from the group consisting
of:

- compounds in which nickel is in oxidation state

zero, such as potassium tetracyanonickelate $K_4[Ni(CN)_4]$, bis(acrylonitrile)nickel zero, bis-(cycloocta-1,5-diene)nickel zero and derivatives containing ligands, such as tetrakis(tri-phenylphosphine)nickel zero;

- compounds of nickel such as carboxylates, carbonate, bicarbonate, borate, bromide, chloride, citrate, thiocyanate, cyanide, formate, hydroxide, hydrophosphite, phosphite, phosphate and derivatives, iodide, nitrate, sulphate, sulphite, aryl- and alkylsulphonates.

9. Process according to one of the preceding claims, characterized in that the organic compounds containing at least one ethylenic double bond are selected from diolefins such as butadiene, isoprene, hexa-1,5-diene, cycloocta-1,5-diene, ethylenically unsaturated aliphatic nitriles, especially linear pentenenitriles such as pent-3-enenitrile and pent-4-enenitrile, monoolefins such as styrene, methylstyrene, vinylnaphthalene, cyclohexene and methylcyclohexene and also mixtures of two or more of these compounds.

10. Process according to one of the preceding claims, characterized in that the amount of compound of nickel or of another transition metal used is selected such that per mole of organic compound to be hydrocyanated or isomerized between 10^{-4} and 1 mol of nickel or of the other transition metal is employed and in that the amount of organic ligand of formula (I) used is selected such that the number of moles of this compound relative to 1 mol of transition metal is from 0.5 to 50.

11. Process according to one of the preceding claims, characterized in that the hydrocyanation reaction is carried out at a temperature from $10^{\circ}C$ to $200^{\circ}C$.

12. Process according to one of the preceding claims for hydrocyanating ethylenically unsaturated nitrile compounds to dinitriles by reaction with hydrogen cyanide, characterized in that it is operated in the presence of a catalyst system comprising at least one transition metal compound, at least one organic compound of formula (I) and a cocatalyst composed of at least one Lewis acid.
13. Process according to Claim 12, characterized in that the ethylenically unsaturated nitrile compounds are selected from ethylenically unsaturated aliphatic nitriles comprising linear pentenenitriles such as pent-3-enenitrile and pent-4-enenitrile and mixtures thereof.
14. Process according to Claim 13, characterized in that the linear pentenenitriles contain amounts of other compounds selected from the group consisting of 2-methylbut-3-enenitrile, 2-methylbut-2-enenitrile, pent-2-enenitrile, valeronitrile, adiponitrile, 2-methylglutaronitrile, 2-ethylsuccinonitrile and butadiene.
15. Process according to one of Claims 12 to 14, characterized in that the Lewis acid employed as cocatalyst is selected from compounds of the elements of groups Ib, IIb, IIIa, IIIb, IVa, IVb, Va, Vb, VIb, VIIb and VIII of the Periodic Table of the Elements.
16. Process according to one of Claims 12 to 15, characterized in that the Lewis acid is selected from salts selected from the group of halides, sulphates, sulphonates, haloalkylsulphonates, perhaloalkylsulphonates, haloalkylacetates, perhaloalkylacetates, carboxylates and phosphates.
17. Process according to one of Claims 12 to 16, characterized in that the Lewis acid is selected from

zinc chloride, zinc bromide, zinc iodide, manganese chloride, manganese bromide, cadmium chloride, cadmium bromide, stannous chloride, stannous bromide, stannous sulphate, stannous tartrate, indium trifluoromethylsulphonate, indium trifluoroacetate, zinc trifluoroacetate, the chlorides or bromides of rare earth elements such as lanthanum, cerium, praseodymium, neodymium, samarium, europium, gadolinium, terbium, dysprosium, hafnium, erbium, thallium, ytterbium and lutetium, and cobalt chloride, ferrous chloride, yttrium chloride and mixtures thereof.

18. Process according to one of Claims 12 to 17, characterized in that the Lewis acid employed represents from 0.01 to 50 mol per mole of transition metal compound.

19. Process according to one of Claims 1 to 18, characterized in that 2-methylbut-3-enenitrile, present in the reaction mixture originating from butadiene hydrocyanation, is isomerized to pentenenitriles in the absence of hydrogen cyanide, in the presence of a catalyst comprising at least one organic ligand of general formula (I) and at least one transition metal compound.

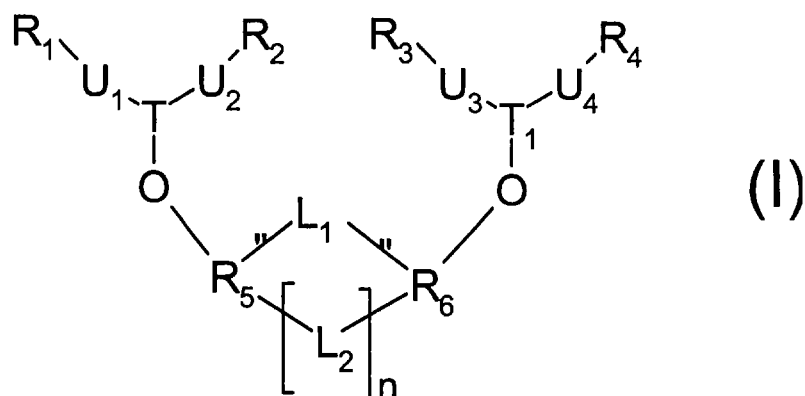
20. Process according to Claim 19, characterized in that the 2-methylbut-3-enenitrile subjected to isomerization is employed alone or in a mixture with 2-methylbut-2-enenitrile, pent-4-enenitrile, pent-3-enenitrile, pent-2-enenitrile, butadiene, adiponitrile, 2-methylglutaronitrile, 2-ethylsuccinonitrile or valeronitrile.

21. Process according to either of Claims 19 and 20, characterized in that the isomerization reaction is carried out at a temperature from 10°C to 200°C.

22. Process according to Claims 19 to 21,

characterized in that the isomerization of 2-methylbut-3-enenitrile to pentenenitriles is carried out in the presence of at least one transition metal compound, at least one organic phosphorous ligand of the formula (I)
5 and a cocatalyst composed of at least one Lewis acid.

23. Organophosphorus compounds corresponding to the general formula I below:



10

in which:

15 T and T₁, which are identical or different, represent a phosphorus, arsenic or antimony atom,

R₁, R₂, R₃ and R₄, which are identical or different, represent a substituted or unsubstituted, aromatic,
20 aliphatic or cycloaliphatic radical comprising one or more rings, which are in fused form or not and which may contain one or more heteroatoms, where the radicals R₁ and R₂ on the one hand and R₃ and R₄ on the other hand may be interconnected by a covalent bond, a hydrocarbon
25 chain or a heteroatom,

U₁, U₂, U₃ and U₄, which are identical or different, represent an oxygen atom or a radical of formula NR in which R denotes a monovalent alkyl, aryl, cycloalkyl,
30 sulphonyl or carbonyl radical,

R_5 and R_6 , which are identical or different, represent an aryl or cycloaliphatic group which may comprise heteroatoms and/or one or more rings, in fused form or not, and which are substituted or unsubstituted,

5

n is an integer equal to 0 or 1,

L_1 , when n is 0, represents a divalent radical selected from the group consisting of the groups NR_7 , PR_8 , SiR_9R_{10} , BR_{11} , S , POR_{12} , SO_2 and CO , in which R_7 is as defined for R above, R_8 and R_{12} may represent the radical OR_{13} , and R_8 , R_9 , R_{10} , R_{11} , R_{12} and R_{13} represent alkyl, aryl or cycloalkyl radicals,

15 L_1 and L_2 , when n is 1, are identical or different and represent a covalent bond or a radical selected from the group consisting of the groups O , NR_7 , PR_8 , SiR_9R_{10} , BR_{11} , S , POR_{12} , SO_2 , CO and $-CR_{14}R_{15}-$, in which R_7 is as defined for R above, R_8 and R_{12} may represent the radical OR_{13} , and R_8 , R_9 , R_{10} , R_{11} , R_{12} , R_{13} , R_{14} and R_{15} represent alkyl, aryl or cycloalkyl radicals, it being possible also for R_{14} and R_{15} to represent a hydrogen atom.

25 24. Compounds according to Claim 23, corresponding to the formulae below:

