

[54] 1-(2-OXYAMINOSULPHONYLPHENYLSULPHONYL)-3-HETEROARYL-UREAS

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[21] Appl. No.: 369,583

[22] Filed: Jun. 21, 1989

Related U.S. Patent Documents

Reissue of:

[64] Patent No.: 4,666,505
Issued: May 19, 1987
Appl. No.: 769,224
Filed: Aug. 23, 1985

[30] Foreign Application Priority Data

Aug. 30, 1984 [DE] Fed. Rep. of Germany 3431929
May 25, 1985 [DE] Fed. Rep. of Germany 3518876

[51] Int. Cl.⁵ C07D 239/69; A01N 43/54

[52] U.S. Cl. 71/92; 71/94;
544/332; 544/320; 544/321; 544/323; 544/327;
546/305

[58] Field of Search 71/92, 94; 544/332,
544/323; 546/305

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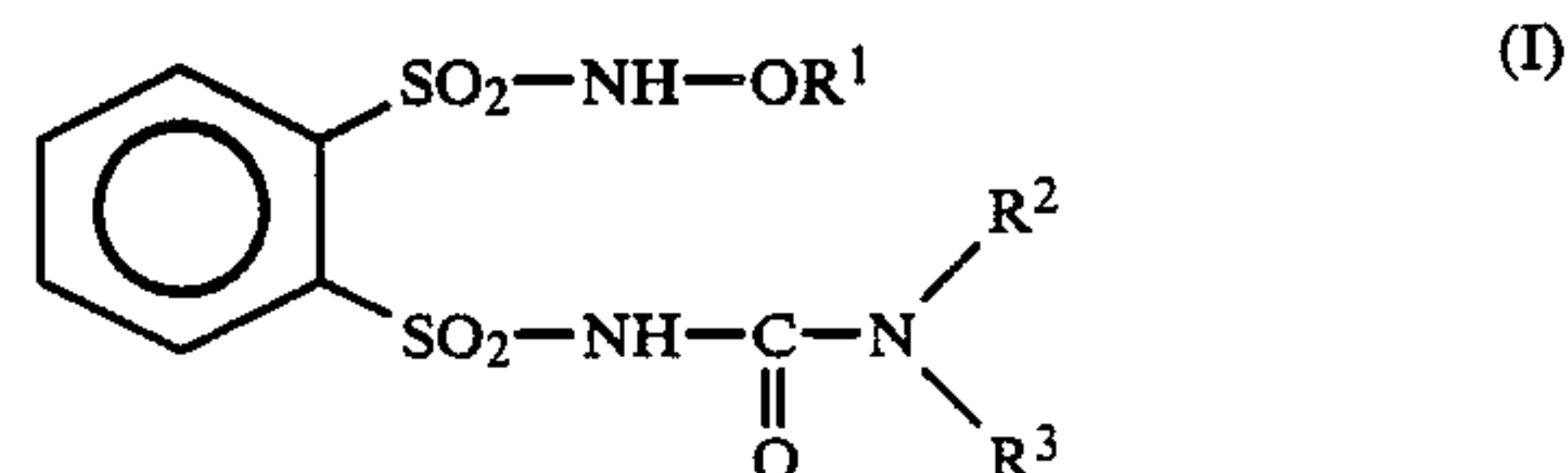
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[57] ABSTRACT

The invention relates to new 1-(2-oxyaminosulphonylphenylsulphonyl)-3-heteroaryl-ureas of the general formula (I)



in which

R¹ represents an optionally substituted radical from the series comprising alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, aralkyl and aryl,

R² represents hydrogen or an optionally substituted radical from the series comprising alkyl, alkenyl, alkynyl and aralkyl and

R³ represents an optionally substituted and/or optionally fused six-membered aromatic heterocyclic radical containing at least one nitrogen atom, the following compounds being excluded: 1-(2-methoxyaminosulphonylphenylsulphonyl)-, 1-(2-ethoxyaminosulphonylphenylsulphonyl)-, 1-(2-propoxyaminosulphonylphenylsulphonyl)-, 1-(2-isopropoxyaminosulphonylphenylsulphonyl)- and 1-(2-butoxyaminosulphonylphenylsulphonyl)-3-(4,6-dimethylpyrimidin-2-yl)-urea -3-(4,6-diethylpyrimidin-2-yl)-urea, -3-(4,6-dipropylpyrimidin-2-yl)-urea, -3-(4,6-diisopropylpyrimidin-2-yl)-urea and -3-(4,6-dibutylpyrimidin-2-yl)-urea, processes for their preparation and their use as herbicides.

11 Claims, No Drawings

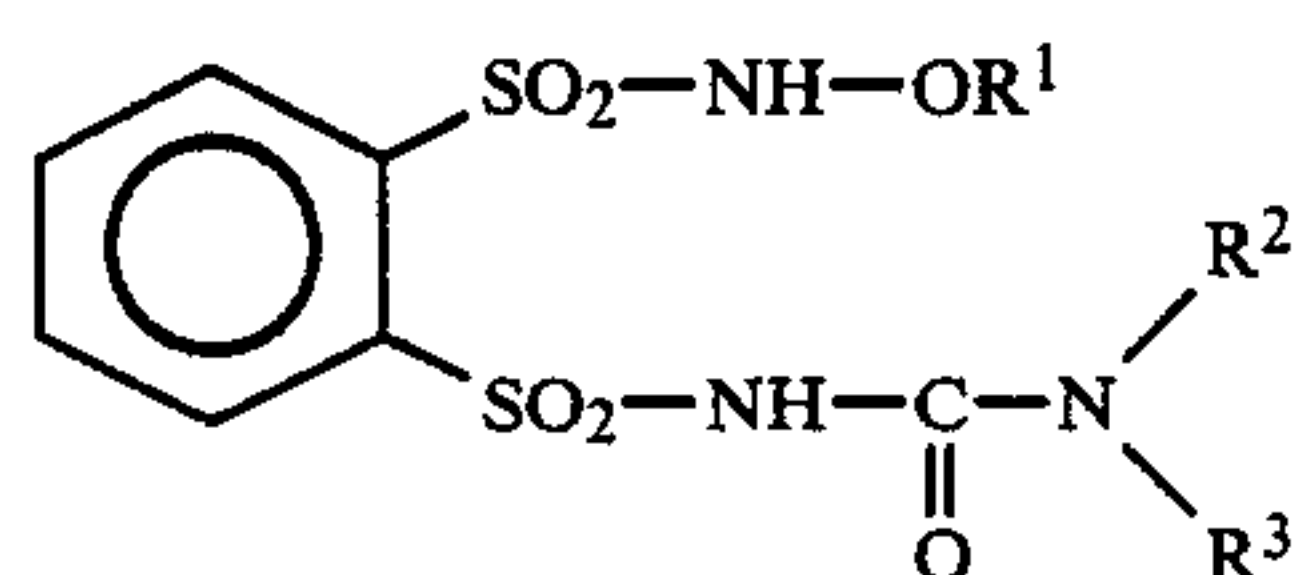
1-(2-OXYAMINOSULPHONYLPHENYLSULPHONYL)-3-HETEROARYL-UREAS

Matter enclosed in heavy brackets [] appears in the original patent but forms no part of this reissue specification; matter printed in italics indicates the additions made by reissue.

The invention relates to new 1-(2-oxyaminosulphonylphenylsulphonyl)-3-heteroaryl-ureas, processes for their preparation and their use as herbicides.

It is known that certain 1-arylsulphonyl-3-heteroarylsureas, such as, for example, 1-(2-methoxyphenylsulphonyl)-3-(4,6-dimethyl-pyrimidin-2-yl)-urea, have a herbicidal action. However, the action of these compounds is not always completely satisfactory (compare U.S. Pat. No. 4,169,719).

New 1-(2-oxyaminosulphonylphenylsulphonyl)-3-heteroaryl-ureas of the general formula (I)



in which

R¹ represents an optionally substituted radical from the series comprising alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, aralkyl and aryl,

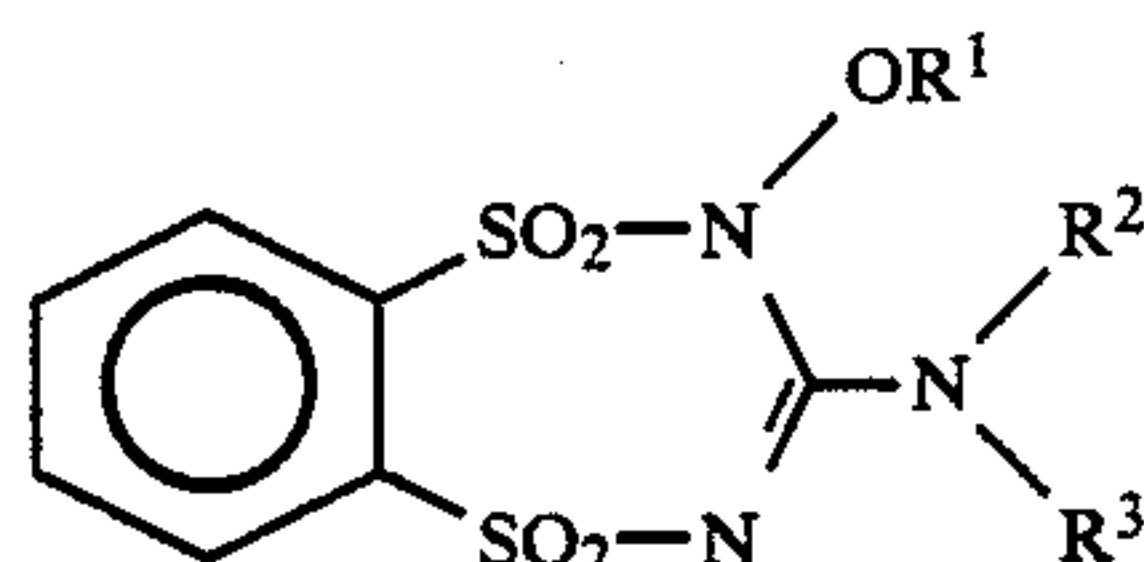
R² represents hydrogen or an optionally substituted radical from the series comprising alkyl, alkenyl, alkynyl and aralkyl and

R³ represents an optionally substituted and/or optionally fused six-membered aromatic heterocyclic radical containing at least one nitrogen atom,

have now been found, the following compounds being excluded: 1-(2-methoxyaminosulphonylphenylsulphonyl)-, 1-(2-ethoxyaminosulphonylphenylsulphonyl)-, 1-(2-propoxyaminosulphonylphenylsulphonyl)-, 1-(2-isopropoxyaminosulphonylphenylsulphonyl)- and 1-(2-butoxyaminosulphonylphenylsulphonyl)-, -3-(4,6-dimethyl-pyrimidin-2-yl)-urea, -3-(4,6-diethyl-pyrimidin-2-yl)-urea, -3-(4,6-dipropylpyrimidin-2-yl)-urea, -3-(4,6-diisopropyl-pyrimidin-2-yl)-urea and -3-(4,6-dibutyl-pyrimidin-2-yl)-urea.

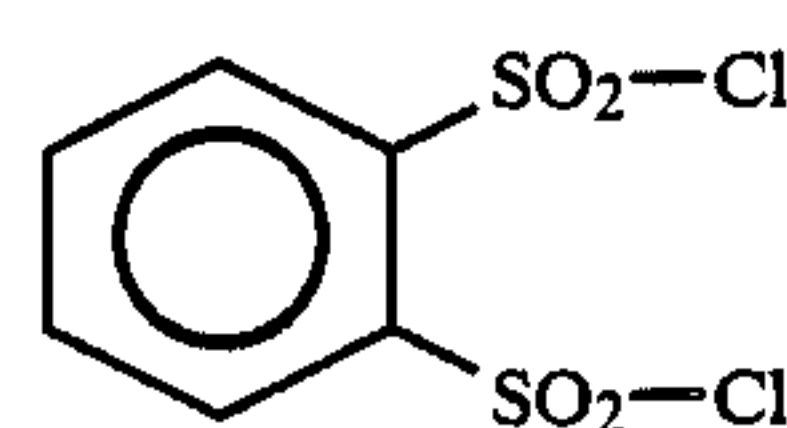
The new compounds of the formula (I) are obtained by a process in which

(a) benzodisultams of the formula (II)

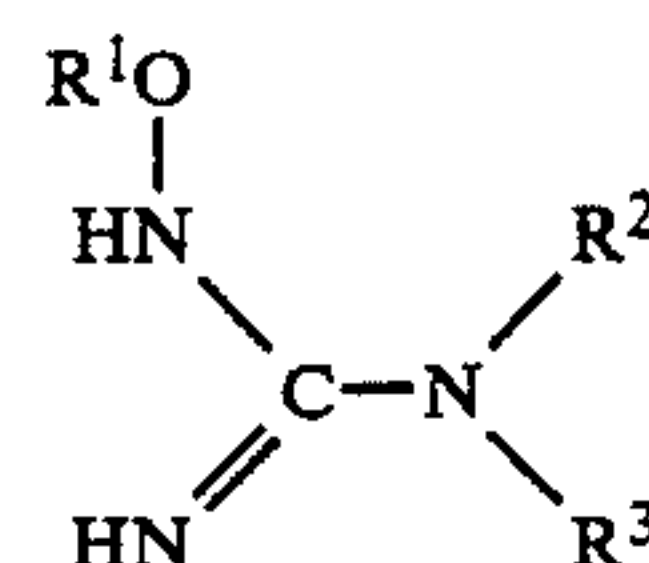


in which R¹, R² and R³ have the abovementioned meanings, are reacted with water, if appropriate in the presence of bases and if appropriate in the presence of diluents, or

(b) benzene-1,2-disulphonic acid dichloride of the formula (III)



is reacted with oxyguanidine derivatives of the formula (IV)



in which R¹, R² and R³ have the abovementioned meaning, in the presence of acid acceptors and if appropriate in the presence of diluents, and the compounds of the formula (II) thereby obtained are reacted—without intermediate isolation—with water, if appropriate in the presence of bases and if appropriate in the presence of diluents.

The new 1-(2-oxyaminosulphonylphenylsulphonyl)-3-heteroaryl-ureas of the formula (I) are distinguished by a powerful herbicidal activity.

Surprisingly, the new compounds of the formula (I) exhibit a considerably better herbicidal action than the already known urea derivatives of the same type of action.

The invention preferably relates to compounds of the formula (I) in which

R¹ represents C₁-C₁₂-alkyl [] (which is optionally substituted by fluorine, chlorine, cyano, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-alkylsulphinyl, C₁-C₄-alkylsulphonyl, C₁-C₄-alkyl-carbonyl, C₁-C₄-alkoxy-carbonyl, C₁-C₄-alkylamino-carbonyl or di-(C₁-C₄-alkyl)-amino-carbonyl []), or represents C₃-C₆-alkenyl [] (which is optionally substituted by fluorine, chlorine or bromine []), C₃-C₆-alkynyl, C₃-C₆-cycloalkyl, C₃-C₆-cycloalkyl-C₁-C₂-alkyl or phenyl-C₁-C₂-alkyl [] (which is optionally substituted by fluorine, chlorine, nitro, cyano, C₁-C₄-alkyl, C₁-C₄-alkoxy or C₁-C₄-alkoxy-carbonyl []), or represents benzhydryl, or represents phenyl [] (which is optionally substituted by fluorine, chlorine, nitro, cyano, C₁-C₄-alkyl, trifluoromethyl, C₁-C₄-alkoxy, C₁-C₂-fluoroalkoxy, C₁-C₄-alkylthio, trifluoromethylthio or C₁-C₄-alkoxycarbonyl []),

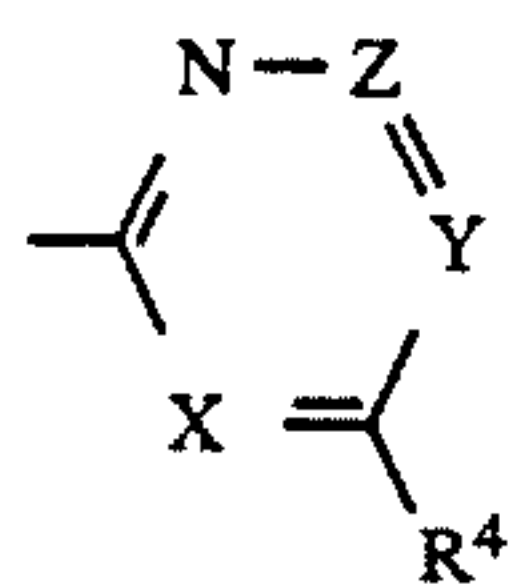
and in which, furthermore,

R² represents hydrogen or C₁-C₄-alkyl [] (which is optionally substituted by fluorine, chlorine, cyano, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-alkylsulphinyl, C₁-C₄-alkylsulphonyl, C₁-C₄-alkyl-carbonyl, C₁-C₄-alkoxy-carbonyl, C₁-C₄-alkylamino-carbonyl or di-(C₁-C₄-alkyl)-aminocarbonyl []), or represents C₃-C₆-alkenyl, C₃-C₆-alkynyl or phenyl-C₁-C₂-alkyl [] (which is optionally substituted by fluorine, chlorine, nitro, cyano, C₁-C₄-alkyl, C₁-C₄-alkoxy or C₁-C₄-alkoxycarbonyl []),

and in which, furthermore,

R³ represents the radical

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wherein

R⁴ represents hydrogen, fluorine, chlorine, bromine, hydroxyl, C₁-C₄-alkyl [] (which is optionally substituted by fluorine and/or chlorine []), C₁-C₄-alkoxy [] (which is optionally substituted by fluorine and/or chlorine []), C₁-C₄-alkylthio [] (which is optionally substituted by fluorine and/or chlorine []), amino, C₁-C₄-alkylamino or di-(C₁-C₄-alkyl)-amino,

X represents nitrogen or a methine bridge (CH),

Y represents nitrogen or an optionally substituted methine bridge C-R⁵,

wherein

R⁵ represents hydrogen, fluorine, chlorine, bromine, cyano, formyl, C₁-C₄-alkyl, C₁-C₃-alkoxy-carbonyl or C₁-C₃-alkyl-carbonyl, and Z represents nitrogen or an optionally substituted methine bridge C-R⁶,

wherein

R⁶ represents hydrogen, fluorine, chlorine, bromine, hydroxyl, C₁-C₄-alkyl [] (which is optionally substituted by fluorine and/or chlorine []), C₁-C₄-alkoxy [] (which is optionally substituted by fluorine and/or chlorine []), C₁-C₄-alkylthio [] (which is optionally substituted by fluorine and/or chlorine []), amino, C₁-C₄-alkylamino or di-(C₁-C₄-alkyl)-amino,

the compounds mentioned above by name being excluded.

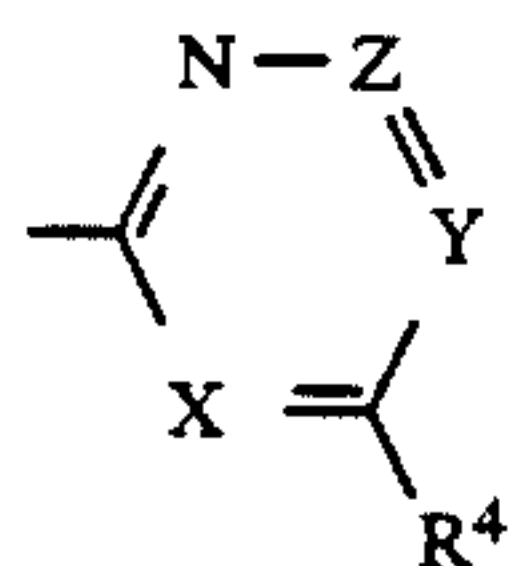
The invention particularly relates to compounds of the formula (I)

in which

R¹ represents C₁-C₈-alkyl [] (which is optionally substituted by fluorine or chlorine []), C₃-C₄-alkenyl, C₁-C₂-alkoxycarbonylmethyl, phenyl, phenethyl or benzyl [] (which is optionally substituted by fluorine, chlorine, nitro, cyano, methyl, methoxy or methoxy-carbonyl []),

R² represents hydrogen and

R³ represents the radical



wherein

R⁴ represents chlorine, methyl, ethyl, methoxy, difluoromethoxy or ethoxy,

X represents nitrogen,

Y represents a methine bridge (CH) and

Z represents an optionally substituted methine bridge C-R⁶,

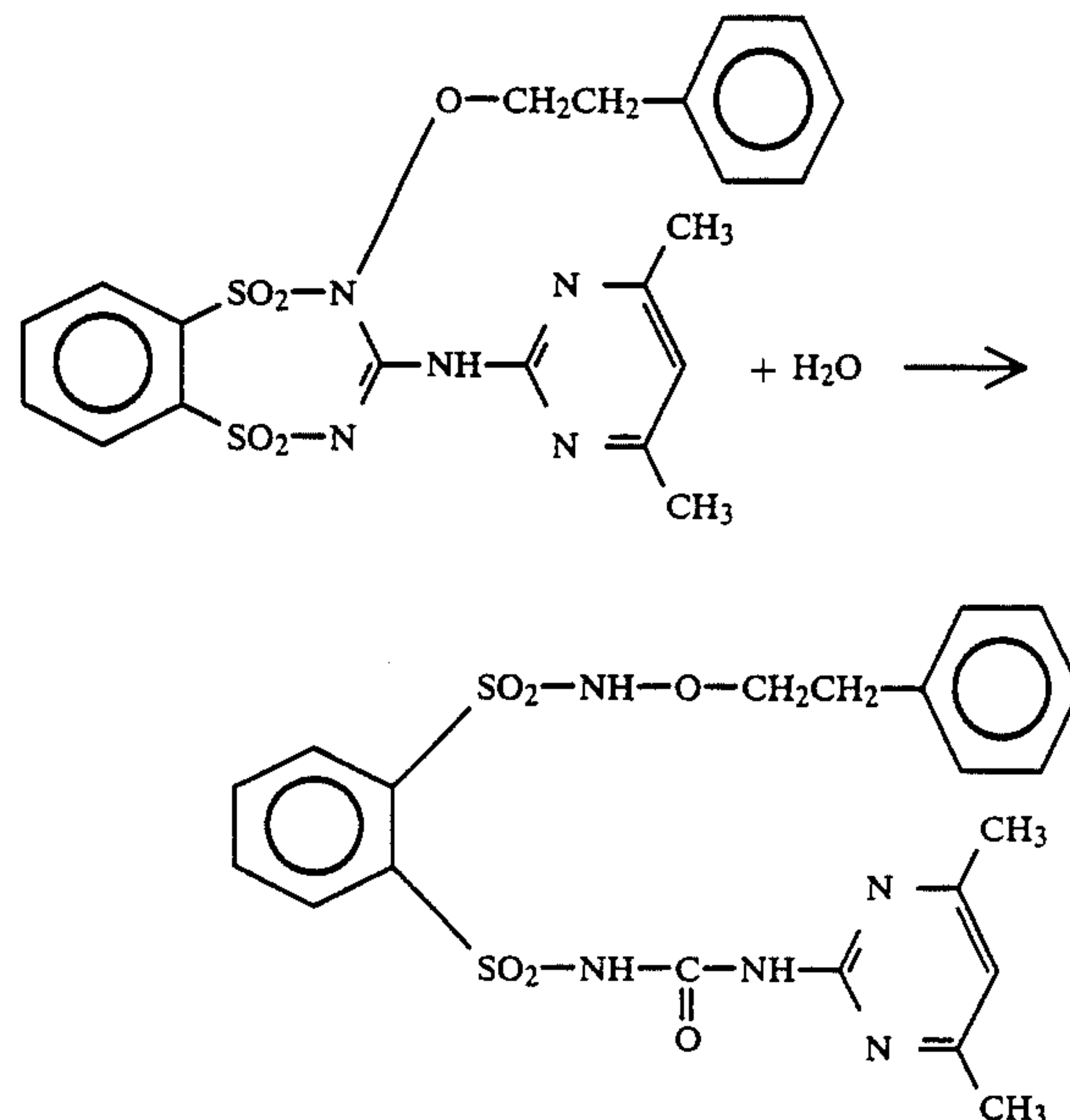
wherein

R⁶ represents hydrogen, chlorine, methyl, methoxy, ethoxy, methylthio, ethylthio, dimethylamino or diethylamino,

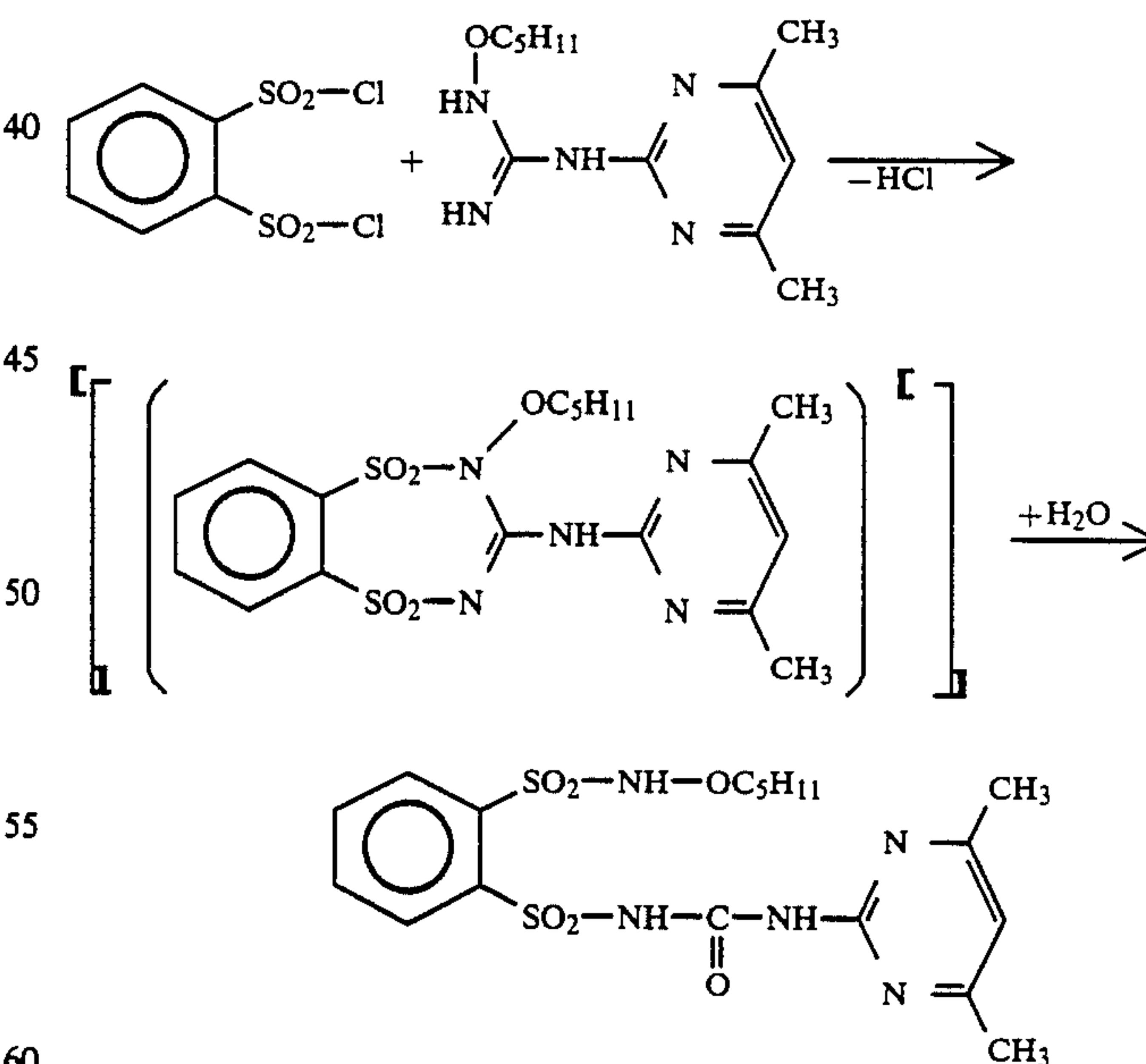
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the compounds mentioned above by name being excluded.

The chemical reaction which takes place in the preparation process according to the invention described above under (a) can be outlined, for example, by the following equation:



The [reaction] reactions which take place in the preparation process according to the invention described above under (b) can be outlined, for example, by the following equation:



Formula (II) provides a general definition of the benzodisultams to be used as starting substances in process (a) according to the invention.

In formula (II), R¹, R² and R³ preferably and particularly have the same meanings as are given above as preferred or as particularly preferred in the context of the definition of the substituents for formula (I).

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Examples of starting substances of formula (II) are listed in the following Table 1.

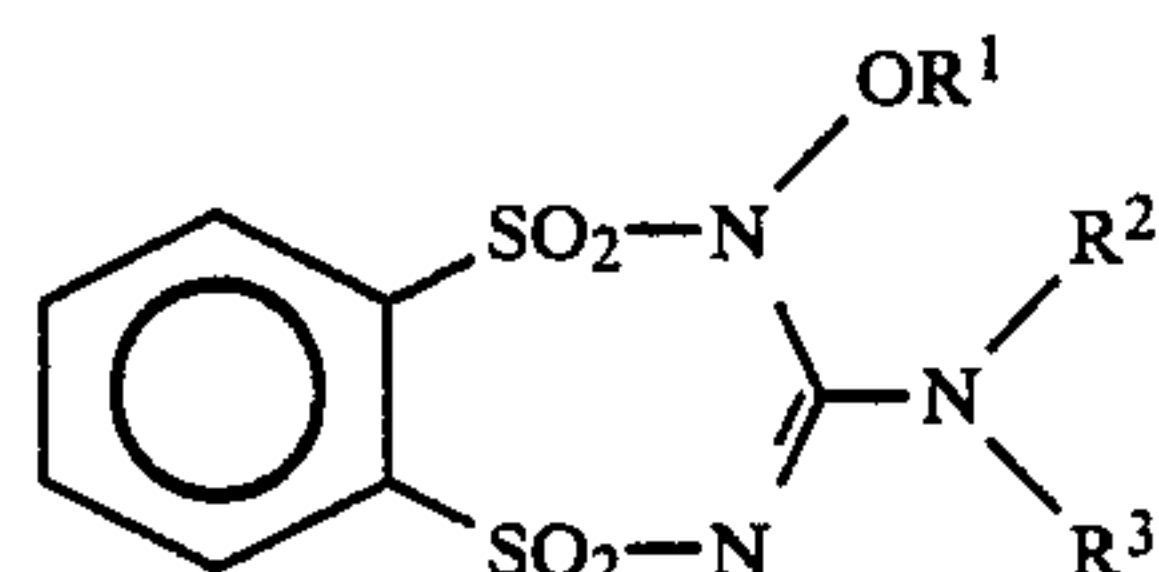


TABLE 1

Examples of starting substances of the formula (II)		
R ¹	R ²	R ³
	H	
-CH ₃	CH ₃	
	H	
-C ₃ H ₇	H	
-CH(CH ₃) ₂	H	
-C ₄ H ₉	H	
-CH ₂ CH(CH ₃) ₂	H	

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TABLE 1-continued

Examples of starting substances of the formula (II)		
R ¹	R ²	R ³
(II) 5 -C ₈ H ₁₇	H	
10	H	
15	H	
20	H	
25	H	
30	H	
35 -CH ₂ CH=CH ₂	H	
40 -CH ₃	H	
45	H	
50 -C ₂ H ₅	H	
55 -CH ₂ CH=CH ₂	H	
60	H	
65	H	

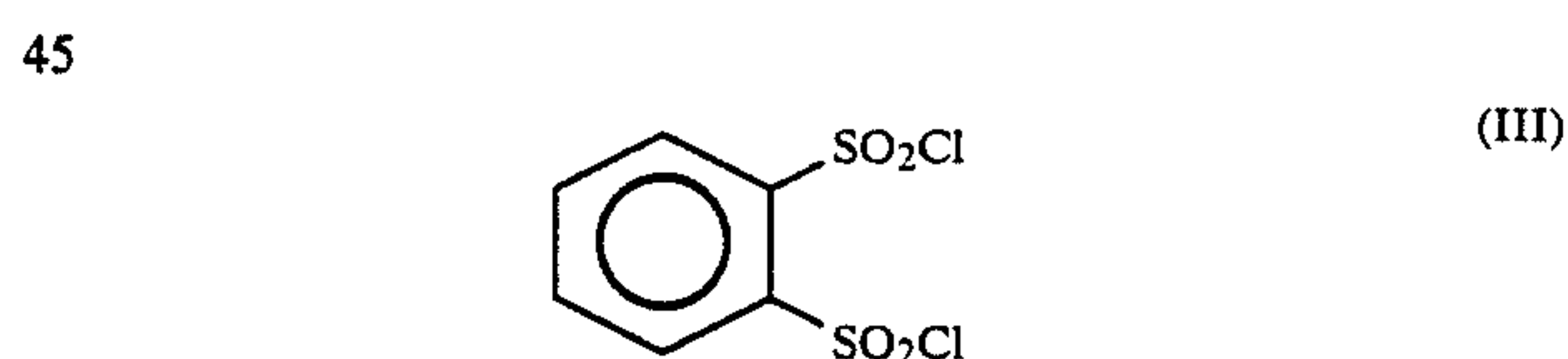
TABLE 1-continued

Examples of starting substances of the formula (II)		
R ¹	R ²	R ³
—CH ₃	H	
—C ₂ H ₅	H	
—CH ₃	H	
	H	
—CH ₃	H	
—C ₂ H ₅	H	
—CH ₃	H	
—CH ₂ CH=CH ₂	H	

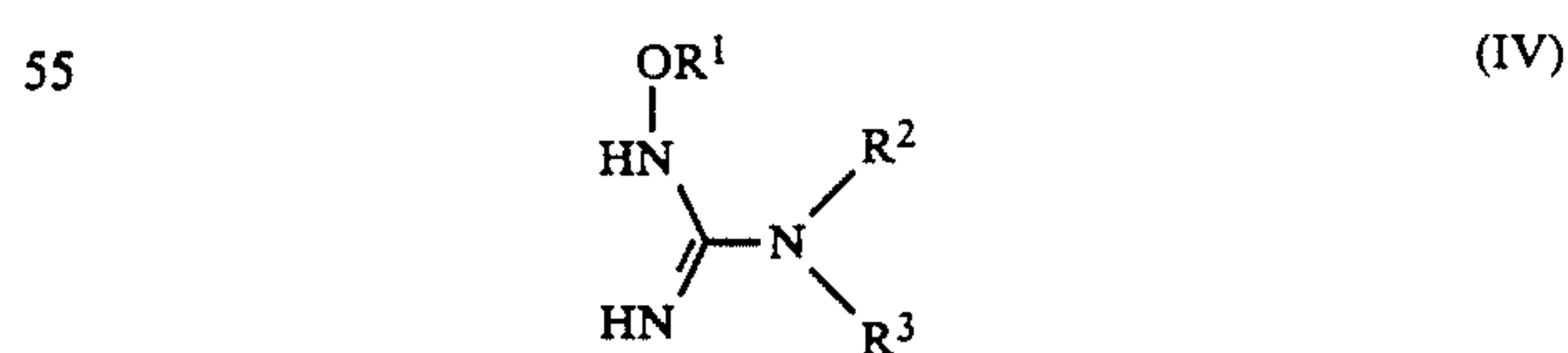
TABLE 1-continued

Examples of starting substances of the formula (II)		
R ¹	R ²	R ³
—CH ₃	H	
—CH ₃	—C ₂ H ₅	
—C ₃ H ₇ (—n)	H	
—CH ₂ COOC ₂ H ₅	H	
—CH ₃	H	

40 The compounds of the formula (II) have not yet been described in the literature. The compounds of the formula (II) are obtained by a process in which benzene-1,2-disulphonic acid dichloride of the formula (III)



50 is reacted with oxyguanidine derivatives of the formula (IV)



60 in which R¹, R² and R³ have the abovementioned meaning, in the presence of acid acceptors, such as, for example, pyridine or diazabicyclooctane (DABCO), and, if appropriate, in the presence of diluents, such as, for example, methylene chloride, chloroform, tetrahydrofuran or dioxane, at temperatures between —30° C. and +50° C.

Working up can be effected by customary methods, for example by concentrating the mixture, taking up the residue in methylene chloride, washing the mixture with dilute hydrochloric acid and with water and separating off, drying, filtering and concentrating the organic phase, the products of the formula (II) remaining in the residue.

The benzene-1,2-disulphonic acid dichloride of the formula (III) to be used as the starting substance is already known (compare J. Org. Chem. 31, (1966), 3289-3292).

Formula (IV) provides a general definition of the oxyguanidine derivatives also to be used as starting substances. In formula (IV), R^1 , R^2 and R^3 preferably and particularly have the same meanings as are given above as preferred or as particularly preferred in the context of the definition of the substituents for formula (I).

Examples which may be mentioned of starting substances of the formula (IV) are: N' -(4-methyl-pyrimidin-2-yl)-, N' -(4-ethyl-pyrimidin-2-yl)-, N' -(4,6-dimethoxypyrimidin-2-yl)-, N' -(2,6-dimethyl-pyrimidin-4-yl)-, N' -(4-difluoromethoxy-6-methyl-pyrimidin-2-yl), N' -(2,6-dimethyl-pyrimidin-4-yl)-, N' -(4,6-dimethyl-pyrimidin-2-yl)-, N' -(4-methoxy-6-methyl-pyrimidin-2-yl)-, N' -(4-ethoxy-6-methyl-pyrimidin-2-yl)-, N' -(4-chloro-6-methoxy-pyrimidin-2-yl)-, N' -(4-chloro-6-ethoxy-pyrimidin-2-yl)-, N' -(4-chloro-6-dimethylamino-pyrimidin-2-yl)-, N' -(4-methyl-6-methylthiopyrimidin-2-yl)- and N' -(4-dimethylamino-6-methyl-pyrimidin-2-yl)- N'' -methoxy-guanidine, $-N''$ -ethoxy-guanidine, $-N''$ -propoxy-guanidine, $-N''$ -isopropoxy-guanidine, $-N''$ -butoxy-guanidine, $-N''$ -isobutoxy-guanidine, $-N''$ -sec.-butoxy-guanidine, $-N''$ -pentoxy-guanidine, $-N''$ -isopentoxy-guanidine, $-N''$ -hexyloxy-guanidine, $-N''$ -octyloxy-guanidine, $-N''$ -allyloxy-guanidine, $-N''$ -(2-chloroethoxy)-guanidine, $-N''$ -(2-fluoroethoxy)-guanidine, $-N''$ -(2-chloro-propoxy)-guanidine, $-N''$ -(2-fluoro-propoxy)-guanidine, $-N''$ -(3-chloro-propoxy)-guanidine, $-N''$ -(4-chloro-butoxy)-guanidine, $-N''$ -methoxycarbonylmethoxy-guanidine, $-N''$ -ethoxycarbonylmethoxy-guanidine, $-N''$ -(1-methoxycarbonyl-ethoxy)-guanidine, $-N''$ -(1-ethoxycarbonyl-ethoxy)-guanidine, $-N''$ -dimethylaminocarbonylmethoxy-guanidine, $-N''$ -(2-phenyl-ethoxy)-guanidine, $-N''$ -phenoxy-guanidine, $-N''$ -(4-methyl-benzyloxy)-guanidine, $-N''$ -(4-fluoro-benzyloxy)-guanidine, $-N''$ -(4-chloro-benzyloxy)-guanidine, $-N''$ -(4-nitrobenzyloxy)-guanidine, $-N''$ -(2,6-dichloro-benzyloxy)-guanidine, $-N''$ -(4-methoxycarbonylbenzyloxy)-guanidine and $-N''$ -(4-ethoxycarbonylbenzyloxy)-guanidine.

The starting substances of the formula (IV) are known in some cases (compare J. Chem. Soc. 1962, page 3915 and DE-OS (German Published Specification) No. 3,334,455).

The compounds of the formula (IV) are obtained by a process in which cyanamide derivatives of the formula (V)



in which R^2 and R^3 have the abovementioned meanings, are reacted with hydroxylamine derivatives of formula (VI)



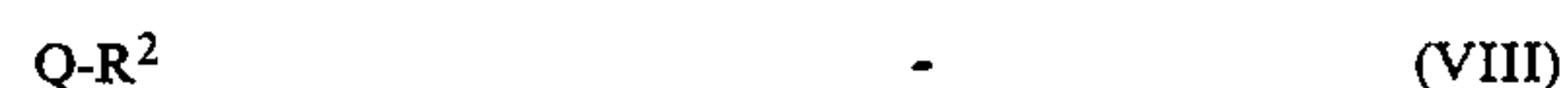
in which R^1 has the abovementioned meanings, or with hydrochlorides of hydroxylamine derivatives of the formula (VI), if appropriate in the presence of diluents, such as, for example, ethanol, propanol or butanol, at temperatures between 20° C. and 120° C. and, if appropriate, the reaction products are treated with acid acceptors, such as, for example, ammonia, potassium carbonate or sodium hydroxide.

The cyanamide derivatives of the formula (V) are known in some cases (compare J. Chem. Soc. 1953, 1725). The compounds of the formula (V) are essentially obtained by the following synthesis routes:

(a) by reaction of alkali metal or alkaline earth metal salts of cyanamide—such as, for example, sodium cyanamide or calcium cyanamide—with chloro-hetarenes of the formula (VII)



in which R^3 has the abovementioned meaning, and, if appropriate, subsequently—if R^2 does not represent hydrogen—with halogen compounds of the formula (VIII)



in which

R^2 represents an optionally substituted radical from the series comprising alkyl, alkenyl, alkynyl and aralkyl and

Q represents chlorine, bromine or iodine, if appropriate in the presence of inert diluents, such as, for example, acetone, acetonitrile or dimethylformamide, at temperatures between 0° C. and 100° C.

After the mixture has been concentrated and the residue has been dissolved in water, the cyanamide derivatives of the formula (V) can be precipitated by acidification, for example with hydrochloric acid, and isolated by filtration with suction.

Alternatively, the compounds of the formula (V) are obtained

(b) in the case where R^3 represents a substituted pyrimidinyl radical, by reaction of cyanoguanidine ("dicyandiamide") with β -dicarbonyl compounds or derivatives thereof, such as acetylacetone (compare J. Chem. Soc. 1953, 1725-1730); acetoacetic acid esters (compare J. Prakt. Chem. 77 (1908), 542 and J. Chem. Soc. 1948, 586) or malonic acid esters (compare German Patent Specification 158,591).

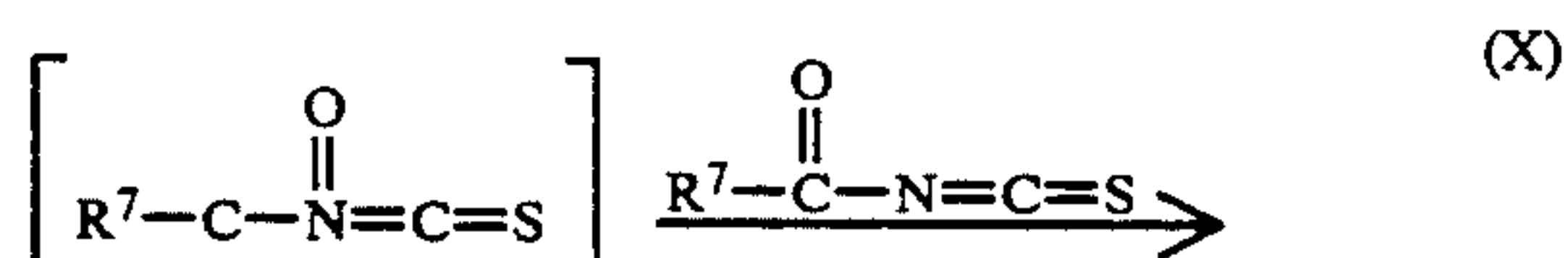
The 2-cyanoamino-4-hydroxy-6-methyl- or -4,6-dihydroxy-pyrimidines obtained from acetoacetic acid esters or malonic acid esters can be converted into corresponding 2-cyanoamino-4-alkoxy-6-methyl- or -4,6-dialkoxy-pyrimidines in a known manner by reaction with alkylating agents, such as, for example, dimethyl sulphate or diethyl sulphate, if appropriate in the presence of diluents, such as, for example, water, methanol, ethanol, n- or iso-propanol, acetone, dioxane or dimethylformamide, and in the presence of acid-binding agents, such as, for example, sodium hydroxide or potassium hydroxide or sodium carbonate or potassium carbonate. To avoid N-alkylation, acylation is carried out, if appropriate, with an acylating agents, such as, for example, acetic anhydride or acetyl chloride, and,

after the alkylation, the product is deacylated again with aqueous acids or bases.

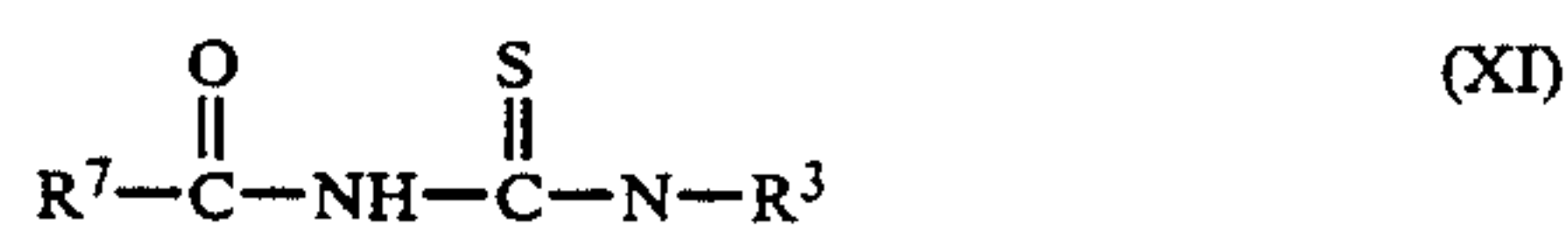
In another alternative process, the compounds of the formula (V) are obtained by a process in which (c) amino-hetarenes of the formula (IX)



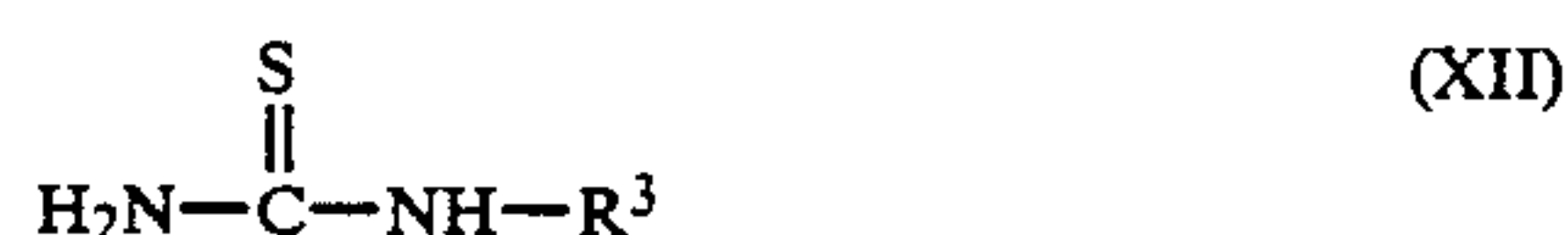
in which R^3 has the abovementioned meaning, are reacted with carbonyl isothiocyanates of the formula (X)



in which R^7 represents ethoxy or phenyl, if appropriate in the presence of an inert diluent, such as, for example, acetone, acetonitrile or toluene, at temperatures between 0°C . and 100°C ., the carbonylthioureas of the formula (XI) thereby formed



in which R^3 and R^7 have the abovementioned meanings, are isolated by filtration with suction, if appropriate after concentration of the mixture, and are reacted with aqueous alkali metal or alkaline earth metal hydroxide solutions, such as, for example, sodium hydroxide solution, if appropriate in the presence of an organic solvent, such as, for example, tetrahydrofuran or dioxane, at temperatures between 0°C . and 120°C ., and the thioureas obtained as crystals after acidification, for example with hydrochloric acid, of the formula (XII)



in which R^3 has the abovementioned meaning, are isolated by filtration with suction and reacted with metal compounds which can bond hydrogen sulphide, such as, for example, lead-II acetate, copper-II acetate, mercury-II acetate or iron-II acetate, in the presence of aqueous alkali metal or alkaline earth metal hydroxide solutions, such as, for example, sodium hydroxide solution, at temperatures between 20°C . and 100°C ., and, when the reaction has ended, the mixture is filtered and the filtrate is acidified with an acid, such as, for example, acetic acid. The products of the formula (V) thereby obtained as crystals can be isolated by filtration with suction.

The starting substances for the preparation processes described above under (a), (b) and (c) for the cyanamide derivatives of the formula (V) are known and/or can be prepared by processes which are known per se.

These include the chloro-hetarenes of the formula (VII) (compare J. Chem. Soc. (c) 1966, 2031; Chem. Pharm. Bull. 11 (1963), 1382-1388 and Arch. Pharm. 295 (1962), 649-657), the halogen compounds of the formula (VIII) (commercially available chemicals), the amino-hetarenes of the formula (IX) (compare Chem. Pharm. Bull. 11, (1963) pages 1382-1388; J. Chem. Soc. 1946, 81 and U.S. Pat. No. 4,299,960) and the carbonyl isothiocyanates of the formula (X) (compare J. Heterocycl. Chem. 5, (1968), 837 and U.S. Pat. No. 4,160,037).

The benzene-1,2-disulphonic acid dichloride of the formula (III) to be used as the starting substance in process (b) according to the invention is already known (compare J. Org. Chem. 31, (1966), 3289-3292).

Formula (IV) provides a general definition of the oxyguanidine derivatives to be used as starting substances in process (b) according to the invention. In formula (IV), R^1 , R^2 and R^3 preferably and particularly have the same meanings as are given above as preferred or as particularly preferred in the context of the definition of the substituents of formula (I).

Examples of compounds of the formula (IV) have already been mentioned above in connection with the description of the starting substances for process (a). The preparation of the starting substances of the formula (IV) has already been described above in connection with the description of the starting substances for process (a).

Process (a) according to the invention for the preparation of the new compounds of the formula (I) is preferably carried out in water as the solvent. Other possible diluents are all the inert organic solvents, but preferably aprotic polar solvents. These include, where appropriate, ketones, such as, for example, acetone and methyl ethyl ketone, nitriles, such as, for example, acetonitrile and propionitrile, dimethylsulphoxide, sulfolane, 1,2-dimethoxyethane and dioxane.

Process (a) is carried out in the presence of bases. Preferred possible bases are alkali metal or alkaline earth metal hydroxides, such as, for example, sodium hydroxide, potassium hydroxide and calcium hydroxide, tertiary amines, such as, for example, triethylamine, N,N-dimethylaniline and N,N-dimethyl-benzylamine, and nitrogen-containing heterocyclic compounds, such as, for example, pyridine or diazabicyclooctane (DABCO).

The reaction temperatures can be varied within a substantial range in process (a) according to the invention. In general, the reaction is carried out between 0°C . and $+100^\circ \text{C}$., preferably between 10°C . and $+80^\circ \text{C}$. Process (a) according to the invention is in general carried out under normal pressure.

For carrying out process (a), in general between 1 and 100 moles, preferably between 5 and 50 moles, of water and, if appropriate, between 1 and 3 moles, preferably between 1 to 2 moles, of a base are employed per mole of benzodisultam of the formula (II).

The reaction components are usually brought together at room temperature and the reaction mixture is stirred until the reaction has ended.

Working up can be carried out in the customary manner; for example, by a procedure in which the mixture is acidified—for example with hydrochloric acid—and concentrated to approximately half the volume, and the product of the formula (I) obtained as crystals is isolated by filtration with suction.

Process (b) according to the invention for the preparation of the new compounds of the formula (I) is preferably carried out using diluents. Possible diluents are virtually all the inert organic solvents, preferably aprotic polar solvents. These include optionally substituted hydrocarbons, such as, for example, methylene chloride, chloroform, 1,2-dichloroethane, toluene, xylene and chlorobenzene, nitriles, such as, for example, acetonitrile and propionitrile, ethers, such as, for example, 1,2-dimethoxyethane, tetrahydrofuran and dioxane, and dimethylformamide, dimethylacetamide, dimethyl-

sulphoxide, sulpholane, pyridine and 2-methyl-5-ethylpyridine.

Virtually all the acid-binding agents which are usually employed can be used as acid acceptors in process (b). These include, in particular, alkali metal and alkaline earth metal hydroxides, alkali metal and alkaline earth metal hydrides, organometallic compounds, such as butyl-lithium, and furthermore aliphatic, aromatic or heterocyclic amines, such as trimethylamine, triethylamine, N,N-dimethylaniline, N,N-dimethyl-benzylamine, diazabicyclooctane (DABCO), diazabicyclononene (DBN), diazabicycloundecene (DBU), pyridine, 2-methyl-5-ethylpyridine and 4-dimethyl-aminopyridine.

The reaction temperatures can be varied within a substantial range in process (b). In general, the reaction is carried out between -80°C . and $+100^{\circ}\text{C}$., preferably between -30°C . and $+50^{\circ}\text{C}$. Process (b) according to the invention is in general carried out under normal pressure.

For carrying out process (b), in general between 1 and 2 moles, preferably between 1.0 and 1.2 moles, of benzene-1,2-disulphonic acid dichloride of the formula (III) and then between 1 to 100 moles, preferably between 5 and 50 moles, of water, and, if appropriate, between 1 and 3 moles, preferably between 1 and 2 moles, of a base are employed per mole of oxyguanidine derivative of the formula (IV).

The reaction components are usually brought together at room temperature or with external cooling and the reaction mixture is stirred until the reaction has ended.

Working up can be carried out in the customary manner; for example by a procedure in which the mixture is concentrated, if appropriate, and/or diluted with an organic solvent which is virtually water-immiscible, such as, for example, methylene chloride, washed with dilute hydrochloric acid and with water, dried, filtered and concentrated. The product of the formula (I) which remains in the residue is isolated by trituration with a suitable organic solvent, such as, for example, ethanol, made to crystallise and isolated by filtration with suction.

The active compounds according to the invention can be used as defoliant, desiccant, agent for destroying broad-leaved plants and, especially, as weedkillers. By weeds, in the broadest sense, there are to be understood all plants which grow in locations where they are undesired. Whether the substances according to the [invention] *invention* act as total or selective herbicides depends essentially on the amount used.

The active compounds according to the invention can be used, for example, in connection with the following plants:

Dicotyledon weeds of the genera: [Sinapis] *Sinapis*, Lepidium, Galium, Stellaria, Matricaria, [Anthemis] *Anthemis*, Galinsoga, Chenopodium, Urtica, Senecio, Amaranthus, Portulaca, Xanthium, Convolvulus, Ipomoea, Polygonum, Sesbania, Ambrosia, Cirsium, Carduus, Sonchus, Solanum, Rorippa, Rotala, Lindernia, Lamium, [Beronica] *Veronica*, Abutilon, Emex, Datura, Viola, Galeopsis, Papaver and Centaurea.

Dicotyledon cultures of the genera: Gossypium, Glycine, Beta, Daucus, Phaseolus, Pisum, Solanum, Linum, Ipomoea, Vicia, Nicotiana, Lycopersicon, Arachis, Brassica, Lactuca, Cucumis and Cucurbita.

Monocotyledon weeds of the genera: Echinochloa, Setaria, Panicum, Digitaria, Phleum, Poa, Festuca,

Eleusine, Brachiaria, Lolium, Bromus, Avena, Cyperus, Sorghum, Agropyron, Cynodon, Monochoria, Fimbristylis, Sagittaria, Eleocharis, Scirpus, Paspalum, Ischaemum, Sphenoclea, Dactyloctenium, Agrostis, Alopecurus and Apera.

Monocotyledon cultures of the genera: Oryza, Zea, Triticum, Hordeum, Avena, Secale, Sorghum, Panicum, Saccharum, Ananas, Asparagus and Allium.

However, the use of the active compounds according to the invention is in no way restricted to these genera, but also extends in the same manner to other plants.

The compounds are suitable, depending on the concentration, for the total combating of weeds, for example on industrial terrain and rail tracks, and on paths and squares with or without tree plantings. Equally, the compounds can be employed for combating weeds in perennial cultures, for example afforestations, decorative tree plantings, orchards, vineyards, citrus groves, nut orchards, banana plantations, coffee plantations, tea plantations, rubber plantations, oil palm plantations, cocoa plantations, soft fruit plantings and hopfields, and for the selective combating of weeds in annual cultures.

The active compounds can be converted to the customary formulations, such as solutions, emulsions, wettable powders, suspensions, powders, dusting agents, pastes, soluble powders, granules, suspension-emulsion concentrates, natural and synthetic materials impregnated with active compound, and very fine capsules in polymeric substances.

These formulations are produced in known manner, for example by mixing the active compounds with extenders, that is liquid solvents and/or solid [carriers] *carriers*, optionally with the use of surface-active agents, that is emulsifying agents and/or dispersing agents and/or foam-forming agents.

In the case of the use of water as an extender, organic solvents can, for example, also be used as auxiliary solvents. As liquid solvents, there are suitable in the main: aromatics, such as xylene, toluene or alkyl naphthalenes, chlorinated aromatics and chlorinated aliphatic hydrocarbons, such as chlorobenzenes, chloroethylenes or methylene chloride, aliphatic hydrocarbons, such as cyclohexane or paraffins, for example petroleum fractions, mineral and vegetable oils, alcohols, such as butanol or glycol as well as their ethers and esters, ketones, such as acetone, methyl ethyl ketone, methyl isobutyl ketone or cyclohexanone, strongly polar solvents, such as dimethylformamide and dimethylsulphoxide, as well as water.

As solid carrier there are suitable: for example ammonium salts and ground natural minerals, such as kaolins, clays, talc, chalk, quartz, attapulgite, montmorillonite or diatomaceous earth, and ground synthetic minerals, such as highly disperse silicic acid, alumina and silicates, as solid carriers for granules there are suitable: for example crushed and fractionated natural rocks such as calcite, marble, pumice, sepiolite and dolomite, as well as synthetic granules of inorganic and organic meals, and granules of organic material such as sawdust, coconut shells, maize cobs and tobacco stalks; as emulsifying and/or foam-forming agents there are suitable: for example non-ionic and anionic emulsifiers, such as polyoxyethylene-fatty esters, polyoxyethylene-fatty alcohol ethers, for example alkylaryl polyglycol ethers, alkylsulphonates, alkylsulphates, arylsulphonates as well as albumin hydrolysis products; as dispersing agents there are suitable: for example lignin-sulphite waste liquors and methylcellulose.

Adhesive such as carboxymethylcellulose and natural and synthetic polymers in the form of powders, granules or latices, such as gum arabic, polyvinyl alcohol and polyvinyl actate, as well as natural phospholipids, such as cephalins and lecithins, and synthetic phospholipids, can be used in the formulations. Further additives can be mineral and vegetable oils.

It is possible to use colorants such as inorganic pigments, for example iron oxide, titanium oxide and Prussian Blue, and organic dyestuffs, such as alizarin dyestuffs, azo dyestuffs and metal phthalocyanine dyestuffs, and trace nutrients such as salts of iron, manganese, boron, copper, cobalt, molybdenum and zinc.

The formulations in general contain between 0.1 and 95 percent by weight of active compound, preferably between 0.5 and 90%.

For the mixtures come known herbicides, such as, for example, N-(2-benzothiazolyl)-N,N'-dimethyl-urea, 3-(3-chloro-4-methylphenyl)-1,1-dimethylurea, 3-(4-isopropylphenyl)-1,1-dimethylurea, 4-amino-6-(1,1-dimethyl ethyl)-3-methylthio-1,2,4-triazin-5(4H)-one, 4-amino-6-(1,1-dimethyl ethyl)-3-ethylthio-1,2,4-triazin-5(4H)-one, 1-amino-6-ethylthio-3-(2,2-dimethylpropyl)-1,3,5-triazine-2,4(1H,3H)-dione, 4-amino-3-methyl-6-phenyl-1,2,4-triazin-5(4H)-one, 2-chloro-4-ethylamino-6-isopropyl-amino-1,3,5-triazine, the R-enantiomer of (trimethylsilyl)-methyl 2-[4-(3,5-dichloropyridin-2-oxy)-phenoxy]-propionate, the R-enantiomer of (2-benzyloxy)-ethyl 2-[4-(3,5-dichloropyridyl-2-oxy)phenoxy]-propionate, 2,4-dichlorophenoxyacetic acid, 2-(2,4-dichlorophenoxy)-propionic acid, 4-chloro-2-methylphenoxy-acetic acid, 2-(2-methyl-4-chloro-phenoxy)-propionic acid, 3,5-diiodo-4-hydroxy-benzonitrile, 3,5-dibromo-4-hydroxy-benzonitrile and diphenyl ethers and phenylpyridazines, such as, for example, pyridates. Surprisingly some mixtures also show a synergistic action.

Mixtures with other known active compounds, such as fungicides, insecticides, acaricides, nematocides, bird repellants, plant nutrients and agents which improve soil structure, are also possible.

The active compounds can be used as such, in the form of their formulations or in the use forms prepared therefrom by further dilution, such as ready-to-use solutions, suspensions, emulsions, powders, pastes and granules. They are used in the customary manner, for example by watering, spraying, atomising or scattering.

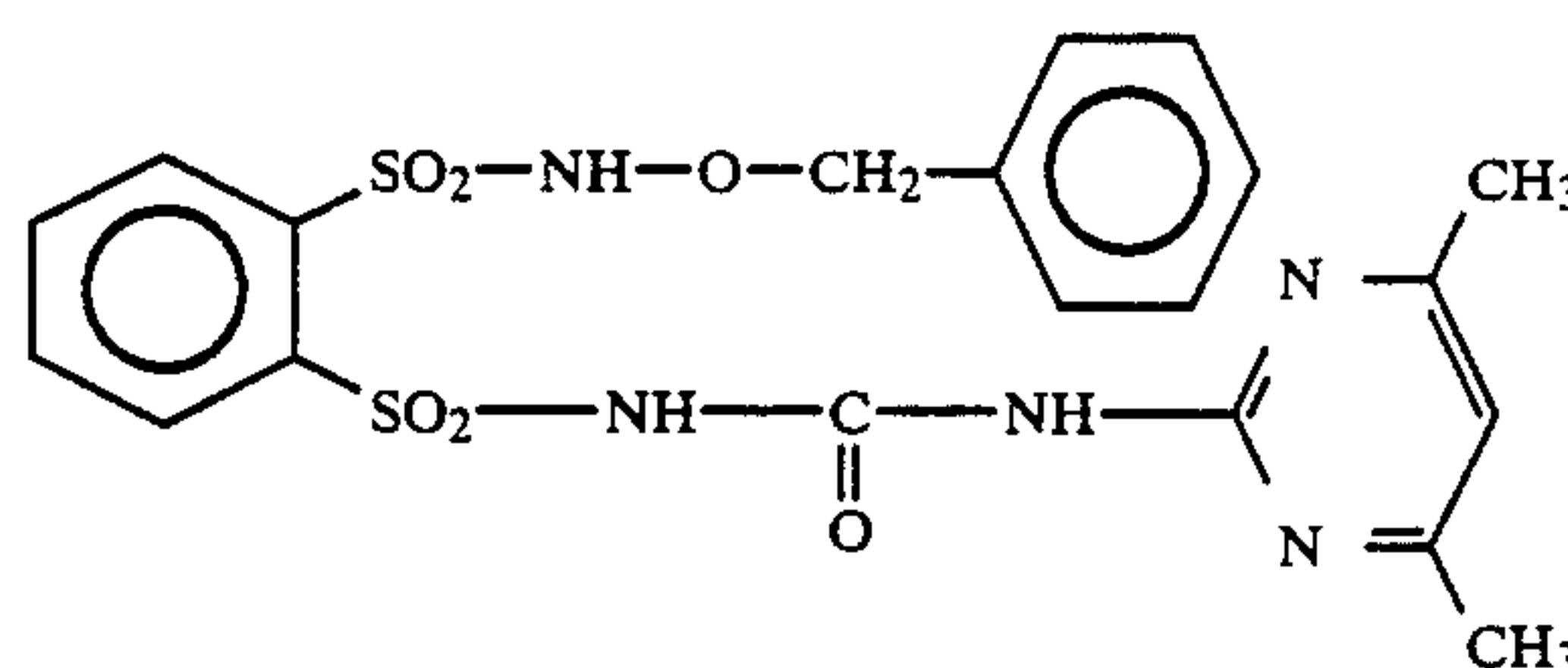
The active compounds according to the invention can be applied either before or after emergence of the plants. They can also be incorporated into the soil before sowing.

The amount of active compound used can vary within a substantial range. It depends essentially on the nature of the desired effect. In general, the amounts used are between 0.001 and 15 kg of active compound per hectare of soil surface, preferably between 0.005 and 10 kg per ha.

The preparation and use of the active compounds according to the invention can be seen from the following examples.

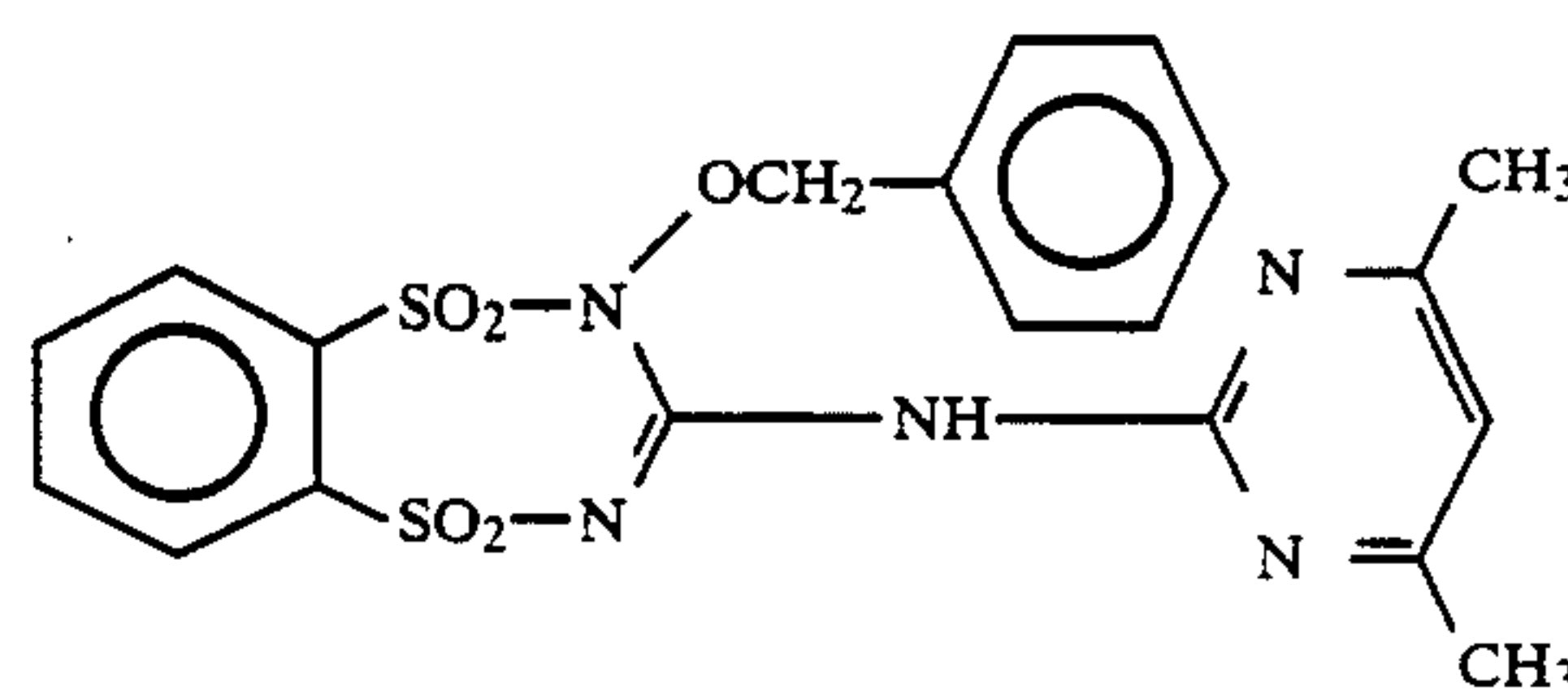
PREPARATION EXAMPLES

EXAMPLE 1



(Process (a))

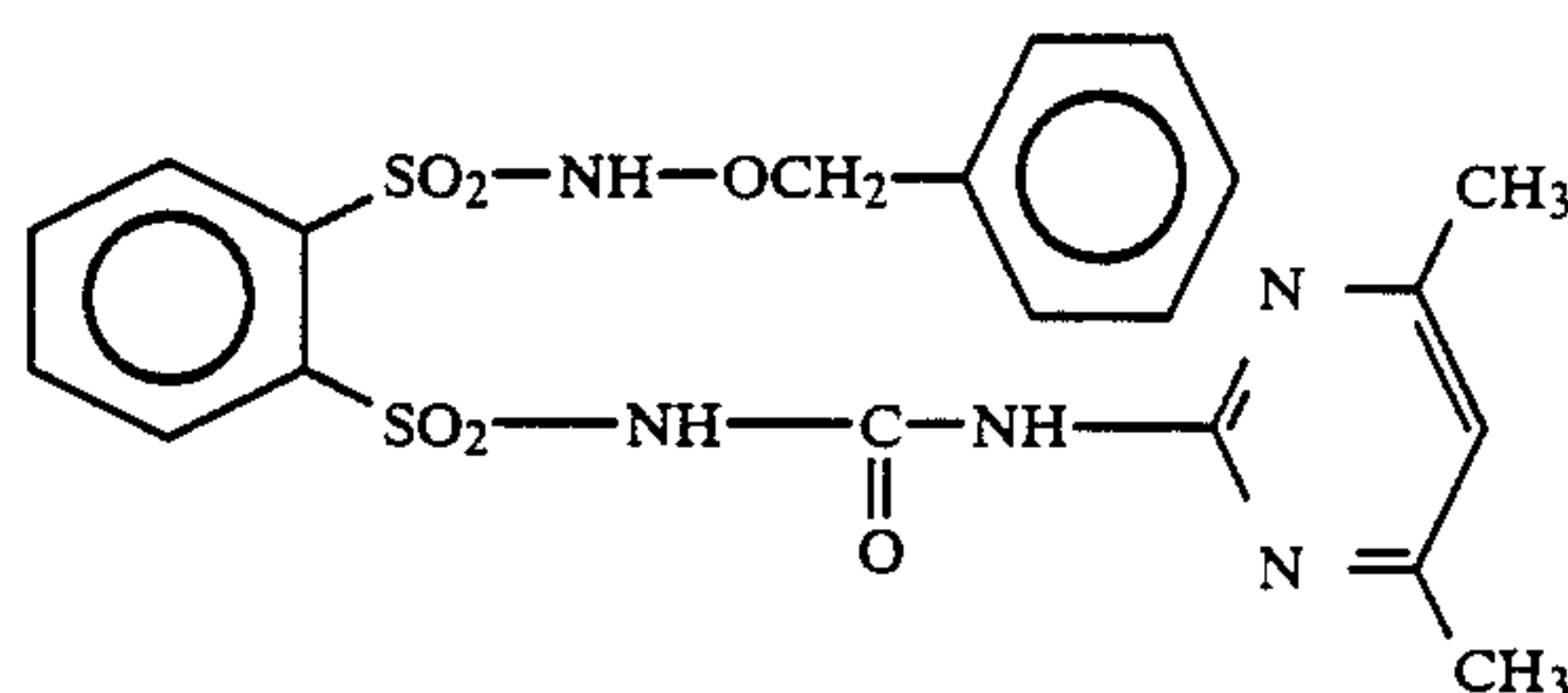
A mixture of 21.9 g (0.05 mole) of the compound of the following structural formula



with 4 g (0.1 mole) of sodium hydroxide, 50 ml of water and 50 ml of dioxane is heated at 40° C. for 2 hours.

The mixture is then acidified with concentrated hydrochloric acid and the solution is concentrated to about half. The product obtained as crystals is filtered off with suction and dried. 12.5 g (54% of theory) of 1-(2-benzyloxyaminosulphonylphenylsulphonyl)-3-(4,6-dimethylpyrimidin-2-yl)-urea of melting point 186° C. (decomposition) are obtained.

EXAMPLE 2



(Process (b))

14 g (0.05 mole) of benzene-1,2-disulphonic acid dichloride are added in portions to a mixture of 13.6 g (0.05 mole) of N'-(4,6-dimethylpyrimidin-2-yl)-N''-benzyloxyguanidine, 12 g (0.15 mole) of pyridine and 200 ml of methylene chloride at -20° C. The mixture is subsequently stirred at -20° C. for 3 hours and at +20° C. for 15 hours.

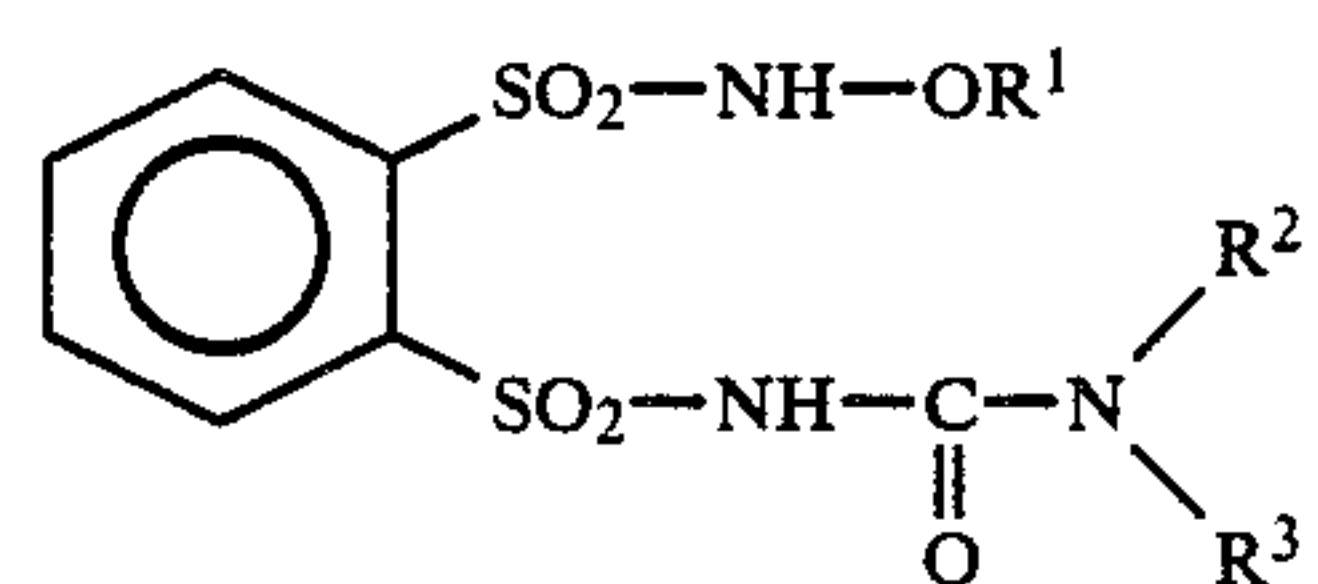
10 ml of water are then added to the reaction mixture and the mixture is stirred at 20° C. for a further 2 hours.

The methylene chloride solution is then washed with dilute hydrochloric acid and water and concentrated. The residue is triturated with ethanol; the product obtained as crystals is isolated by filtration with suction.

6.2 g (27% of theory) of 1-(2-benzyloxyaminosulphonylphenylsulphonyl)-3-(4,6-dimethylpyrimidin-2-

yl)urea of melting point 188° C. (decomposition) are obtained.

The compounds of the formula (I) listed in the following Table 2 can also be prepared by the process described by way of example in the preceding Examples 1 and 2:



(I)

TABLE 2

Example No.	R ¹	R ²	R ³	Melting point (° C.)
3	$-\text{CH}_2-\text{CH}=\text{CH}_2$	H		182 (decomp.)
4	$-\text{C}_8\text{H}_{17}(-n)$	H		145
5		H		
6		H		
7		H		
8		H		
9		H		

TABLE 2-continued

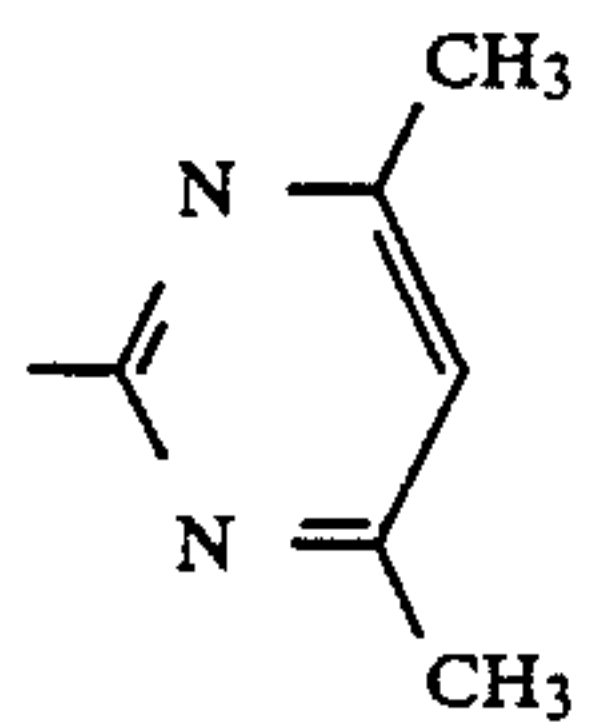
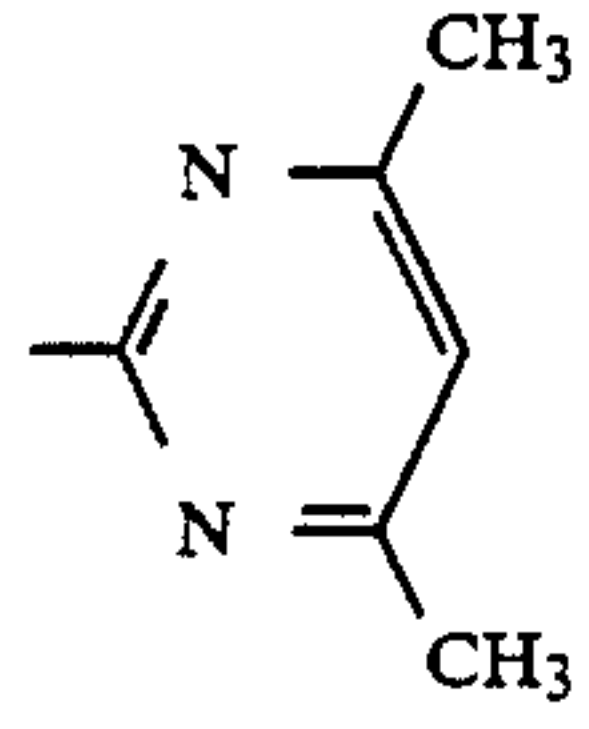
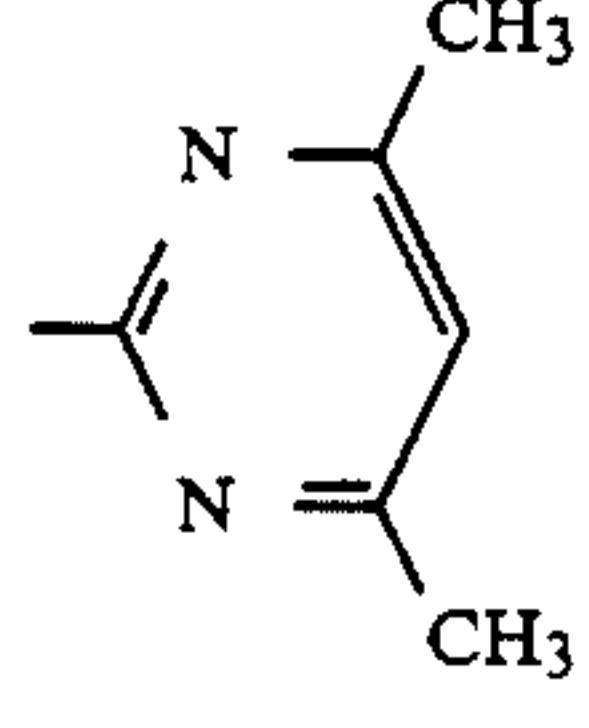
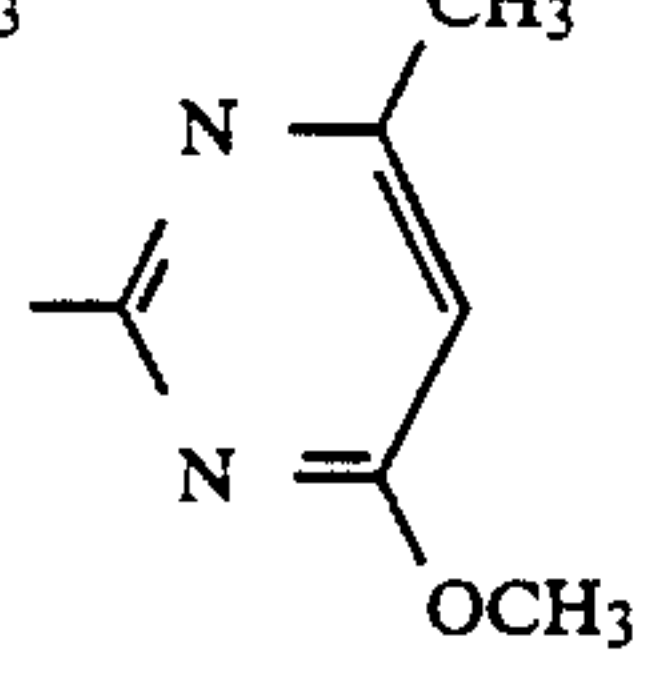
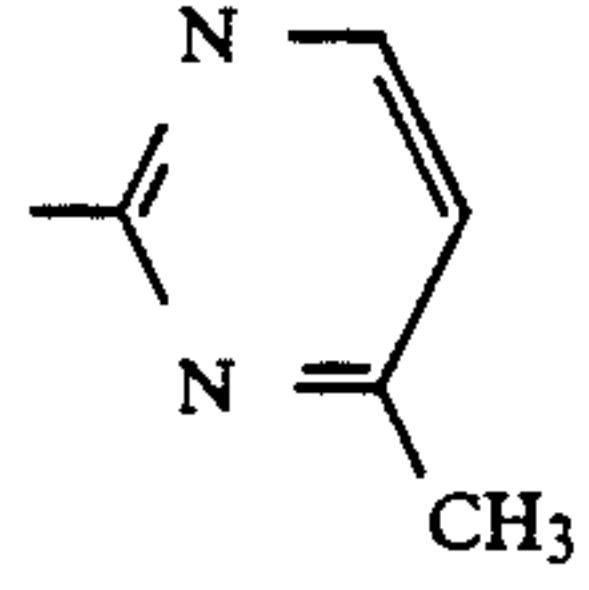
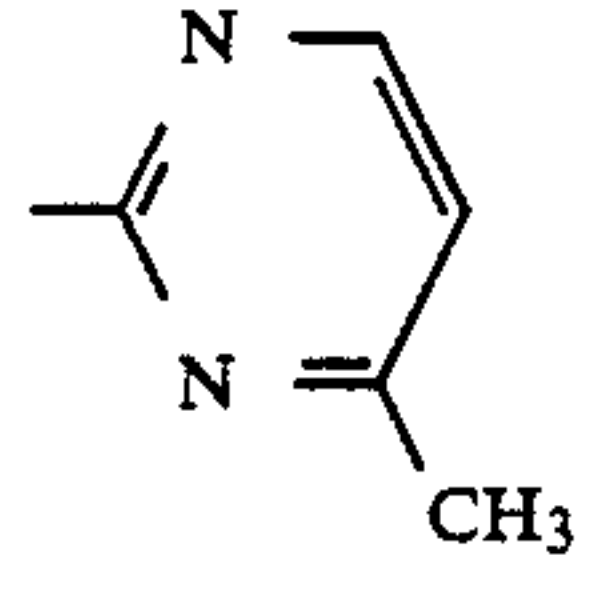
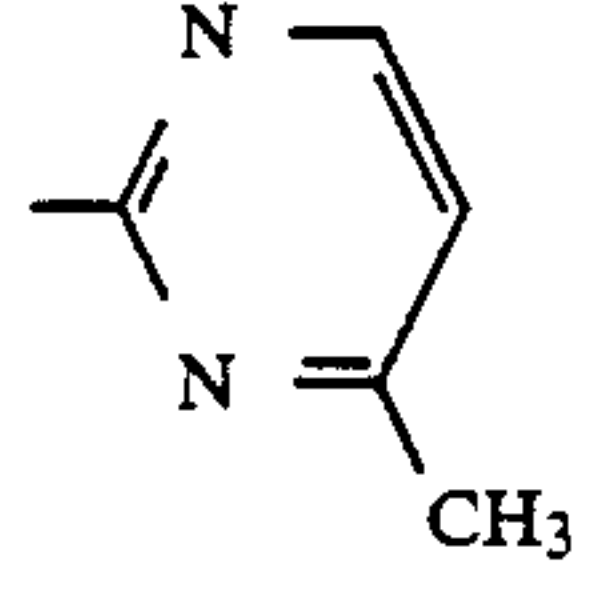
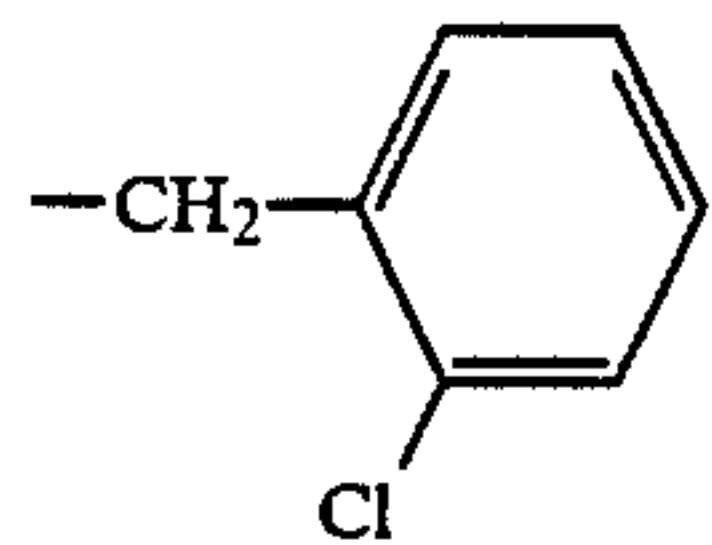
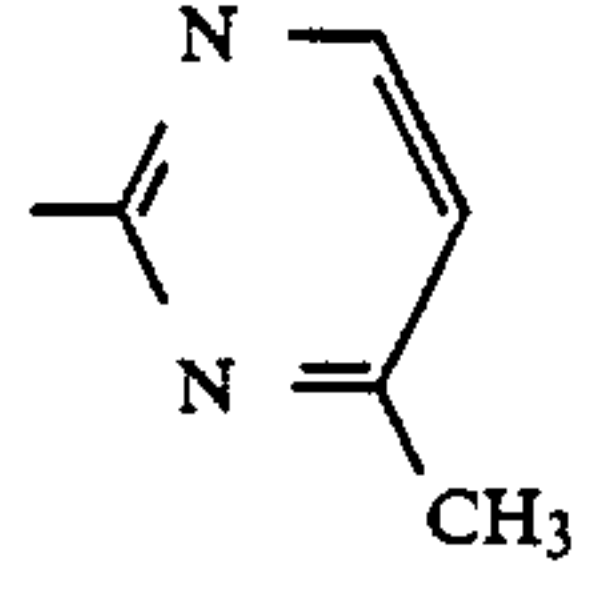
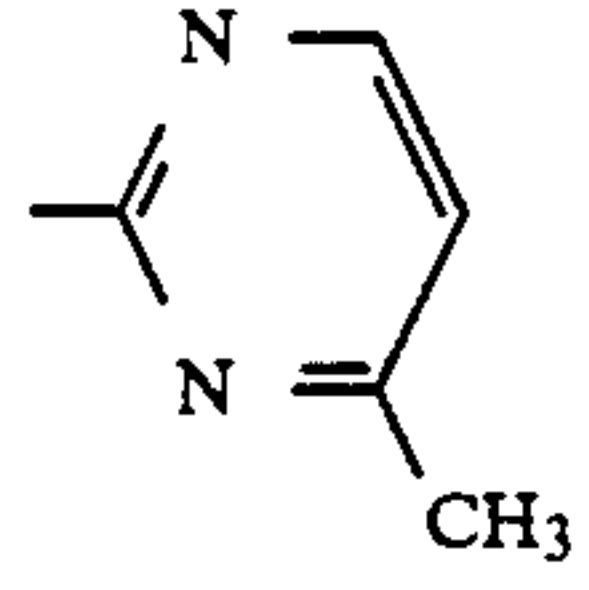
Example No.	R ¹	R ²	R ³	Melting point (° C.)
10	$-\text{CH}_2-\text{COOC}_2\text{H}_5$	H		
11	$-\text{CH}(\text{CH}_3)-\text{COOCH}_3$	H		
12	$-\text{CH}_2\text{CH}(\text{CH}_3)_2$	H		
13	$-\text{CH}_3$	CH ₃		
14	$-\text{CH}_3$	H		194-195
15	$-\text{C}_2\text{H}_5$	H		
16	$-\text{CH}_2-\text{CH}=\text{CH}_2$	H		
17		H		
18	$-\text{CH}_2\text{COOC}_2\text{H}_5$	H		

TABLE 2-continued

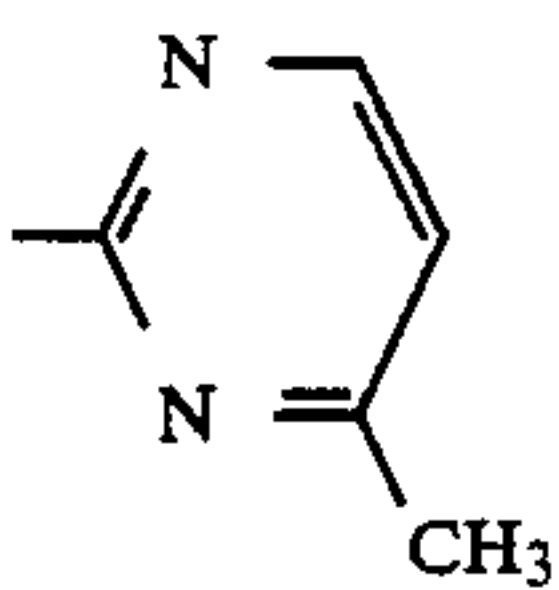
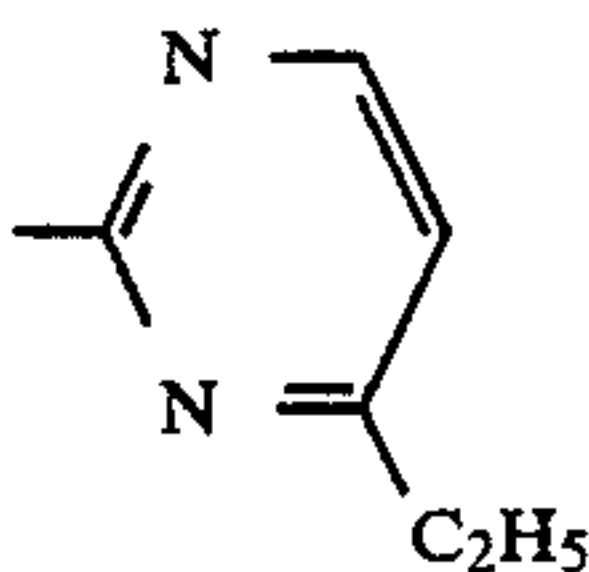
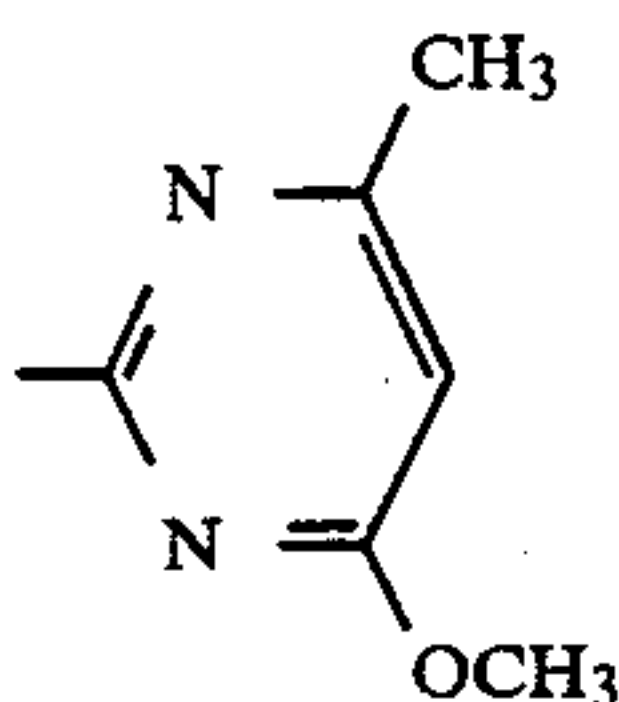
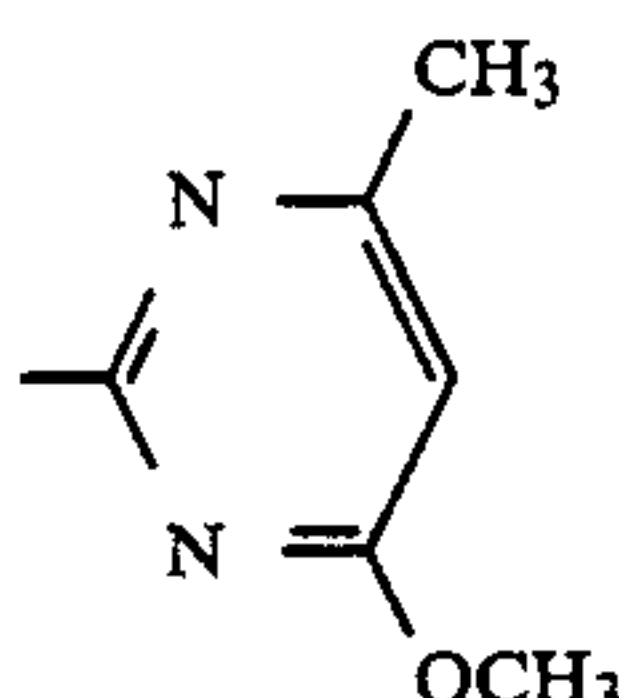
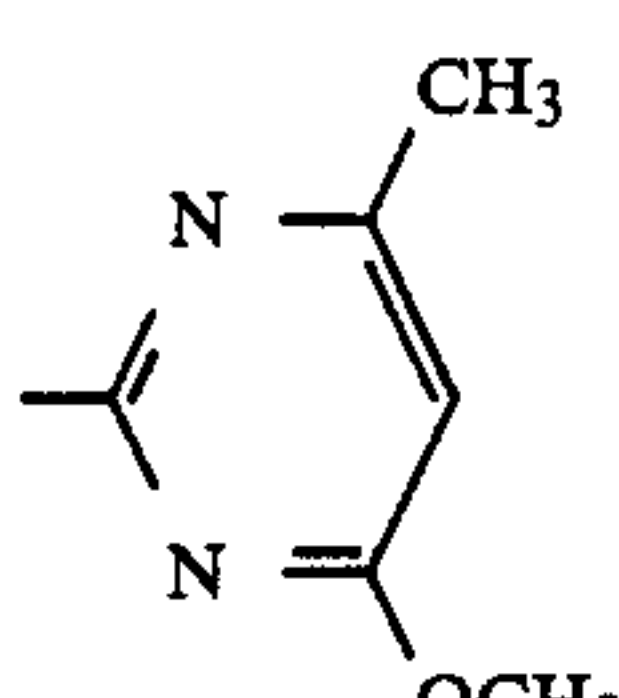
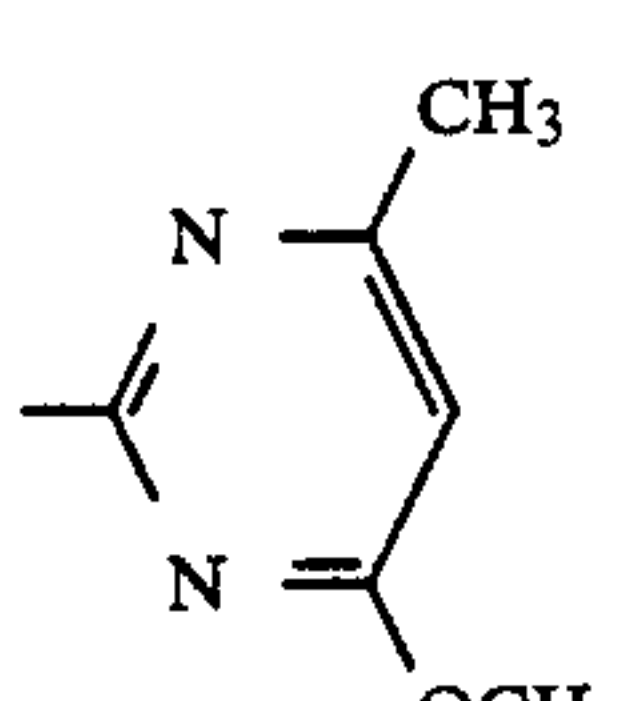
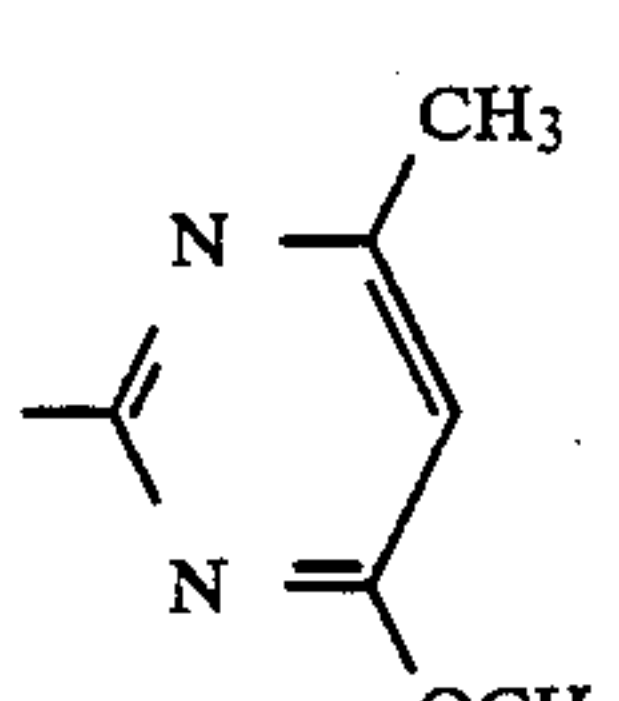
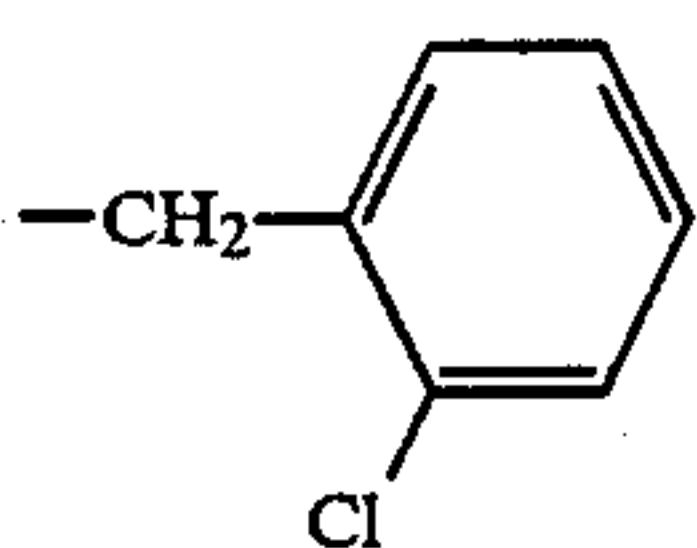
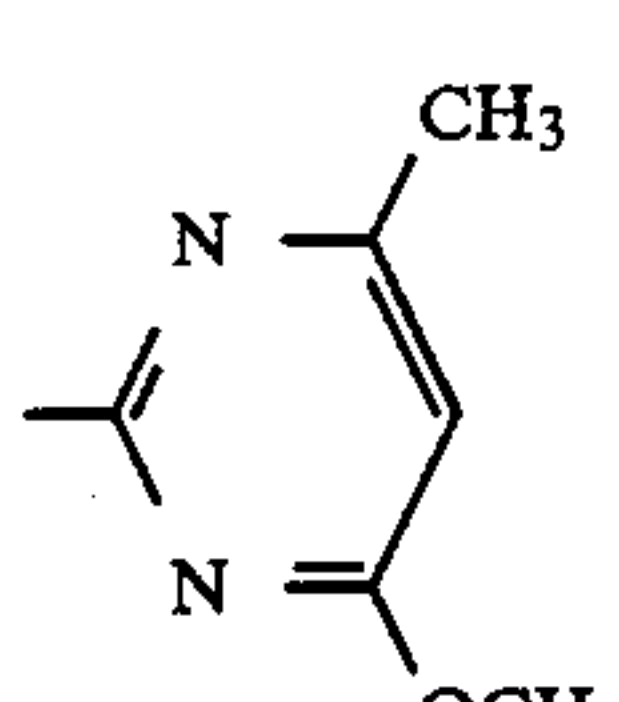
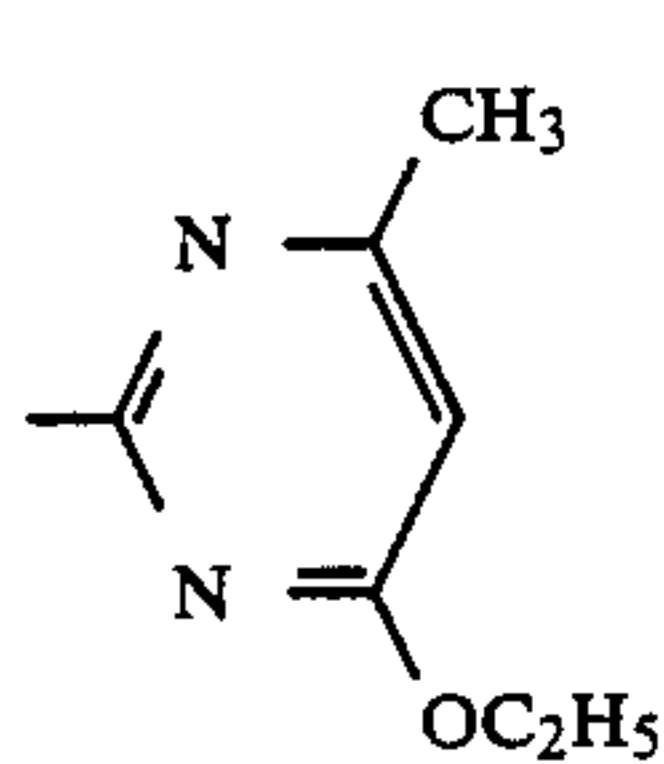
Example No.	R ¹	R ²	R ³	Melting point (° C.)
19	$-\text{CH}_2\text{CH}(\text{CH}_3)_2$	H		
20	$-\text{CH}_3$	H		
21	$-\text{CH}_3$	H		218 (decomp.)
22	$-\text{C}_2\text{H}_5$	H		
23	$-\text{C}_3\text{H}_7$	H		
24	$-\text{CH}_2-\text{CH}=\text{CH}_2$	H		
25	$-\text{CH}_2\text{CH}(\text{CH}_3)_2$	H		
26		H		
27	$-\text{CH}_3$	H		

TABLE 2-continued

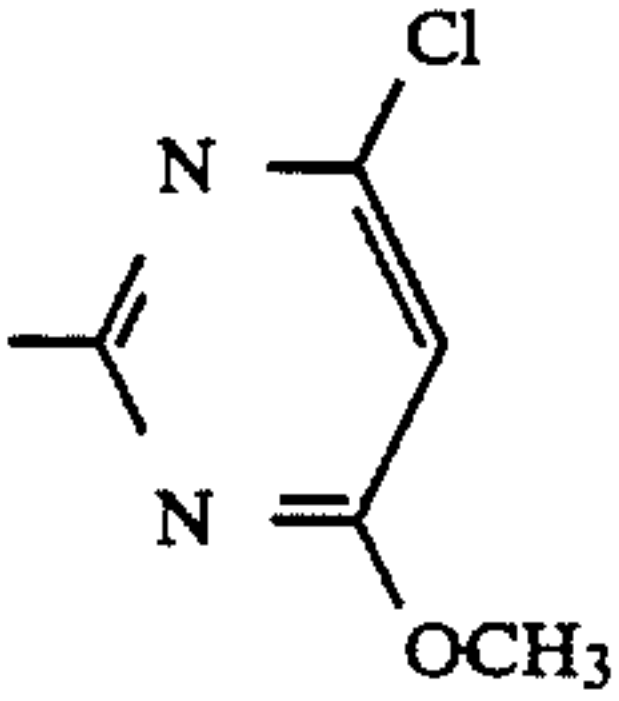
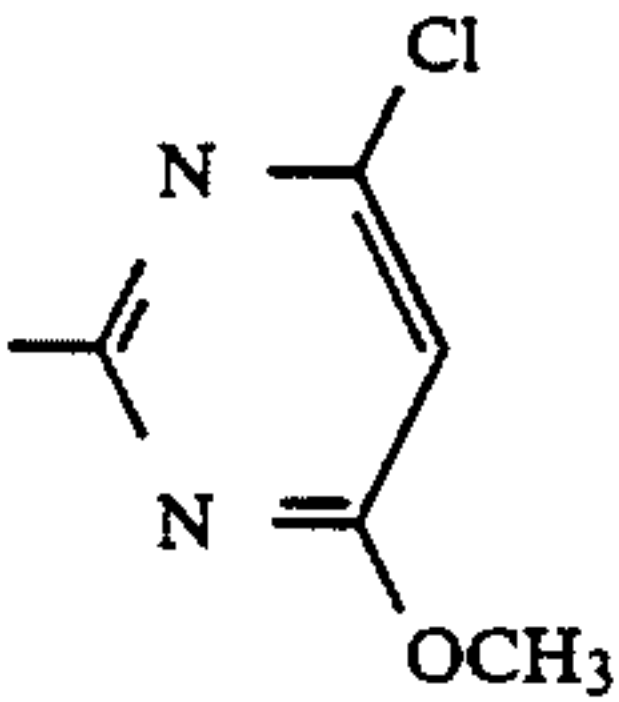
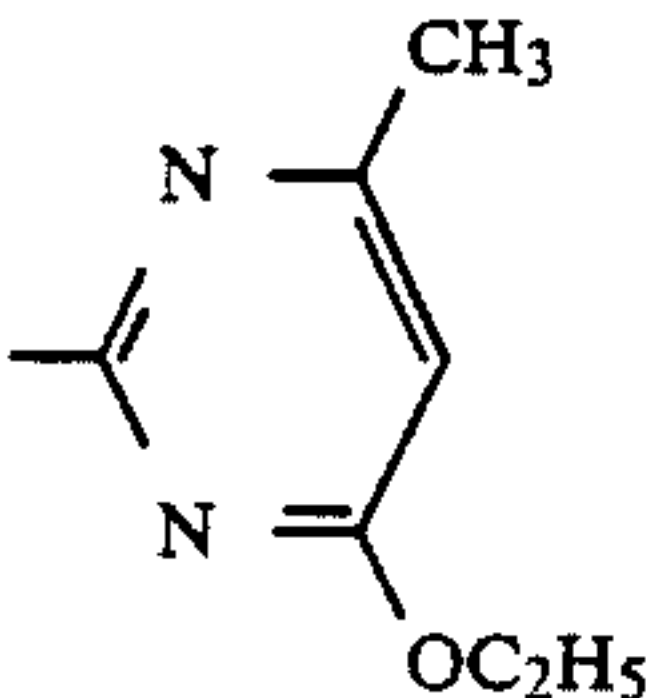
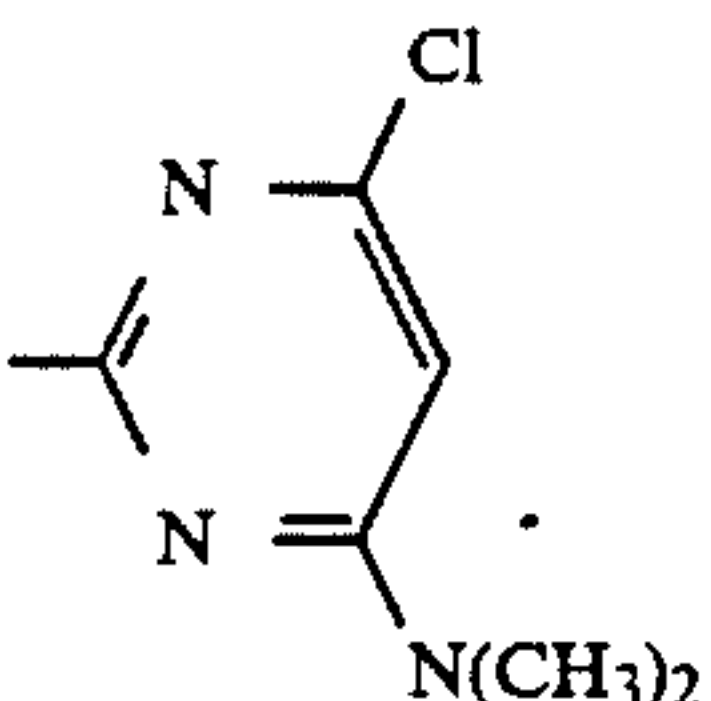
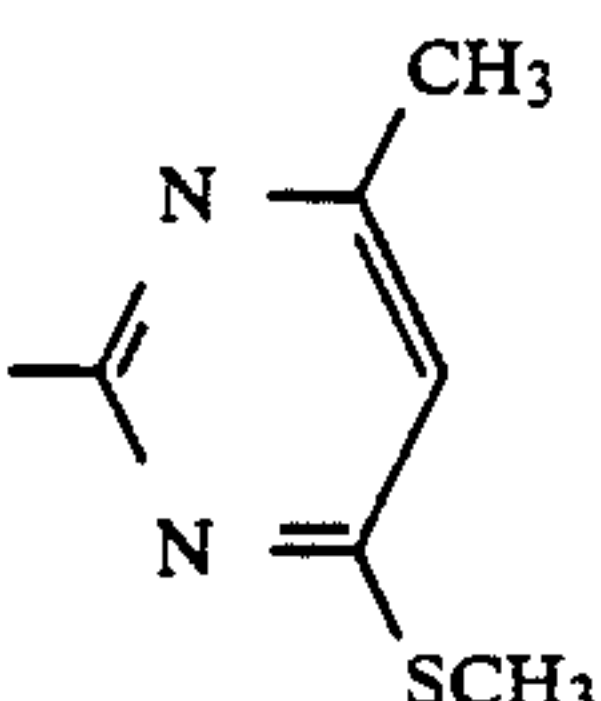
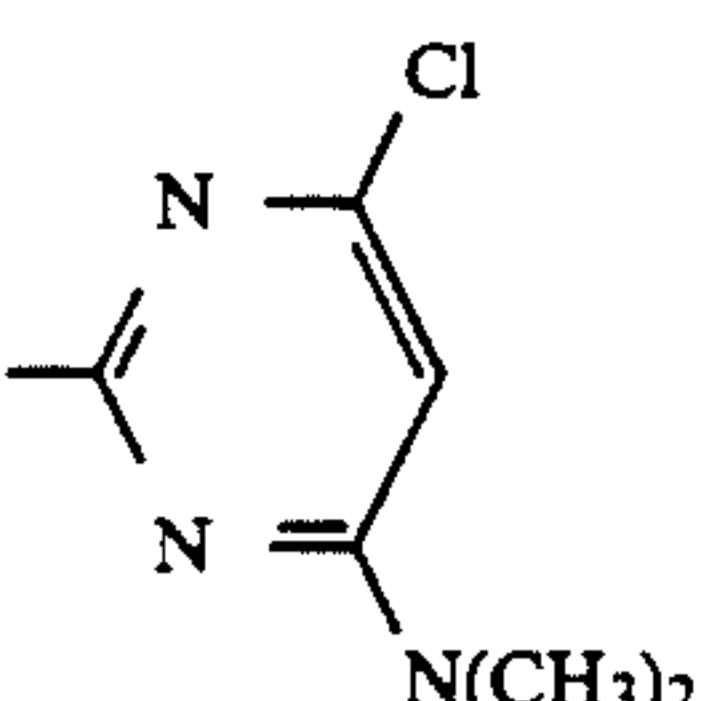
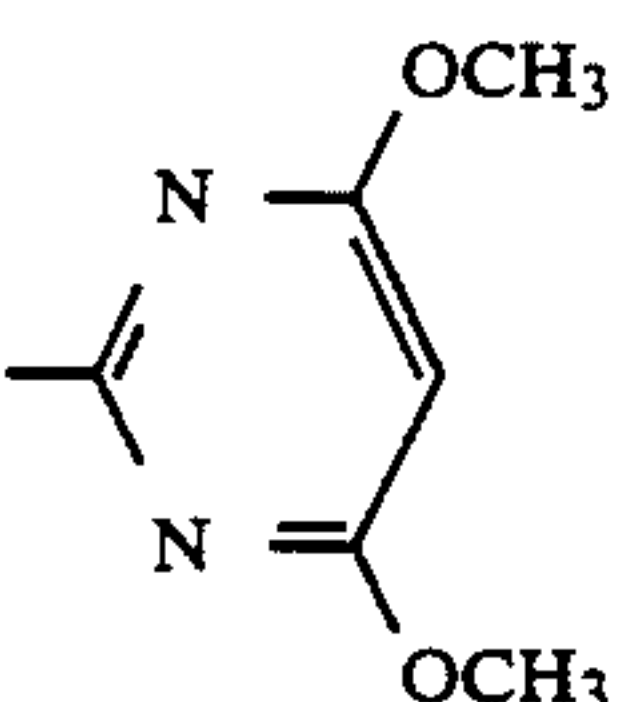
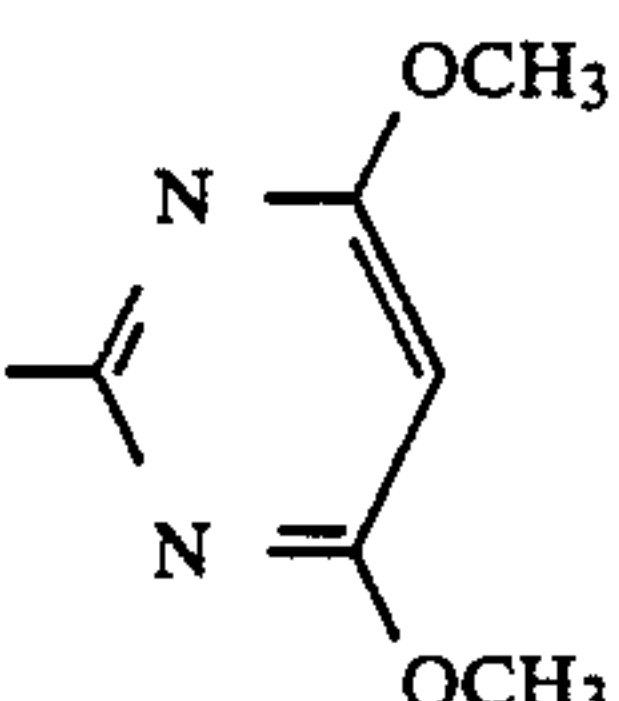
Example No.	R ¹	R ²	R ³	Melting point (° C.)
28	—CH ₃	H		
29	—C ₂ H ₅	H		
30	—CH ₃	H		
31	—CH ₃	H		
32	—CH ₃	H		
33	—CH ₃	H		
34	—CH ₃	H		
35	—C ₂ H ₅	H		

TABLE 2-continued

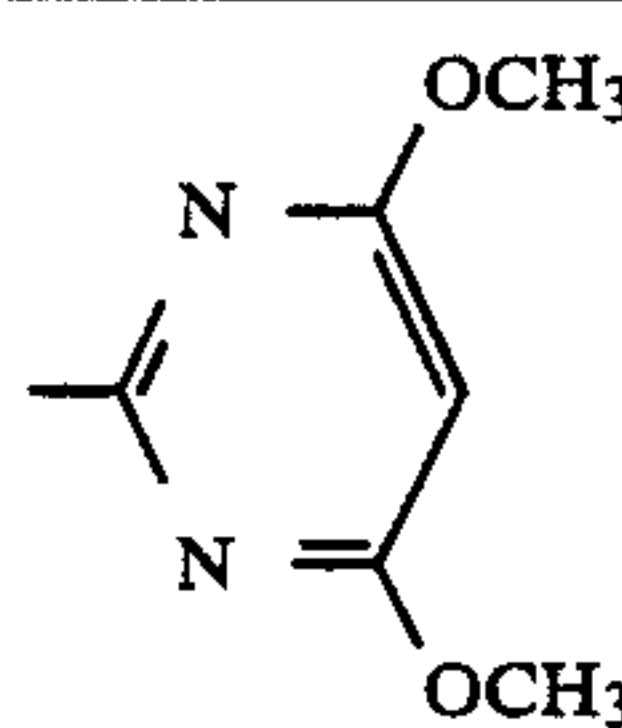
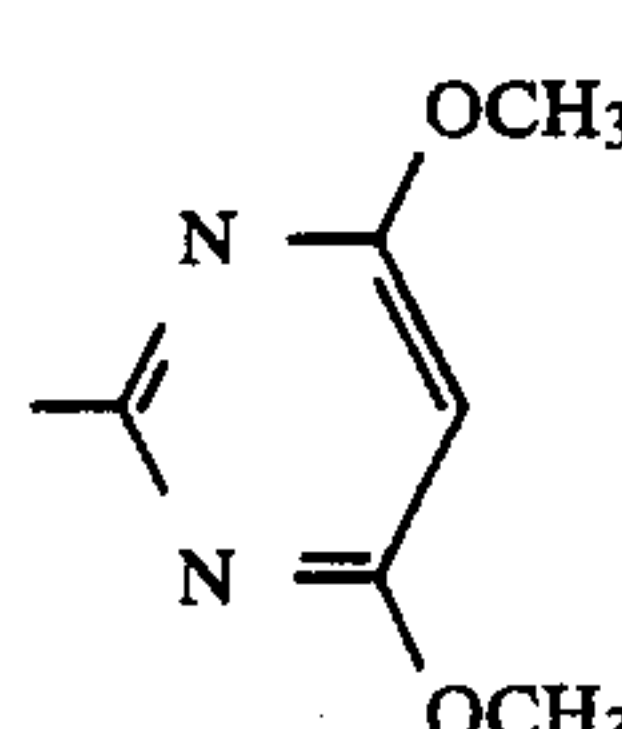
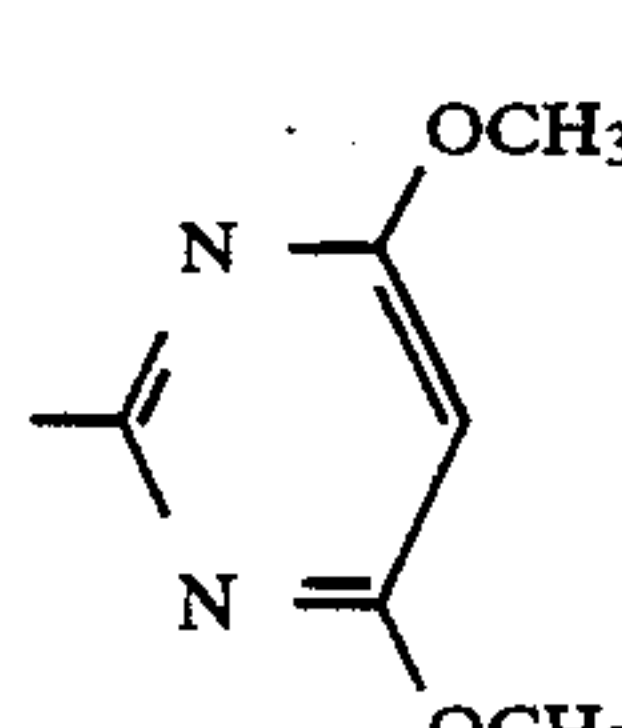
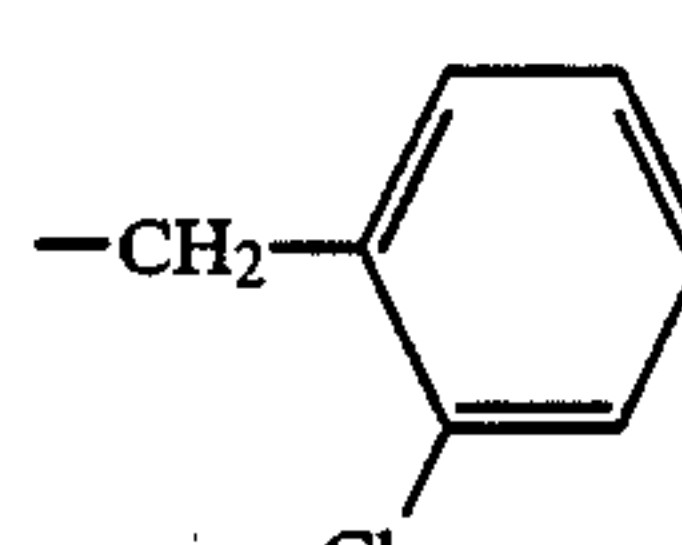
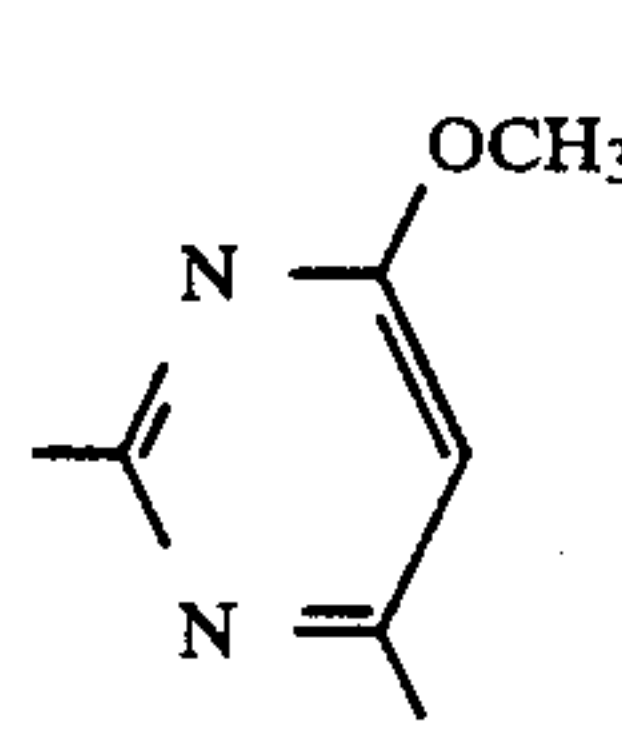
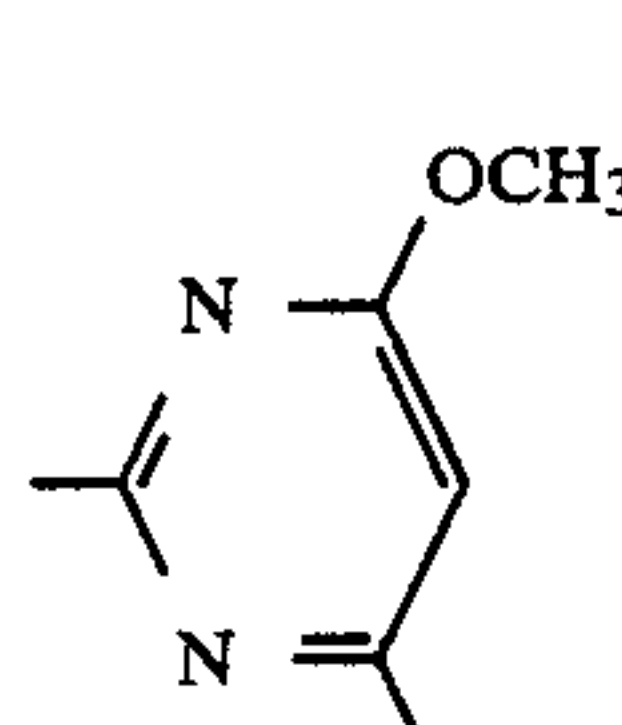
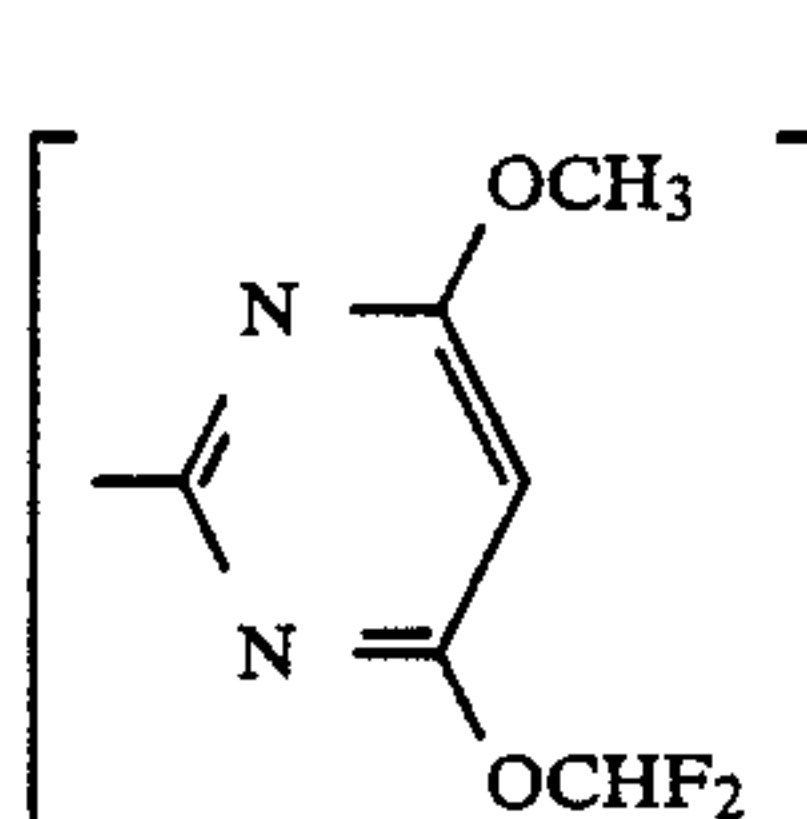
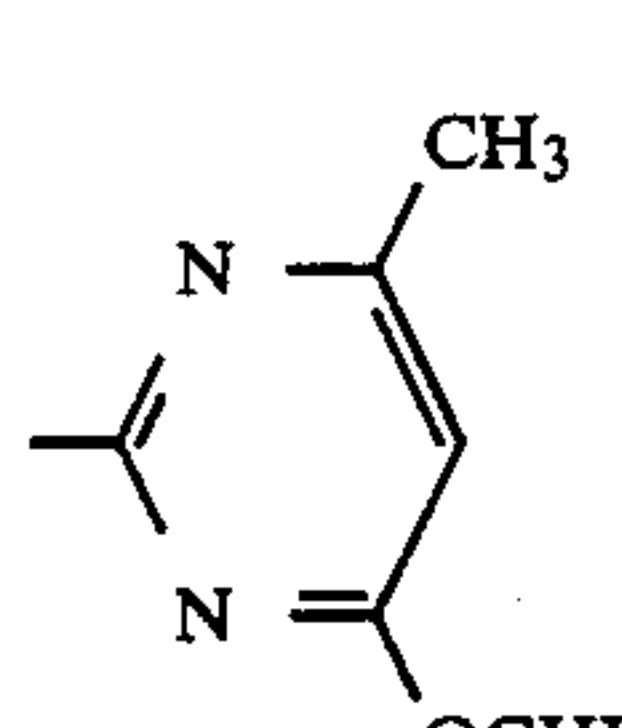
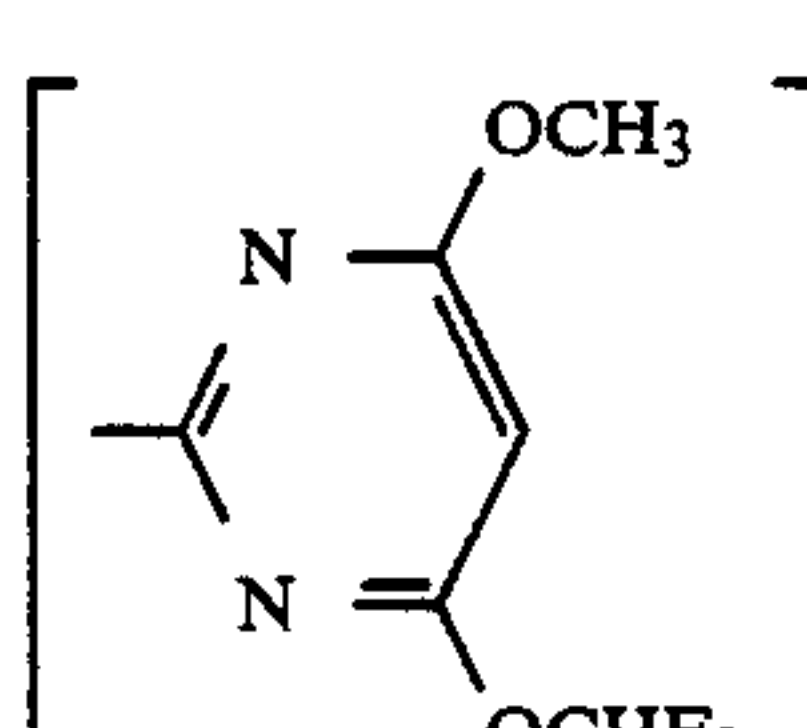
Example No.	R ¹	R ²	R ³	Melting point (° C.)
36	—C ₃ H ₇	H		
37	—CH ₂ —CH=CH ₂	H		
38	—CH ₂ CH(CH ₃) ₂	H		
39		H		
40	—C ₄ H ₉	H		
41	—CH ₃	H	 	
42	—C ₂ H ₅	H		

TABLE 2-continued

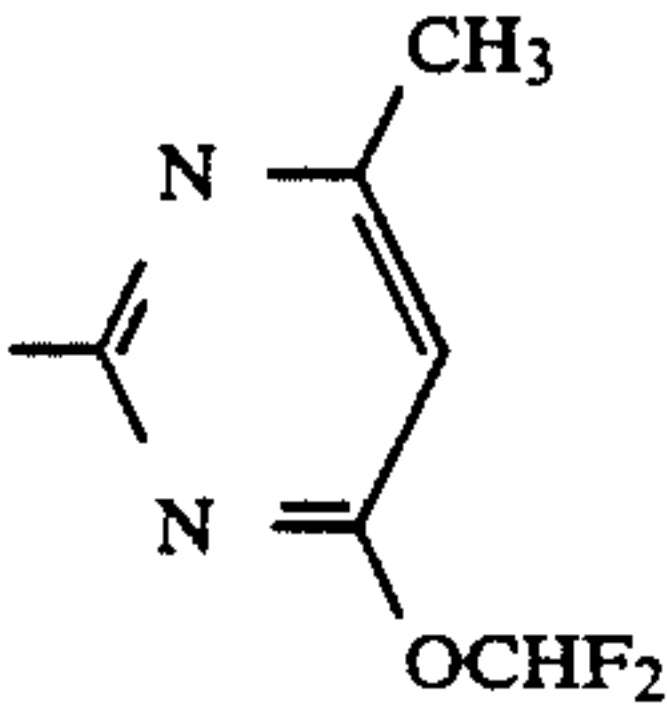
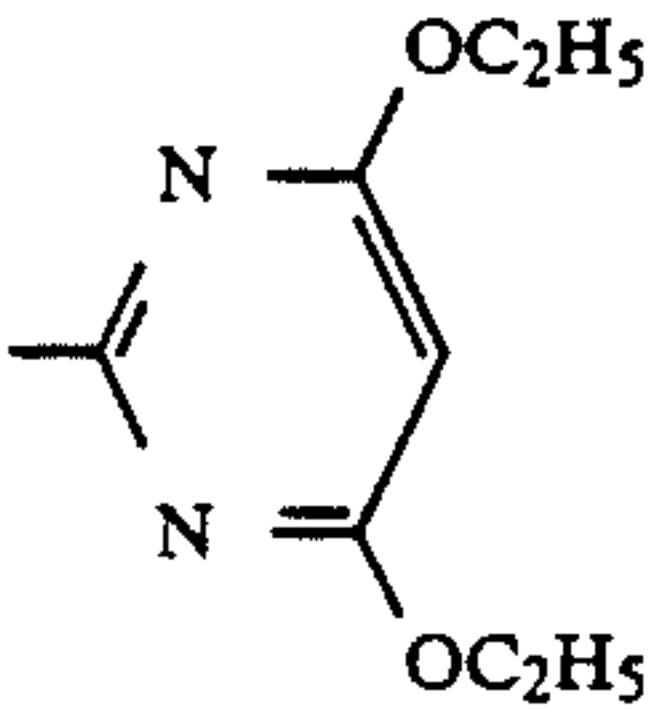
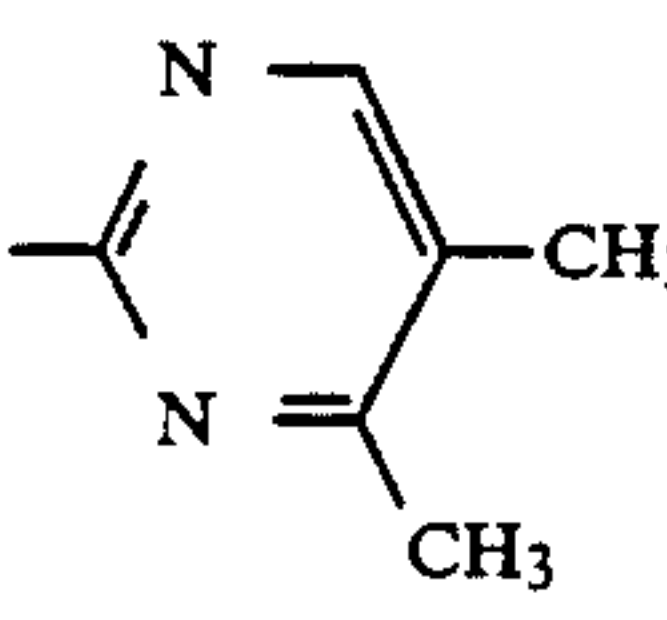
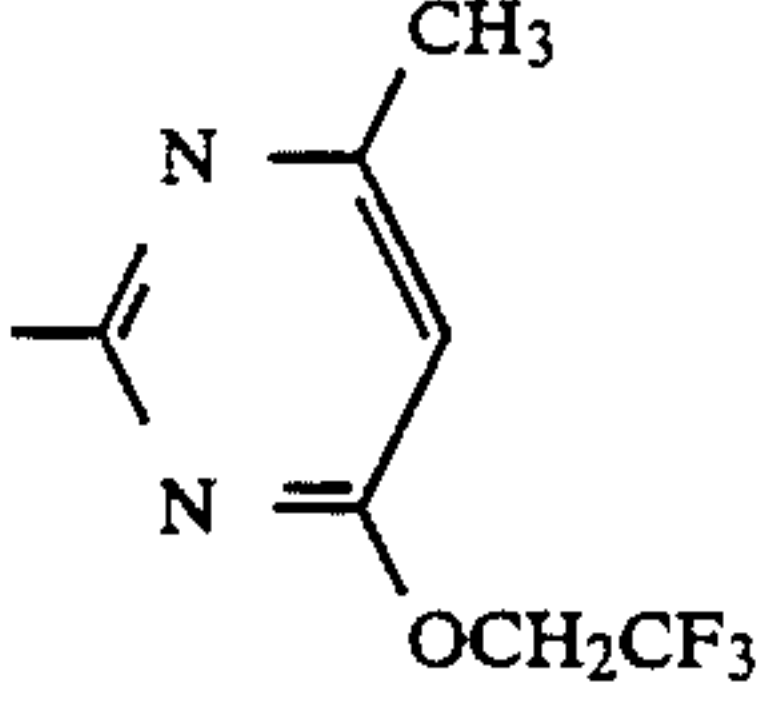
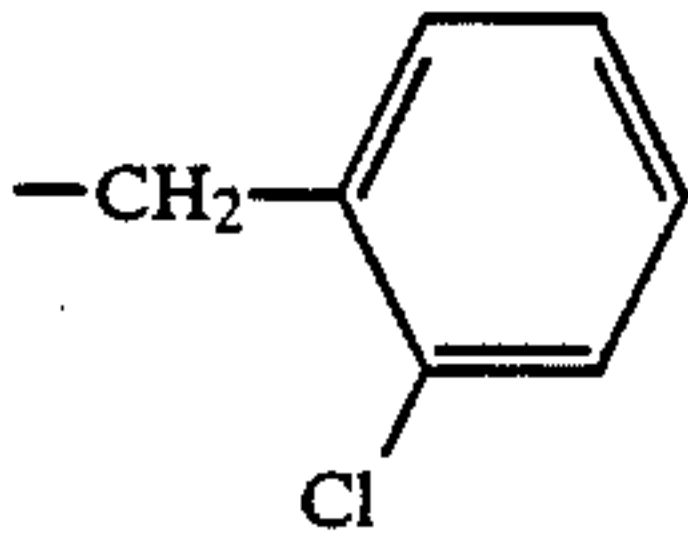
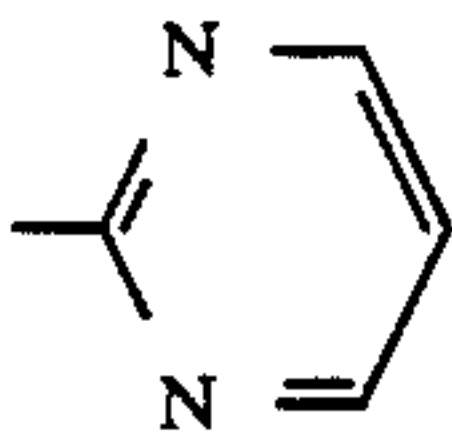
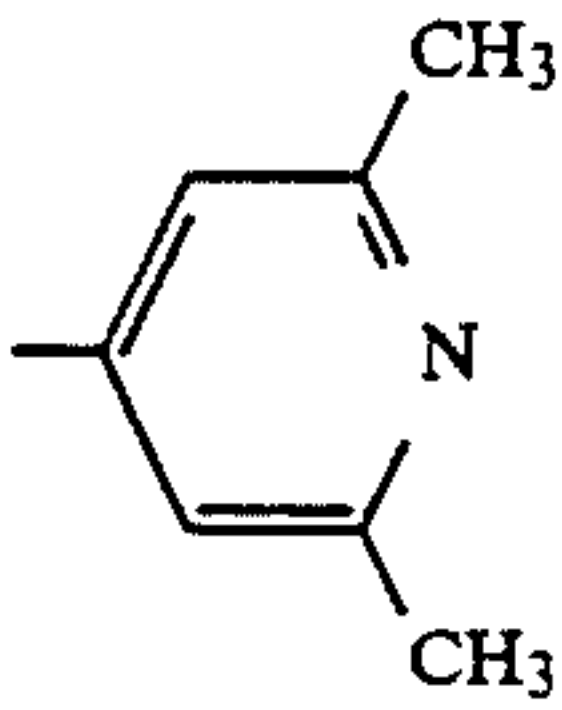
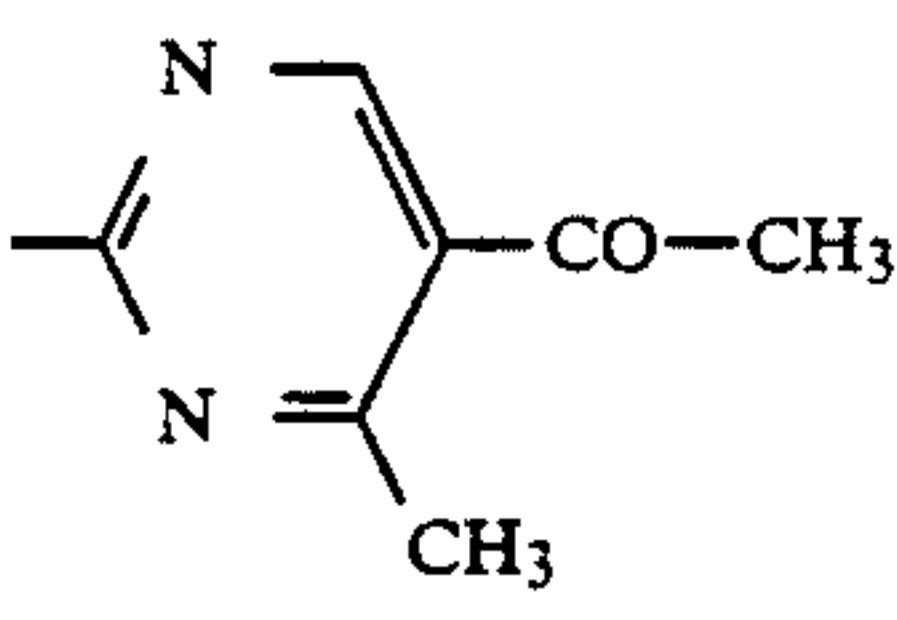
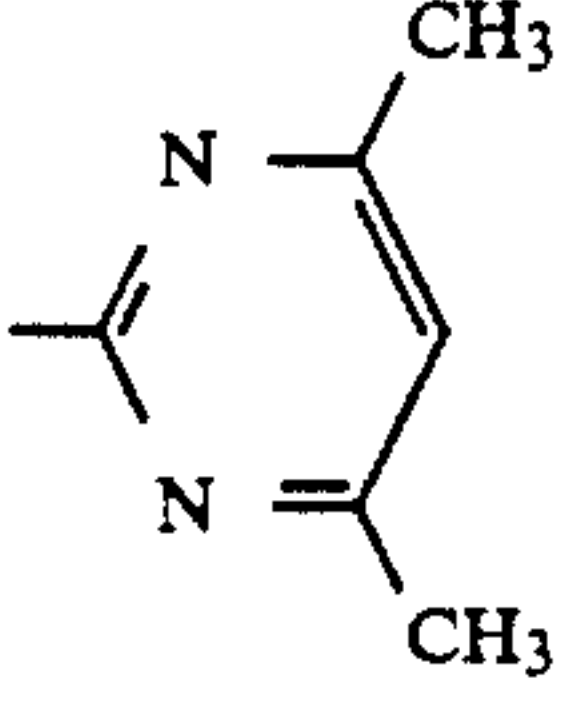
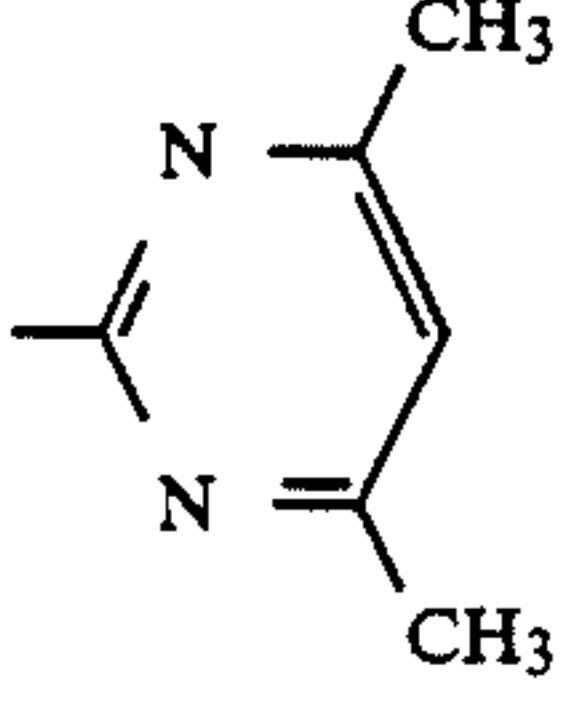
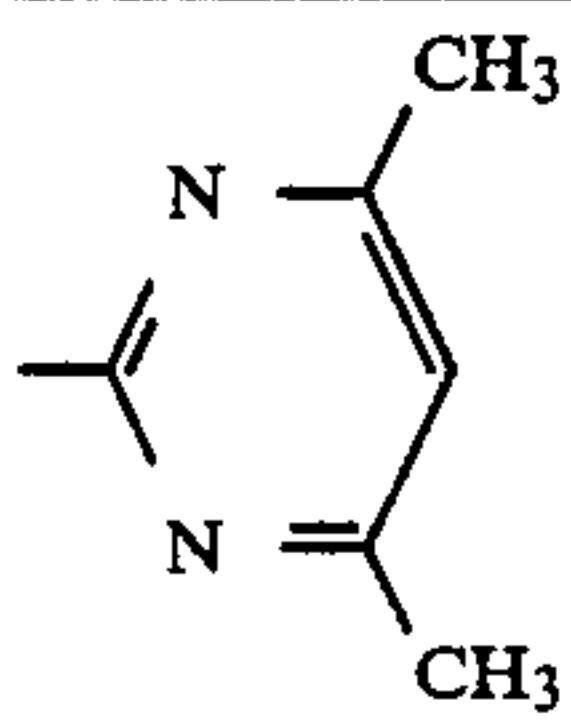
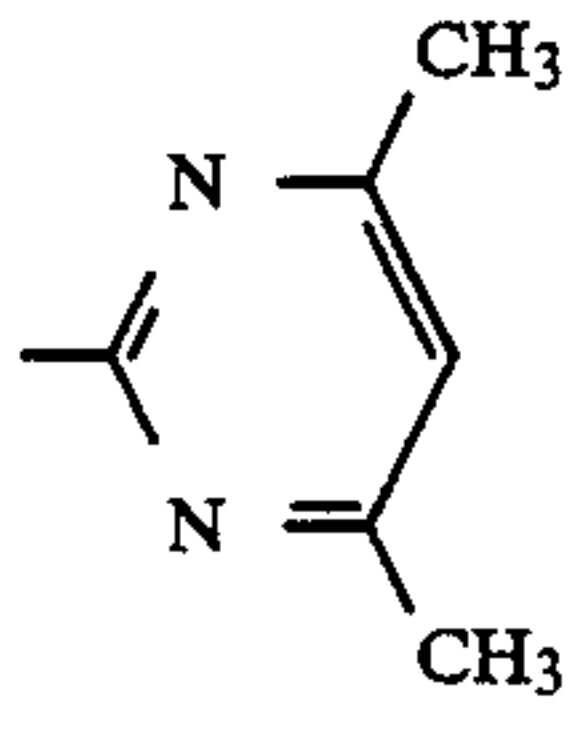
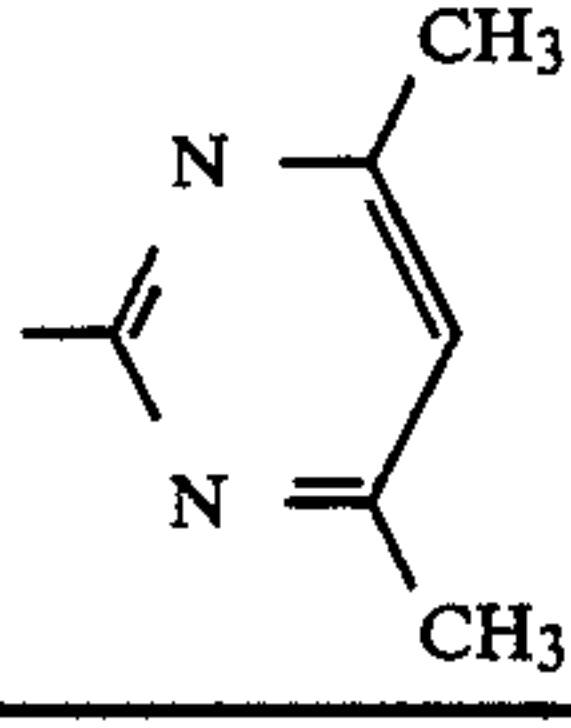
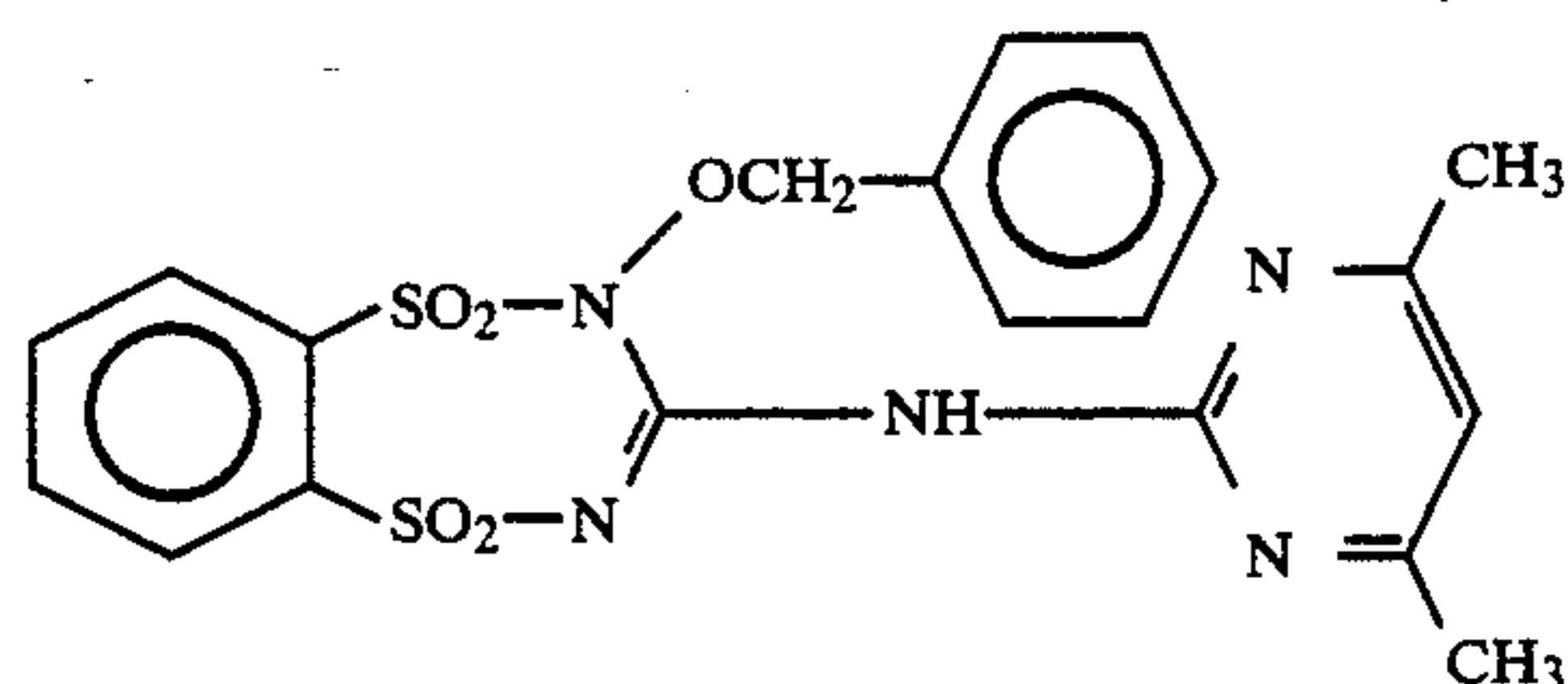
Example No.	R ¹	R ²	R ³	Melting point (° C.)
				
43	-CH ₃	H		
44	-CH ₃	H		
45	-CH ₃	H		
46		H		
47	-CH ₃	H		
48	-CH ₃	H		
49	-C ₂ H ₅	H		218
50	-C ₃ H ₇ (-n)	H		212 (decomp.)

TABLE 2-continued

Example No.	R ¹	R ²	R ³	Melting point (° C.)
51	—C ₃ H ₇ (—i)	H		218 (decomp.)
52	—C ₄ H ₉ (—n)	H		109
53	—CH ₃	H		218

Preparation of the starting compounds of the formula
(II)

EXAMPLE (II-1)



14 g (0.05 mole) of benzene-1,2-disulphonic acid dichloride are added in portions to a mixture of 13.6 g (0.05 mole) of N'-(4,6-dimethylpyrimidin-2-yl)-N''-benzyloxy-guanidine, 12 g (0.15 mole) of pyridine and 100 ml of methylene chloride at —20° C. The mixture is

subsequently stirred at —20° C. for 3 hours and at +20° C. for 15 hours. The reaction mixture is then evaporated and 70 ml of dioxane are added to the residue. The mixture is filtered. The filtrate is concentrated, the residue is triturated with ethanol and the product which has precipitated is isolated by filtration with suction.

15 g (68% of theory) of the compound of the above-mentioned structural formula of melting point 199° C. are obtained.

The compounds of the formula (II) listed in the following Table 3 can be prepared by the process described by way of example in the preceding example:

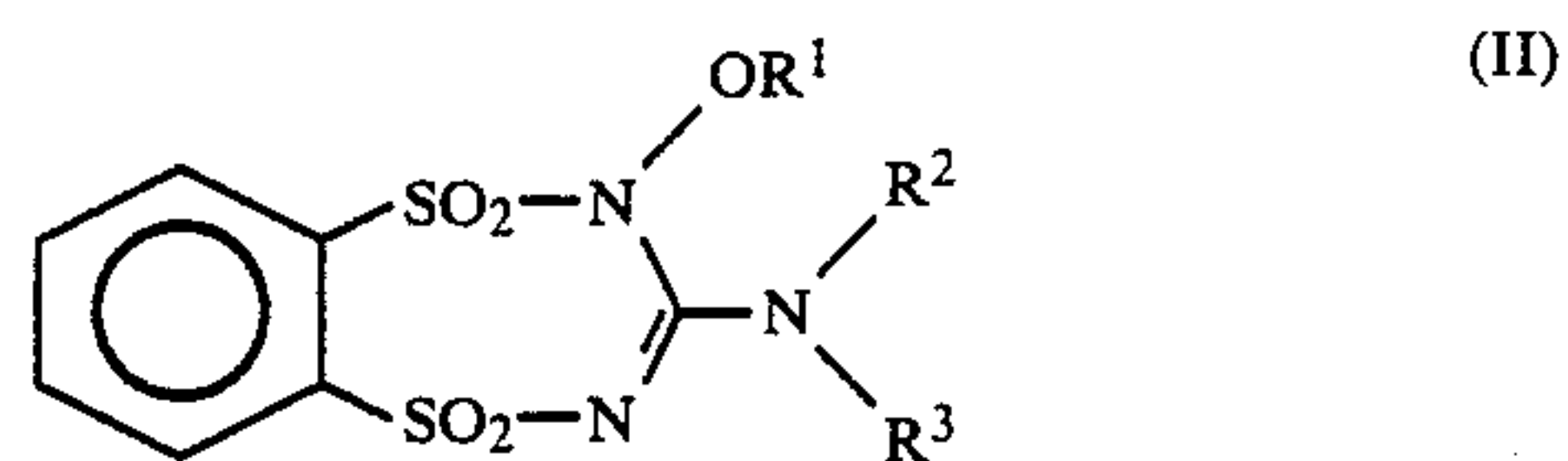


TABLE 3

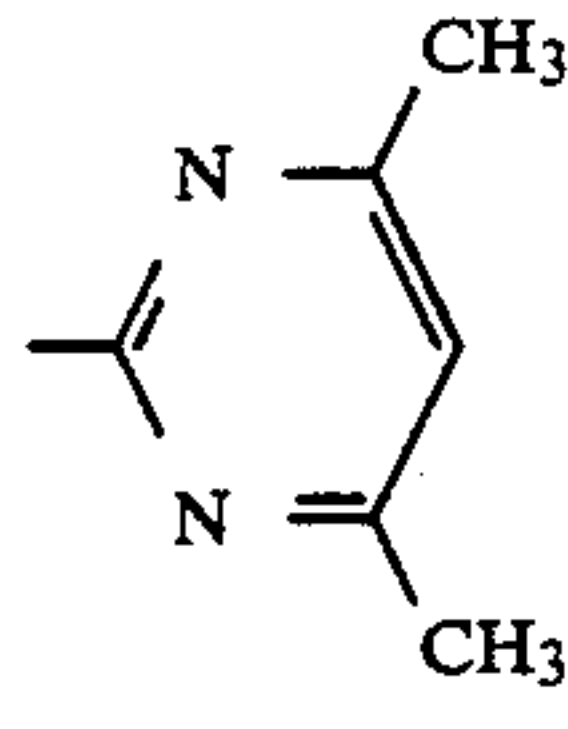
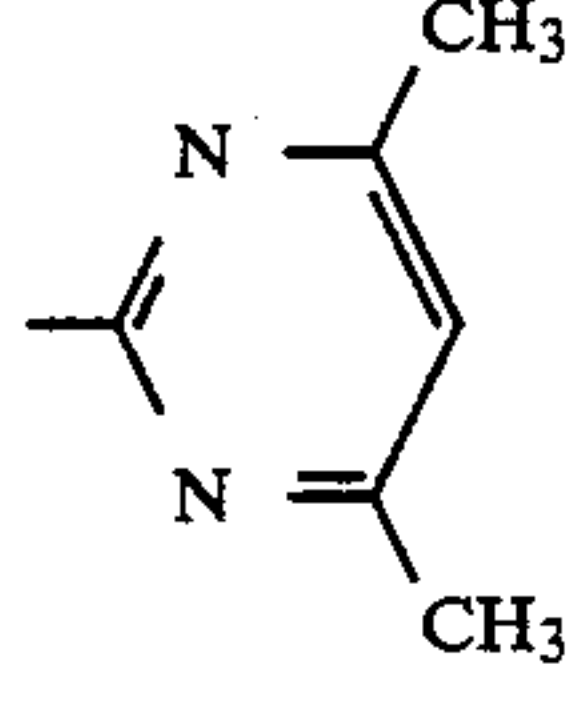
Example No.	R ¹	R ²	R ³	Melting point (° C.)
(II-2)	—CH ₂ —CH=CH ₂	H		180 (decomp.)
(II-3)	—C ₈ H ₁₇ (—n)	H		164

TABLE 3-continued

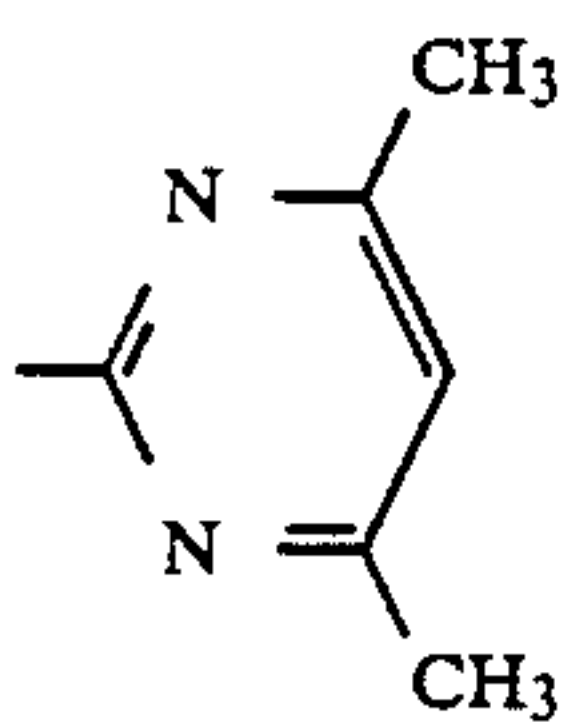
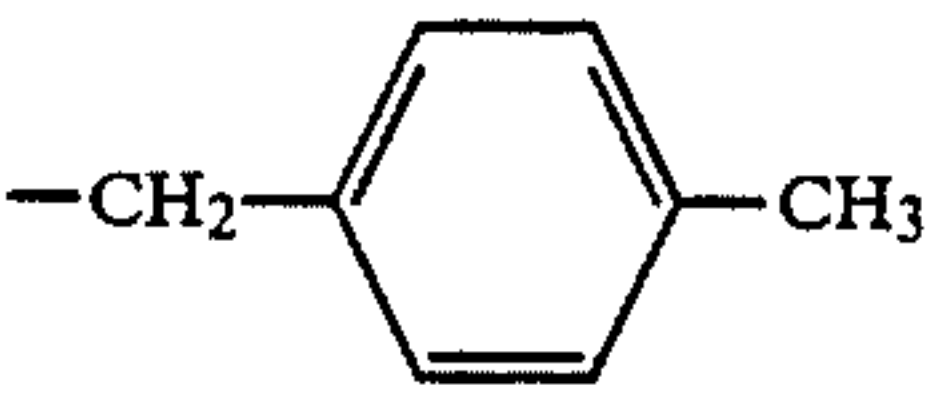
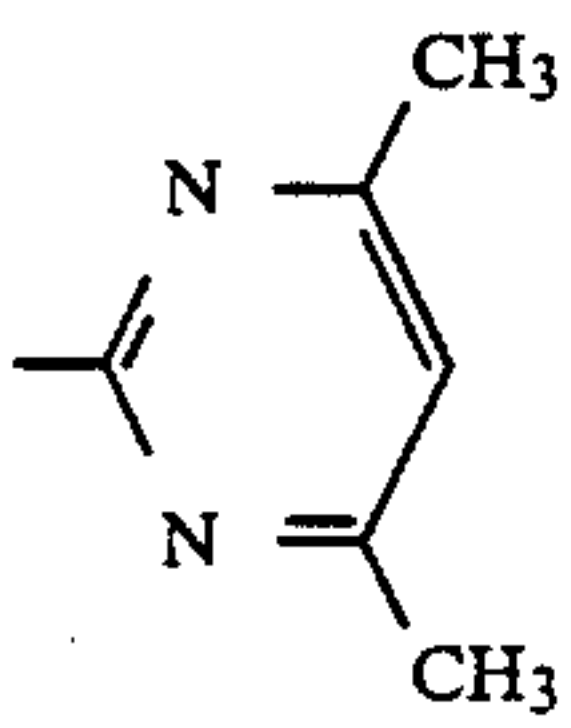
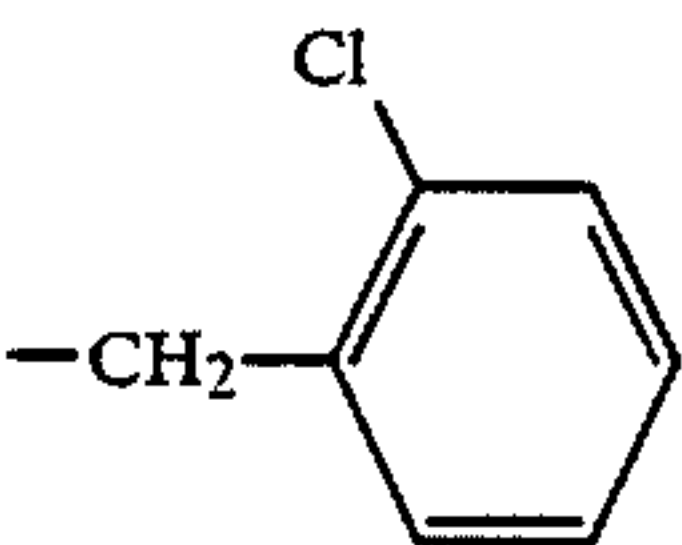
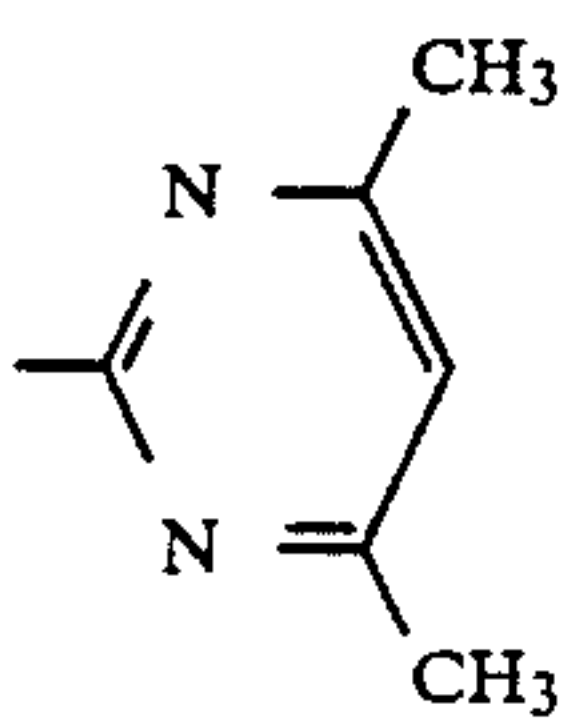
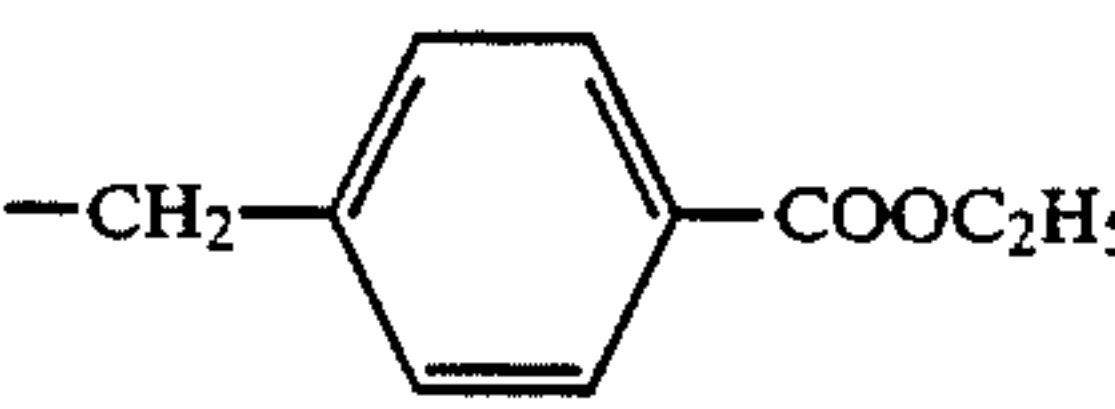
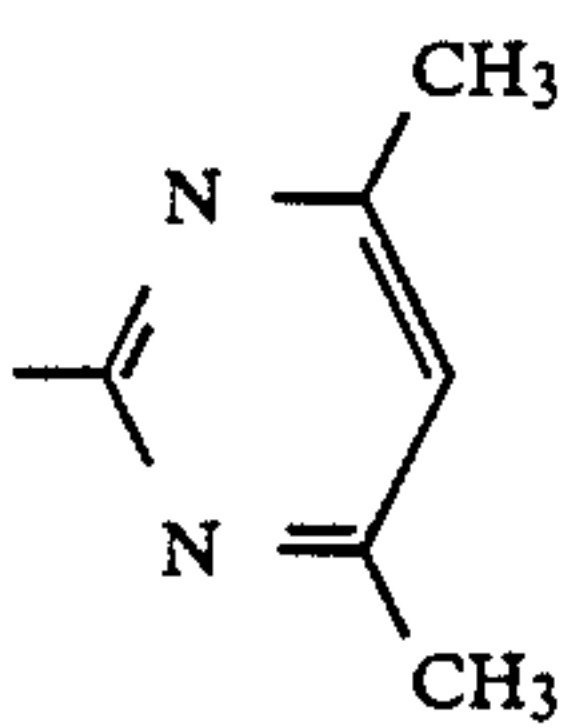
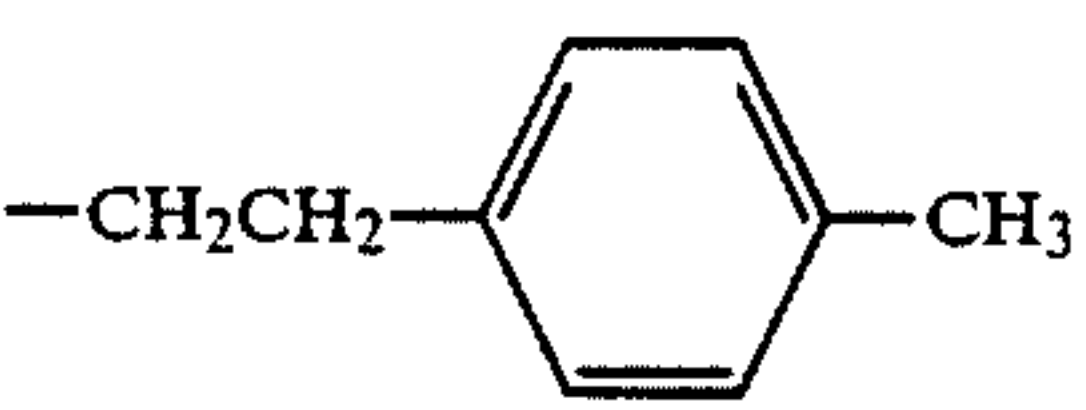
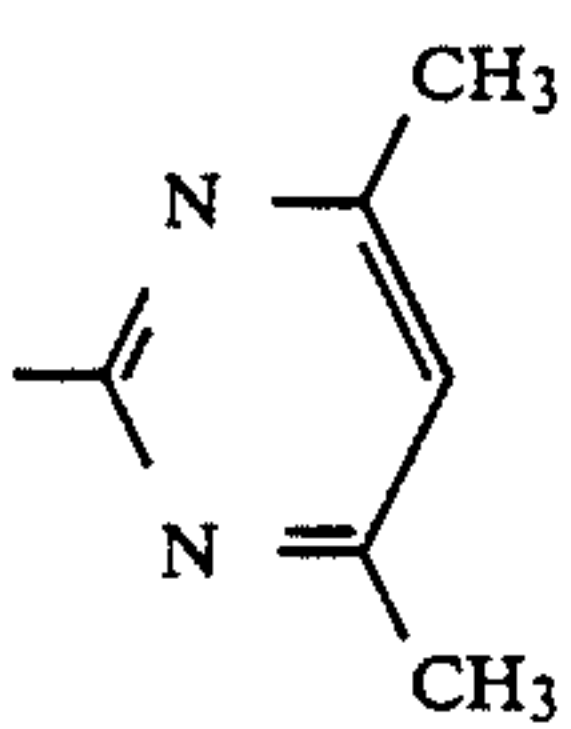
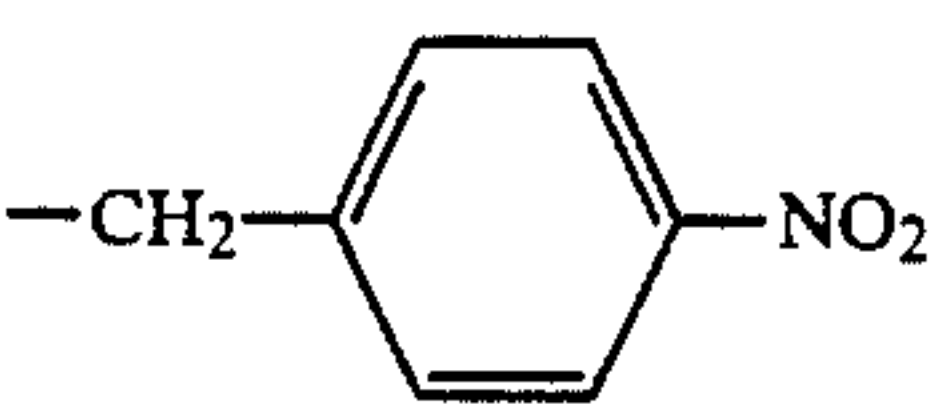
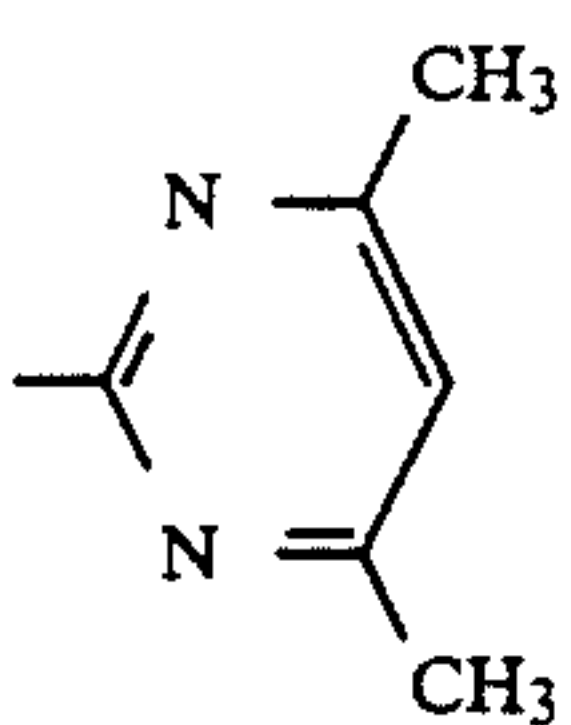
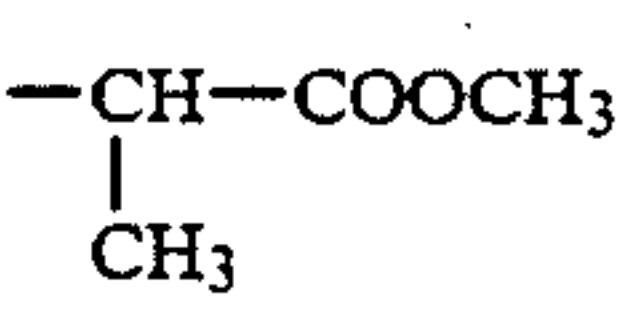
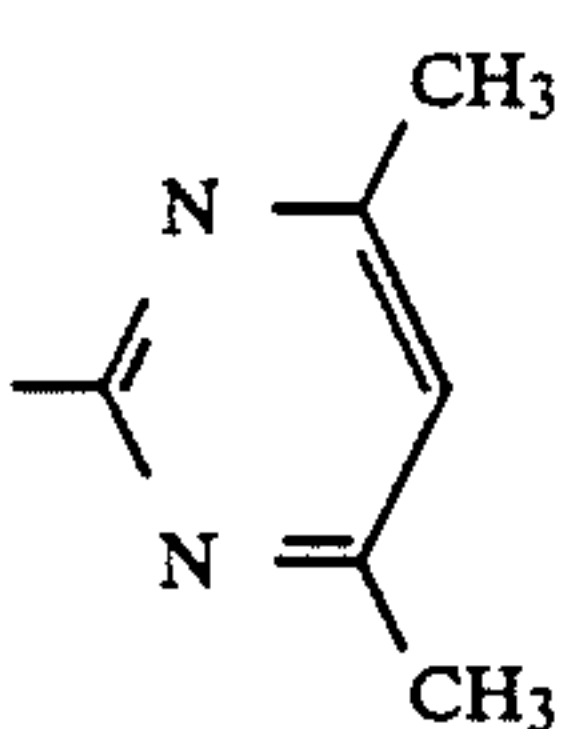
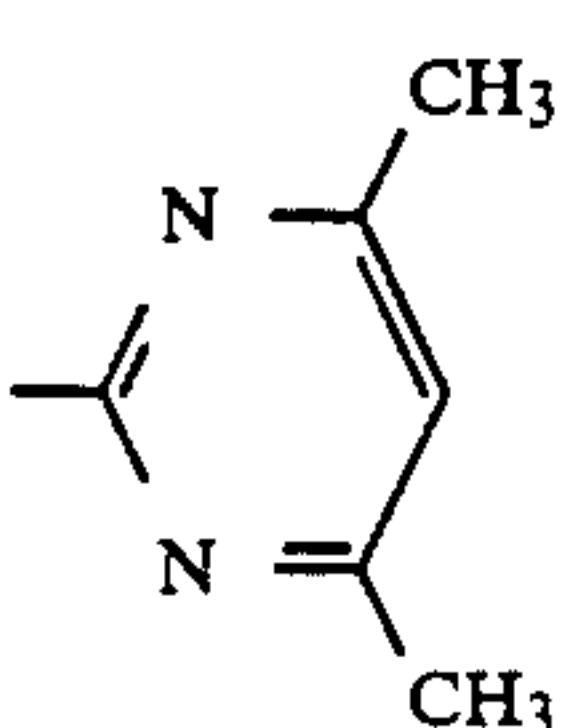
Example No.	R ¹	R ²	R ³	Melting point (°C.)
(II-4)	$-\text{CH}_2-\text{COOC}_2\text{H}_5$	H		210 (decomp.)
(II-5)		H		
(II-6)		H		
(II-7)		H		
(II-8)		H		
(II-9)		H		
(II-10)		H		
(II-11)	$-\text{CH}_2-\text{CH}(\text{CH}_3)_2$	H		amorphous

TABLE 3-continued

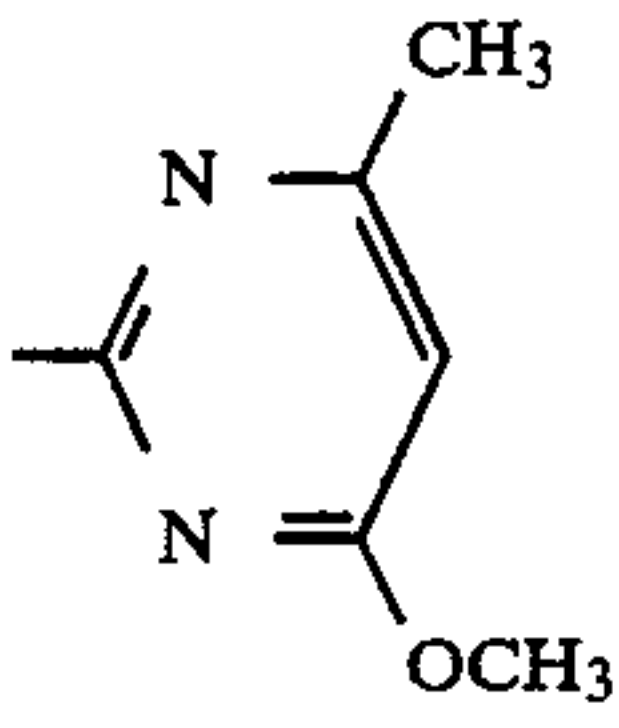
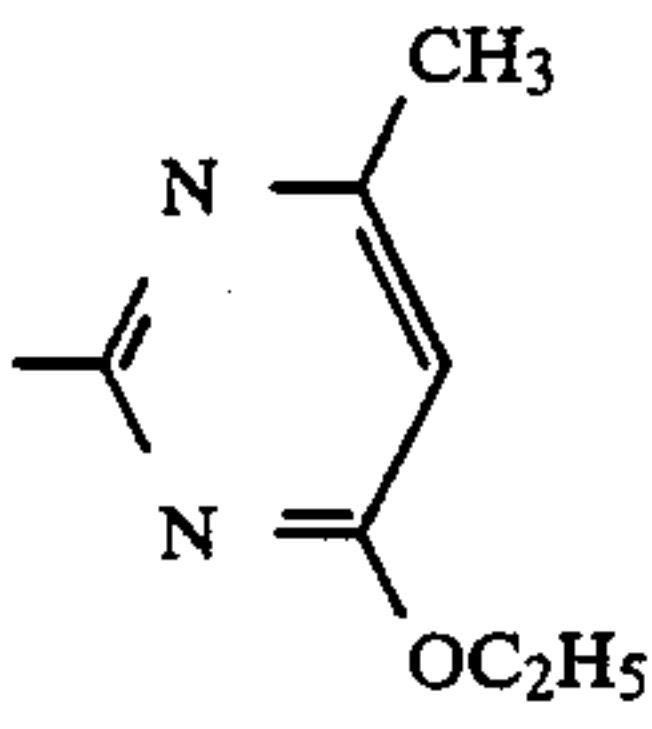
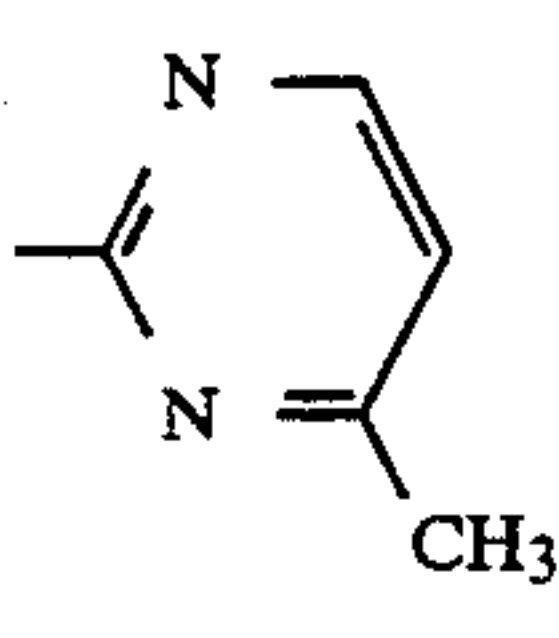
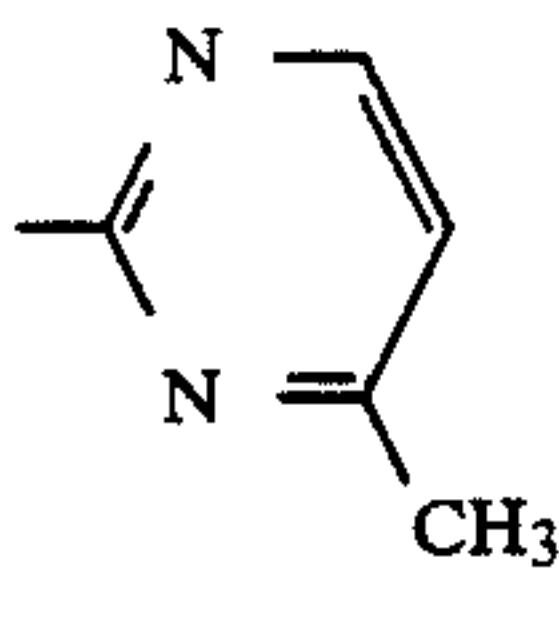
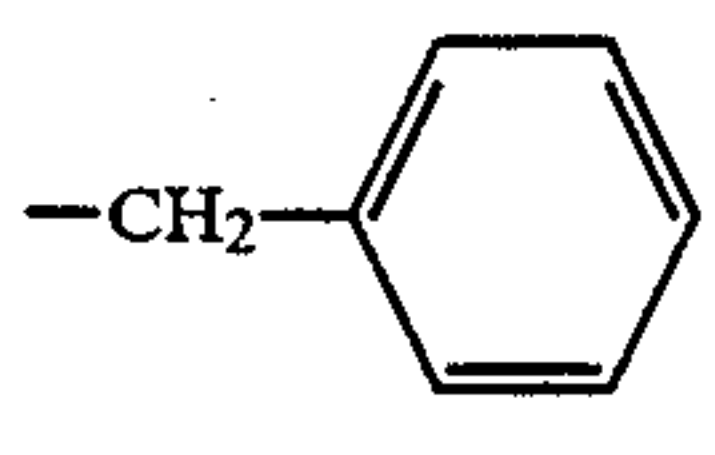
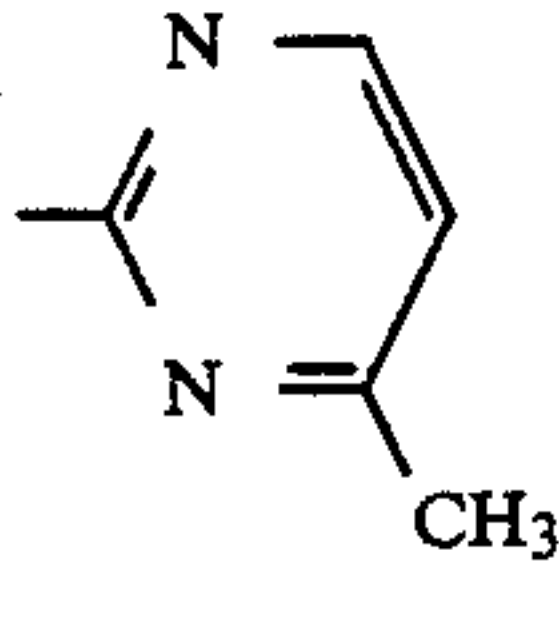
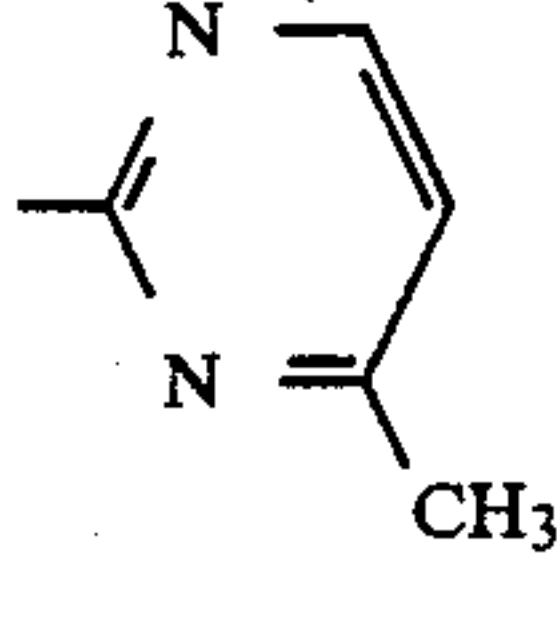
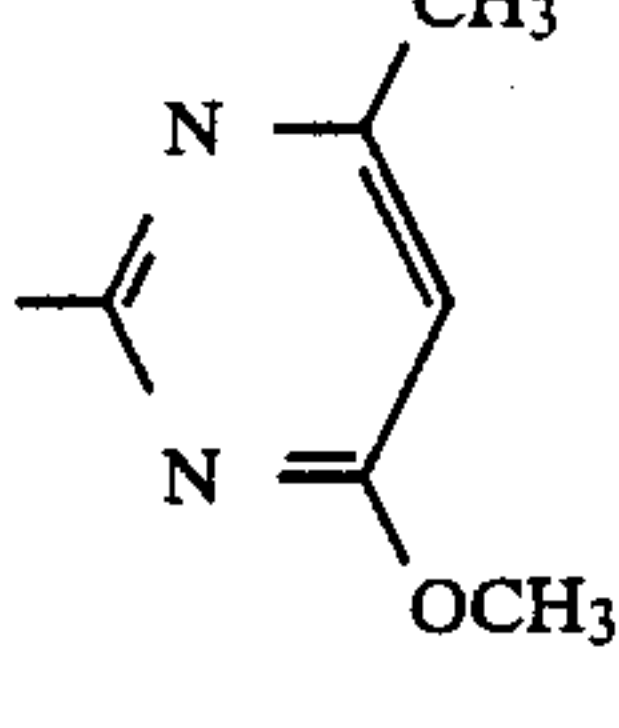
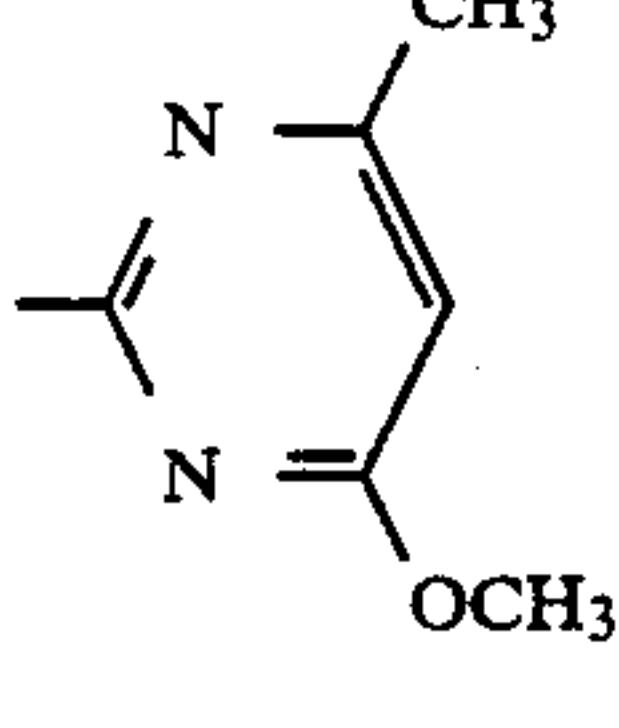
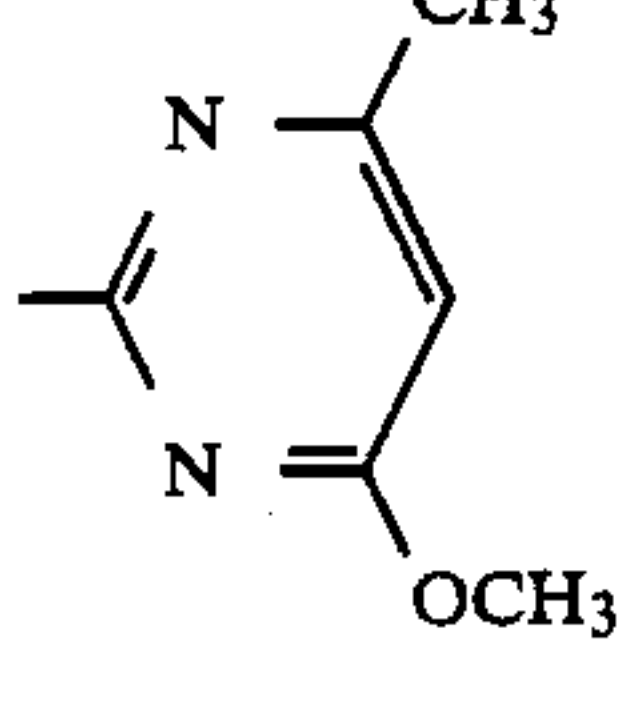
Example No.	R ¹	R ²	R ³	Melting point (°C.)
(II-12)	—CH ₃	CH ₃		
(II-13)	—CH ₃	C ₂ H ₅		
(II-14)	—CH ₃	H		
(II-15)	—CH ₂ CH(CH ₃) ₂	H		
(II-16)		H		
(II-17)	—CH ₂ —COOC ₂ H ₅	H		
(II-18)	—CH ₃	H		151 (decomp.)
(II-19)	—C ₂ H ₅	H		
(II-20)	—C ₃ H ₇	H		

TABLE 3-continued

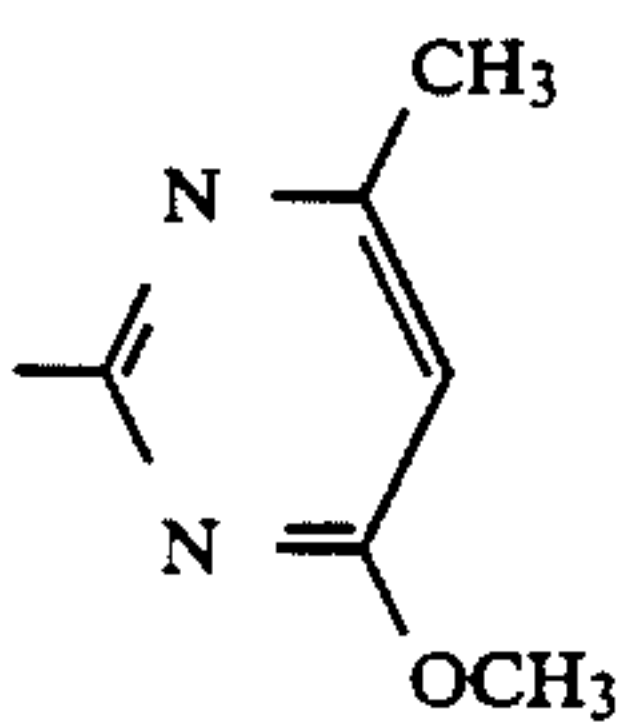
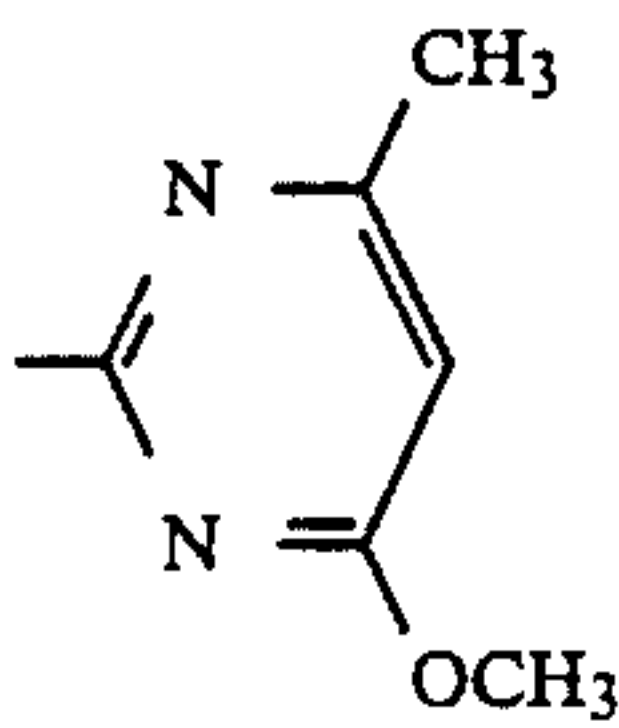
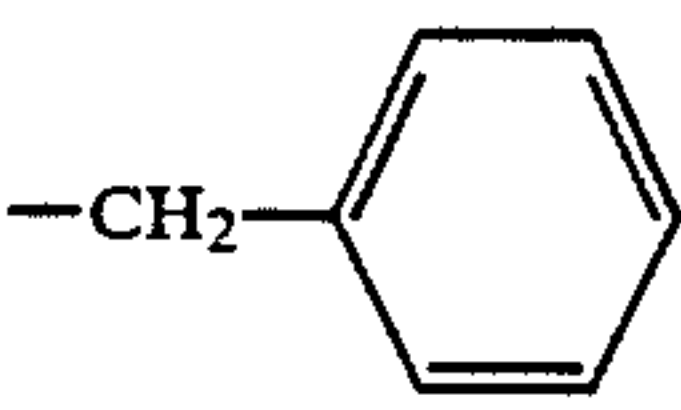
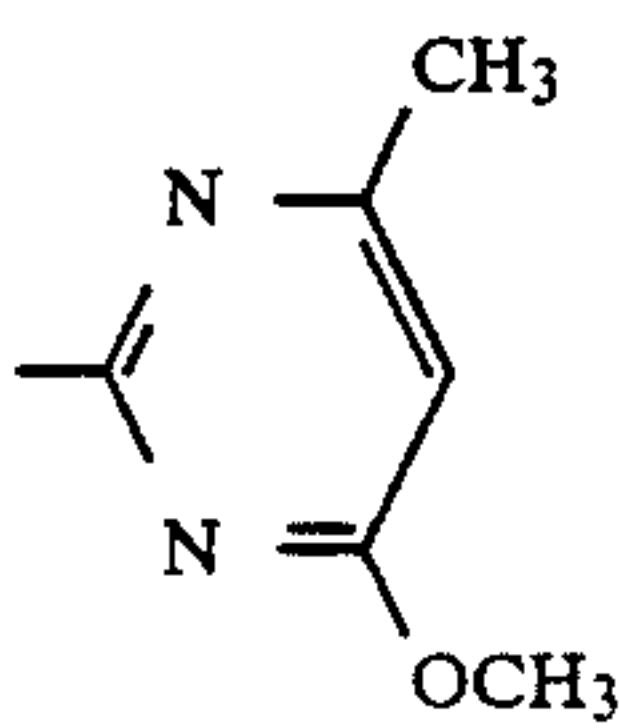
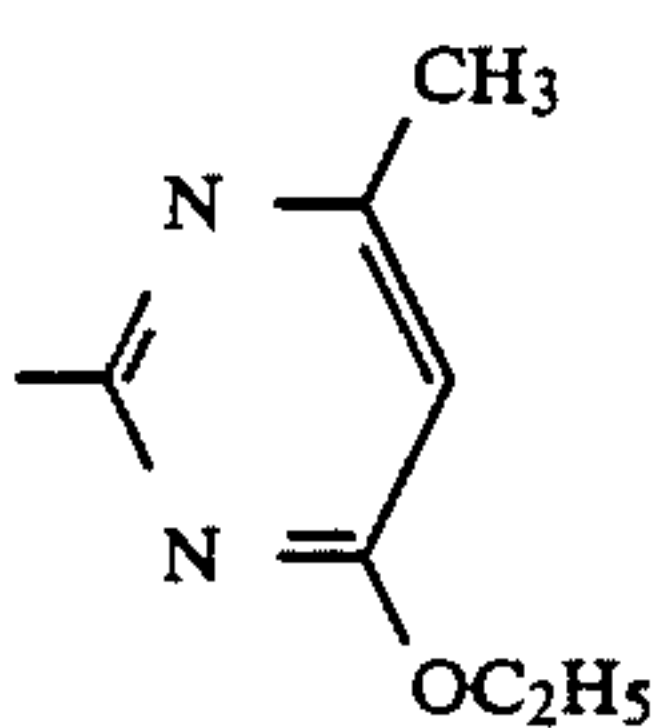
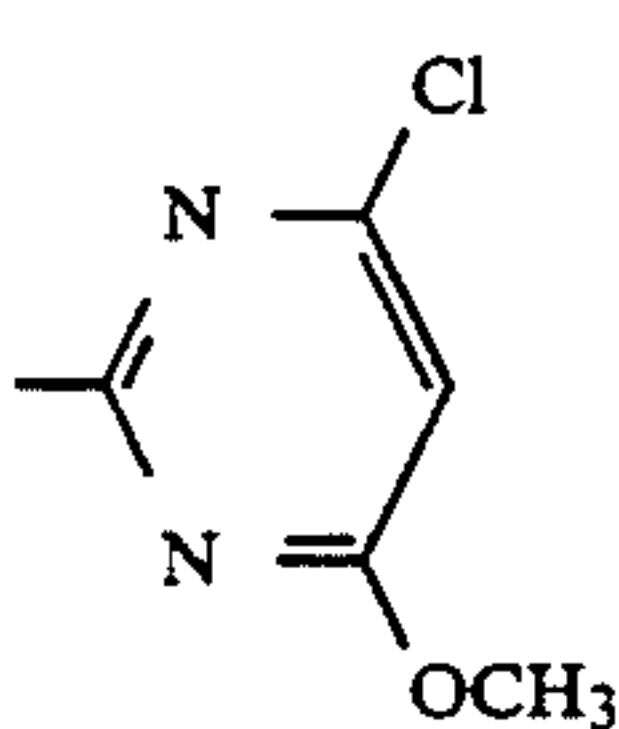
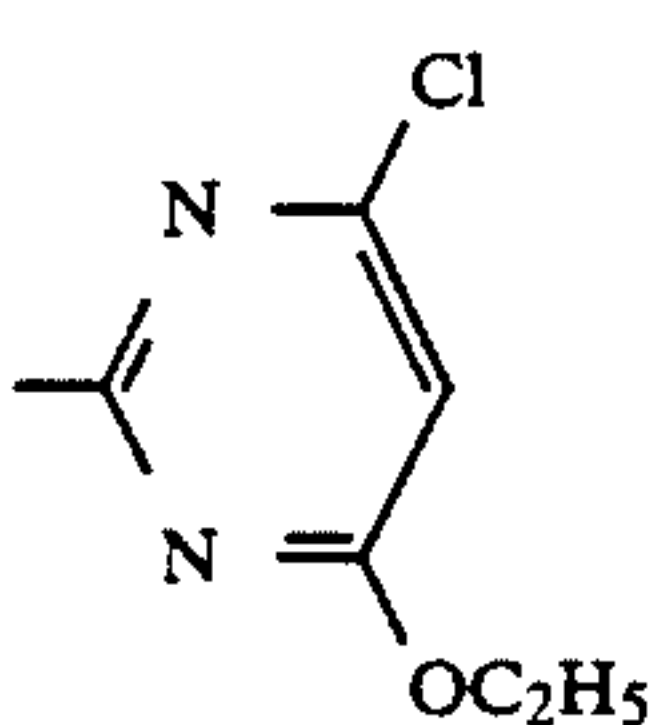
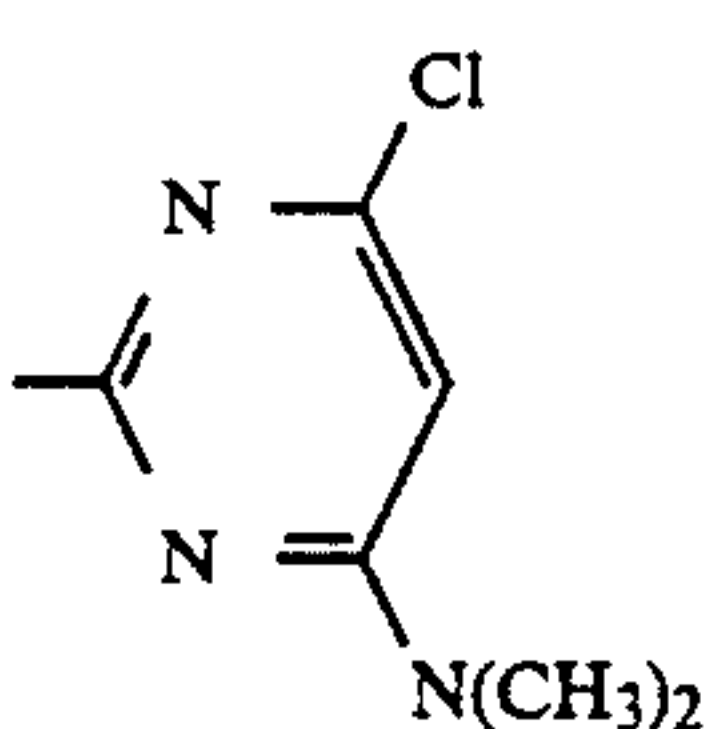
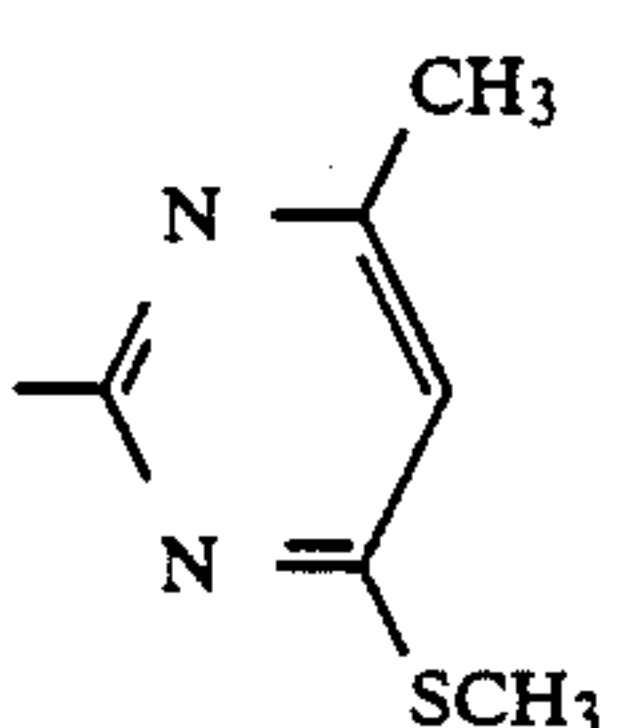
Example No.	R ¹	R ²	R ³	Melting point (°C.)
(II-21)	$-\text{CH}_2-\text{CH}=\text{CH}_2$	H		
(II-22)	$-\text{CH}_2\text{CH}(\text{CH}_3)_2$	H		
(II-23)		H		
(II-24)	$-\text{CH}_3$	H		
(II-25)	$-\text{CH}_3$	H		
(II-26)	$-\text{CH}_3$	H		
(II-27)	$-\text{CH}_3$	H		
(II-28)	$-\text{CH}_3$	H		

TABLE 3-continued

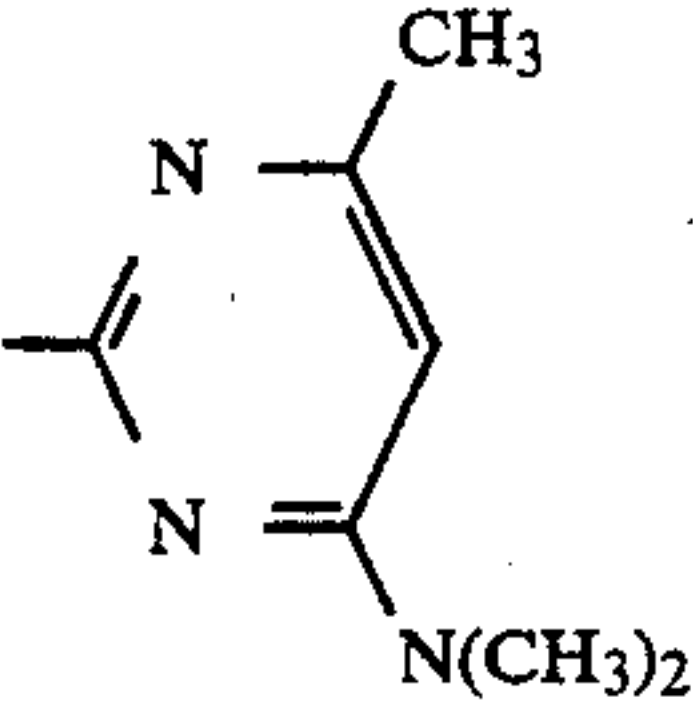
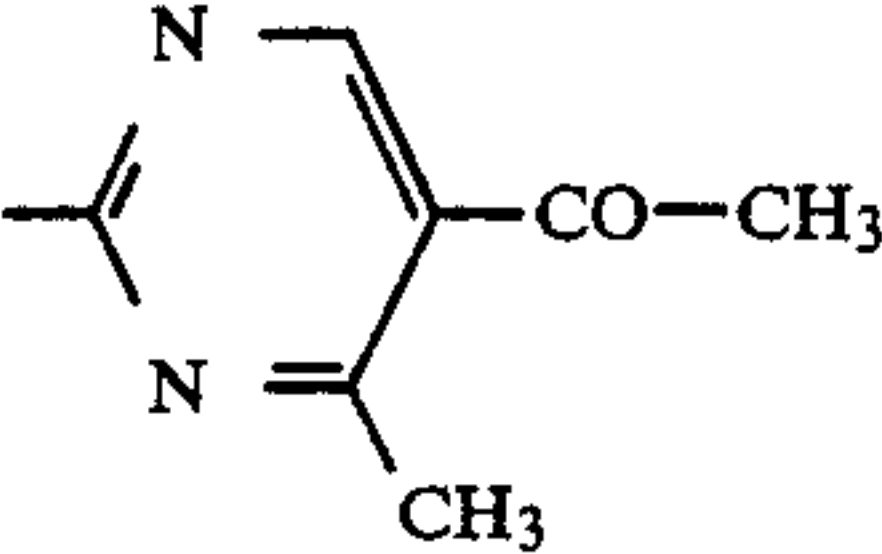
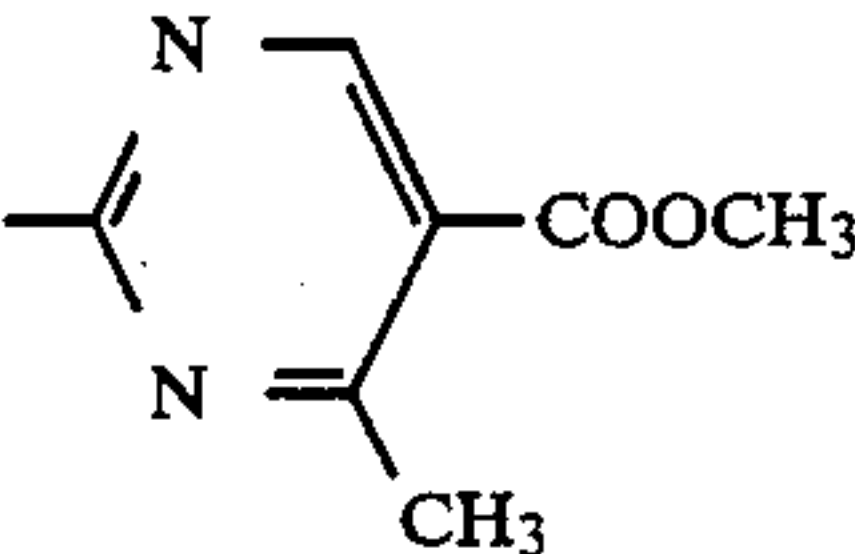
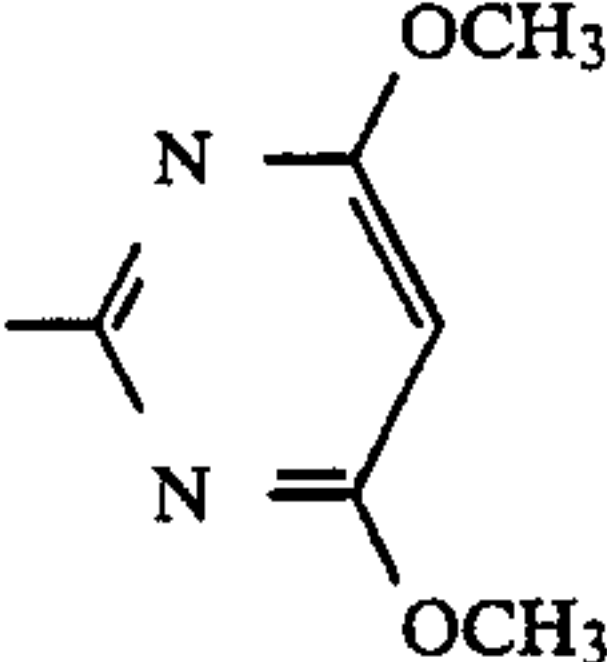
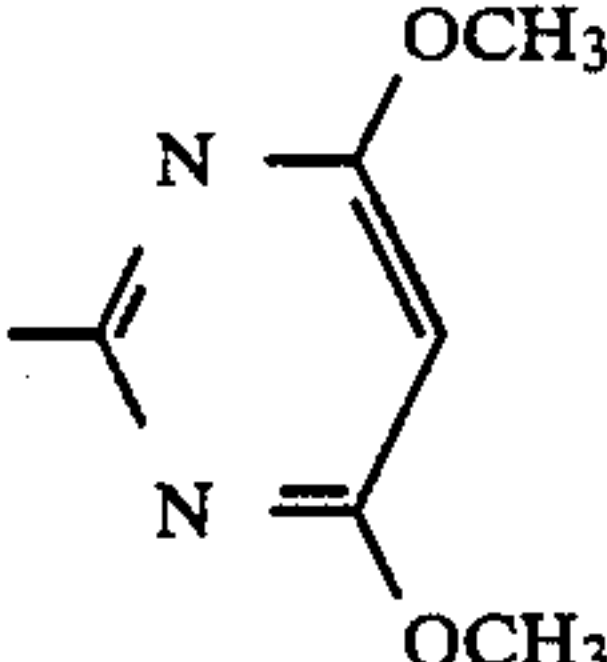
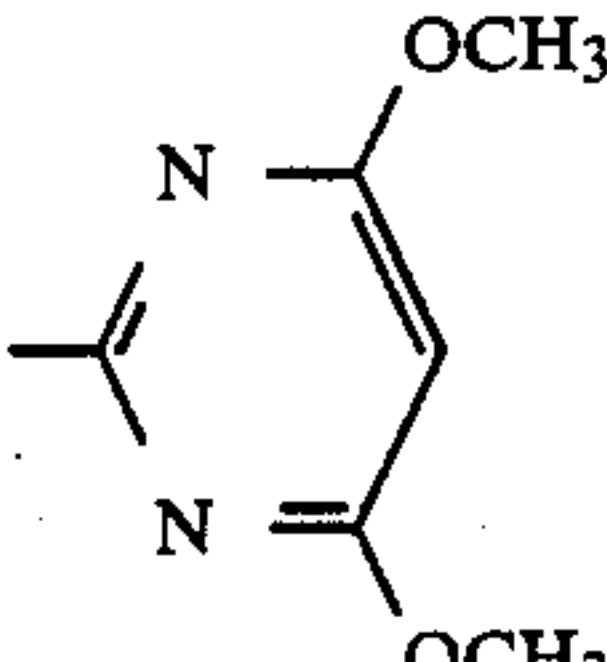
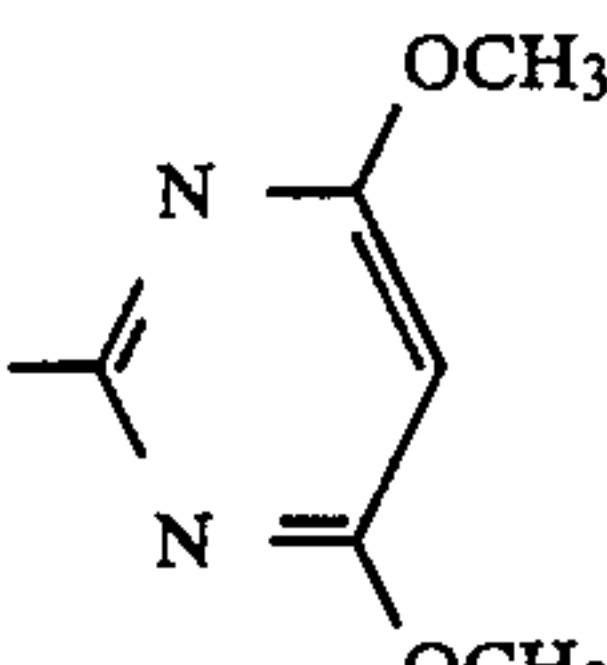
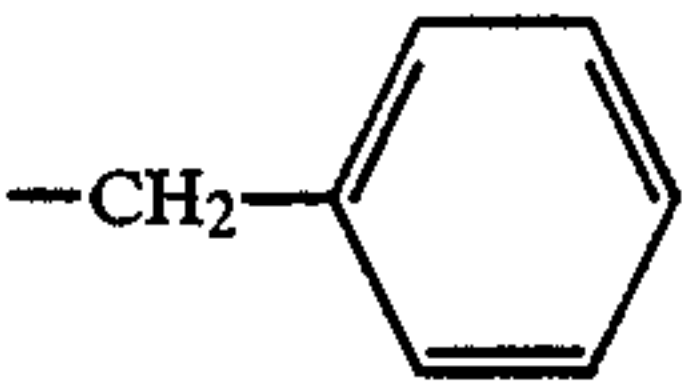
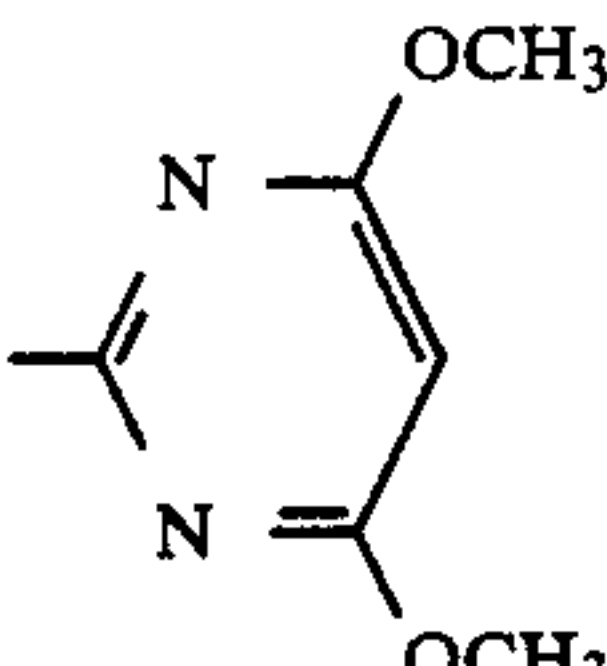
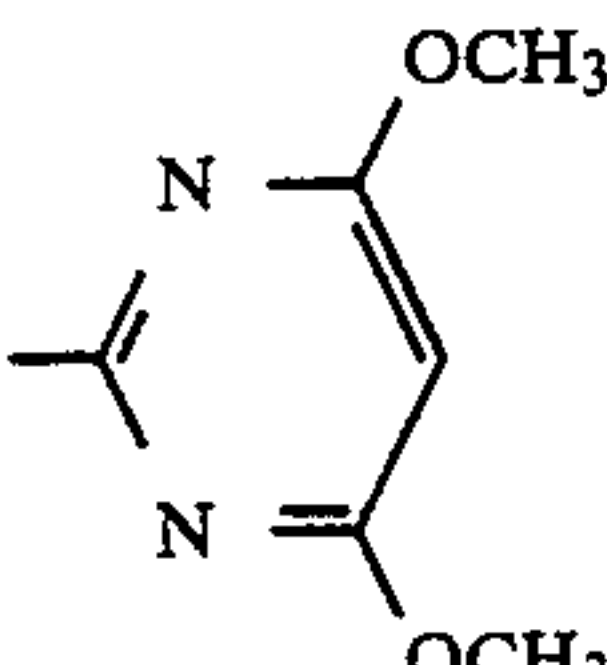
Example No.	R ¹	R ²	R ³	Melting point (°C.)
(II-29)	—CH ₃	H		
(II-30)	—CH ₃	H		
(II-31)	—CH ₃	H		
(II-32)	—CH ₃	H		
(II-33)	—C ₂ H ₅	H		
(II-34)	—C ₃ H _{7(-n)}	H		
(II-35)	—C ₄ H _{9(-n)}	H		
(II-36)		H		
(II-37)	—CH ₂ —CH=CH ₂	H		

TABLE 3-continued

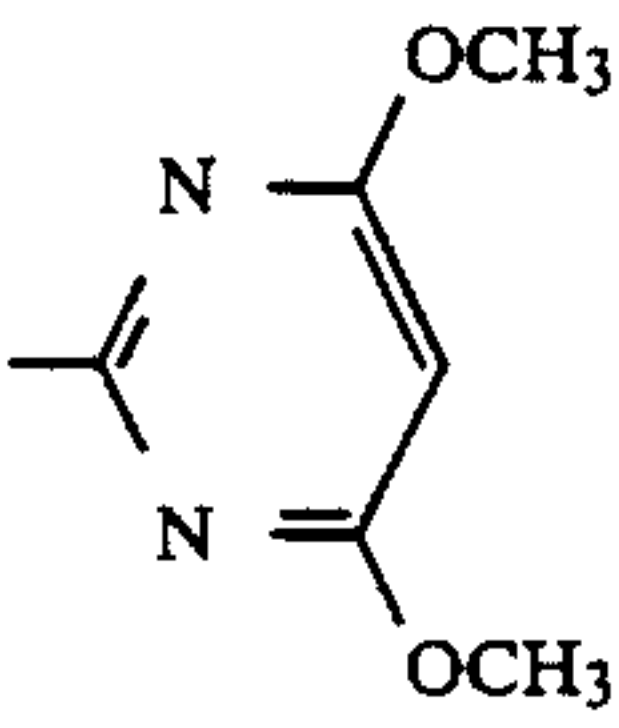
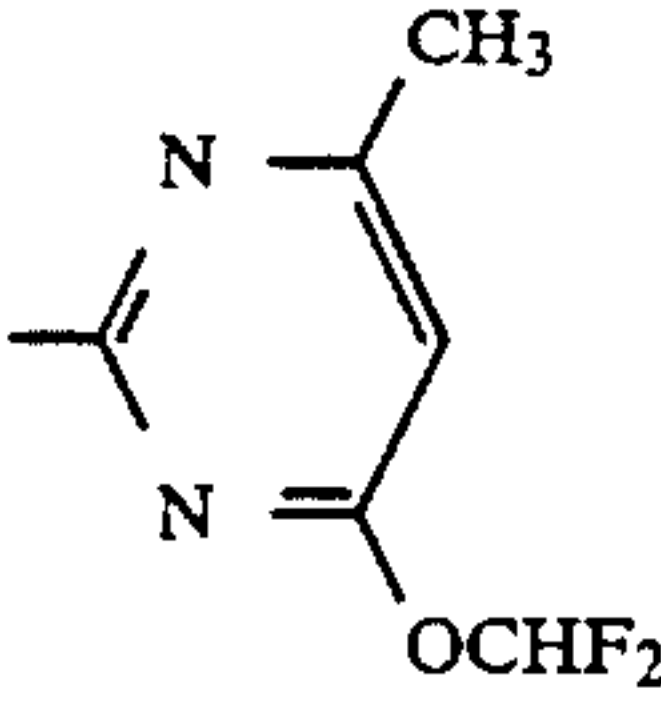
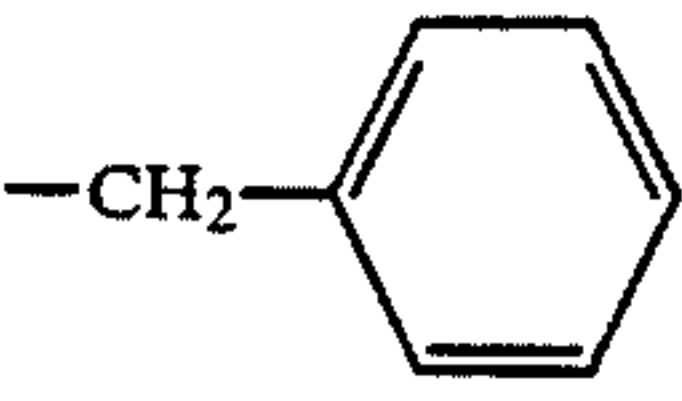
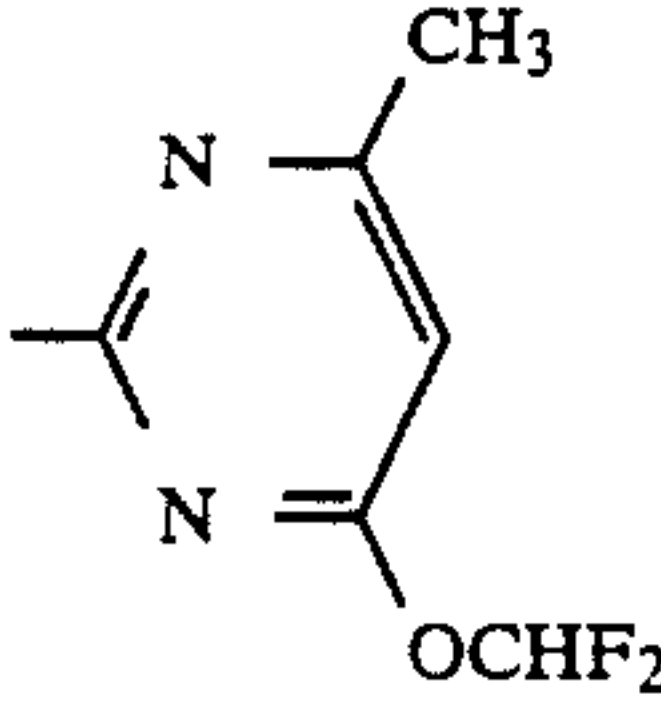
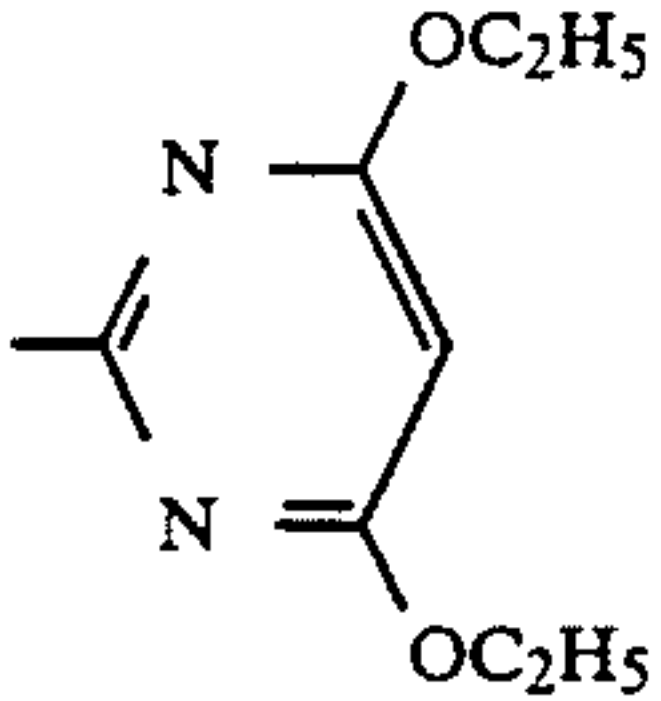
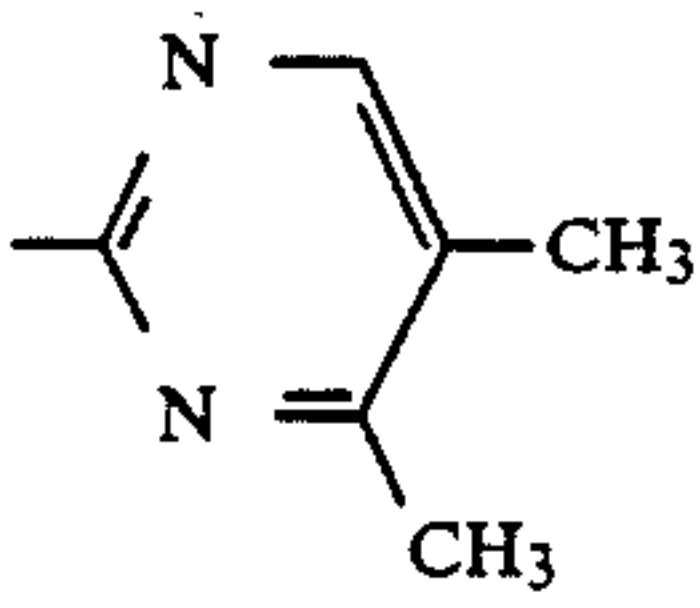
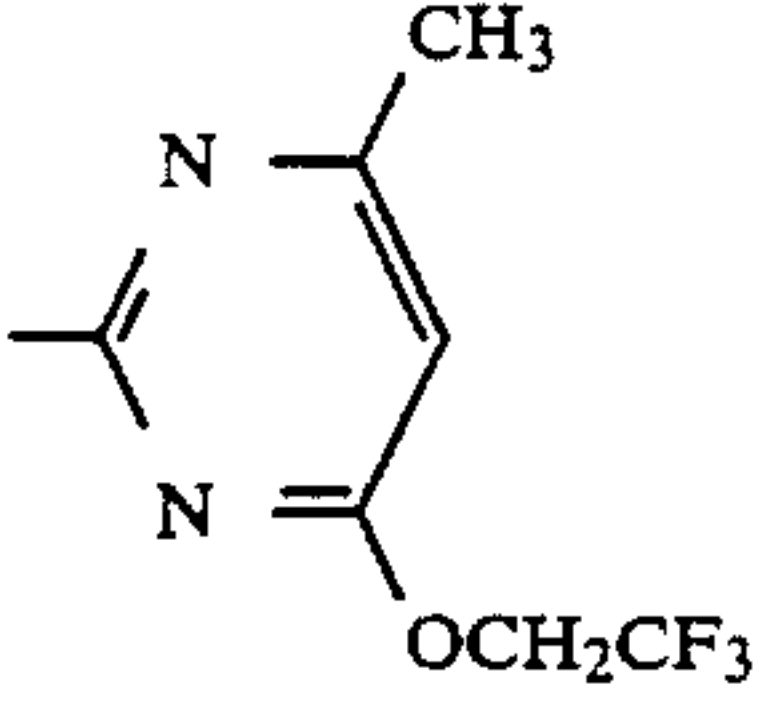
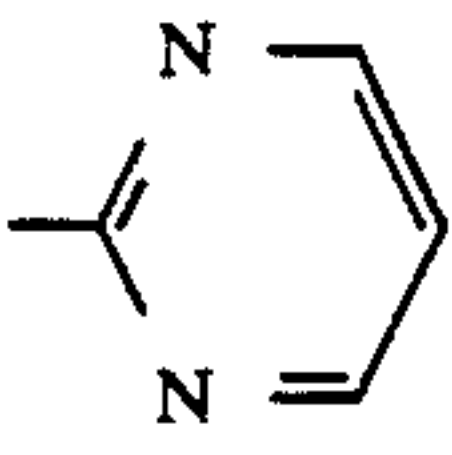
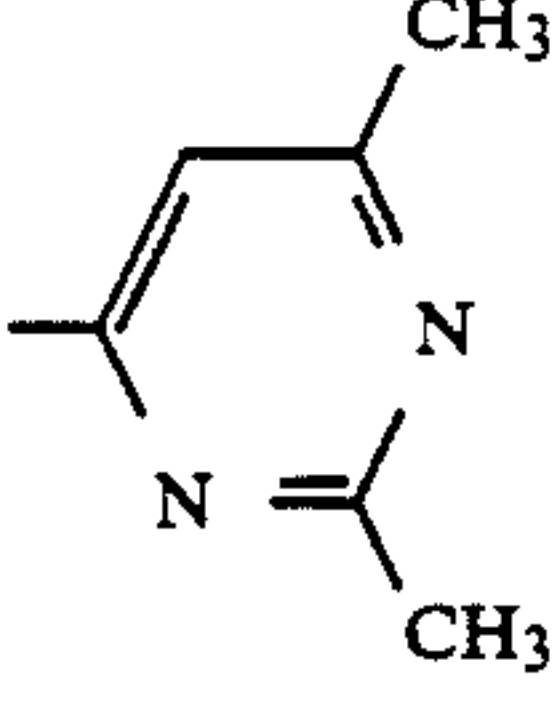
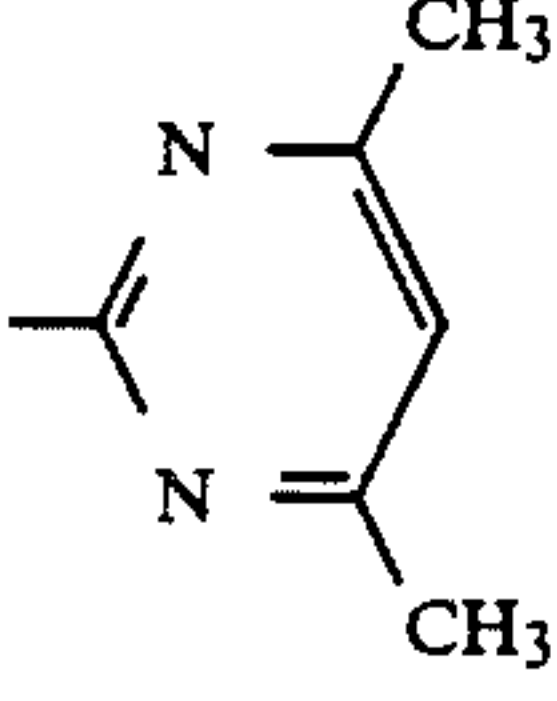
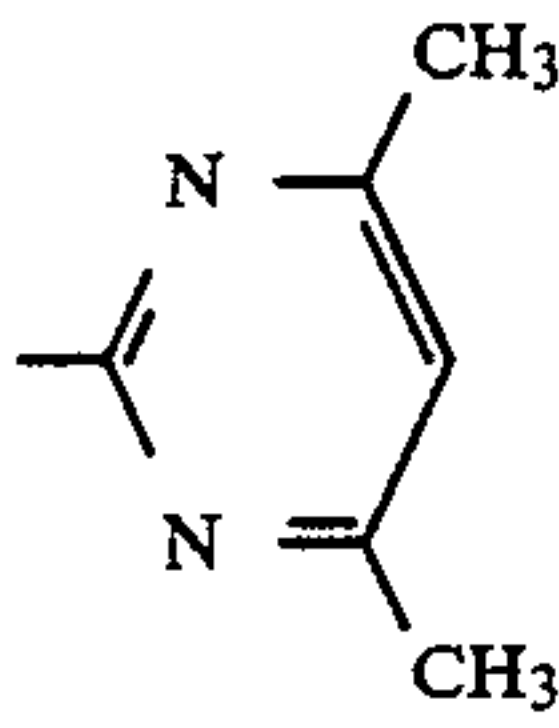
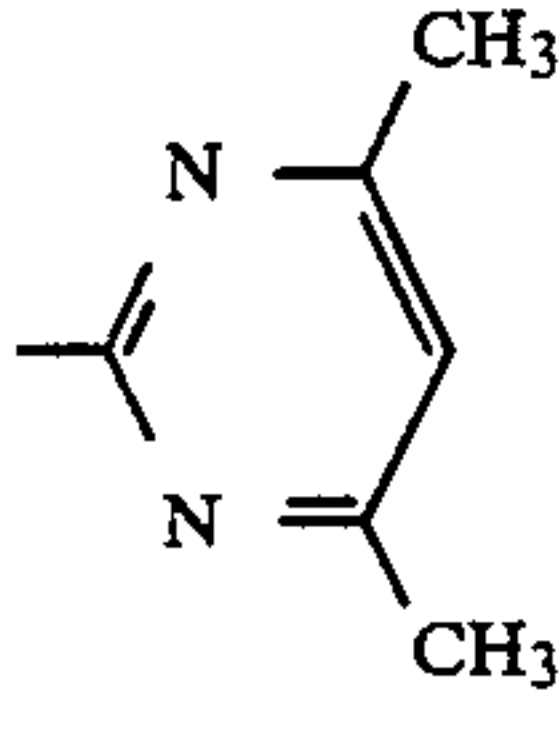
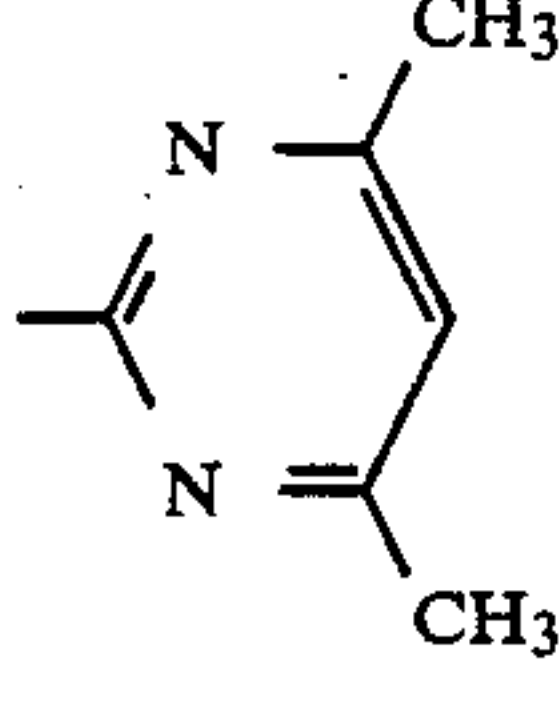
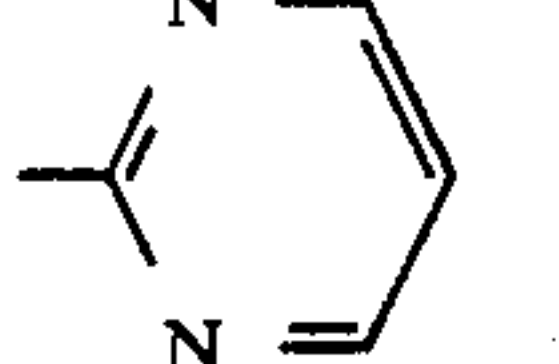
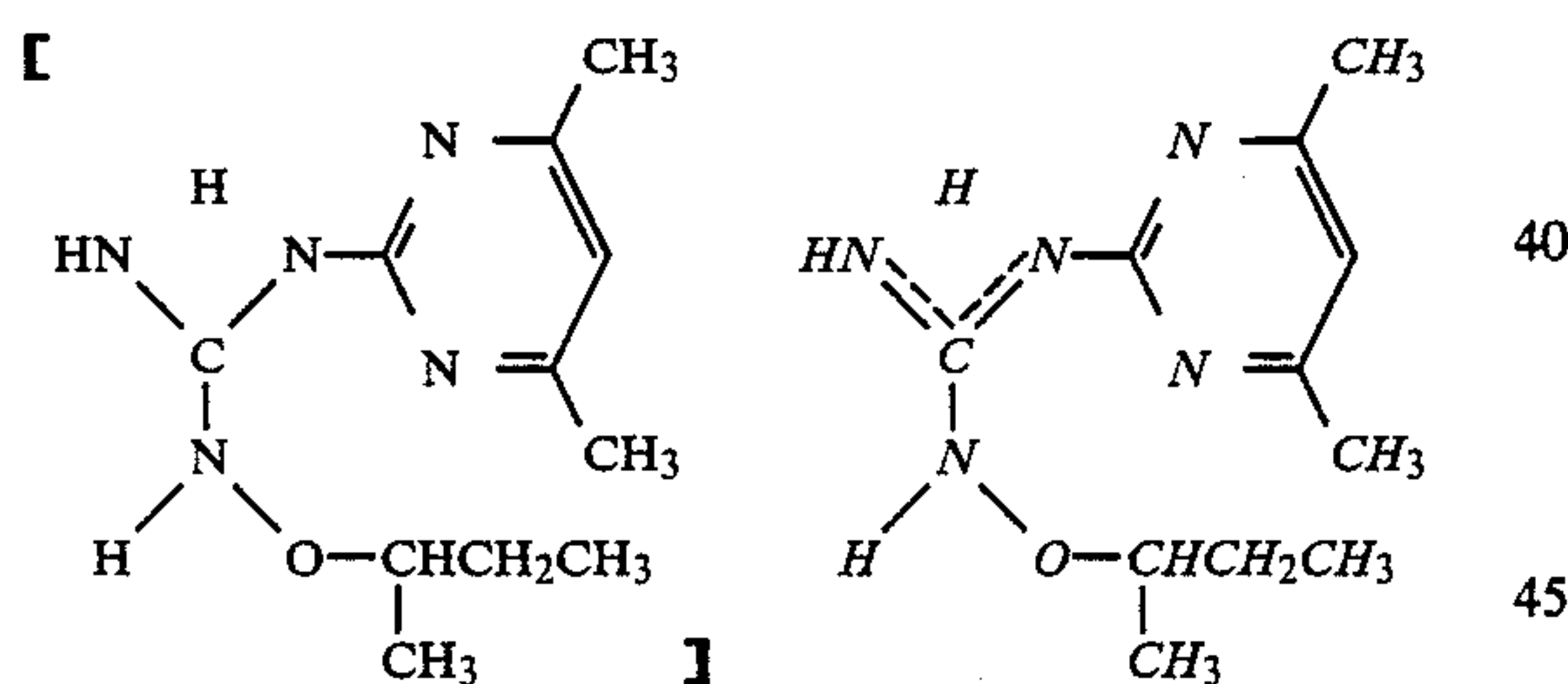
Example No.	R ¹	R ²	R ³	Melting point (°C.)
(II-38)	$-\text{CH}_2\text{CH}(\text{CH}_3)_2$	H		
(II-39)	$-\text{CH}_3$	H		
(II-40)		H		
(II-41)	$-\text{CH}_3$	H		
(II-42)	$-\text{CH}_3$	H		
(II-43)	$-\text{CH}_3$	H		
(II-44)	$-\text{CH}_2\text{CH}(\text{CH}_3)_2$	H		
(II-45)	$-\text{CH}_3$	H		
(II-46)	$-\text{CH}_3$	H		158

TABLE 3-continued

Example No.	R ¹	R ²	R ³	Melting point (°C.)
(II-47)	-C ₂ H ₅	H		104
(II-48)	-C ₃ H ₇ (-n)	H		134
(II-49)	-C ₄ H ₉ (-n)	H		179
(II-50)	-CH ₃	H		187

Preparation of the starting substances of the formula (IV)

EXAMPLE (IV-1)



A mixture of 143 g (0.97 mole) of 2-cyanoamino-4,6-dimethylpyrimidine, 94.3 g (1.06 moles) of O-sec-butyl-hydroxylamine and 190 ml of ethanol is heated at

the boiling point under reflux for 6 hours. The mixture is then filtered with suction, the filtrate is concentrated and 500 ml of water are added to the residue. The product thereby obtained as crystals is isolated by filtration with suction.

131 g (57% of theory) of N'-(4,6-dimethylpyrimidin-2-yl)-N''-sec-butoxy-guanidine of melting point 78° C. are obtained.

The compounds of the formula (IV) listed in the following Table 4 can be prepared analogously:

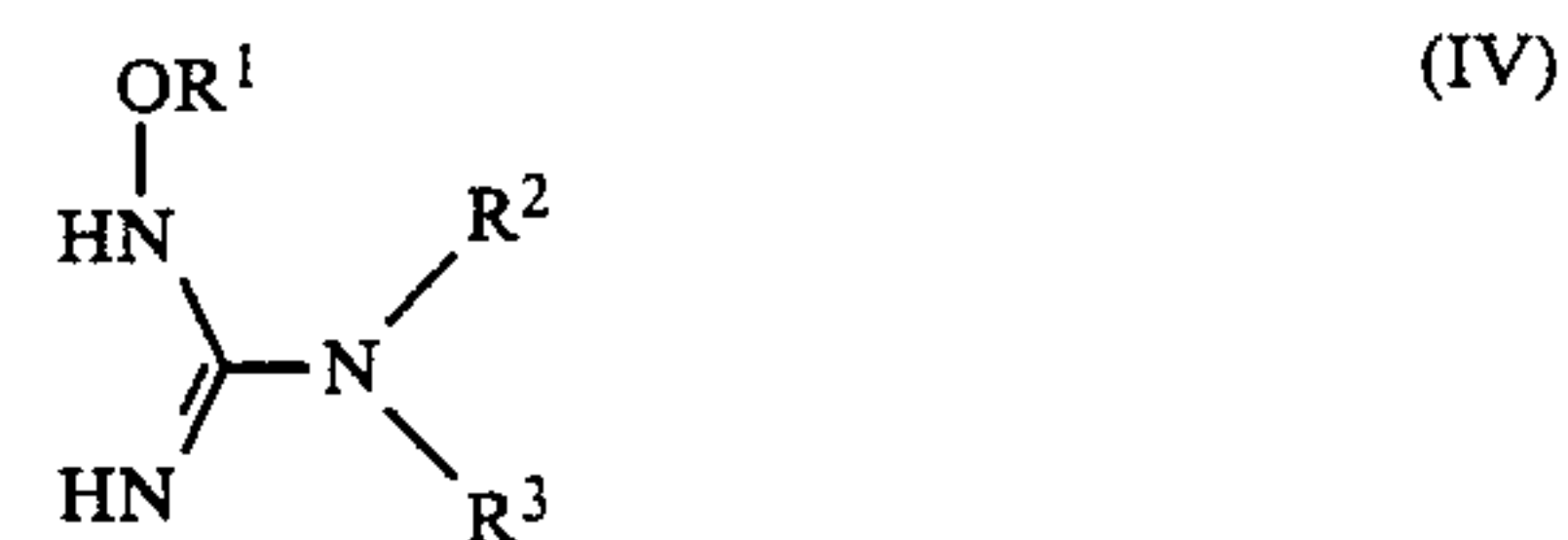


TABLE 4

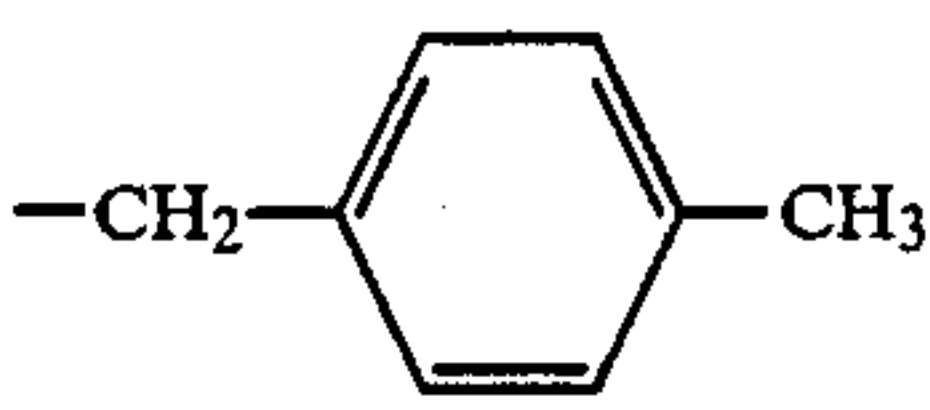
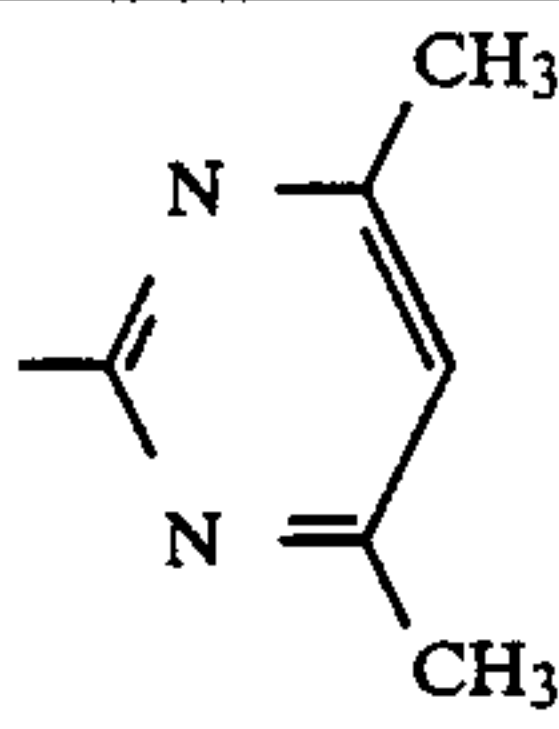
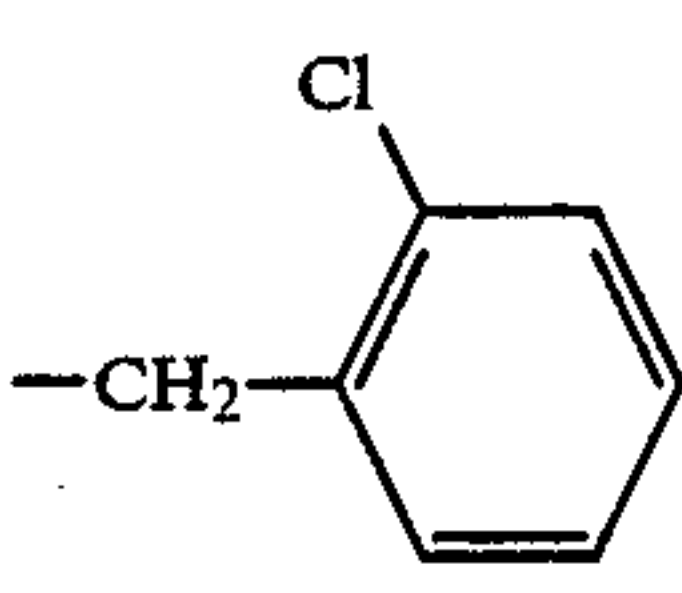
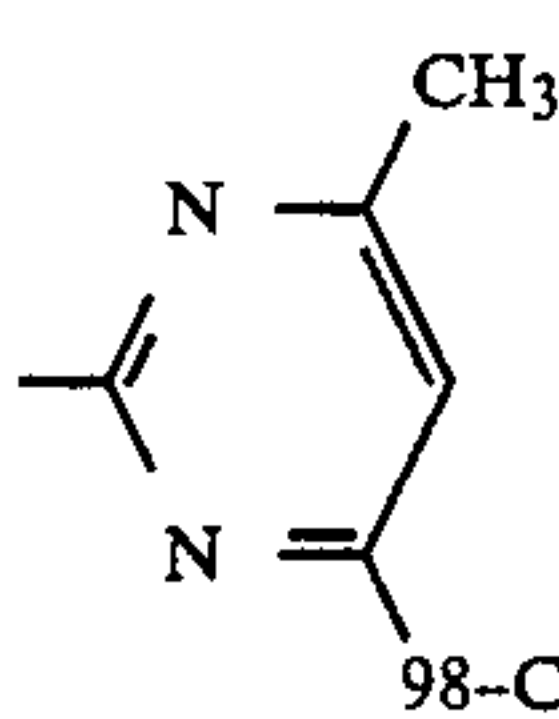
Example No.	R ¹	R ²	R ³	Melting point (°C.)
(IV-2)		H		85-86
(IV-3)		H		102-103

TABLE 4-continued

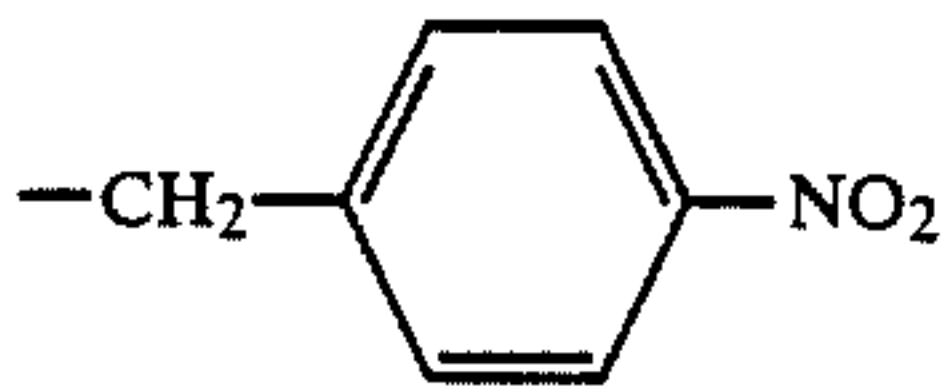
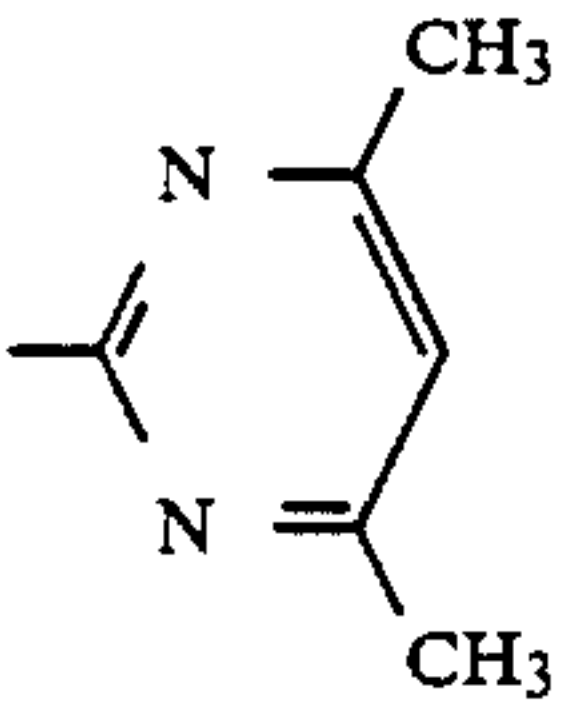
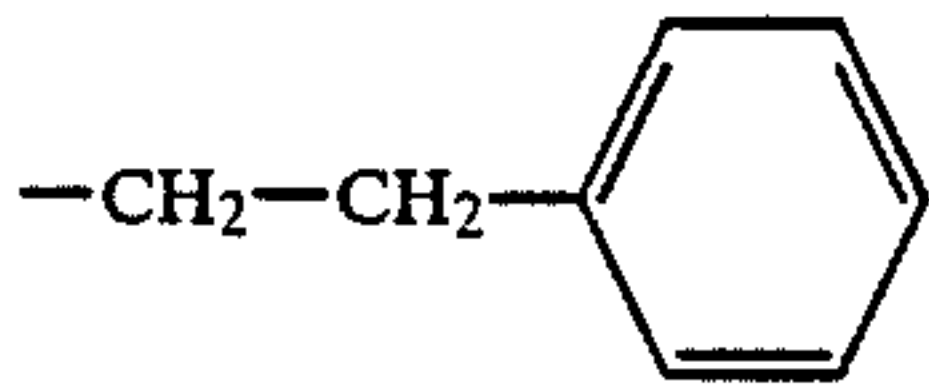
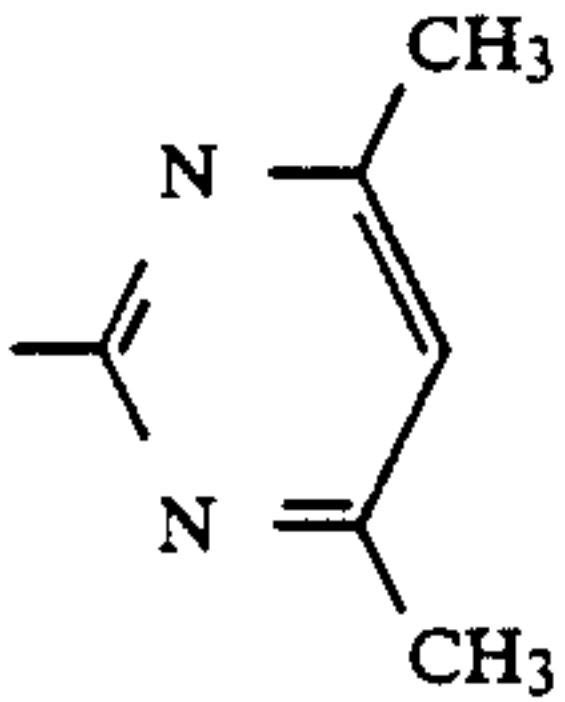
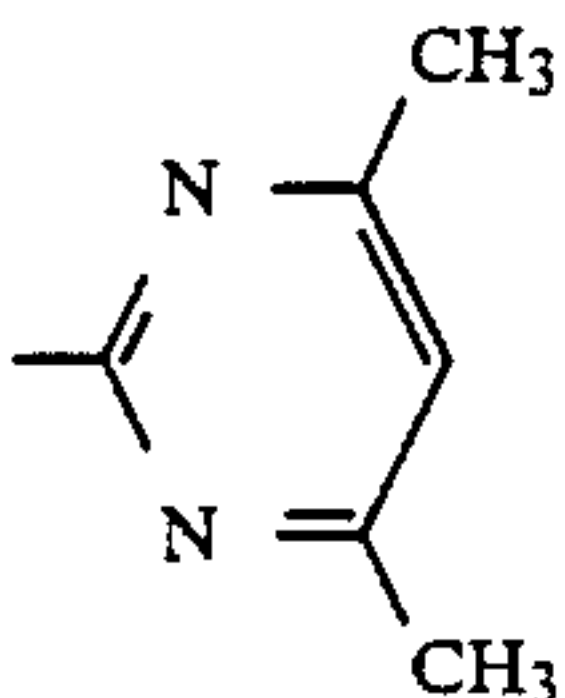
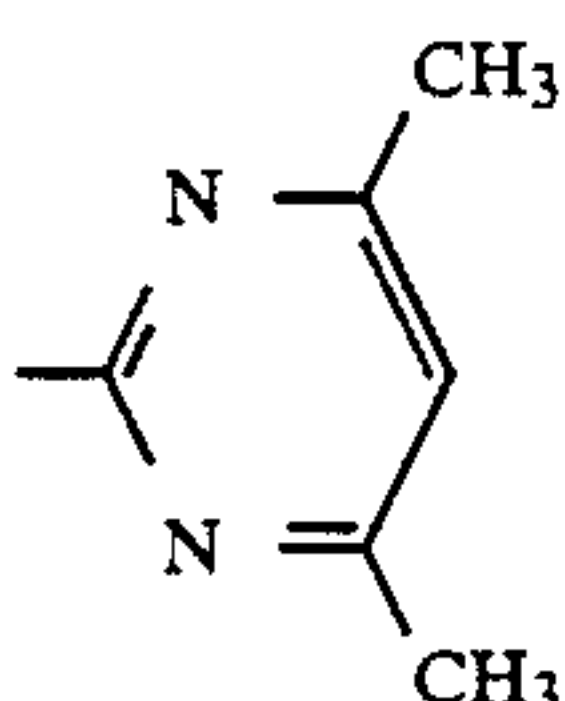
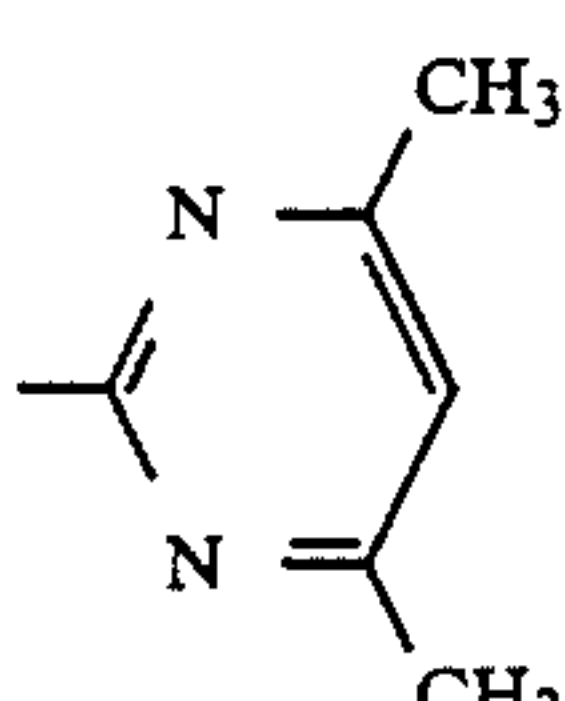
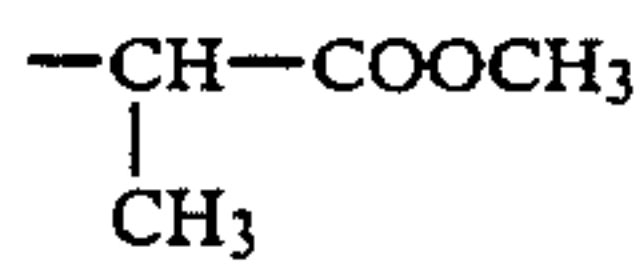
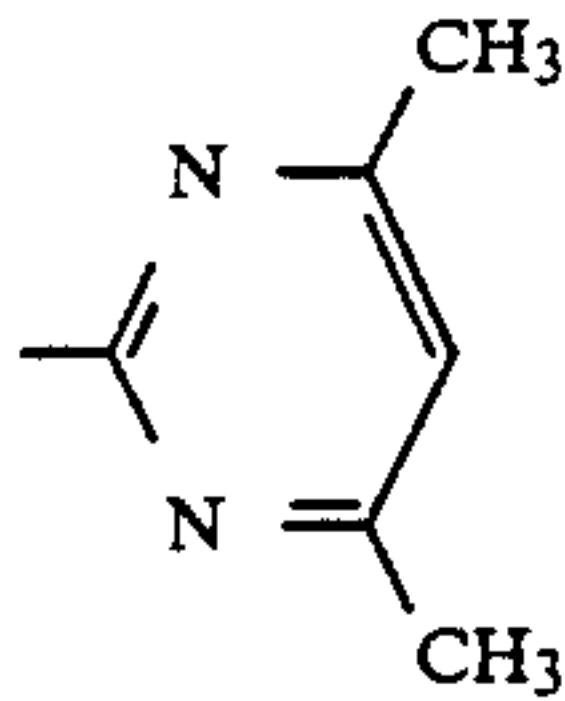
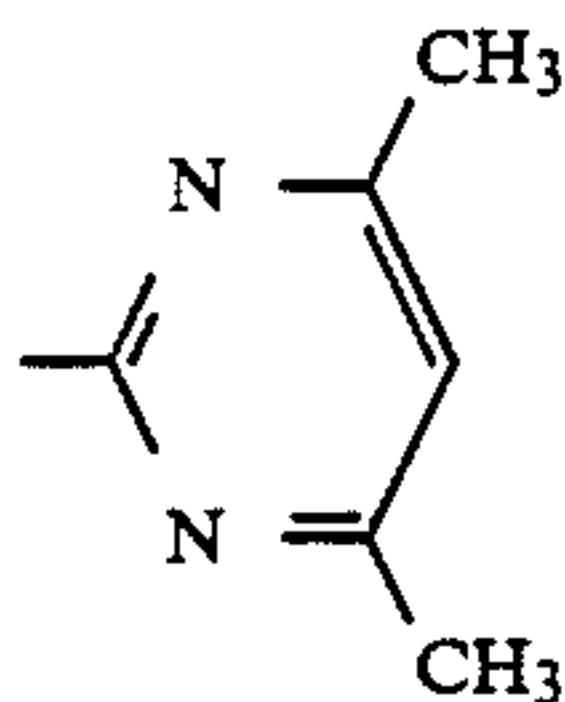
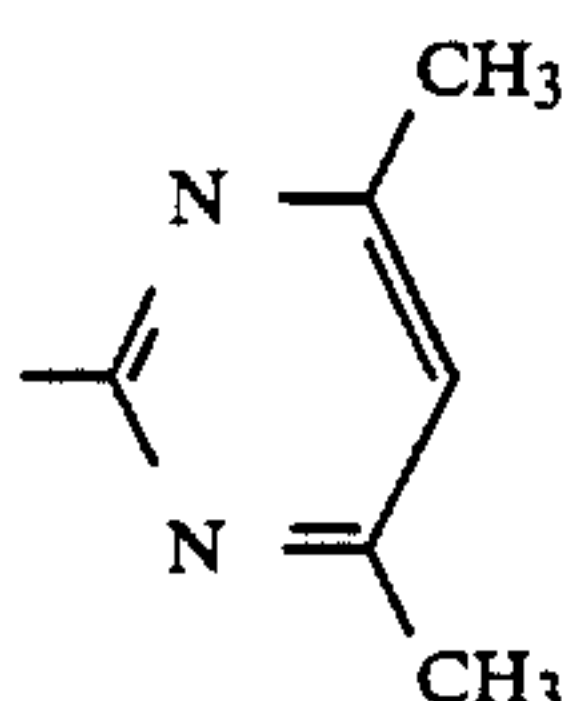
Example No.	R ¹	R ²	R ³	Melting point (°C.)
(IV-4)		H		170-172
(IV-5)		H		$n_D^{24.5} = 1.5776$
(IV-6)	$-\text{CH}_2-\text{CH}=\text{CH}_2$	H		103
(IV-7)	$-\text{C}_8\text{H}_{17}(-n)$	H		58
(IV-8)	$-\text{CH}_2-\text{COOC}_2\text{H}_5$	H		98''99
(IV-9)		H		147-148
(IV-10)	$-\text{CH}_3$	$-\text{CH}_3$		95
(IV-11)	$-\text{CH}_2\text{CH}(\text{CH}_3)_2$	H		52

TABLE 4-continued

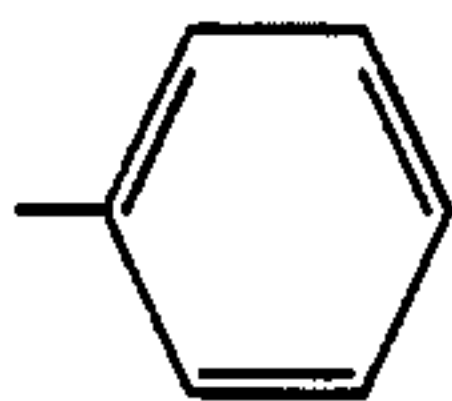
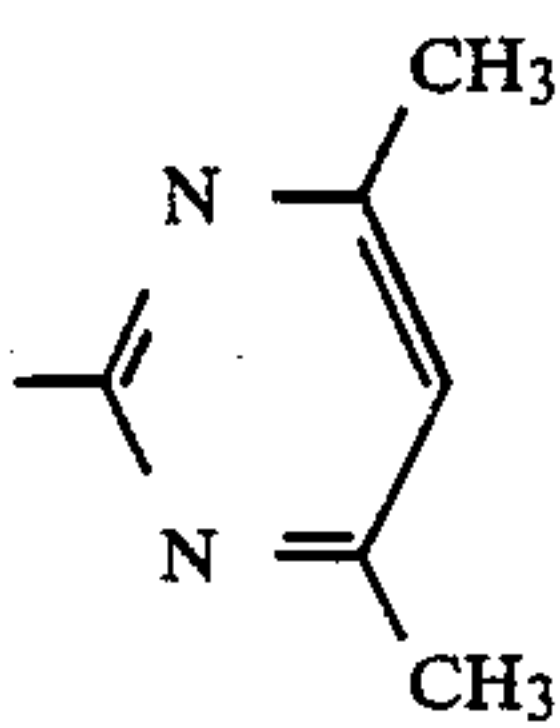
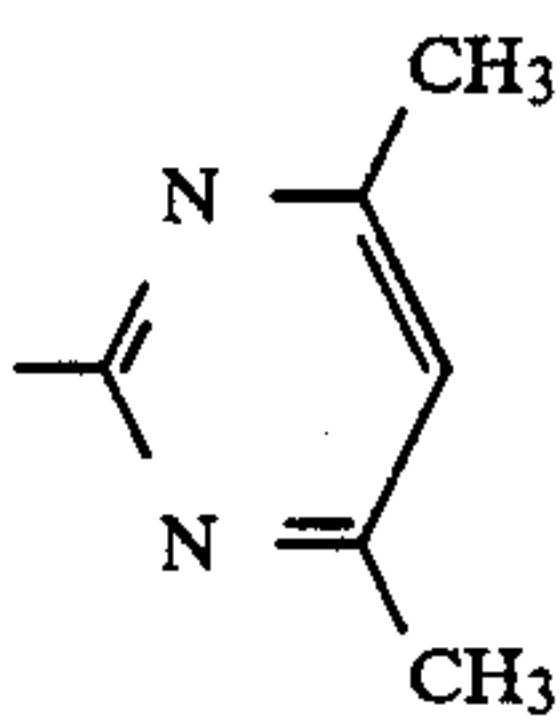
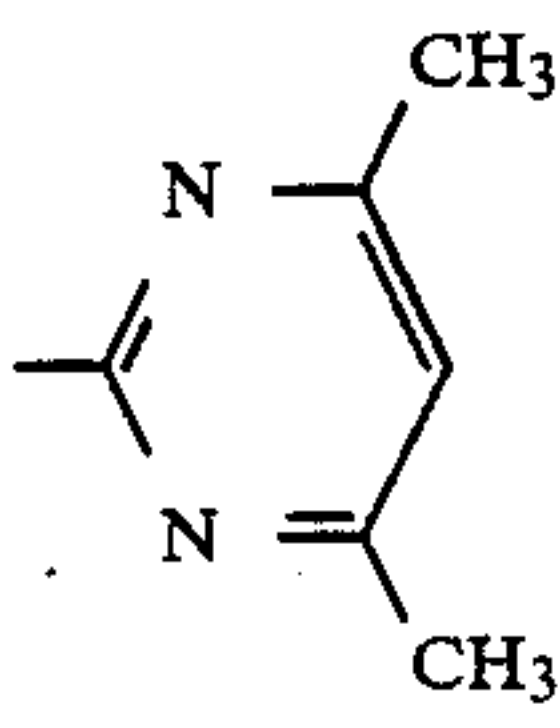
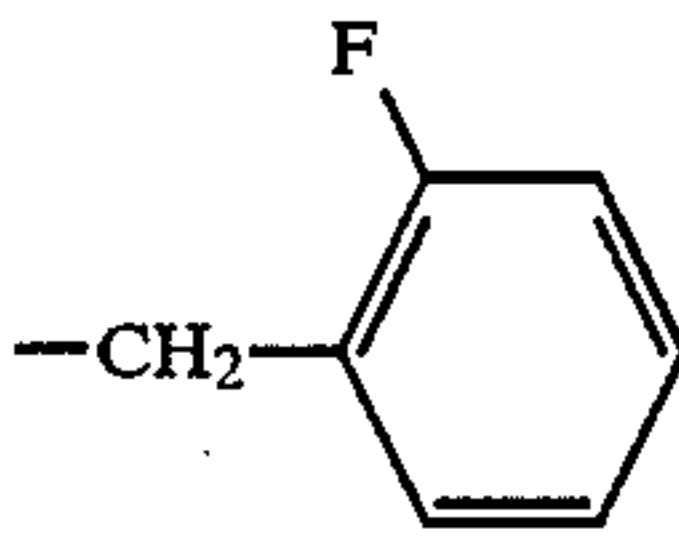
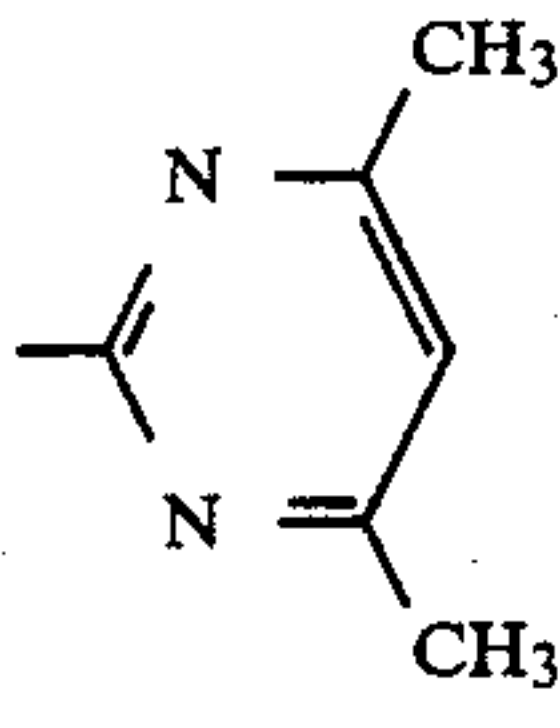
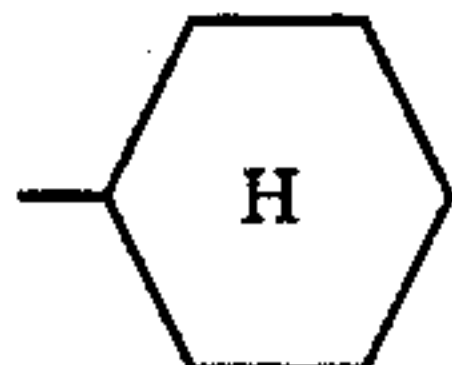
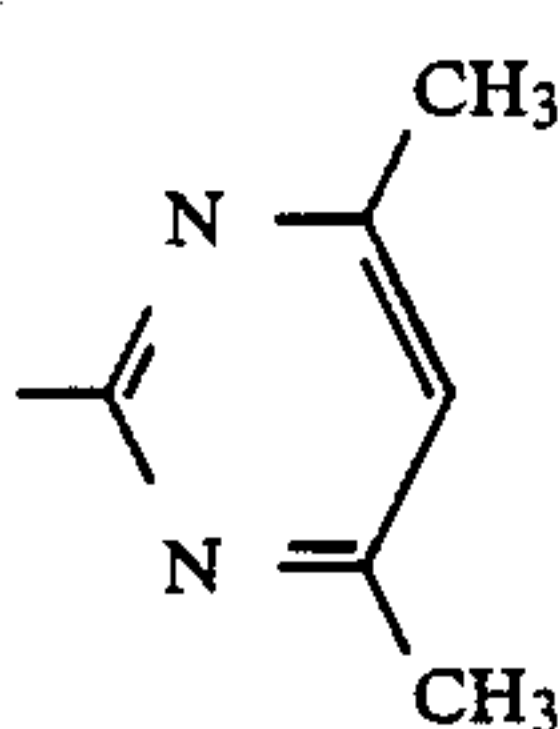
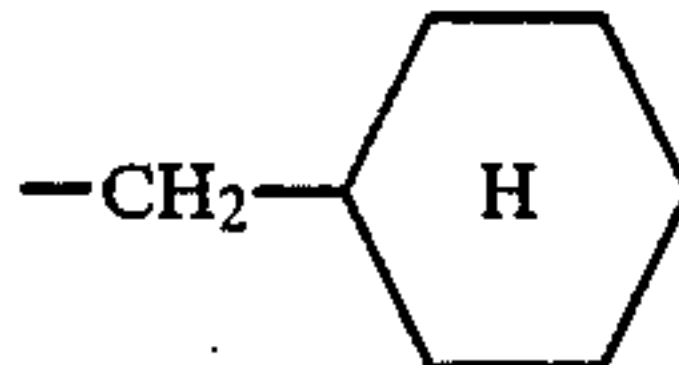
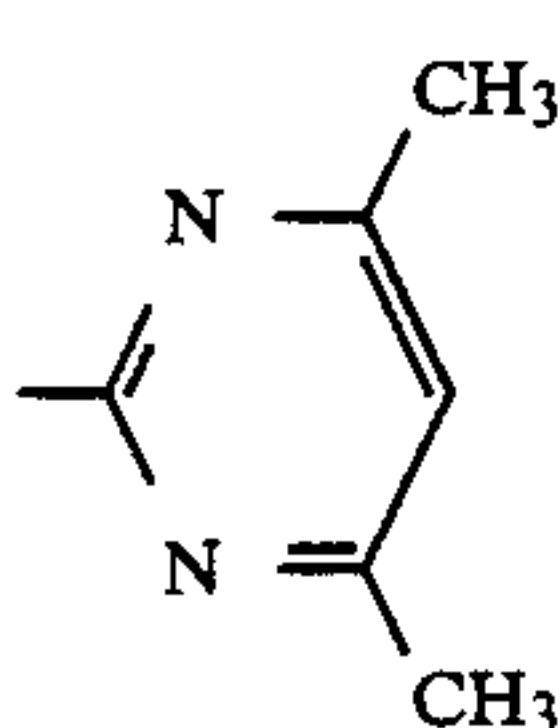
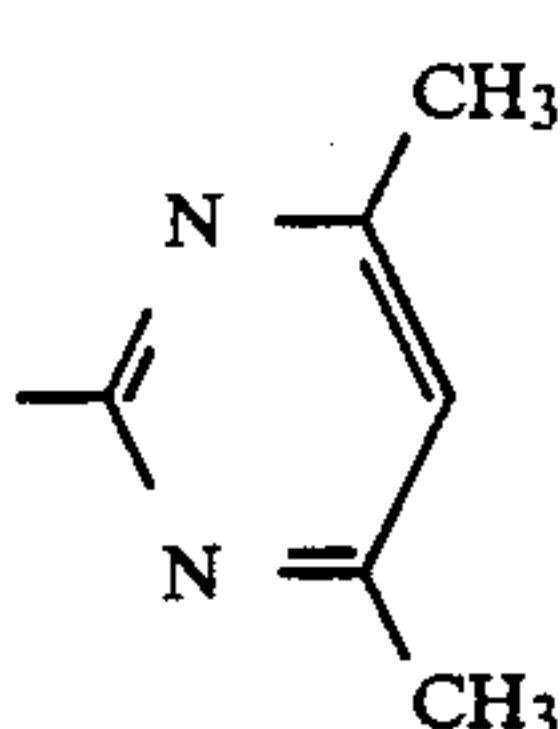
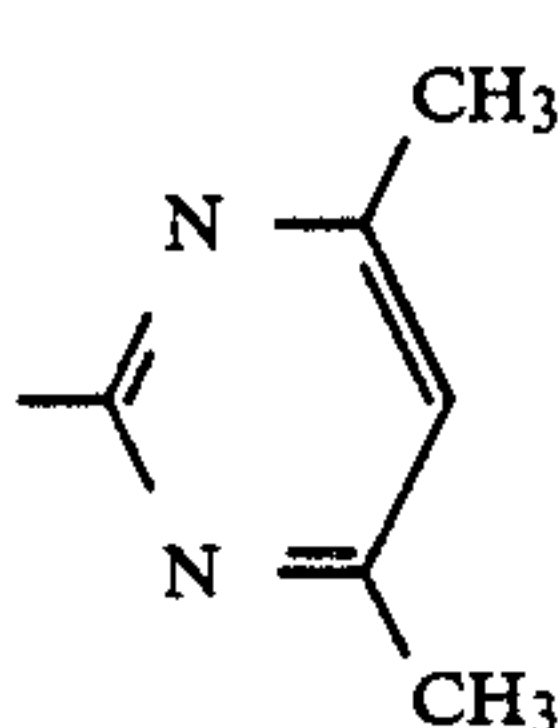
Example No.	R ¹	R ²	R ³	Melting point (°C.)
(IV-12)		H		189-192 (decomp.)
(IV-13)	$-\text{CH}_2\text{CH}_2\text{CH}_2\text{Cl}$	H		137
(IV-14)	$-\text{CH}_2\text{COOCH}_3$	H		148-149
(IV-15)		H		114-116
(IV-16)		H		
(IV-17)		H		
(IV-18)	$-\text{CH}_2-\text{CO}-\text{N}(\text{CH}_3)_2$	H		
(IV-19)	$-\text{CH}_2\text{OCH}_3$	H		

TABLE 4-continued

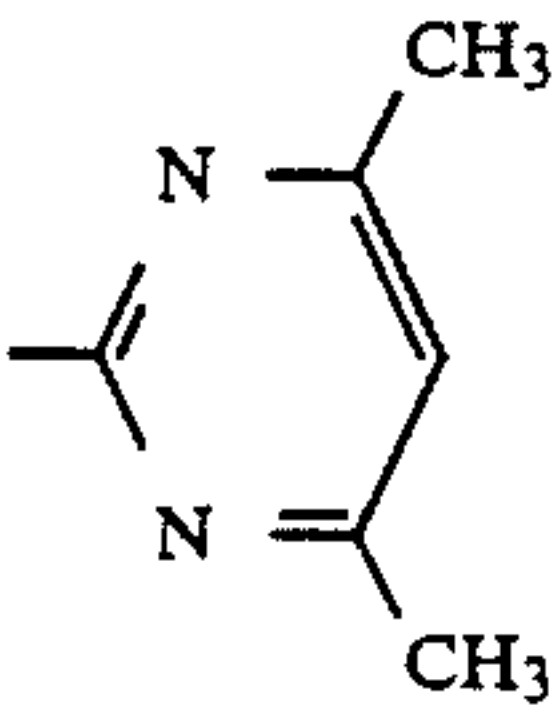
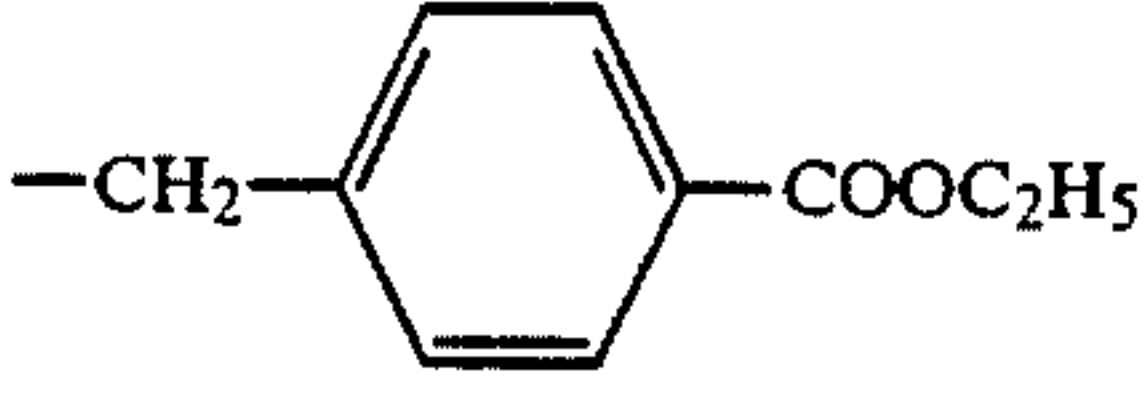
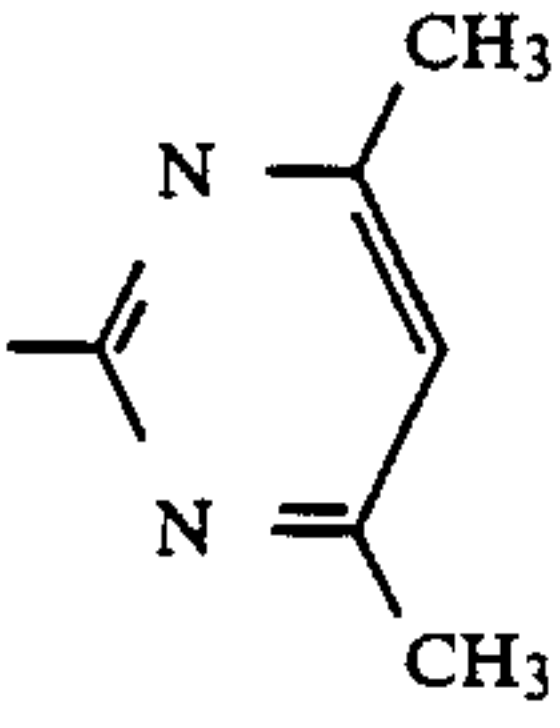
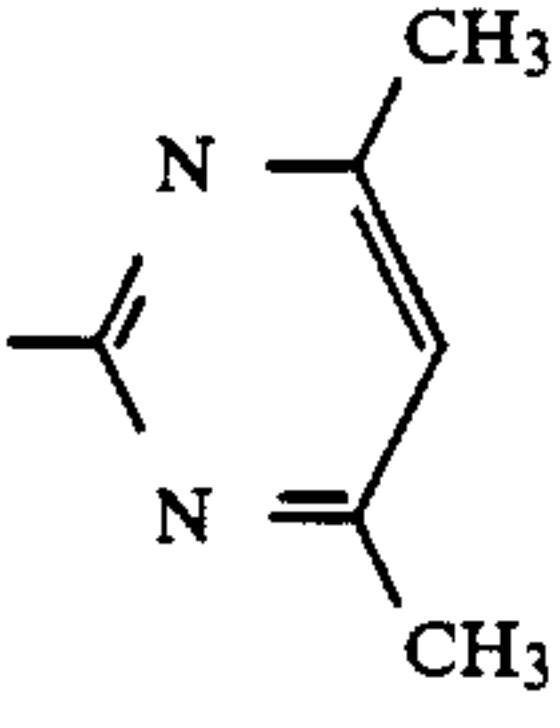
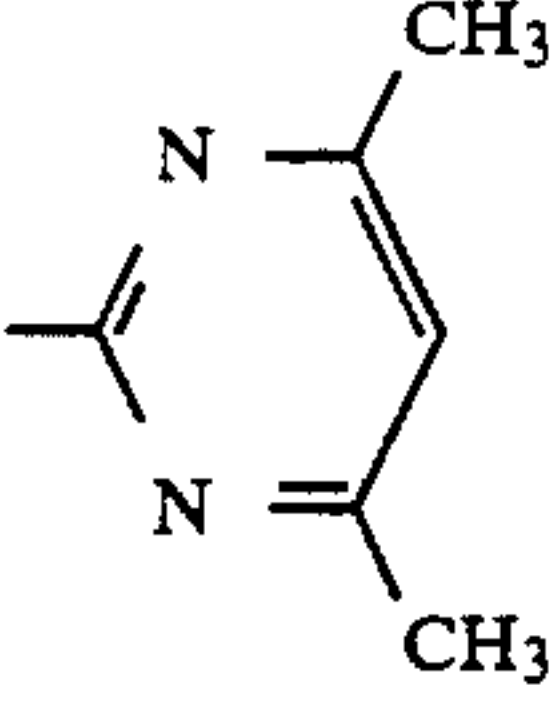
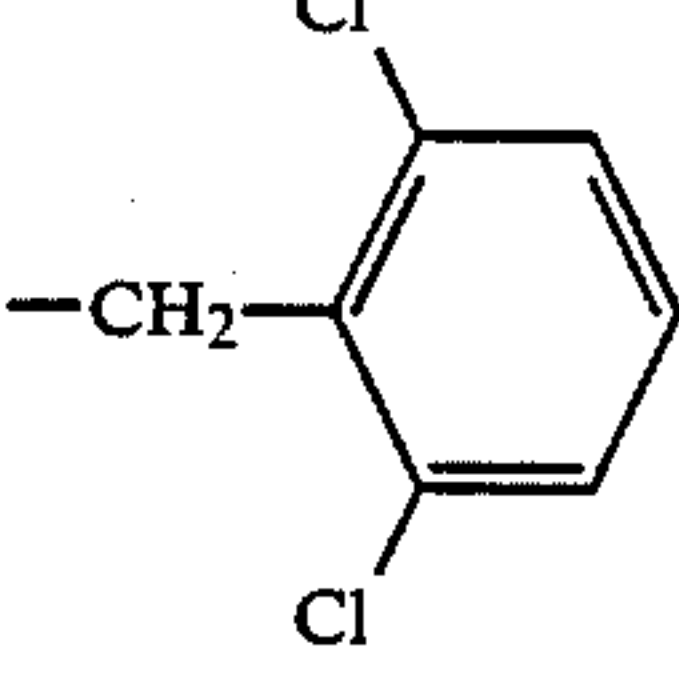
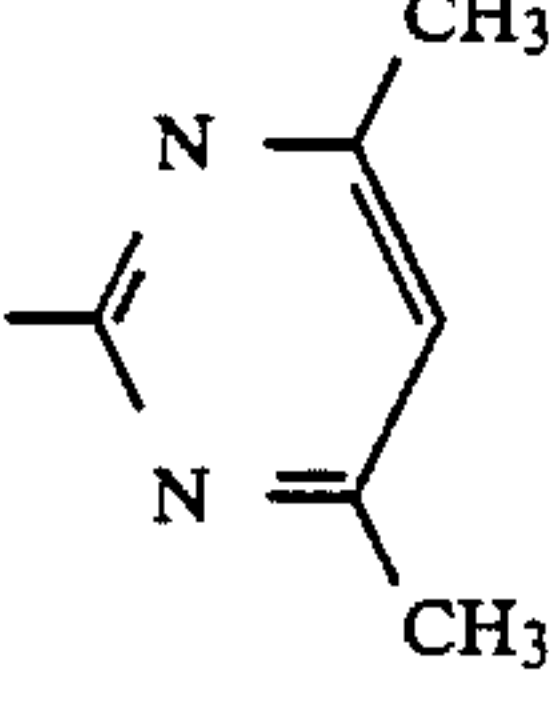
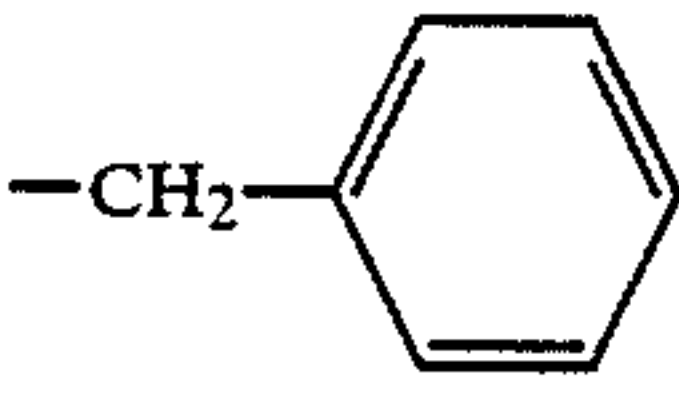
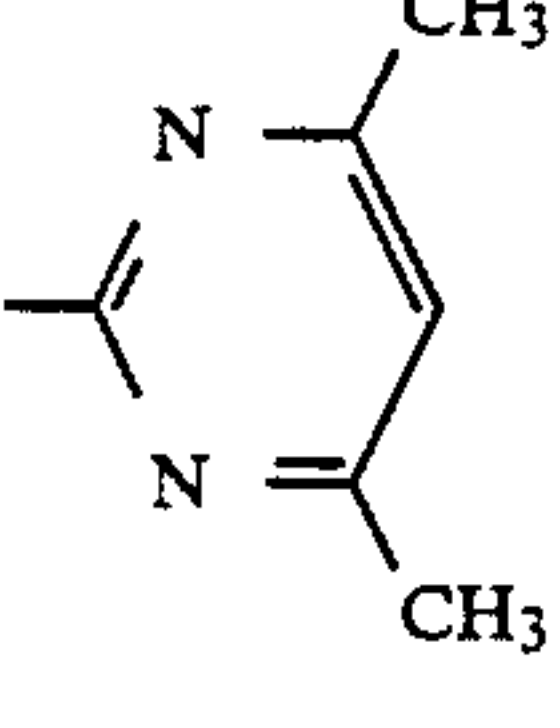
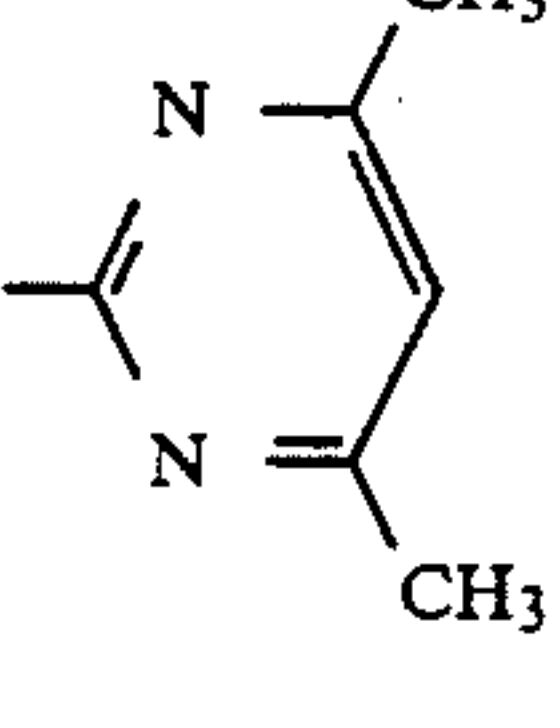
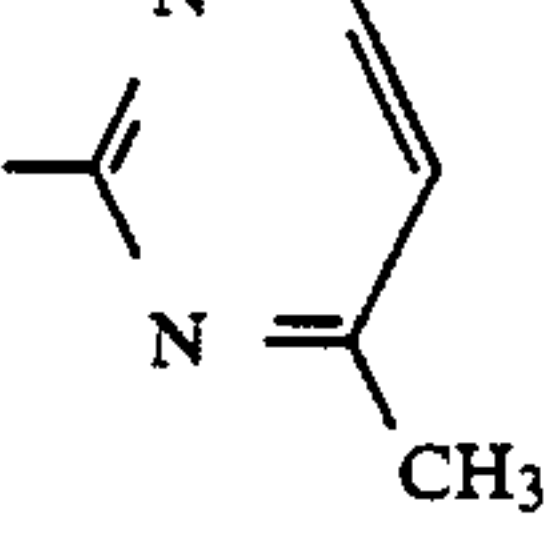
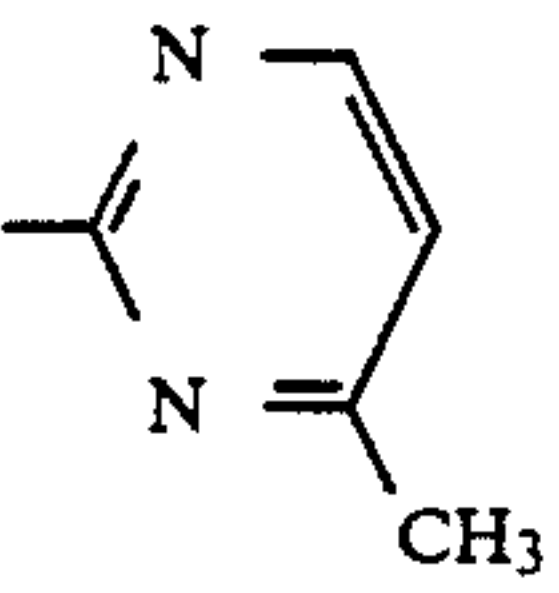
Example No.	R ¹	R ²	R ³	Melting point (°C.)
(IV-20)	$-\text{CH}_2\text{SCH}_3$	H		
(IV-21)		H		138
(IV-22)	$-\text{CH}_2\text{CH}_2\text{OCH}_3$	H		
(IV-23)	$-\text{CH}_2\text{CF}_3$	H		
(IV-24)		H		152
(IV-25)		H		102
(IV-26)	$-\text{CH}_2-\text{COOCH}(\text{CH}_3)_2$	H		112
(IV-27)	$-\text{CH}_3$	H		152
(IV-28)	$-\text{C}_2\text{H}_5$	H		95

TABLE 4-continued

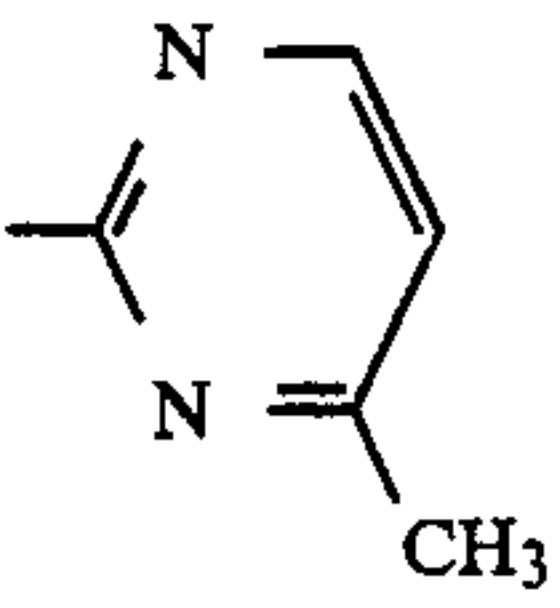
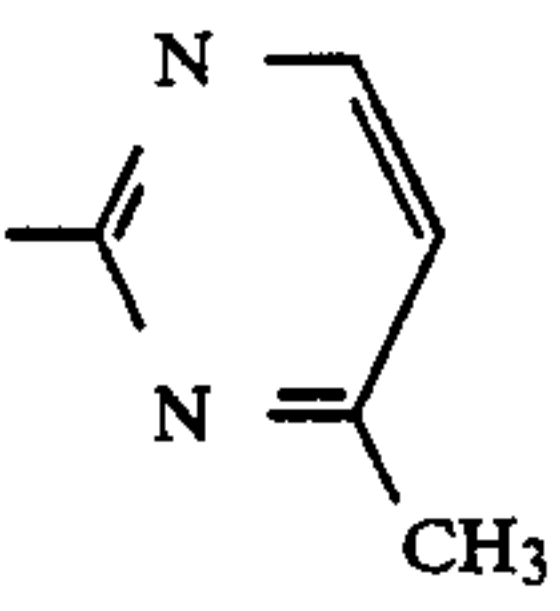
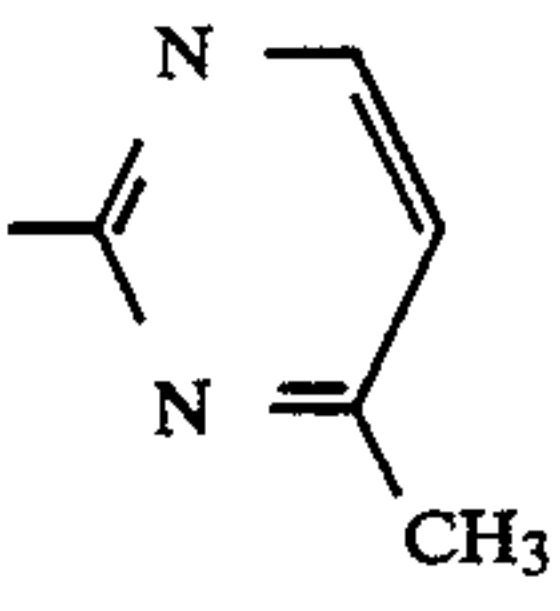
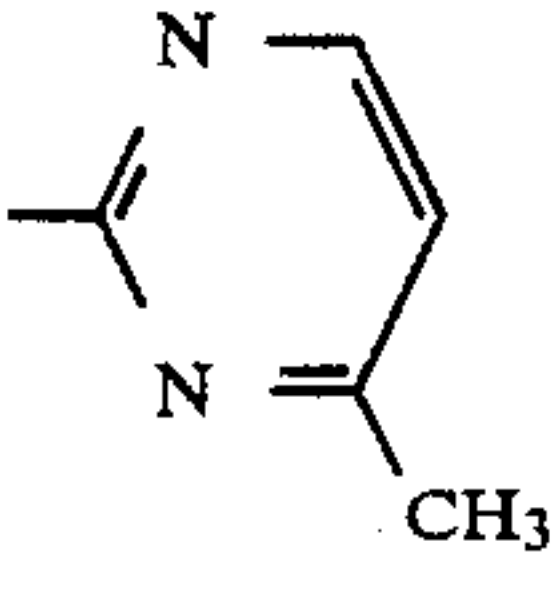
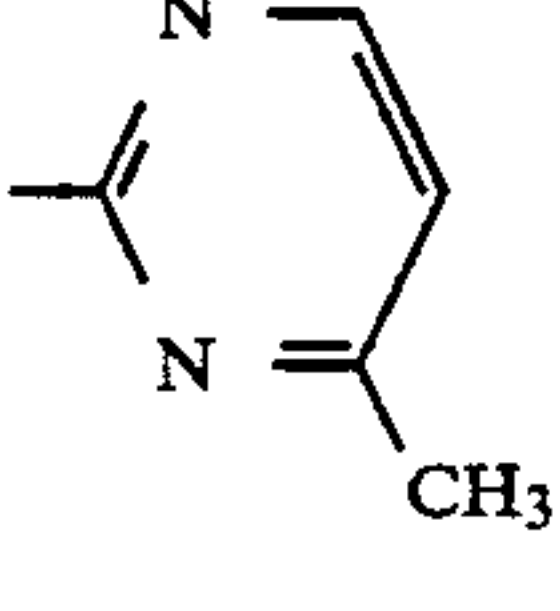
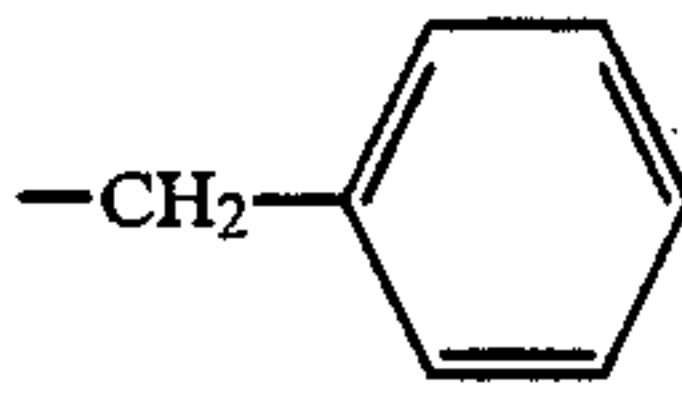
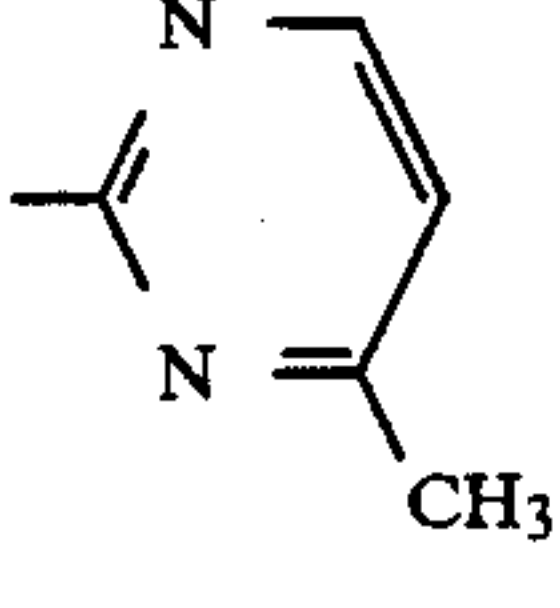
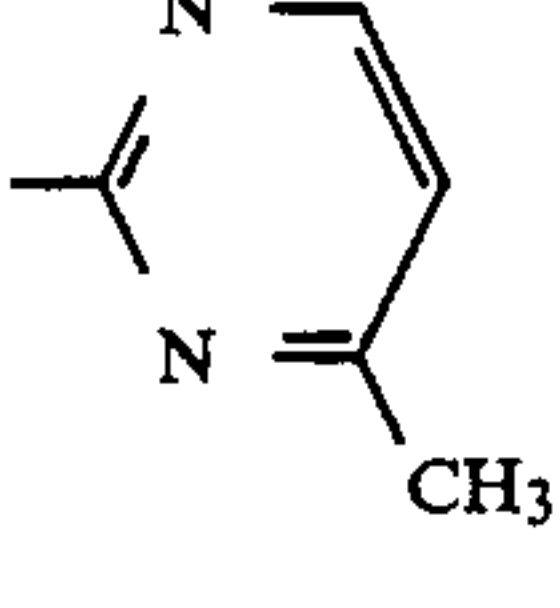
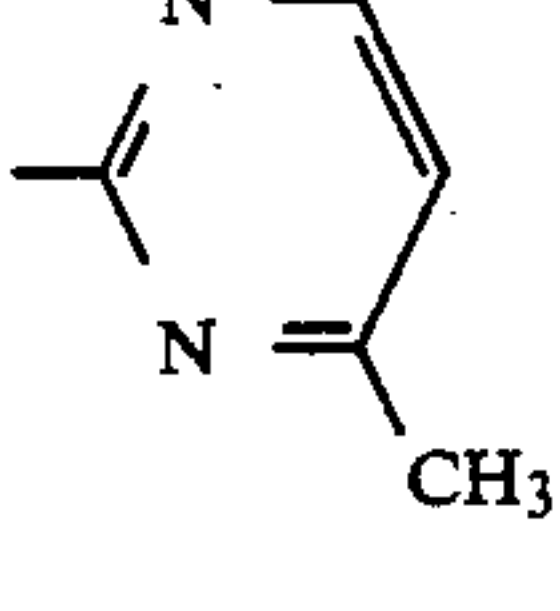
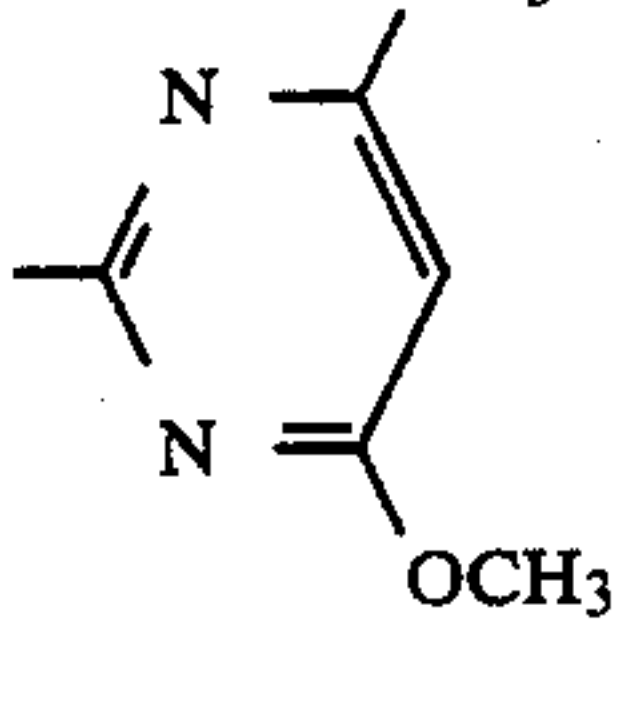
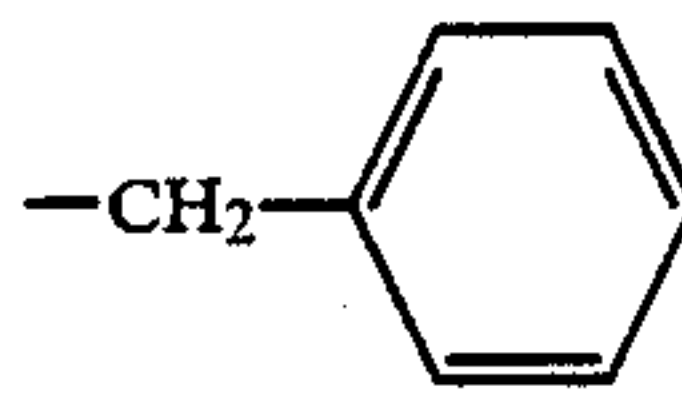
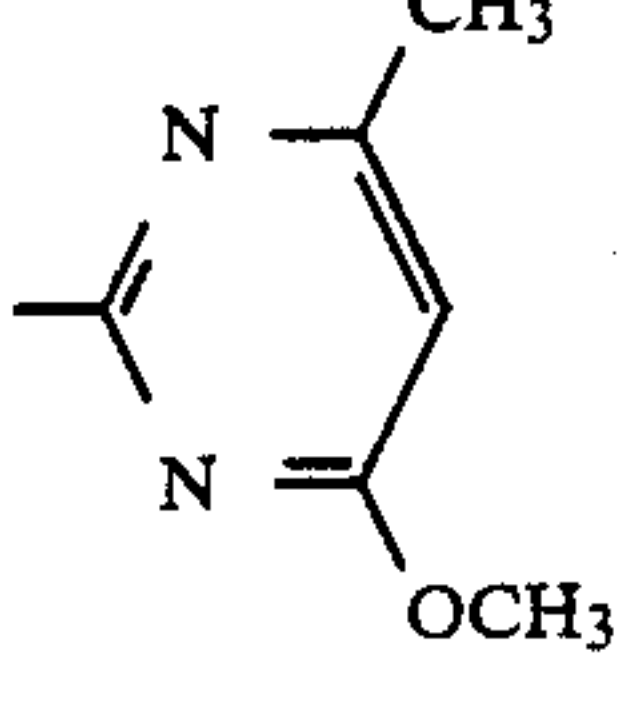
Example No.	R ¹	R ²	R ³	Melting point (°C.)
(IV-29)	-C ₃ H ₇ (-n)	H		
(IV-30)	-CH(CH ₃) ₂	H		
(IV-31)	-C ₄ H ₉ (-n)	H		
(IV-32)	-CH ₂ CH(CH ₃) ₂	H		
(IV-33)	-CH ₂ -CH=CH ₂	H		
(IV-34)		H		150
(IV-35)	-CH ₂ -COOC ₂ H ₅	H		
(IV-36)	-CH ₃	H		98
(IV-37)	-CH ₃	H		126
(IV-38)		H		(amorphous)

TABLE 4-continued

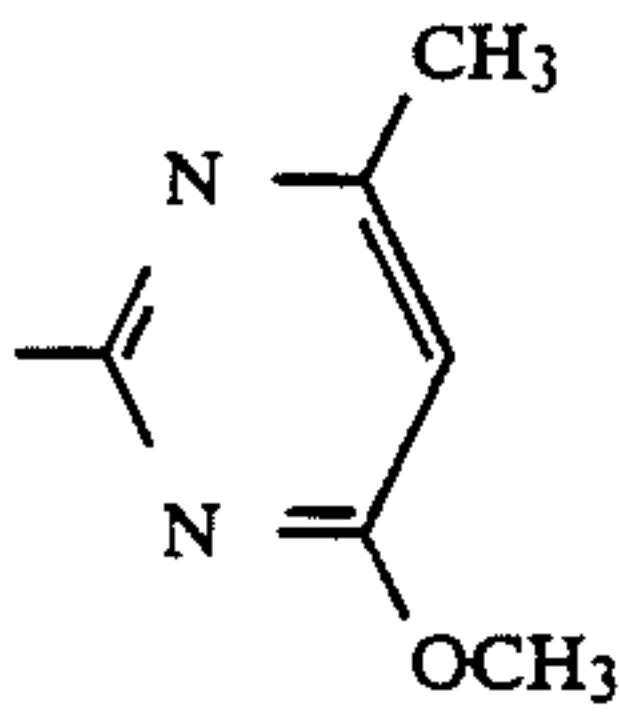
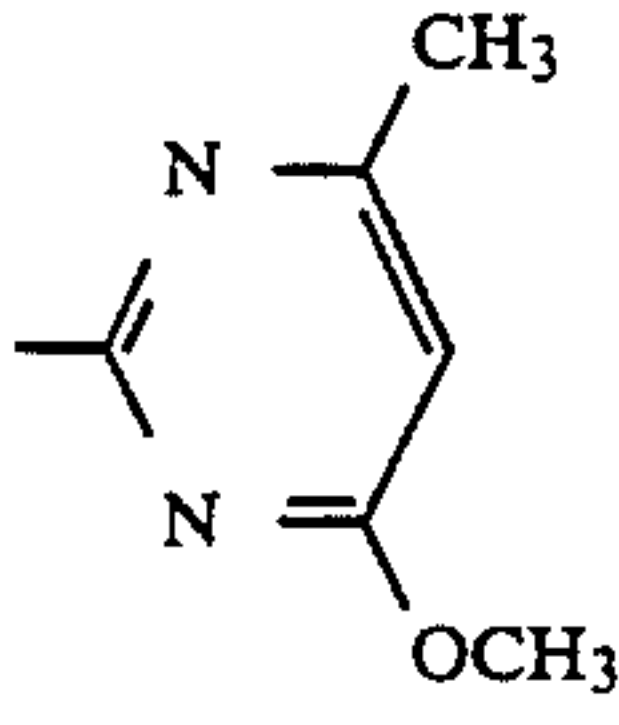
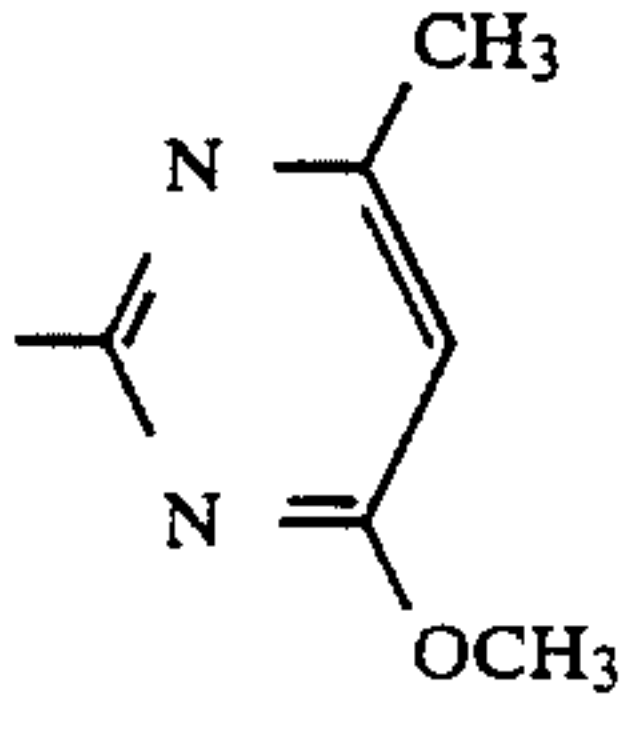
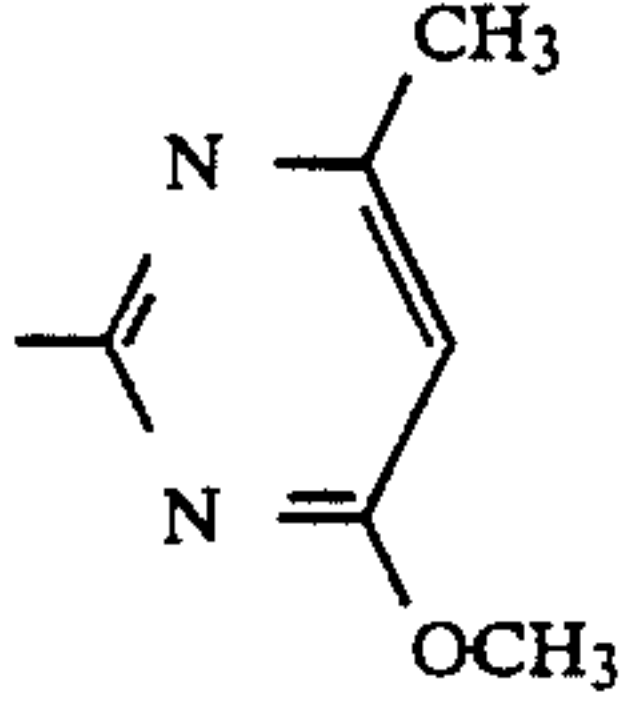
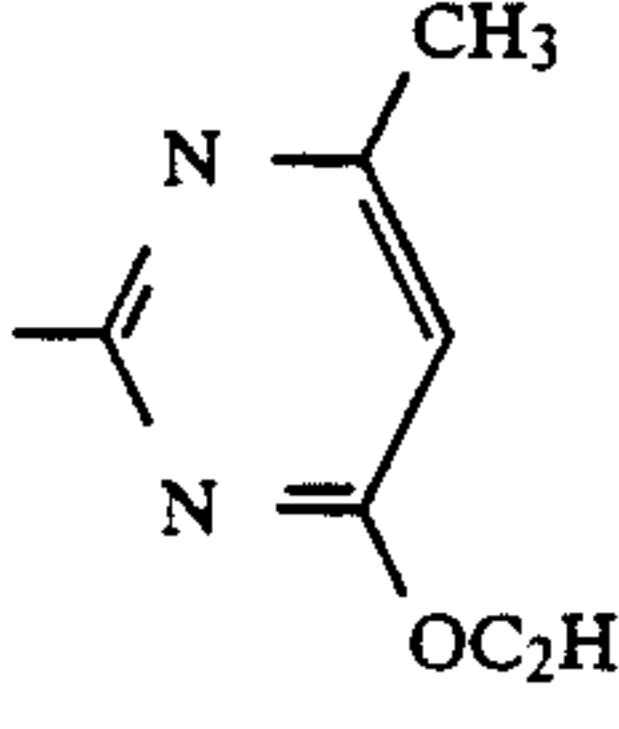
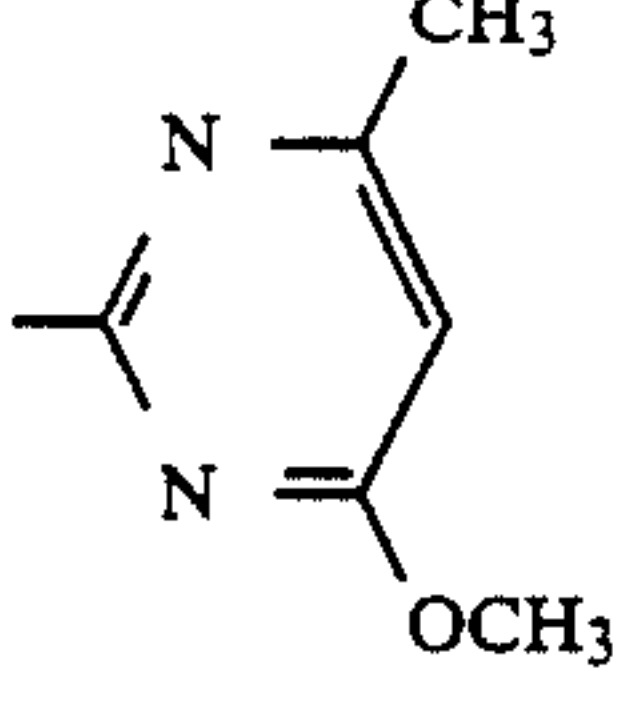
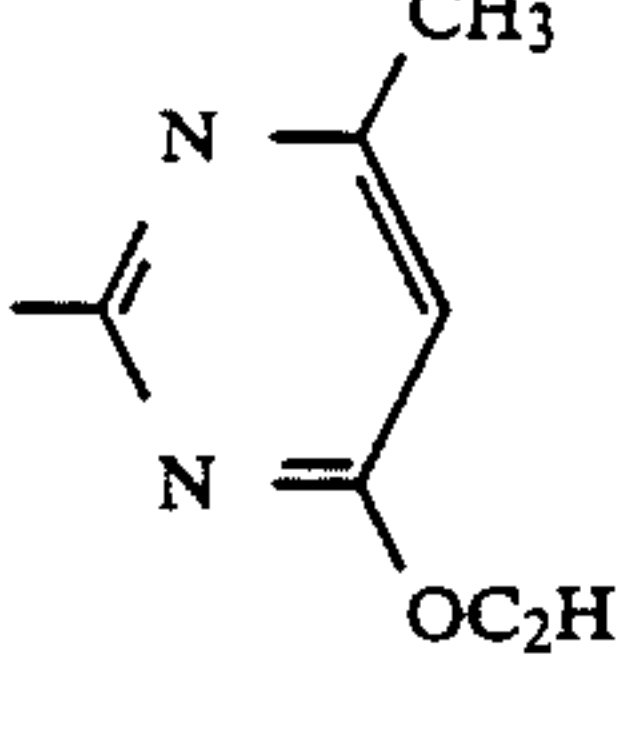
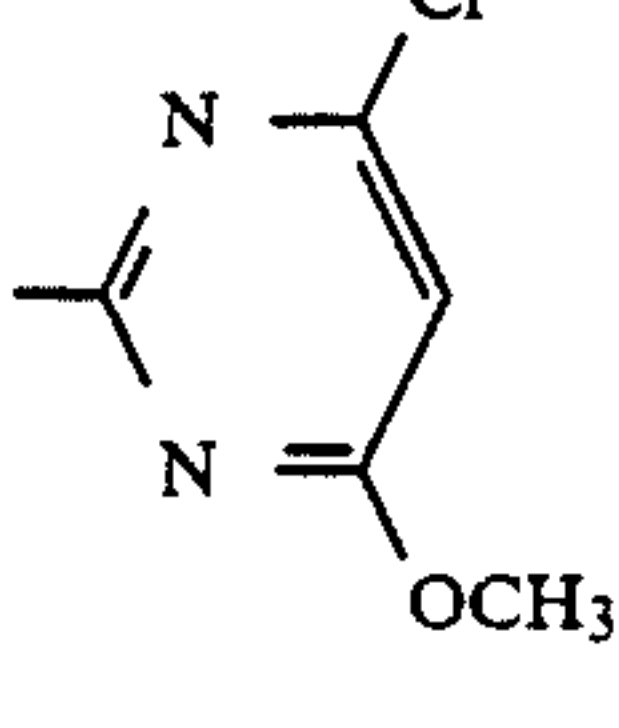
Example No.	R ¹	R ²	R ³	Melting point (°C.)
(IV-39)	—C ₂ H ₅	H		(amorphous)
(IV-40)	—C ₃ H ₇	H		
(IV-41)	—CH ₂ CH(CH ₃) ₂	H		
(IV-42)	—CH ₂ —CH=CH ₂	H		
(IV-43)	—CH ₃	H		
(IV-44)	—CH ₃	CH ₃		135
(IV-45)	—CH ₃	C ₂ H ₅		
(IV-46)	—CH ₃	H		112

TABLE 4-continued

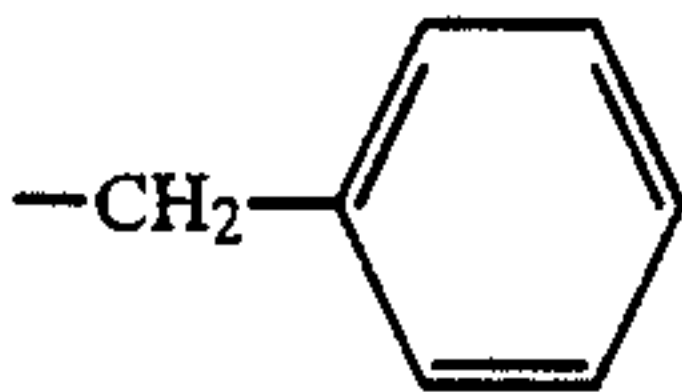
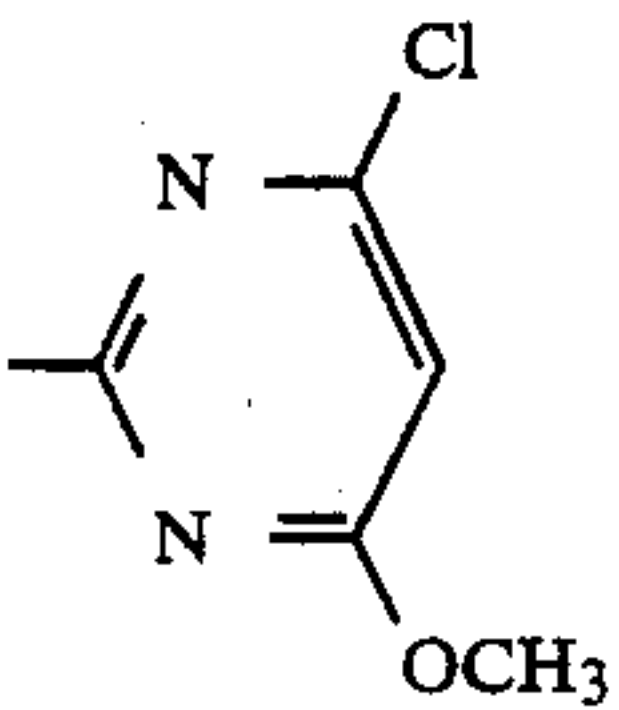
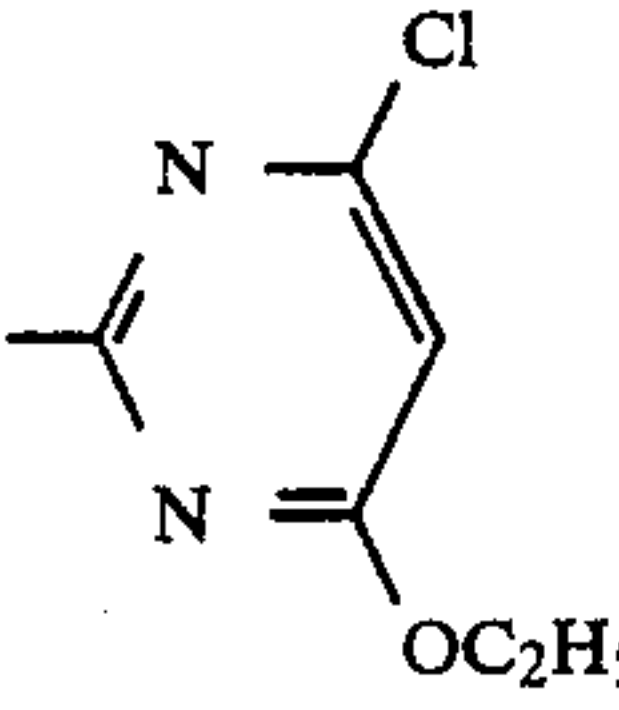
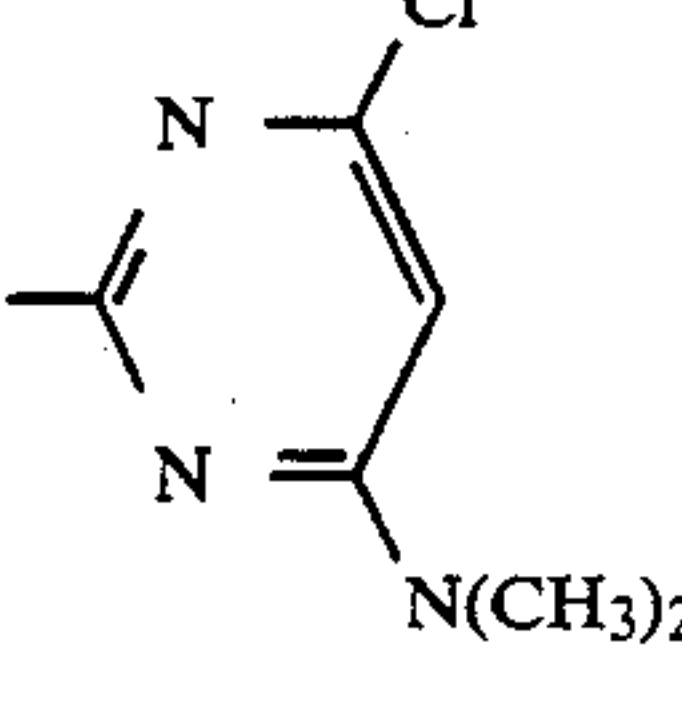
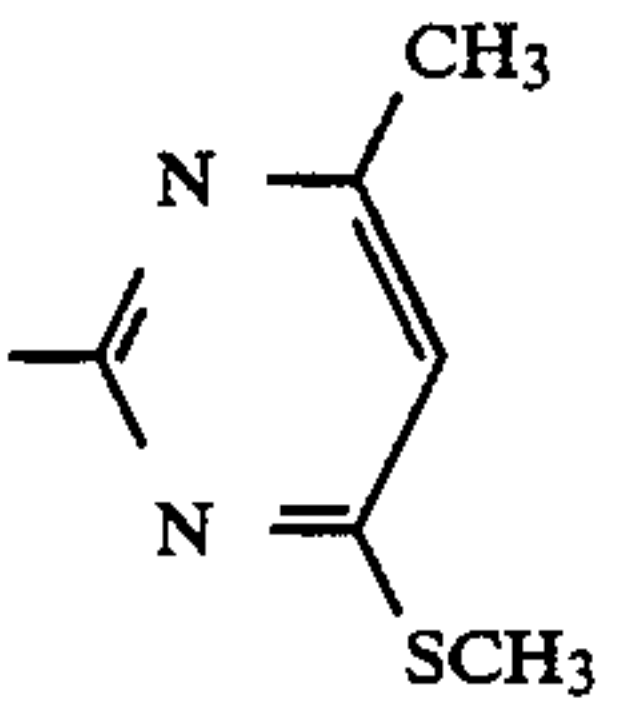
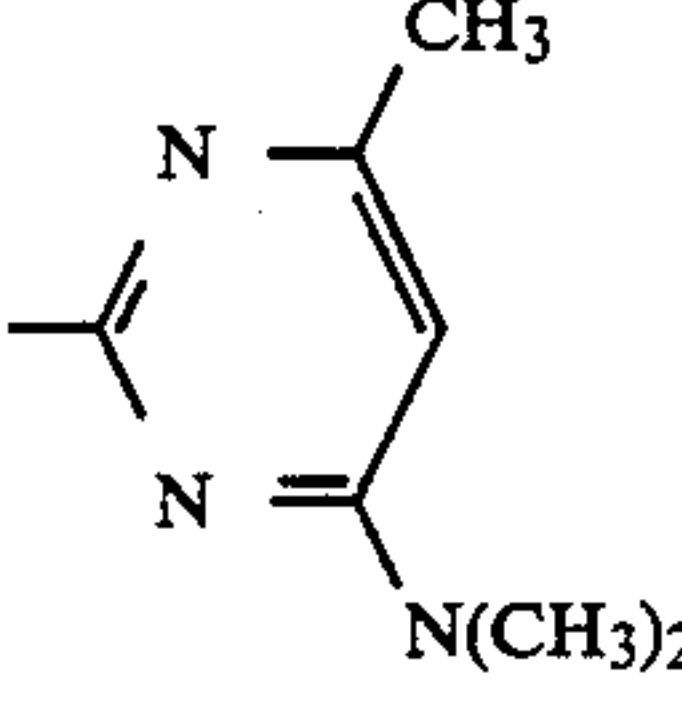
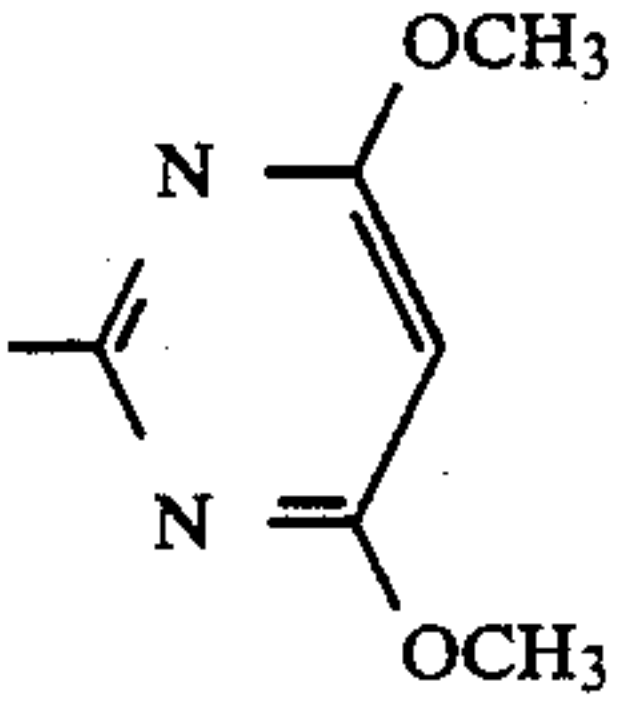
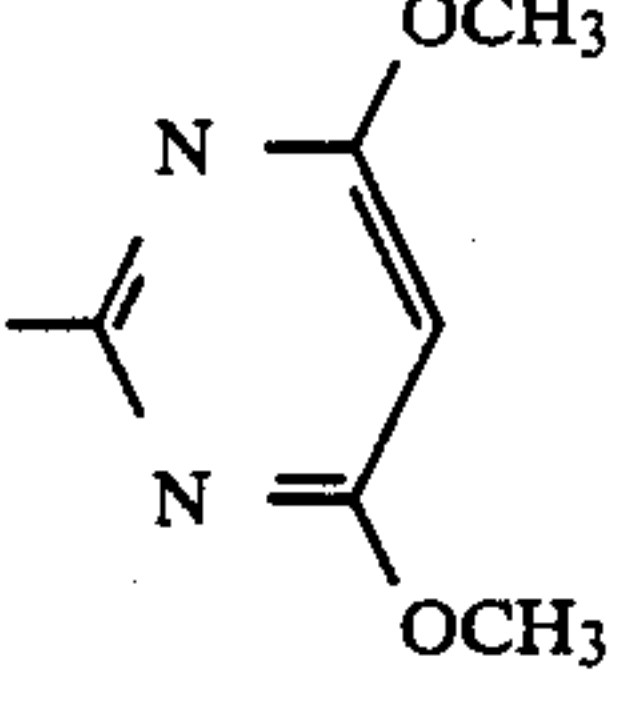
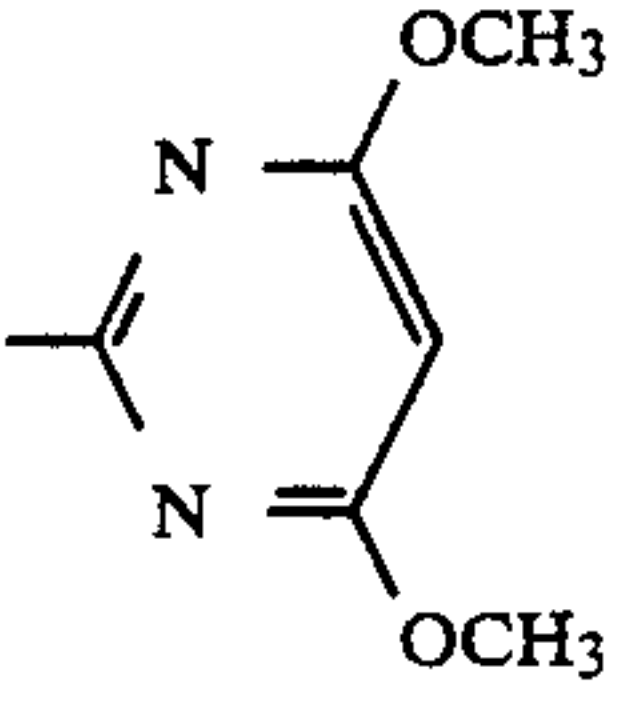
Example No.	R ¹	R ²	R ³	Melting point (°C.)
(IV-47)		H		
(IV-48)	-CH ₃	H		
(IV-49)	-CH ₃	H		
(IV-50)	-CH ₃	H		
(IV-51)	-CH ₃	H		
(IV-52)	-CH ₃	H		122
(IV-53)	-C ₂ H ₅	H		
(IV-54)	-C ₃ H ₇	H		

TABLE 4-continued

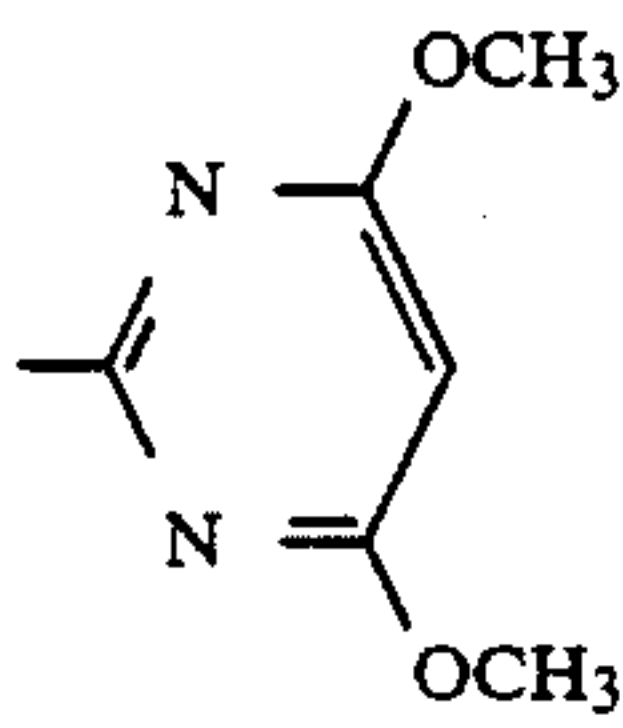
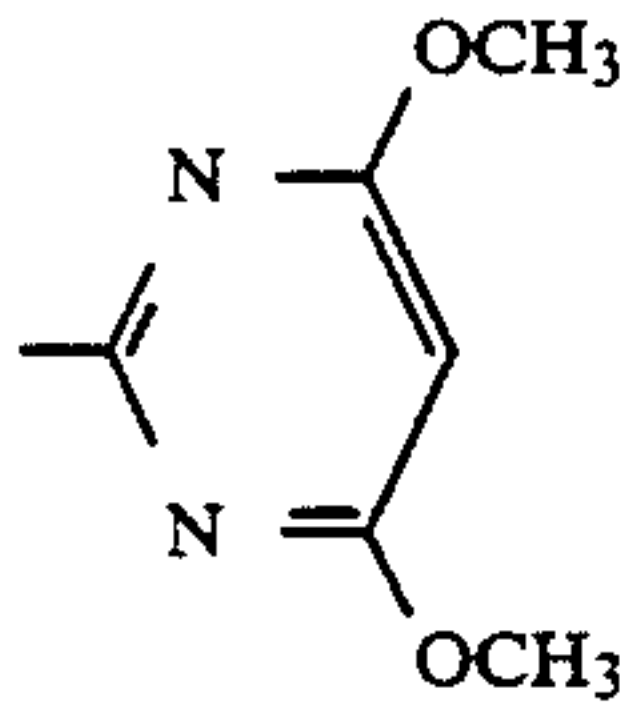
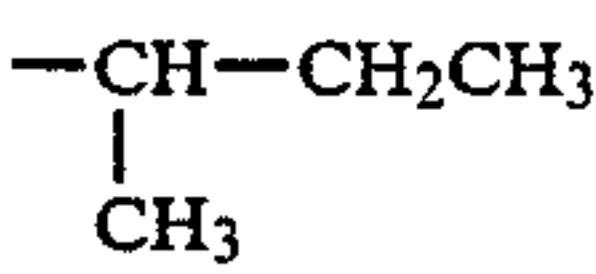
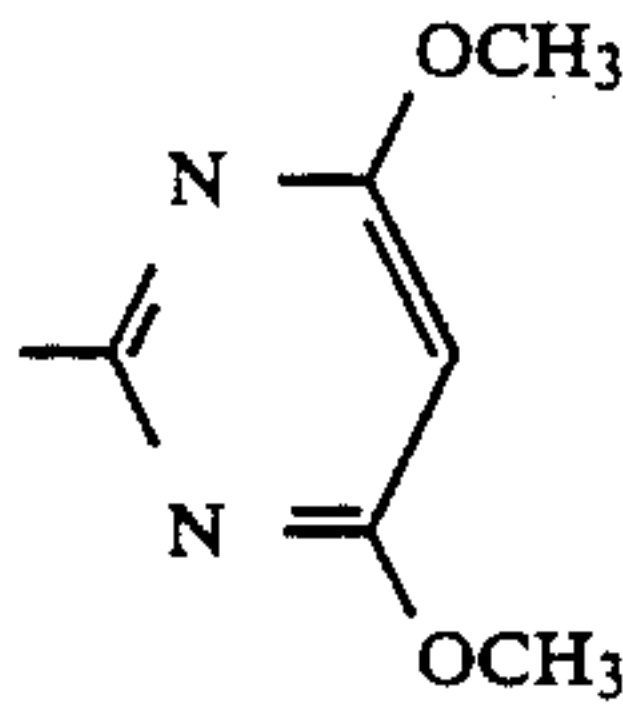
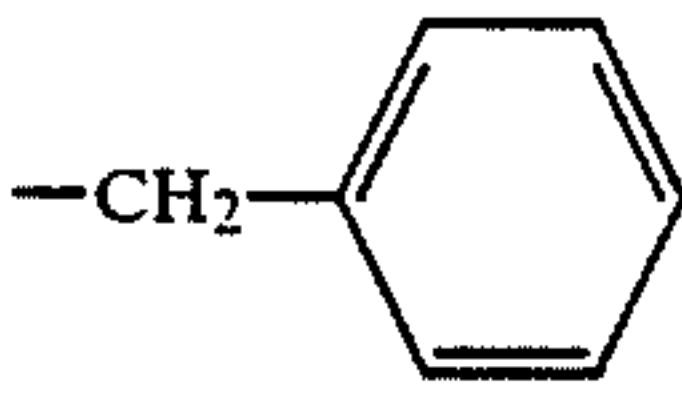
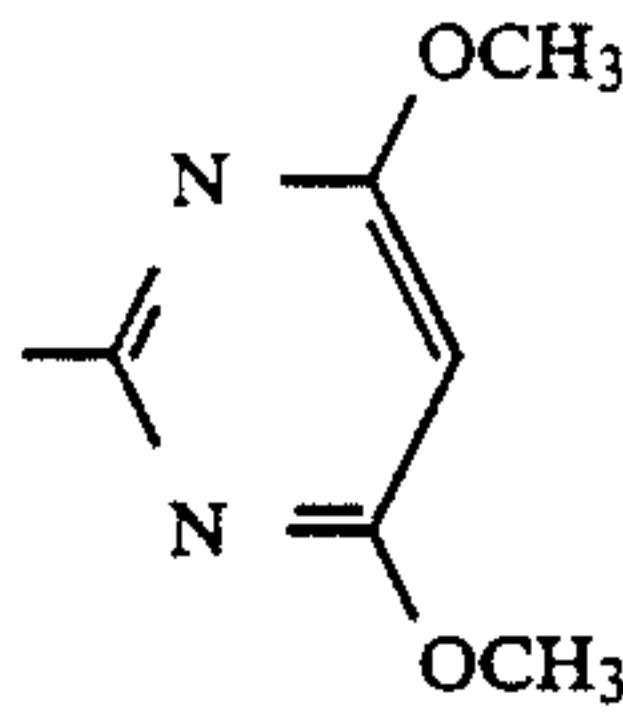
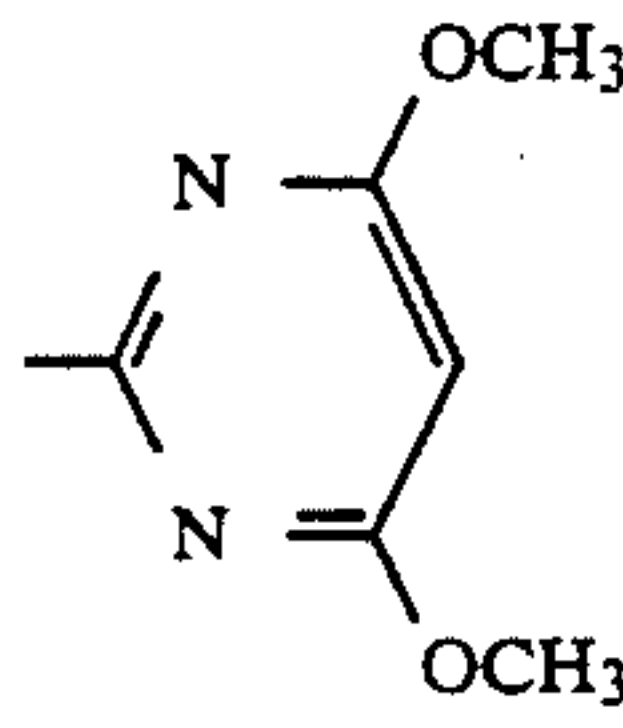
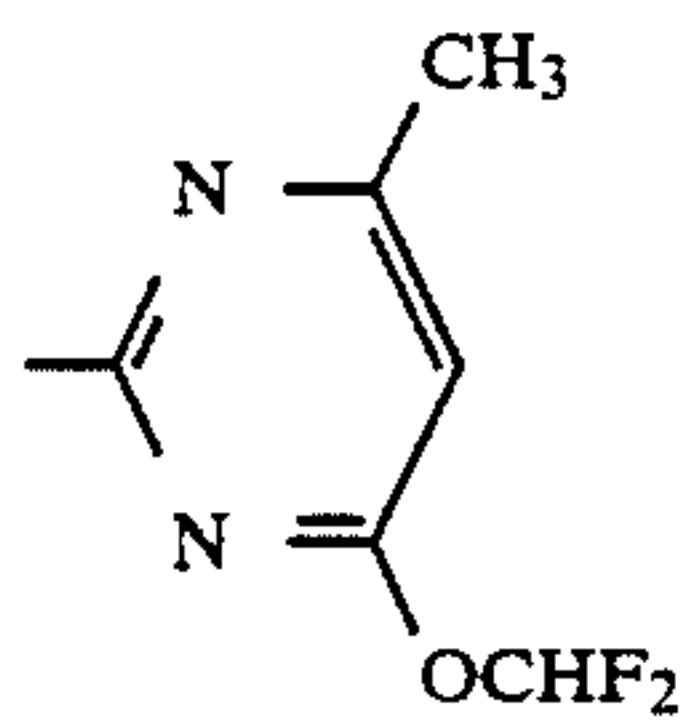
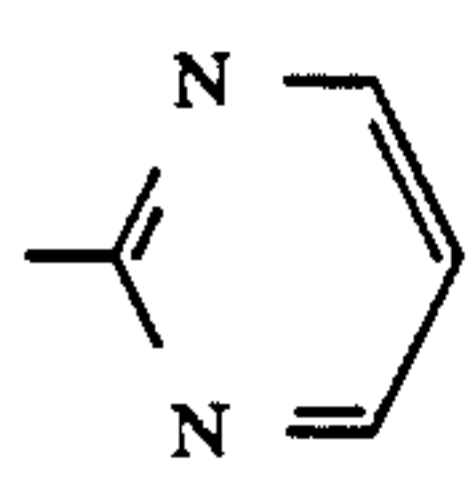
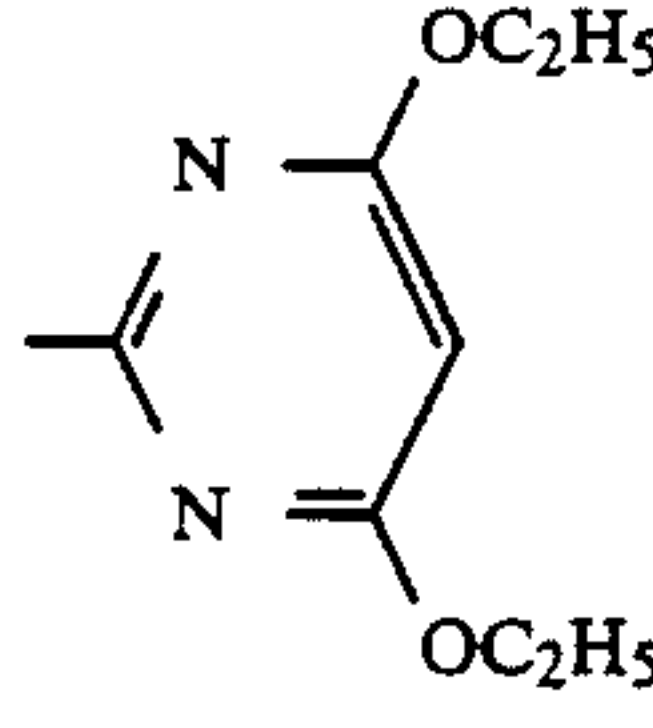
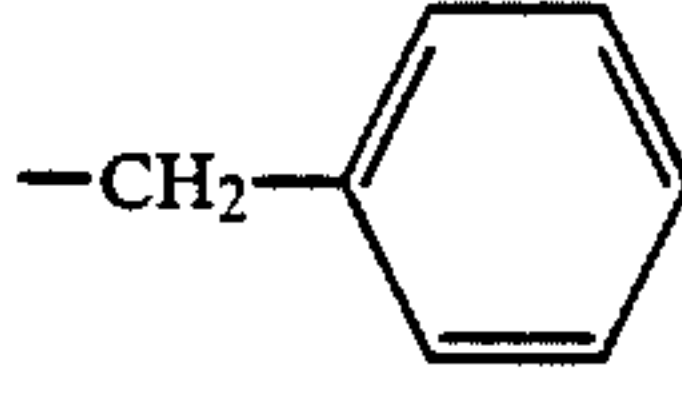
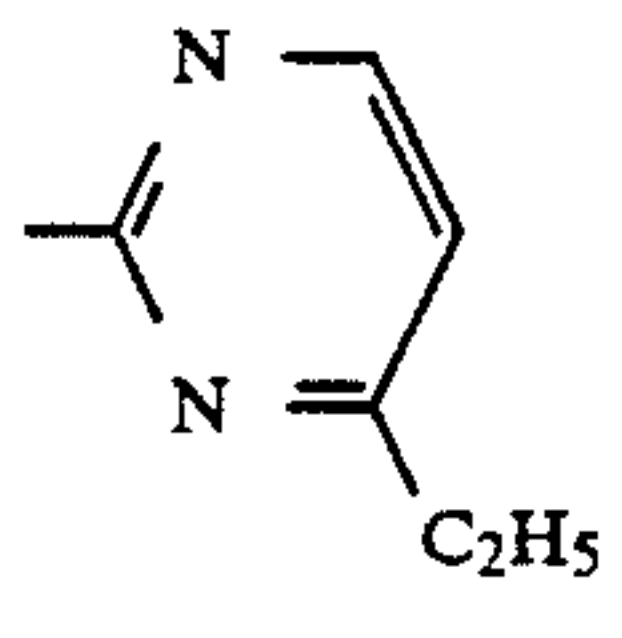
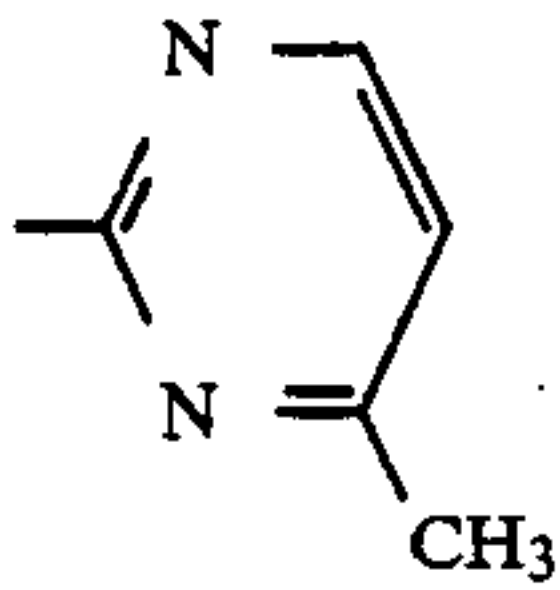
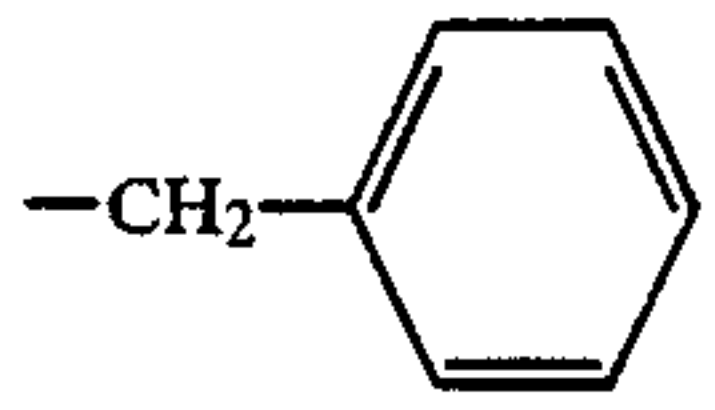
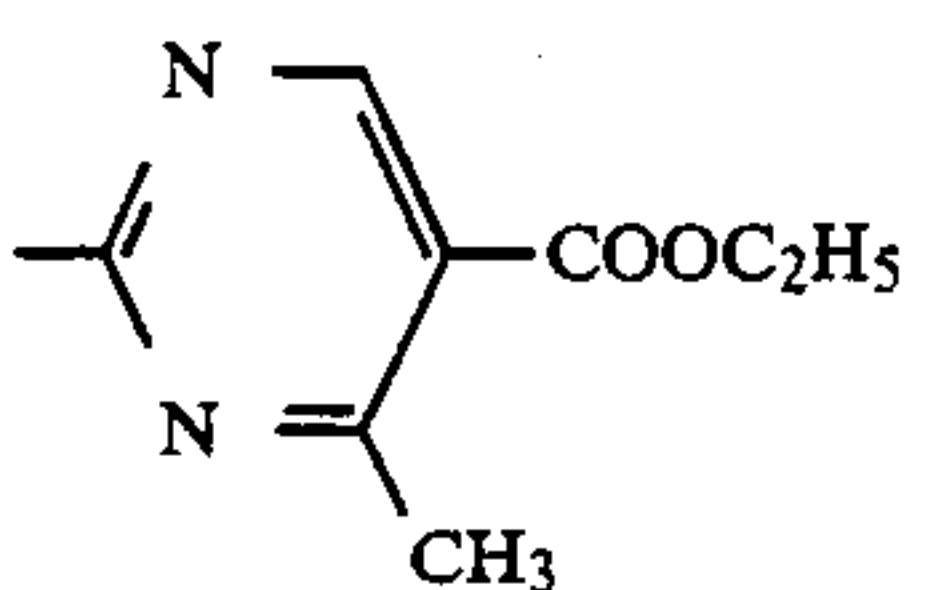
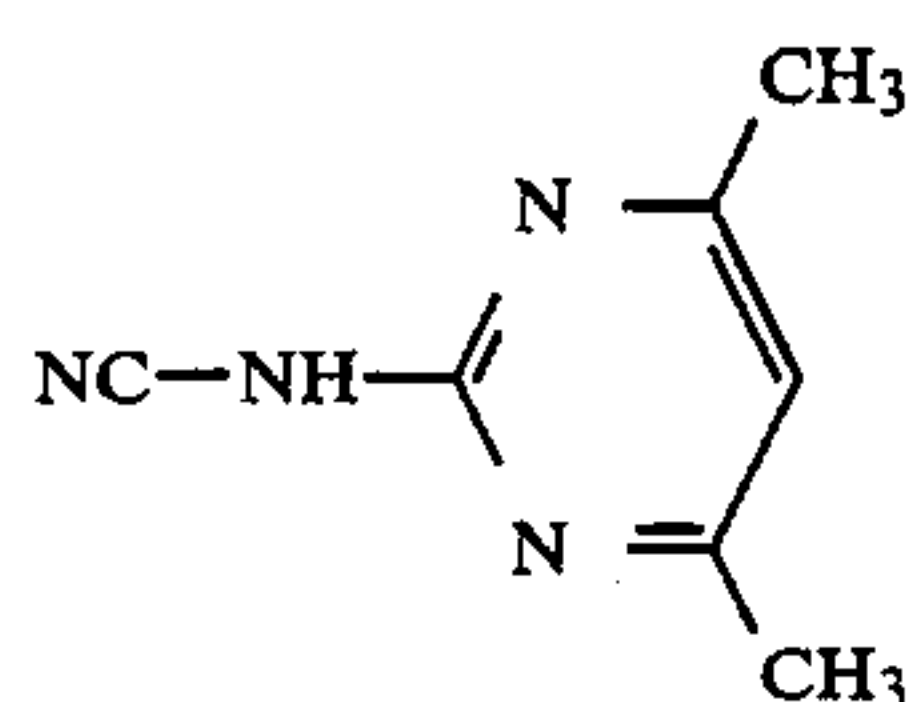
Example No.	R ¹	R ²	R ³	Melting point (°C.)
(IV-55)	—C ₄ H ₉	H		
(IV-56)	—CH ₂ CH(CH ₃) ₂	H		76
(IV-57)		H		68
(IV-58)		H		74
(IV-59)	—CH ₂ —COOC ₂ H ₅	H		
(IV-60)	—CH ₃	H		
(IV-61)	—CH ₃	H		107-109
(IV-62)	—CH ₃	H		
(IV-63)		H		112

TABLE 4-continued

Example No.	R ¹	R ²	R ³	Melting point (°C.)
(IV-64)	$-\text{CH}(\text{C}_6\text{H}_5)_2$	H		165
(IV-65)		H		130

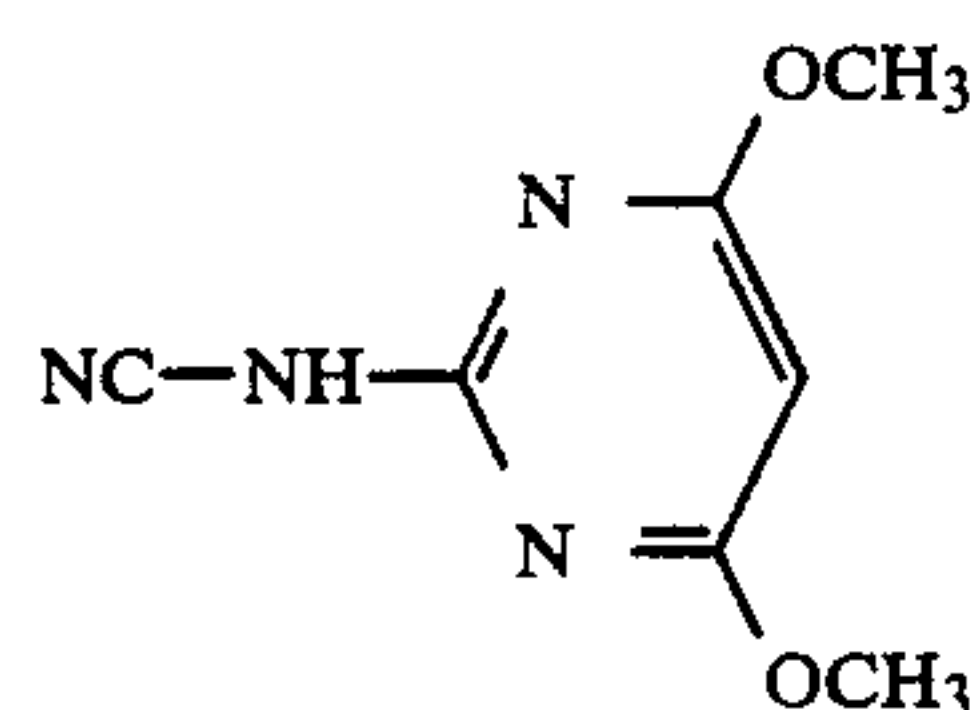
Preparation of the starting substances of the formula
(V)

EXAMPLE (V-1)



A mixture of 42 g (0.5 mole) of cyanoguanidine ("di-cyandiamide") and 50 g (0.5 mole) of 2,4-pentanedione ("acetylacetone") is heated at 120° C. for 15 hours. After cooling, 500 ml of water are then added to the reaction mixture and the solution is acidified with hydrochloric acid at 0° C. to 10° C. The product thereby obtained as crystals is isolated by filtration with suction. 51.8 g (70% of theory) of 2-cyanoamino-4,6-dimethylpyrimidine of melting point 205° C. are obtained.

EXAMPLE (V-2)



A solution, heated to 100° C., of 24 g (0.427 mole) of potassium hydroxide in 100 ml of water is added to a mixture of 9.2 g (0.043 mole) of 4,6-dimethoxypyrimidin-2-yl-thiourea and 70 ml of water at 100° C., with stirring. The mixture is subsequently stirred at 100° C. for 2 minutes and a solution, warmed to 100° C., of 16.2 g (0.05 mole) of lead-II acetate in 30 ml of water is then added. The mixture is heated under reflux for a further 5 minutes and then cooled to 0° C. to 5° C., and 30 ml of glacial acetic acid are added to the aqueous solution. The product thereby obtained as crystals is isolated by filtration with suction.

6.3 g (81.5% of theory) of 2-cyanoamino-4,6-dimethoxy-pyrimidine of melting point 202° C. are obtained.

The compounds of the formula (V) listed in the following Table 5 can be prepared by the process described by way of example in the preceding example:

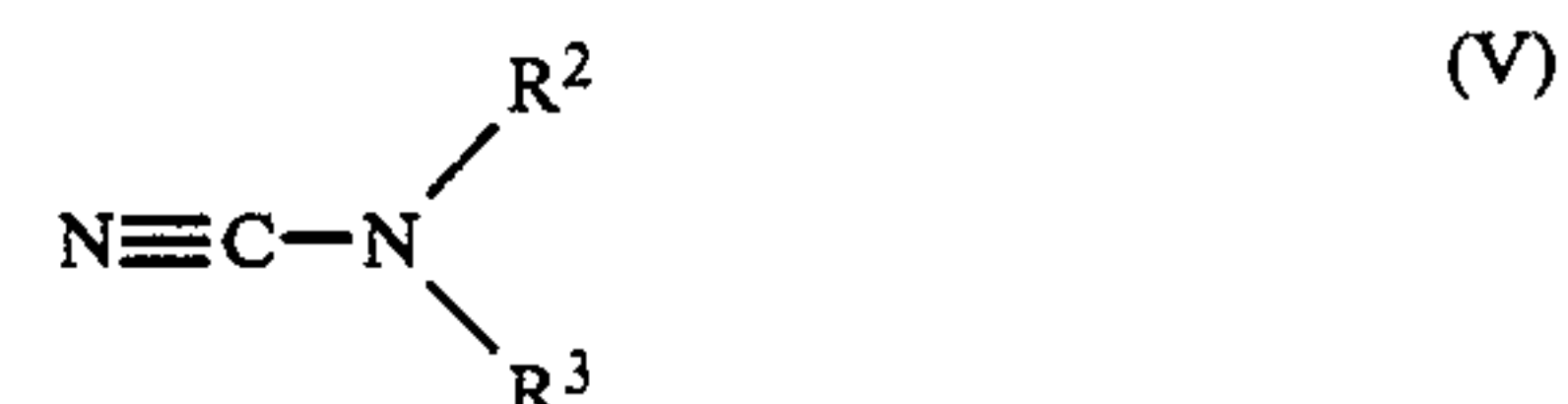


TABLE 5

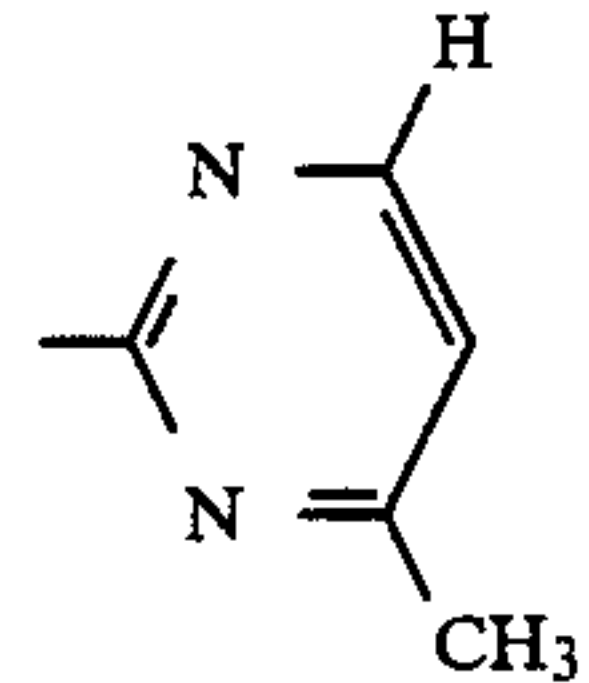
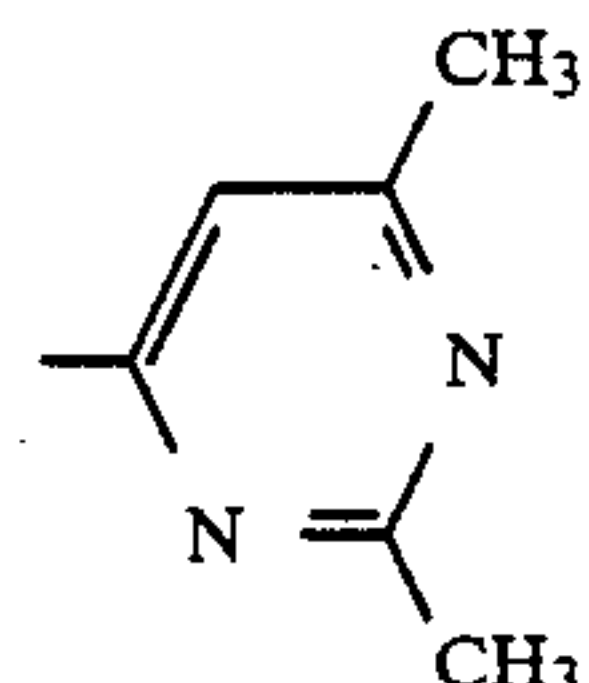
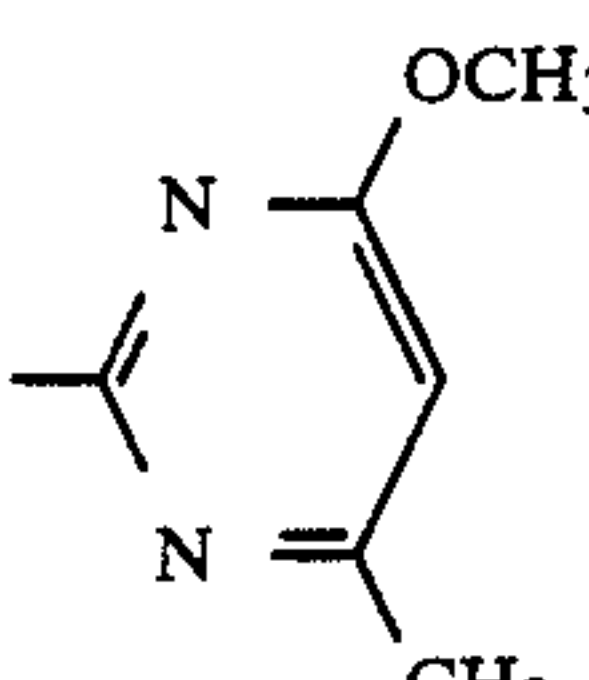
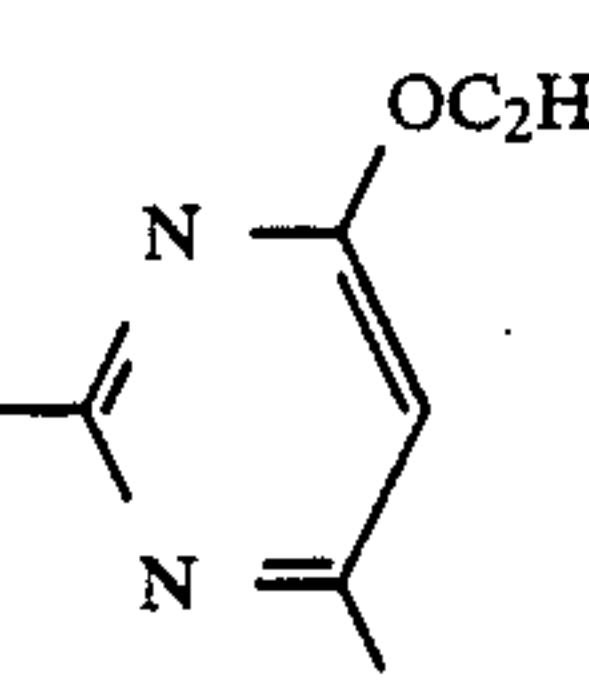
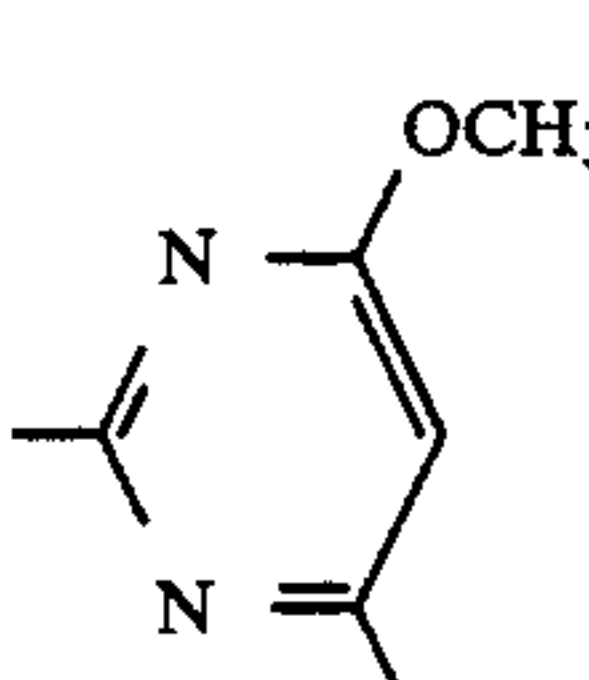
Example No.	R ²	R ³	Melting point (°C.)
(V-3)	H		203 (decomp.)
(V-4)	H		234
(V-5)	H		258
(V-6)	H		
(V-7)	H		200

TABLE 5-continued

Example No.	R ²	R ³	Melting point (°C.)
(V-8)	H		
(V-9)	H		
(V-10)	H		
(V-11)	H		
(V-12)	H		174
(V-13)	H		174
(V-14)	H		146
(V-15)	H		> 300
(V-16)	H		126

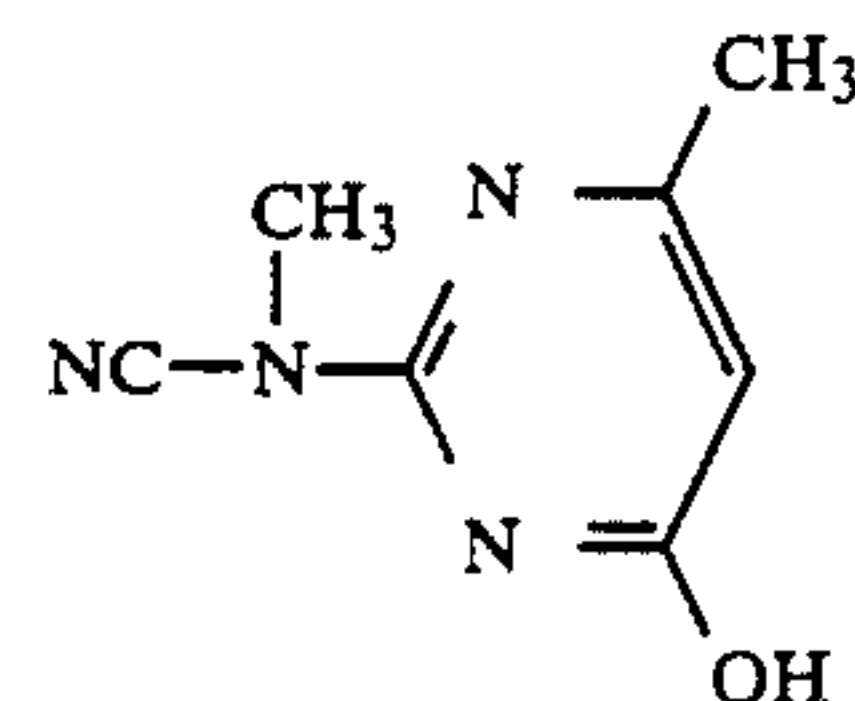
TABLE 5-continued

Example No.	R ²	R ³	Melting point (°C.)
(V-17)	H		186

10 2-(Alkyl-cyano-amino)-pyrimidines of the formula (V) can be prepared, for example, as follows:

EXAMPLE (V-18)

15



20

25 12.6 g (0.1 mole) of dimethyl sulphate are added dropwise to a solution of 15 g (0.1 mole) of 2-cyanoamino-4-hydroxy-6-methyl-pyrimidine and 4.1 g (0.1 mole) of sodium hydroxide in 60 ml of water, whereupon the reaction temperature rises from 20° C. to 40° C. After the mixture has been stirred at 20° C. for two hours, the product obtained as crystals is isolated by filtration with suction.

30

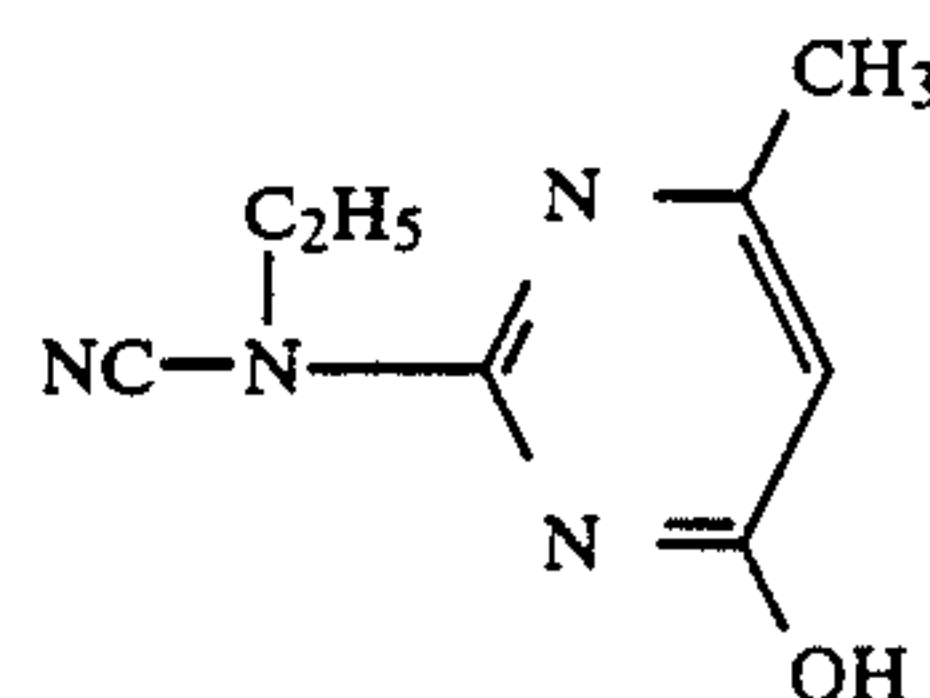
11.1 g (68% of theory) of 2-(methyl-cyano-amino)-4-hydroxy-6-methyl-pyrimide of melting point 290° C. are obtained.

35

The following compound is obtained analogously:

EXAMPLE (V-19)

40

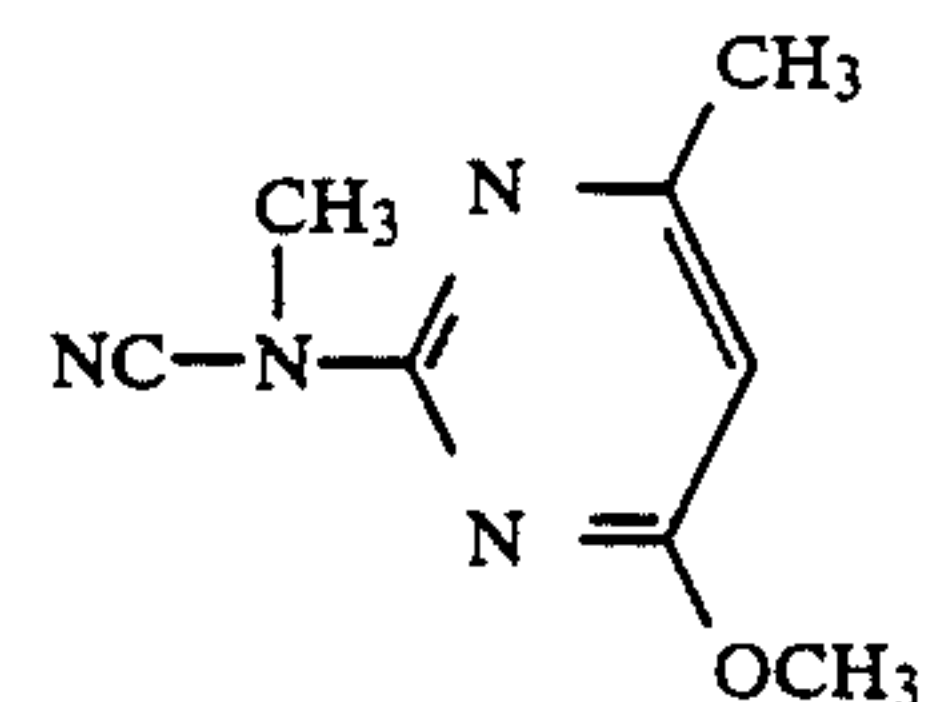


45

Melting point: 215° C. to 220° C.

(EXAMPLE (V-20))

50



55

60 127.5 g (1 mole) of dimethyl sulphate are added dropwise to a solution of 75 g (0.5 mole) of 2-cyanoamino-4-hydroxy-6-methyl-pyrimidine—prepared according to process (b)—and 44 g (1.1 moles) of sodium hydroxide in 750 ml of water, whereupon the reaction temperature rises from 20° C. to 35° C. After the mixture has been stirred at 20° C. for twelve hours, the pH value is brought to between 9 and 10 by addition of sodium hydroxide solution and the product obtained as crystals is isolated by filtration with suction.

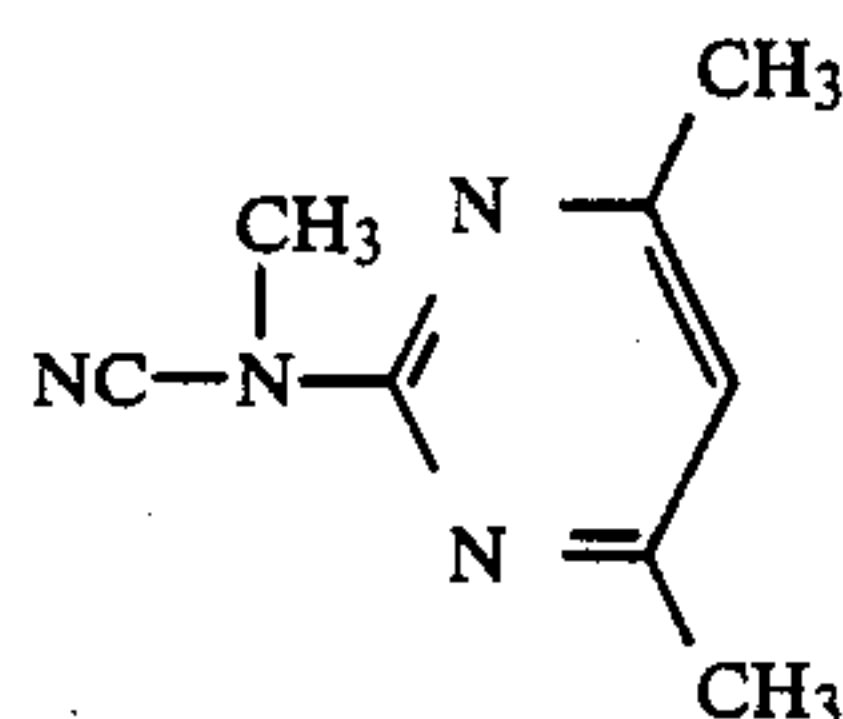
65

61

13 g (15% of theory) of 2-(methyl-cyanoamino)-4-methoxy-6-methyl-pyrimidine of melting point 123° C. are obtained.

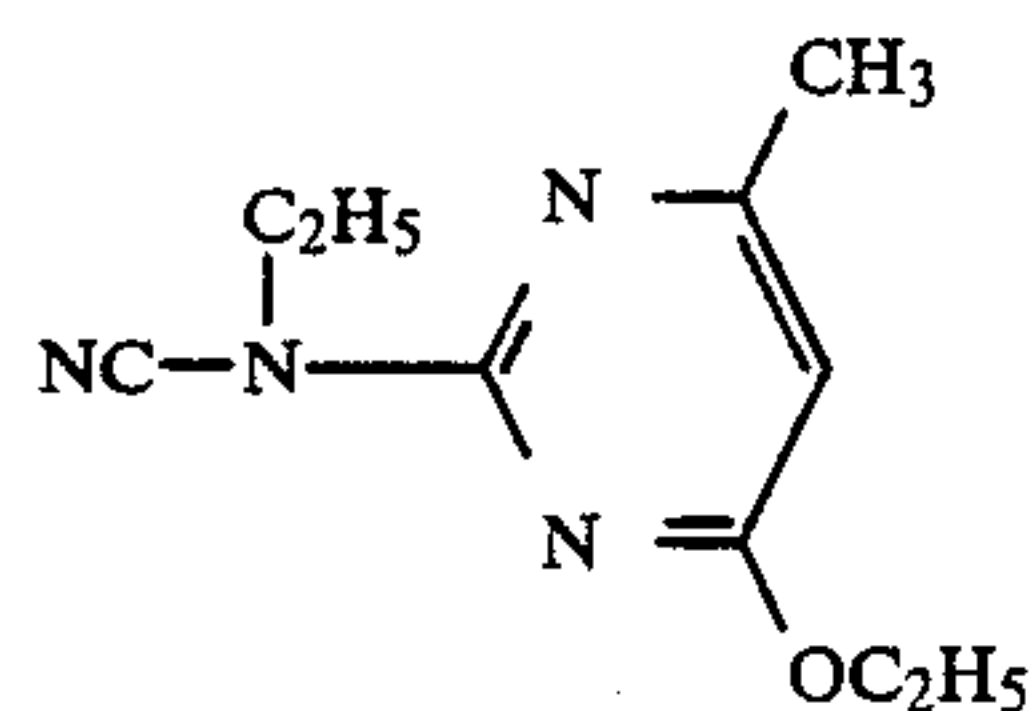
The following compounds are obtained analogously:

EXAMPLE (V-21)



Melting point: 104° C.

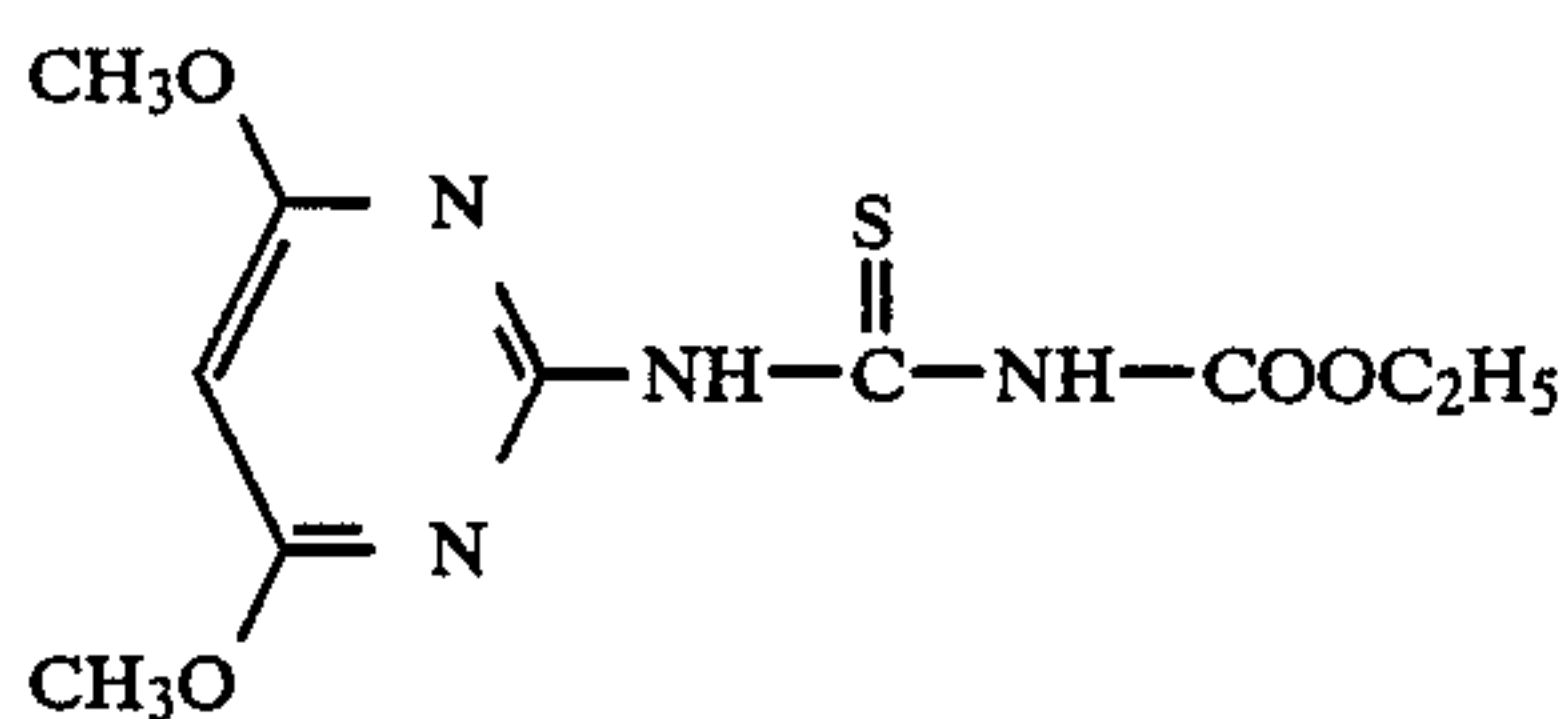
EXAMPLE (V-22)



Melting point: 71° C.

Preparation of the starting substances of the formula (XI)

EXAMPLE (XI-1)



A mixture of 15.5 g (0.1 mole) of 2-amino-4,6-dimethoxy-pyrimidine, 13.1 g (0.1 mole) of ethoxycarbonyl isothiocyanate and 200 ml of acetonitrile is stirred at 60° C. for 2 hours. It is then cooled at 10° C. and the product obtained as crystals is isolated by filtration with suction.

22.5 g (79% of theory) of 1-(ethoxycarbonyl)-3-(4,6-dimethoxy-pyrimidin-2-yl)-thiourea of melting point 194° C. (decomposition) are obtained.

The compounds of the formula (XI) listed in the following Table 6 can be prepared by the process described by way of example in the preceding example:

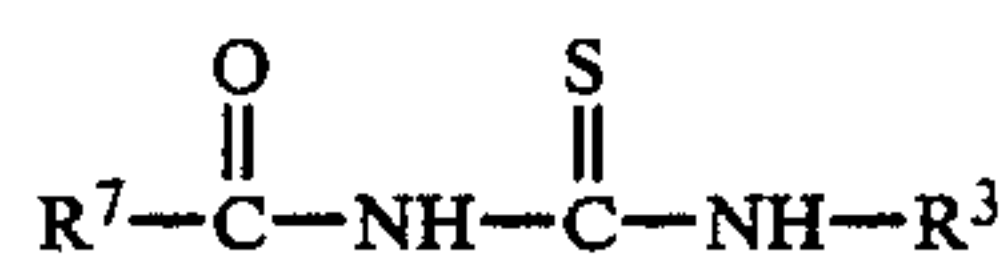


TABLE 6

Example No.	R ⁷	R ³	Melting point (°C.)
(XI-2)			189

62

TABLE 6-continued

Example No.	R ⁷	R ³	Melting point (°C.)
5 (XI-3)			198-9 (decomp.)
10 (XI-4)	-OC ₂ H ₅		217
15 (XI-5)			190
20 (XI-6)	-OC ₂ H ₅		140
25 (XI-7)			145
30 (XI-8)			161
35 (XI-9)	-OC ₂ H ₅		119
40 (XI-10)			182
45 (XI-11)	-OC ₂ H ₅		184-185
50 (XI-12)			189
55 (XI-13)	-OC ₂ H ₅		184-185

TABLE 6-continued

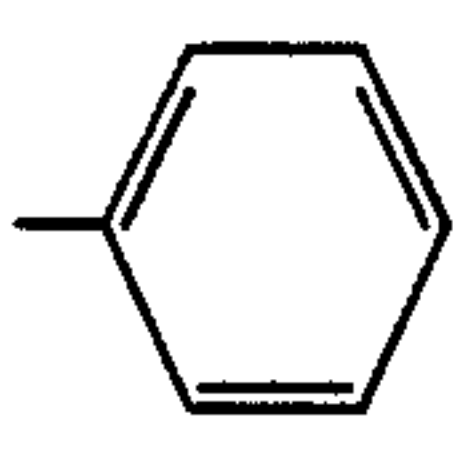
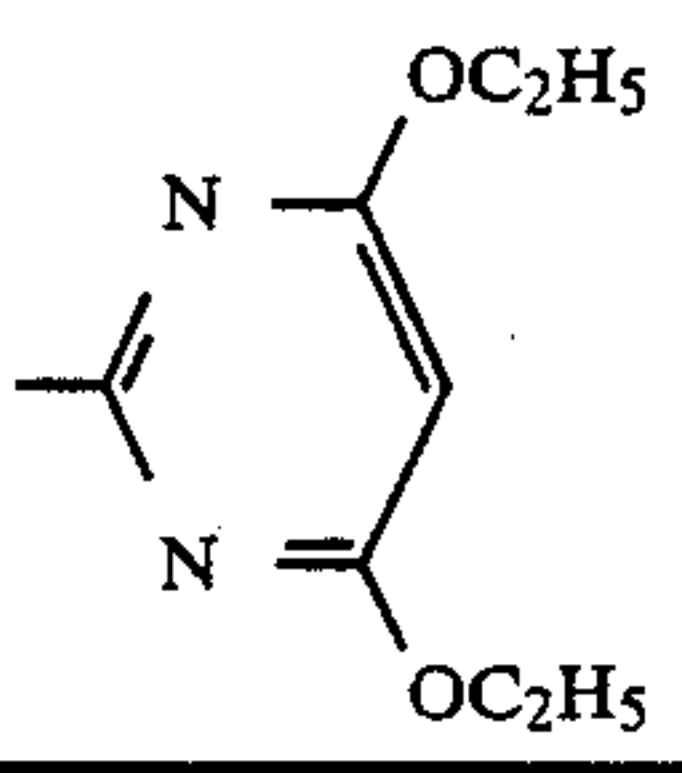
Example No.	R ⁷	R ³	Melting point (°C.)
(XI-12)	-OC ₂ H ₅		173
(XI-13)	-OC ₂ H ₅		160-162
(XI-14)	-OC ₂ H ₅		132-136
(XI-15)	-OC ₂ H ₅		169
(XI-16)			156
(XI-17)	-OC ₂ H ₅		
(XI-18)	-OC ₂ H ₅		
(XI-19)			

TABLE 6-continued

Example No.	R ⁷	R ³	Melting point (°C.)
(XI-20)	-OC ₂ H ₅		168
(XI-21)	-OC ₂ H ₅		
(XI-22)			
(XI-23)	-OC ₂ H ₅		
(XI-24)			
(XI-25)	-OC ₂ H ₅		
(XI-26)			
(XI-27)	-OC ₂ H ₅		
(XI-28)			173

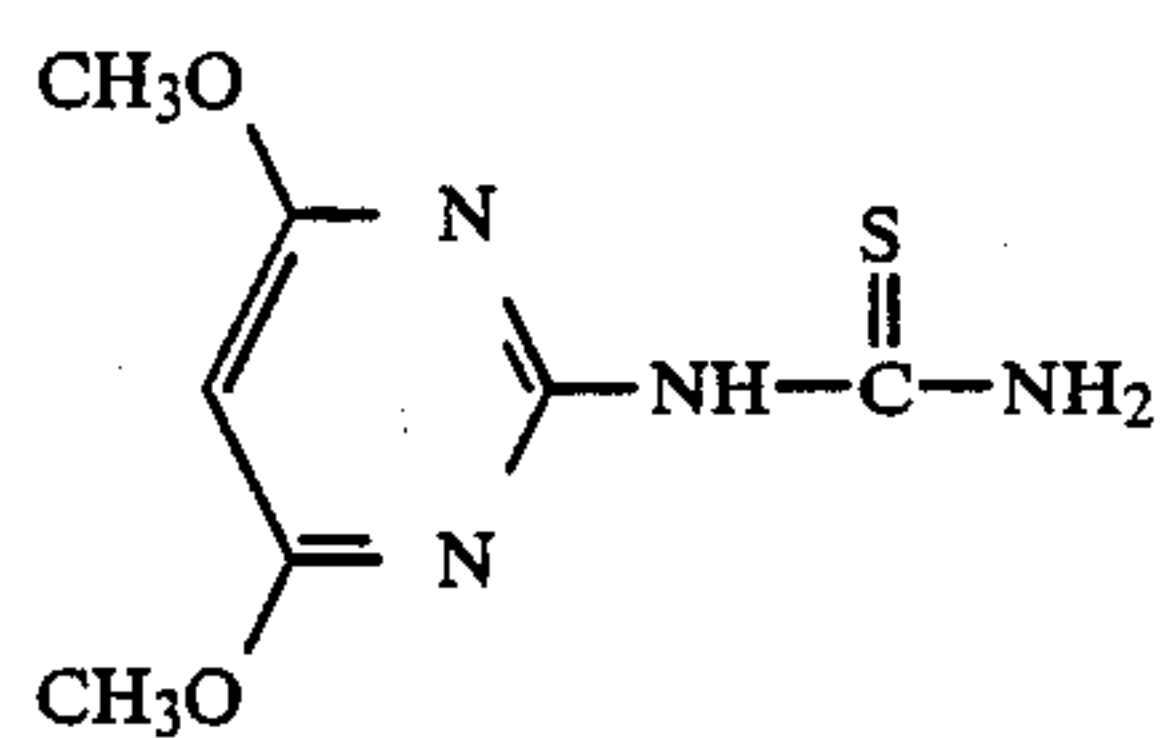
65

TABLE 6-continued

Example No.	R ⁷	R ³	Melting point (°C.)
(XI-29)			179

Preparation of the starting substances of the formula (XII)

EXAMPLE (XII-1)



A mixture of 5.0 g (0.0175 mole) of 1-(ethoxycarbonyl)-3-(4,6-dimethoxy-pyrimidin-2-yl)-thiourea, 4.0 g (0.1 mole) of sodium hydroxide and 100 ml of water is stirred at 20° C. for 2 days. Dilute hydrochloric acid is then added dropwise, with stirring, until the solution has been rendered acid and the evolution of CO₂ has ended. The product obtained as crystals is isolated by filtration with suction.

3.5 g (94% of theory) of 4,6-dimethoxy-pyrimidin-2-yl-thiourea of melting point 245°-8° C. (decomposition) are obtained.

The compounds of the formula (XII) listed in the following Table 7 can be prepared by the process described by way of example in the preceding example:

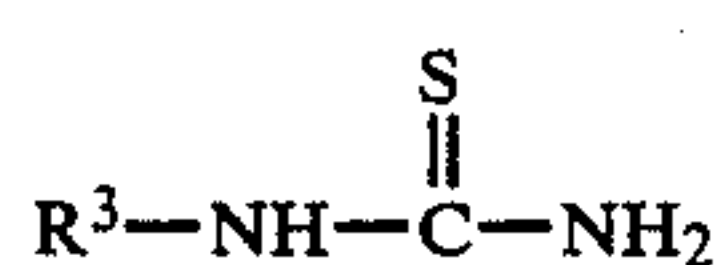
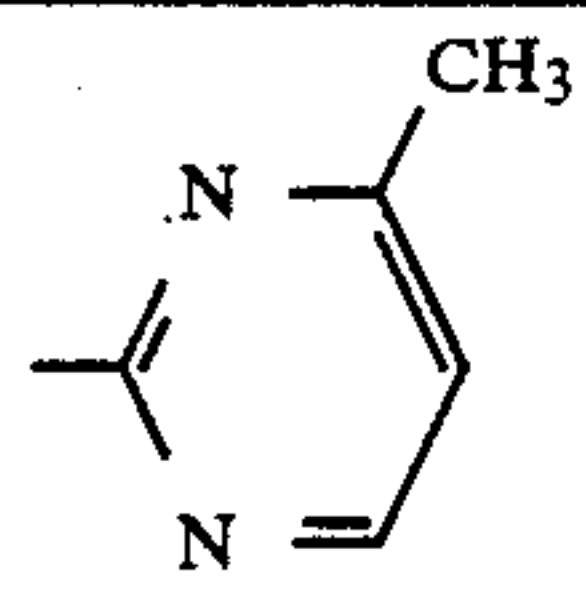
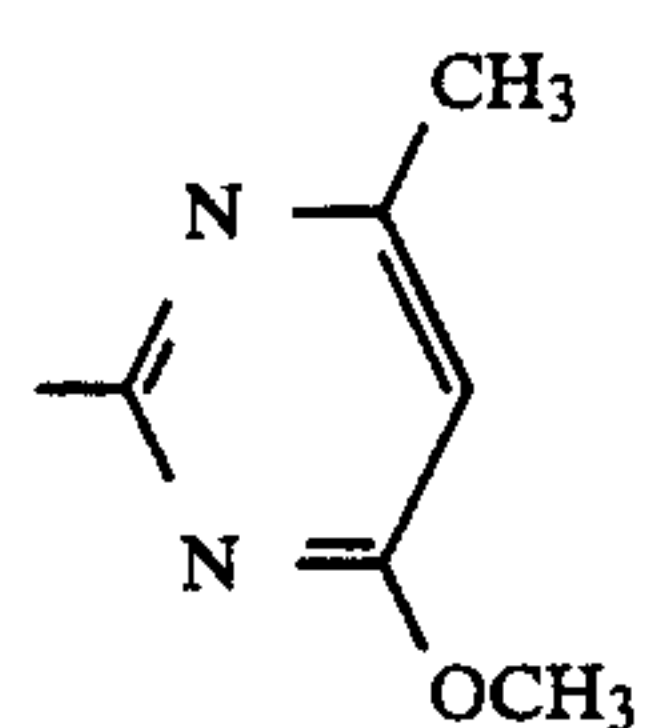
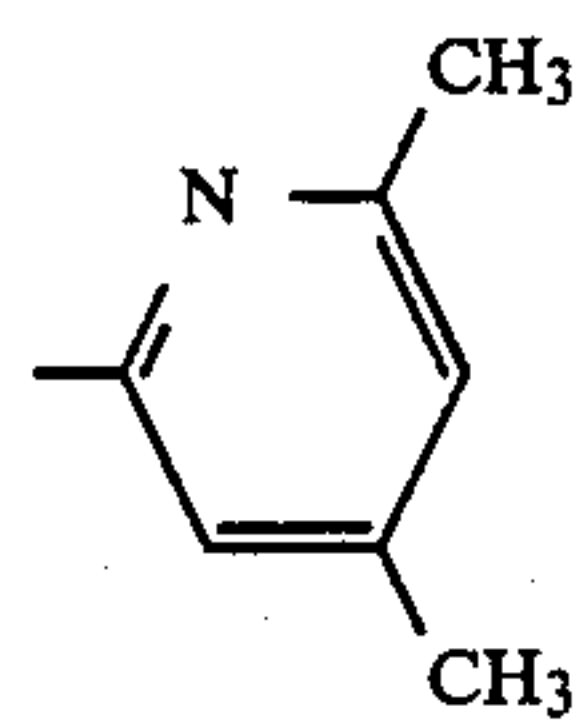


TABLE 7

Example No.	R ³	Melting point (°C.)
(XII-2)		264-265 (decomp.)
(XII-3)		231 (decomp.)
(XII-4)		259-260 (decomp.)

66

TABLE 7-continued

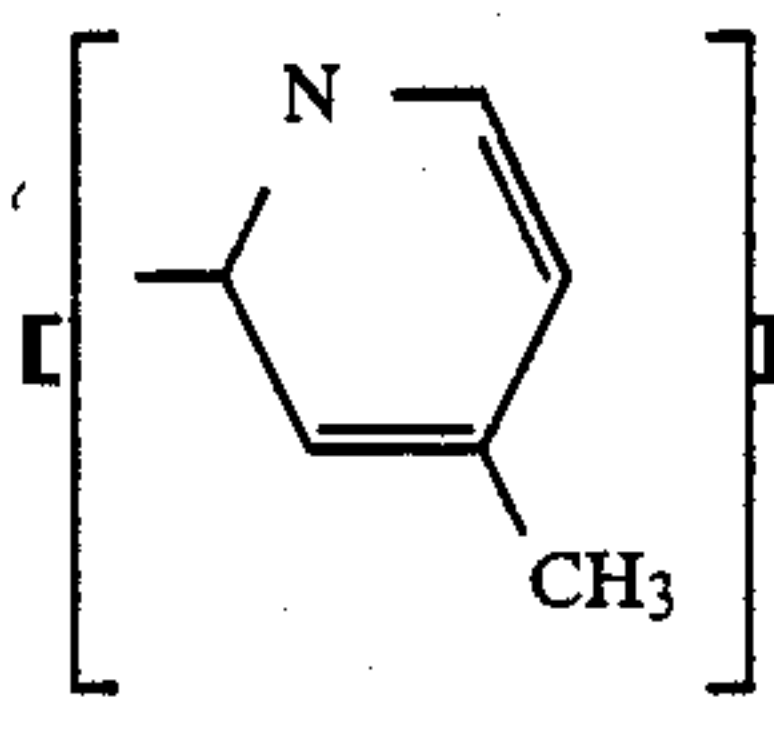
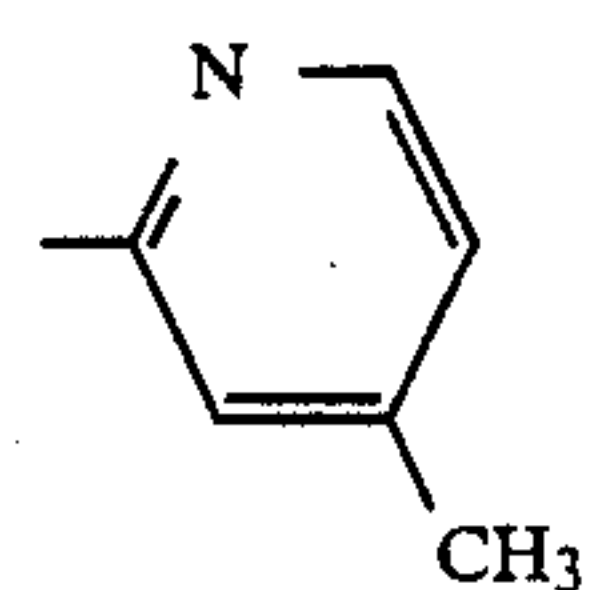
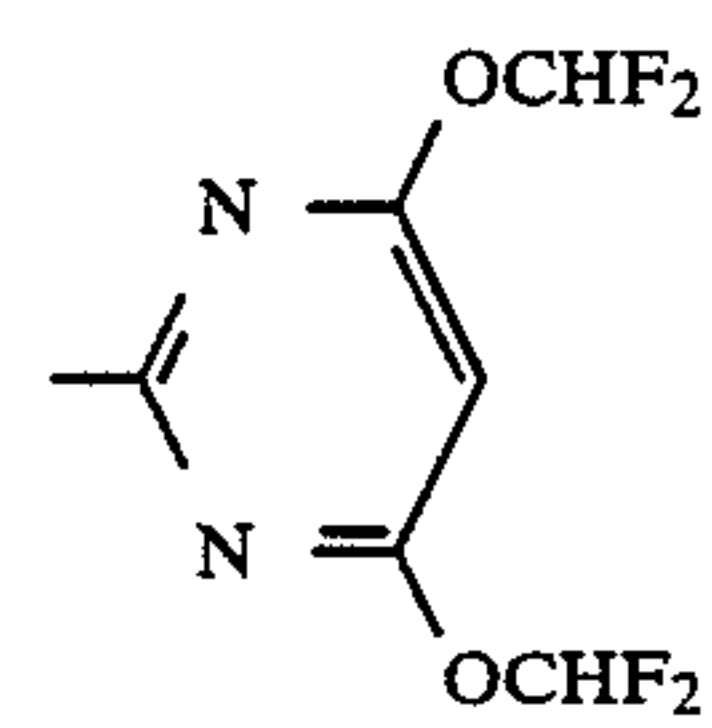
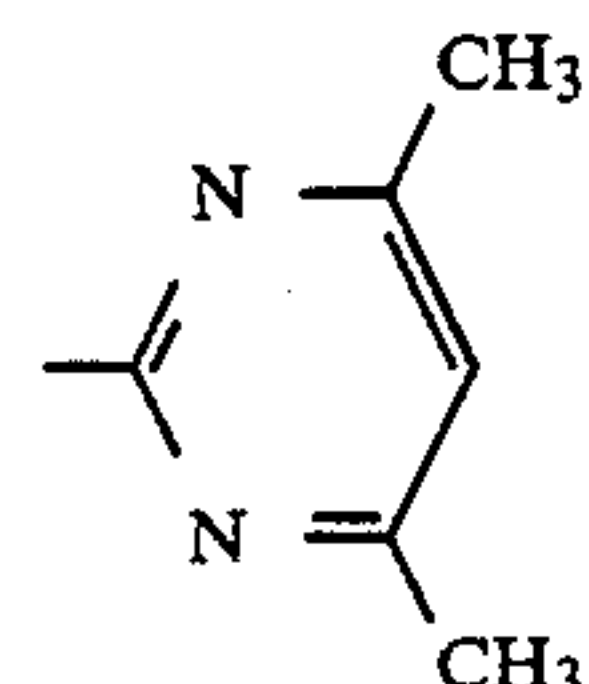
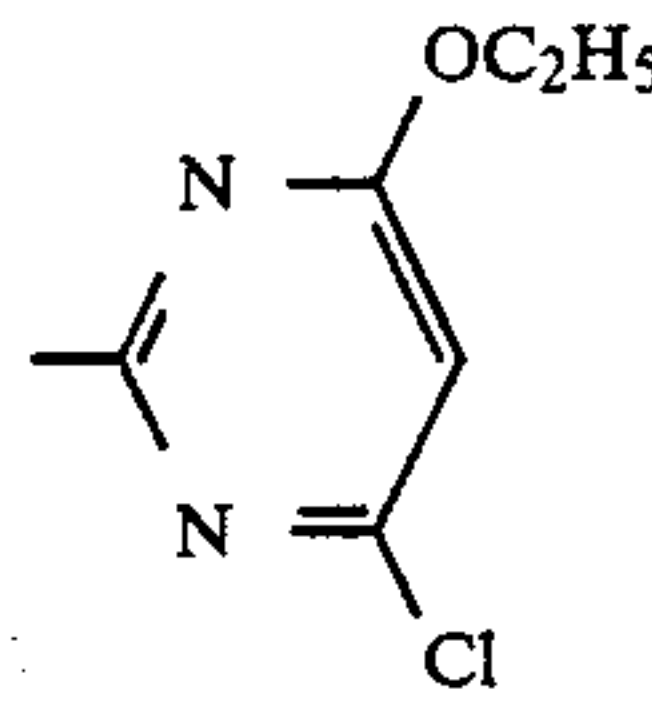
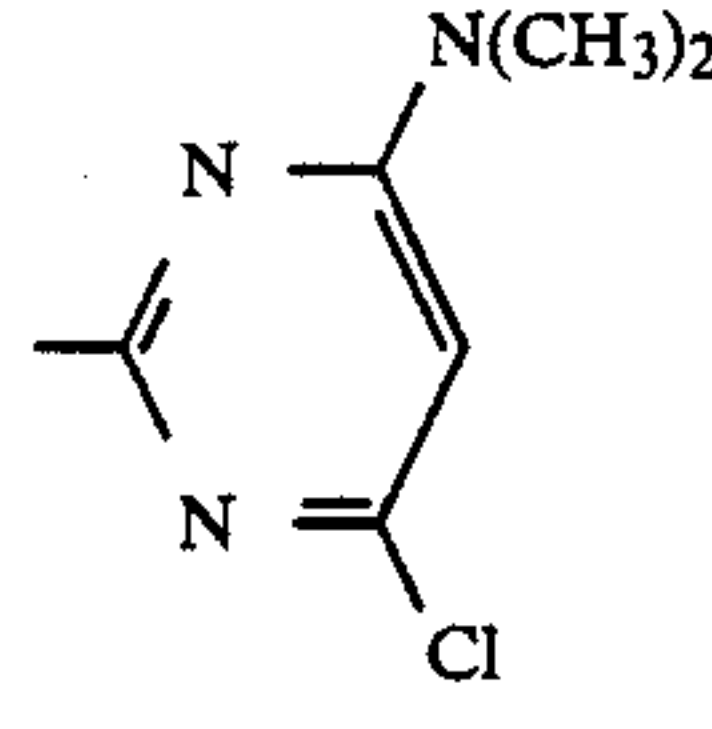
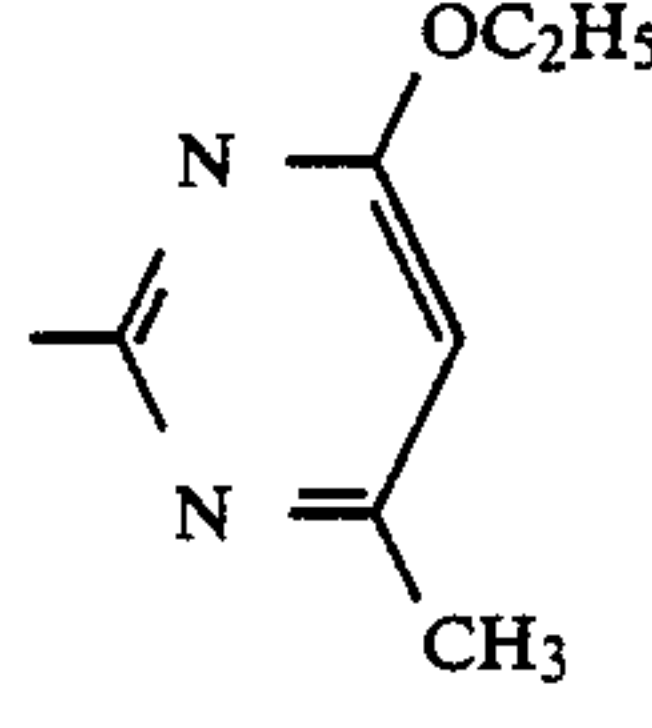
Example No.	R ³	Melting point (°C.)
(XII-5)		214-215
(XII-6)		192-194
(XII-7)		225-227 (decomp.)
(XII-8)		225-227 (decomp.)
(XII-9)		156
(XII-10)		
(XII-11)		

TABLE 7-continued

Example No.	R ³	Melting point (°C.)
(XII-12)		
(XII-13)		
(XII-14)		263
(XII-15)		166
(XII-16)		248

EXAMPLE A

Pre-emergence test

Solvent: 5 parts by weight of acetone

Emulsifier: 1 part by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, 1 part by weight of active compound is mixed with the stated amount of solvent, the stated amount of emulsifier is added and the concentrate is diluted with water to the desired concentration.

Seeds of the test plants are sown in normal soil and, after 24 hours, watered with the preparation of the active compound. It is expedient to keep constant the amount of water per unit area. The concentration of the active compound in the preparation is no importance, only the amount of active compound applied per unit area being decisive. After three weeks, the degree of damage to the plants is rated in % damage in comparison to the development of the untreated control. The figures denote:

0% = no action (like untreated control)

100% = total destruction

In this test, the active compounds according to the invention exhibit a very good herbicidal activity.

EXAMPLE B

Pre-emergence test

Solvent: 5 parts by weight of acetone

Emulsifier: 1 part by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, 1 part by weight of active compound is mixed with the stated amount of solvent, the stated amount of emulsifier is added and the concentrate is diluted with water to the desired concentration.

Test plants which have a height of 5-15 cm are sprayed with the preparation of the active compound in such a way as to apply the particular amount of active compound desired per unit area. The concentration of the spray liquor is so chosen that the particular amounts of active compound desired are applied in 2,000 l of water/ha. After three weeks, the degree of damage to the plants is rated in % damage in comparison to the development of the untreated control. The figures denote:

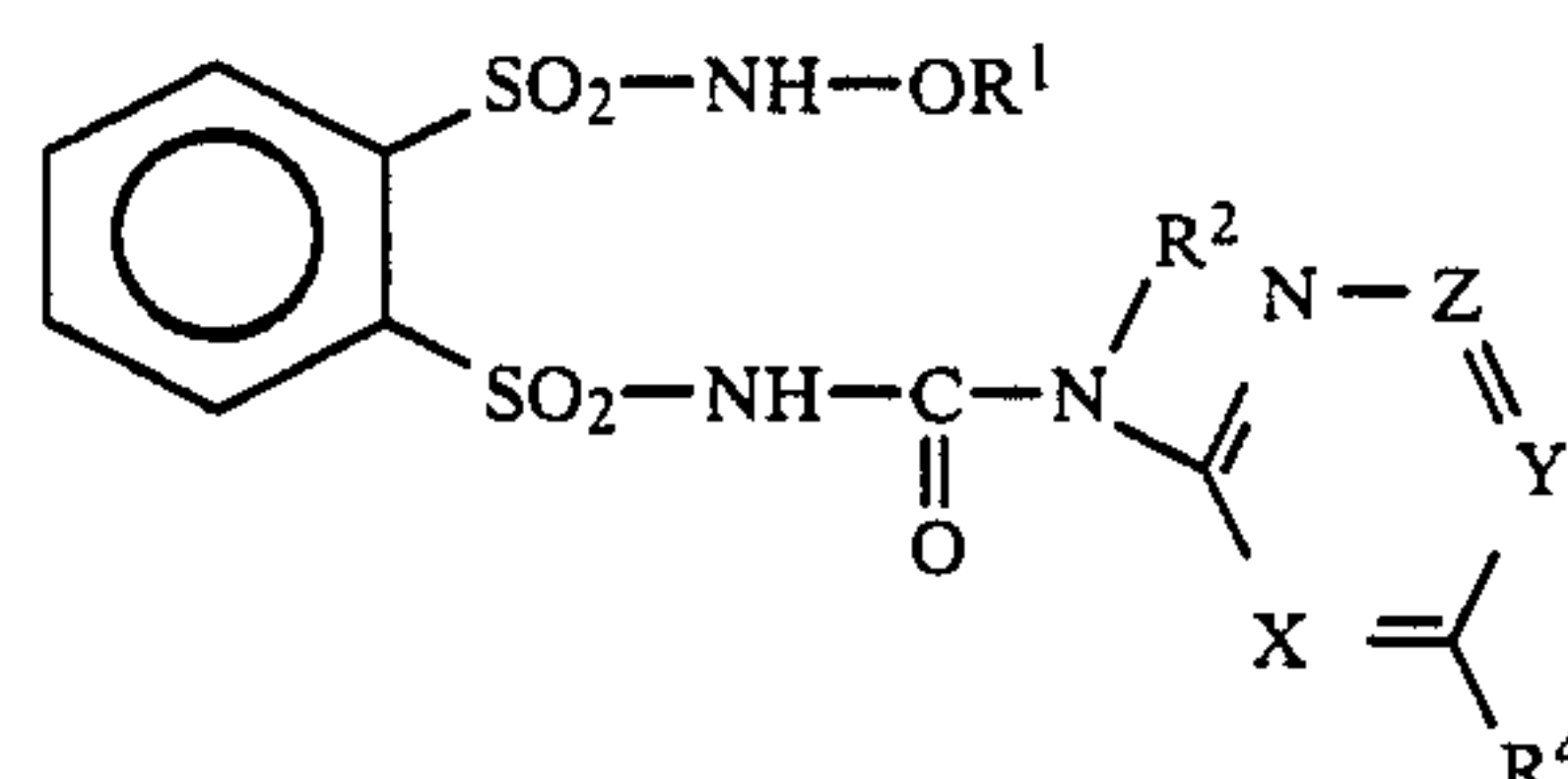
0% = no action (like untreated control)

100% = total destruction

In this test, the active compounds according to the invention exhibit a very good herbicidal activity.

We claim:

1. A 1-(2-oxyaminosulphonylphenylsulphonyl)-3-heteroaryl-urea of the formula



in which

R¹ represents C₁-C₁₂-alkyl which is optionally substituted by fluorine, chlorine, cyano, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-alkylsulphinyl, C₁-C₄-alkylsulphonyl, C₁-C₄-alkyl-carbonyl, C₁-C₄-alkoxy-carbonyl, C₁-C₄-alkylamino-carbonyl or di-[(C₁-C₄-alkyl)](C₁-C₄-alkyl)-amino-carbonyl, or represents C₃-C₆-alkenyl which is optionally substituted by fluorine, chlorine or bromine, C₃-C₆-alkinyl, C₃-C₆-cycloalkyl, C₃-C₆-cycloalkyl-C₁-C₂-alkyl or phenyl-C₁-C₂-alkyl which is optionally substituted by fluorine, chlorine, nitro, cyano, C₁-C₄-alkyl, C₁-C₄-alkoxy or C₁-C₄-alkoxy-carbonyl, or represents benzhydryl, or represents phenyl which is optionally substituted by fluorine, chlorine, nitro, cyano, C₁-C₄-alkyl, trifluoromethyl, C₁-C₄-alkoxy, C₁-C₂-fluoroalkoxy, C₁-C₄-alkylthio, trifluoromethylthio or C₁-C₄-alkoxy-carbonyl,

R² represents hydrogen or C₁-C₄-alkyl which is optionally substituted by fluorine, chlorine, cyano, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-alkylsulphinyl, C₁-C₄-alkylsulphonyl, C₁-C₄-alkyl-carbonyl, C₁-C₄-alkoxy-carbonyl, C₁-C₄-alkylamino-carbonyl or di-(C₁-C₄-alkyl)-aminocarbonyl, or represents C₃-C₆-alkenyl, C₃-C₆-alkinyl or phenyl-C₁-C₂-alkyl which is optionally substituted by fluorine, chlorine, nitro, cyano, C₁-C₄-alkyl, C₁-C₄-alkoxy or C₁-C₄-alkoxycarbonyl,

R⁴ represents hydrogen, fluorine, chlorine, bromine, hydroxyl, C₁-C₄-alkyl which is optionally substituted by fluorine and/or chlorine, C₁-C₄-alkoxy [which is optionally substituted by fluorine and/or chlorine], C₁-C₄-alkylthio which is optionally

substituted by fluorine and/or chlorine, amino, C₁-C₄-alkylamino or di-(C₁-C₄-alkyl)-amino,

X represents nitrogen,

Y represents an optionally substituted methine bridge C-R⁵,

R⁵ represents hydrogen, fluorine, chlorine, bromine, cyano, formyl, C₁-C₄-alkyl, C₁-C₃-alkoxy-carbonyl or C₁-C₃-alkyl-carbonyl, and

Z represents an optionally substituted methine bridge C-R⁶, and

R⁶ represents hydrogen, fluorine, chlorine, bromine, hydroxyl, C₁-C₄-alkyl [which is optionally substituted by fluorine and/or chlorine], C₁-C₄-alkylthio which is optionally substituted by fluorine and/or chlorine, amino, C₁-C₄-alkylamino or di-(C₁-C₄-alkyl)-amino.

2. A compound according to claim 1, in which

R¹ represents C₁-C₈-alkyl which is optionally substituted by fluorine or chlorine, C₃-C₄-alkenyl, C₁-C₂-alkoxy-carbonmethyl, phenyl, phenethyl or benzyl which is optionally substituted by fluorine, chlorine, nitro, cyano, methyl, methoxy or methoxy-carbonyl,

R² represents hydrogen,

R⁴ represents chlorine, methyl, ethyl, methoxy, di-fluoromethoxy or ethoxy,

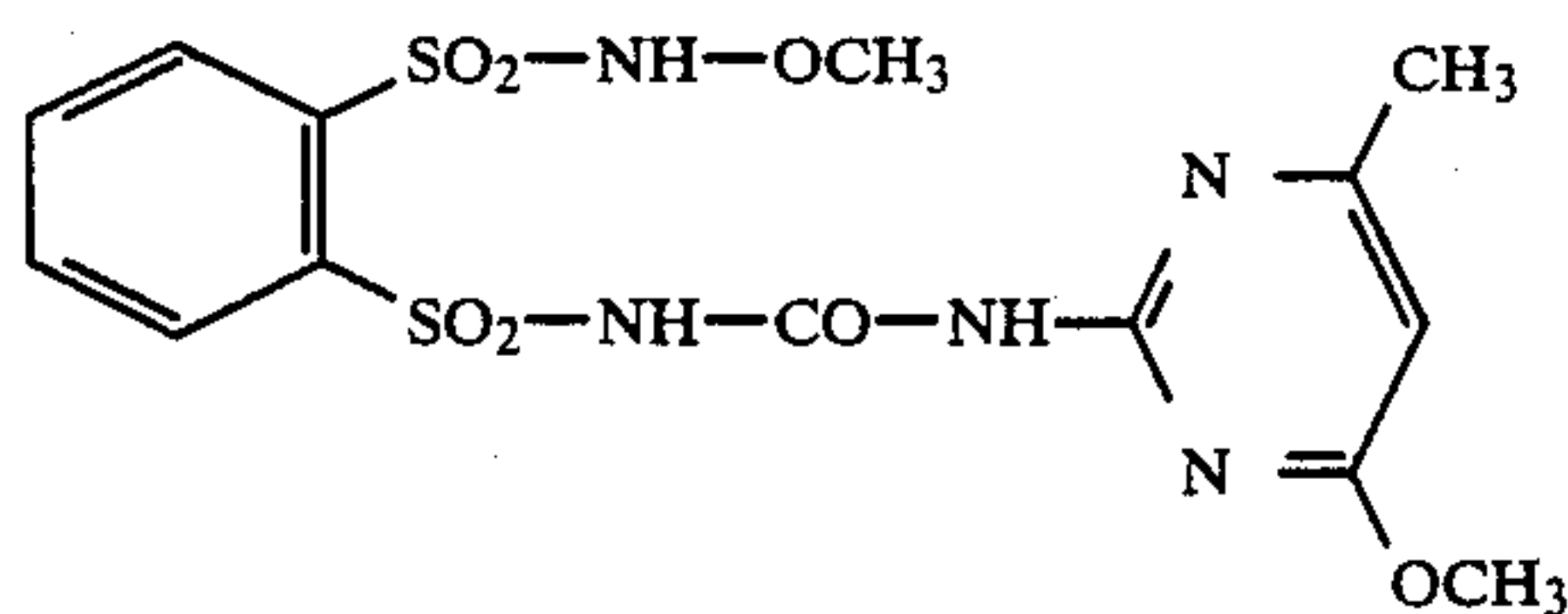
X represents nitrogen,

Y represents a methine bridge (CH) and

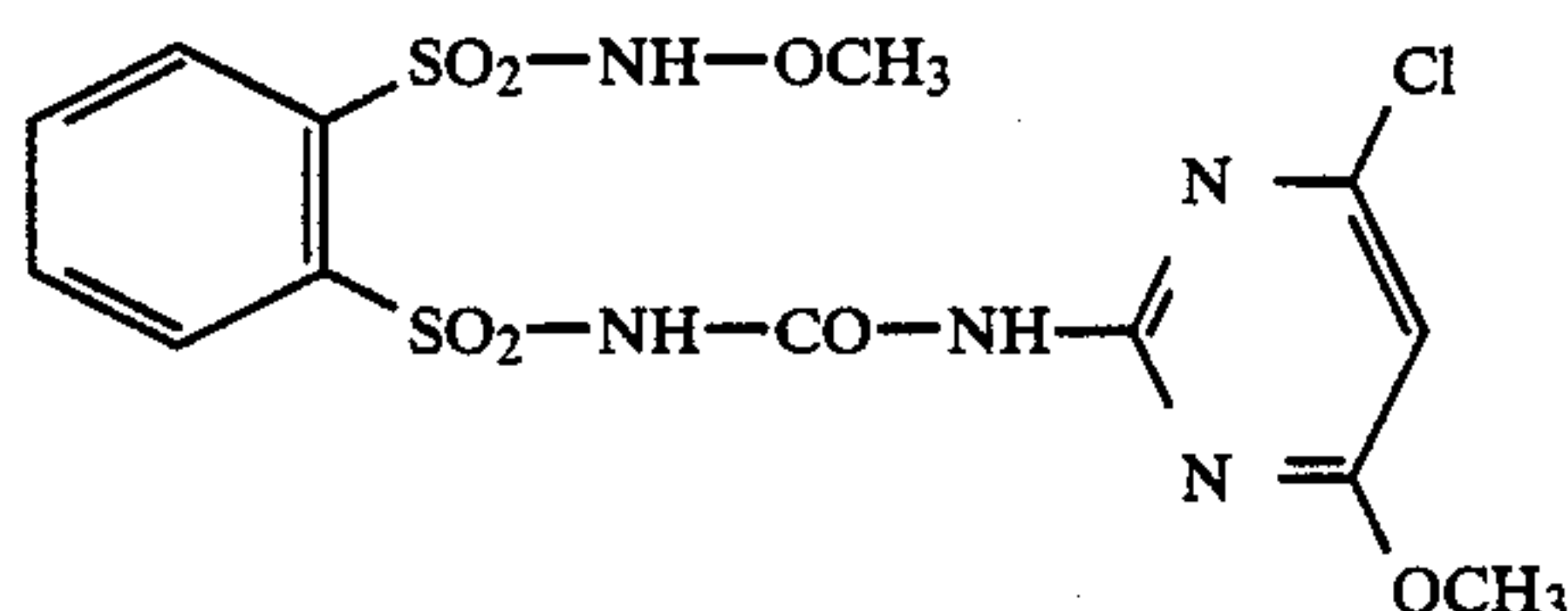
Z represents an optionally substituted methine bridge C-R⁶, and

R⁶ represents hydrogen, chlorine, methoxy, ethoxy, methylthio, ethylthio, [dimethylamino] dimethyl-amino or diethylamino.

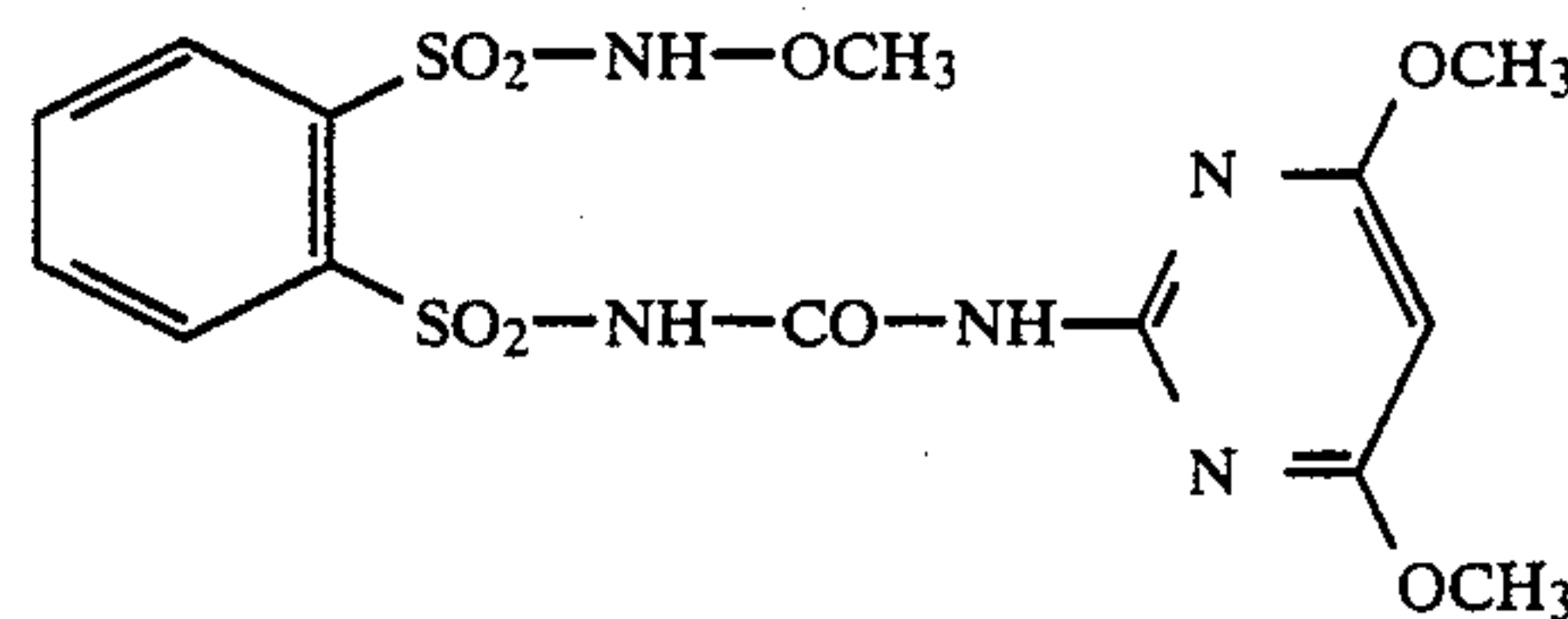
3. A compound according to claim 1, wherein such compound is 1-(2-methoxyaminosulphonyl-phenylsulphonyl)-3-(4-methoxy-6-methyl-pyrimidin-2-yl)-urea of the formula



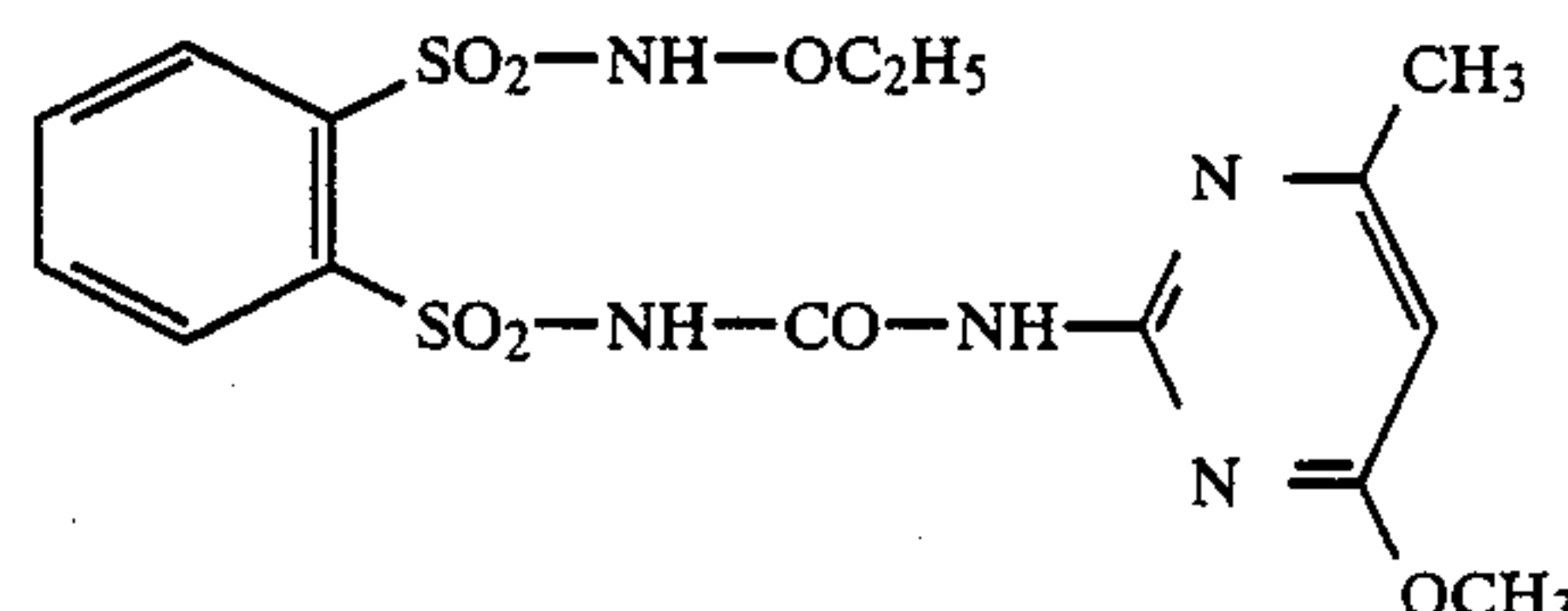
4. A compound according to claim 1, wherein such compound is 1-(2-methoxyaminosulphonyl-phenylsulphonyl)-3-(4-chloro-6-methoxy-pyrimidin-2-yl)-urea of the formula



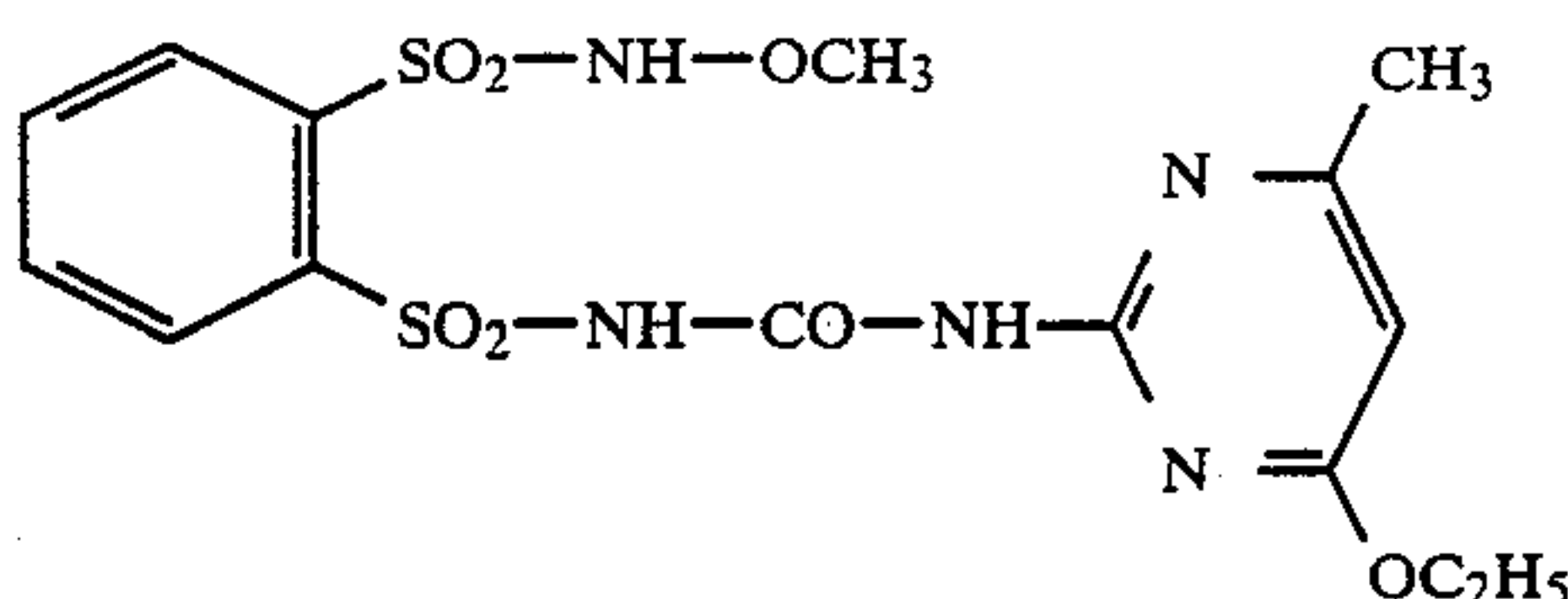
5. A compound according to claim 1, wherein such compound is 1-(2-methoxyaminosulphonyl-phenylsulphonyl)-3-(4,6-dimethoxy-pyrimidin-2-yl)-urea of the formula



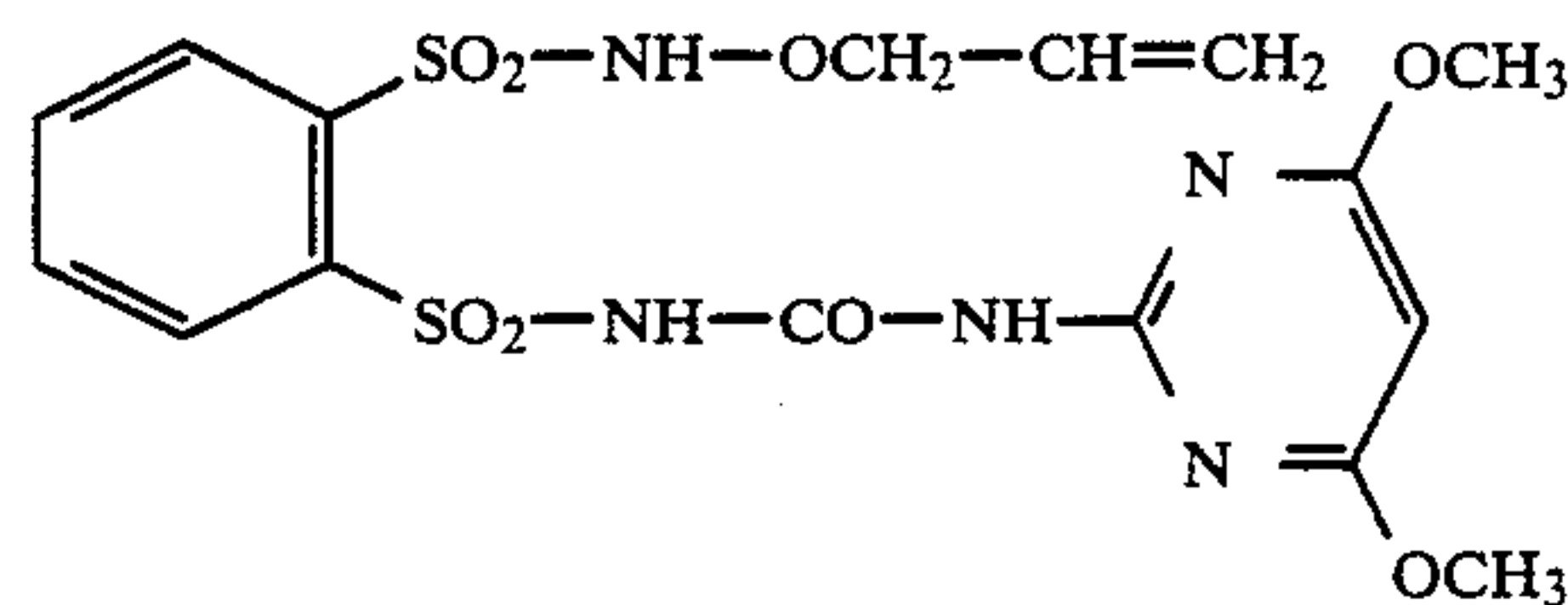
6. A compound according to claim 1, wherein such compound is 1-(2-ethoxyaminosulphonyl-phenylsulphonyl)-3-(4-methoxy-6-methyl-pyrimidin-2-yl)-urea of the formula



7. A compound according to claim 1, wherein such compound is 1-(2-methoxyaminosulphonyl-phenylsulphonyl)-3-(4-ethoxy-6-methyl-pyrimidin-2-yl)-urea of the formula



8. A compound according to claim 1, wherein such compound is 1-(2-allyloxyaminosulphonyl-phenylsulphonyl)-3-(4,6-dimethoxy-pyrimidin-2-yl)-urea of the formula



9. A herbicidal composition comprising a herbicidally effective amount of a compound according to claim 1 and a diluent.

10. A method of combating unwanted vegetation which comprises applying to such vegetation or to a locus from which it is desired to exclude such vegetation a herbicidally effective amount of a compound according to claim 1 and a diluent.

11. The method according to claim 10, wherein such compound is

1-(2-methoxyaminosulphonyl-phenylsulphonyl)-3-(4-methoxy-6-methyl-pyrimidin-2-yl)-urea,

1-(2-methoxyaminosulphonyl-phenylsulphonyl)-3-(4-chloro-6-methoxy-pyrimidin-2-yl)-urea,

1-(2-methoxyaminosulphonyl-phenylsulphonyl)-3-(4,6-dimethoxy-pyrimidin-2-yl)-urea,

1-(2-ethoxyaminosulphonyl-phenylsulphonyl)-3-(4-methoxy-6-methyl-pyrimidin-2-yl)-urea,

1-(2-methoxyaminosulphonyl-phenylsulphonyl)-3-(4-ethoxy-6-methyl-pyrimidin-2-yl)-urea, or

1-(2-allyloxyaminosulphonyl-phenylsulphonyl)-3-(4,6-dimethoxy-pyrimidin-2-yl)-urea.

* * * * *

UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

PATENT NO. : Re 33,298

DATED : August 14, 1990

INVENTOR(S) : Diehr et al.

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Col. 69, claim 2	Delete " carbonmethyl " and substitute -- carbonyl-
line 22	methyl --

Signed and Sealed this
Twenty-sixth Day of May, 1992

Attest:

DOUGLAS B. COMER

Attesting Officer

Acting Commissioner of Patents and Trademarks