



US00RE34069E

United States Patent [19]

[11] E

Patent Number: Re. 34,069

Köster et al.

[45] Reissued Date of Patent: Sep. 15, 1992

[54] PROCESS FOR THE PREPARATION OF OLIGONUCLEOTIDES

[75] Inventors: Hubert Köster, Concord, Mass.; Nanda D. Sinha, San Rafael, Calif.

[73] Assignee: Biosyntech GmbH, Hamburg, Fed. Rep. of Germany

[21] Appl. No.: 481,572

[22] Filed: Feb. 16, 1990

Related U.S. Patent Documents

Reissue of:

[64] Patent No.: 4,725,677
Issued: Feb. 16, 1988
Appl. No.: 752,178
Filed: Aug. 10, 1984[51] Int. Cl.⁵ C07H 15/12; C07H 17/00[52] U.S. Cl. 536/27; 536/28;
536/29

[58] Field of Search 536/27, 28, 29

[56] References Cited

U.S. PATENT DOCUMENTS

4,310,662 1/1982 Crea 536/27
4,415,732 11/1983 Caruthers et al. 536/27
4,419,509 12/1983 Hsiung 536/27
4,458,066 7/1984 Caruthers et al. 536/27
4,476,301 10/1984 Imbach et al. 536/29
4,500,707 2/1985 Caruthers et al. 536/29
4,591,614 5/1986 Miller et al. 536/27
4,605,735 8/1986 Miyoshi et al. 536/27

FOREIGN PATENT DOCUMENTS

0040099 11/1981 European Pat. Off. .
81302110.2 11/1981 European Pat. Off. .
0061746 10/1982 European Pat. Off. .
0064796 11/1982 European Pat. Off. .
82200564.1 11/1982 European Pat. Off. .
0090789 10/1983 European Pat. Off. .
83870031.8 10/1983 European Pat. Off. .
0131993 1/1985 European Pat. Off. .
8420951.6 1/1985 European Pat. Off. .

OTHER PUBLICATIONS

Clesen et al., "Tetrahedron Letters", vol. 25, No. 12, pp. 1307-1310, 1984.

Lelsing et al., "Jour. of the Amer. Chem. Soc." vol. 98, No. 12, Jun. 1976, pp. 3655-3661.

Narang, "Tetrahedron", vol. 39, No. 1, pp. 3-22, 1983.
Marugg et al., "Recl. Trav. Chim. Pays-Bas" vol. 103, pp. 97-98, 1984.Ogilvie, K. K. et al., *Can J. Chem* 58:2686 (1980).V. Amarnath and A. D. Broom, *Chemical Reviews* 77(2):183 (1977).H. Koster et al., *Nucleic Acids Research Symposium Series No. 7* (1980) pp. 39-61.Caruthers, M. H., *Science* 230:281 (1985).Zon, G. et al., *Nucleic Acids Research* 13(22):8181 (1985).Urdea, M. S. et al. *Nucleic Acids Research Symposium Ser. 16* (1985) pp. 257-260.Gao, X. et al. *Nucleic Acids Research* 13(2):573 (1985).Beaucage and Caruthers (1981) *Tet. Lett.* 22:1859-1862.

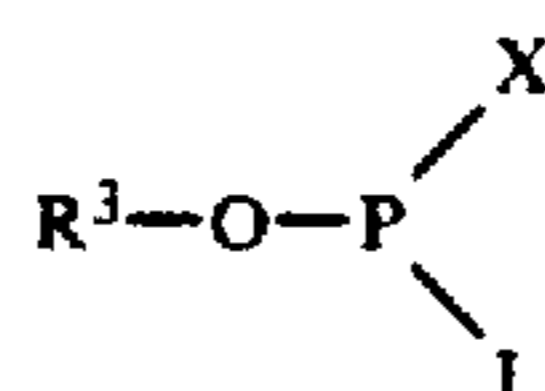
Primary Examiner—Ronald W. Griffin

Attorney, Agent, or Firm—Hamilton, Brook, Smith & Reynolds

[57]

ABSTRACT

The invention relates to a process for the preparation of oligonucleotides by the following steps: reaction of a nucleoside with a phosphine derivative, reaction of the nucleotide derivative thus obtained with a nucleoside bonded to a polymeric carrier, oxidation of the carrier-bound nucleoside-nucleotide thus obtained with formation of phosphotriester groups, blocking of free primary 5'-OH groups, elimination of a protective group from the terminal 5'-OH group, where appropriate single or multiple repetition of the abovementioned steps to introduce further nucleoside phosphate or oligonucleoside phosphate units, and cleavage of the nucleoside-carrier bond and, where appropriate, elimination of all protective groups present in the oligonucleoside phosphates. The phosphine derivative used is a compound of the general formula III



(III)

in which X and L can react with OH groups of the sugar units in the oligonucleotides, and R³ is a protective group which can be liberated by β-elimination.

19 Claims, 5 Drawing Sheets

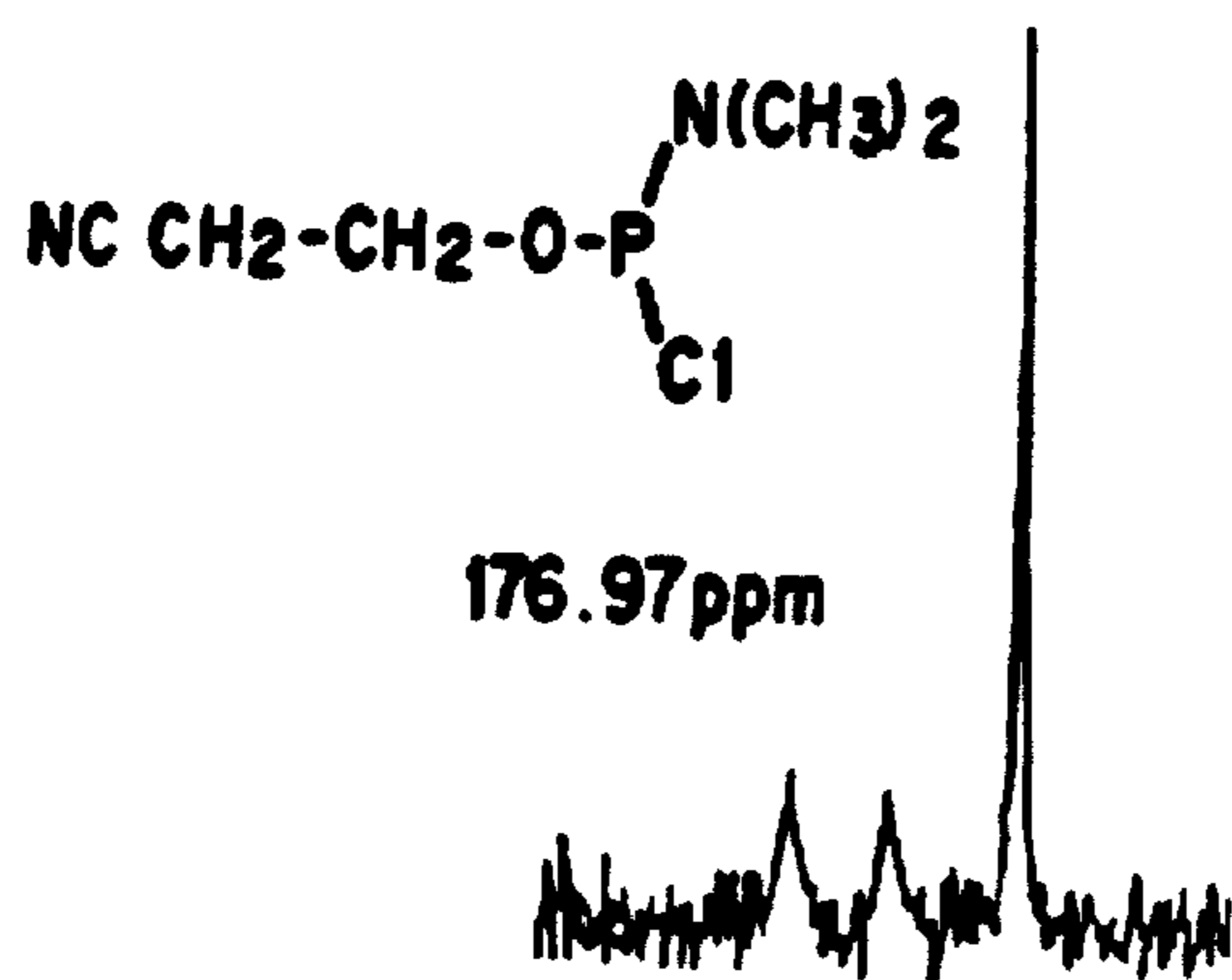


FIG. 1a

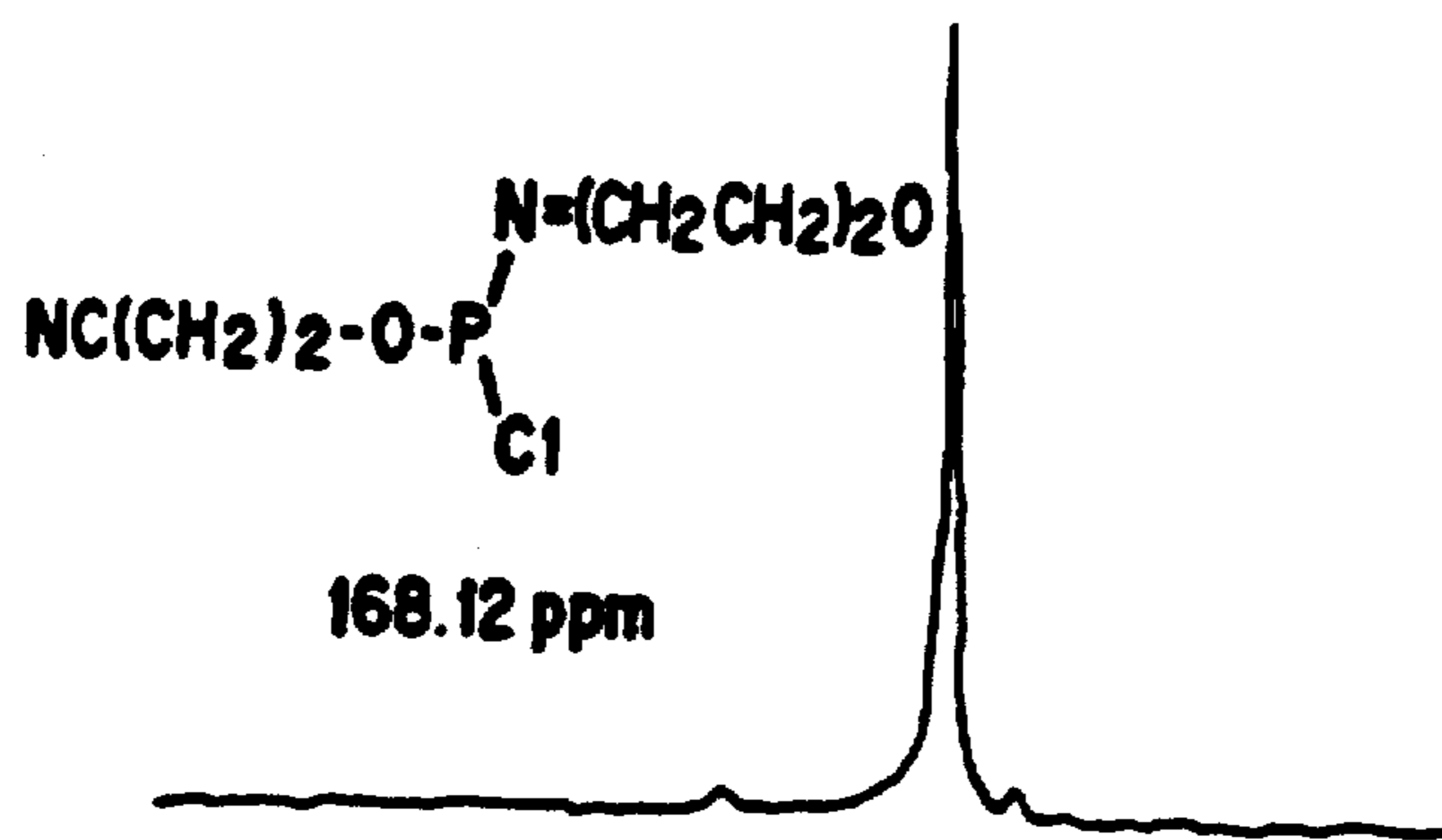


FIG. 1b

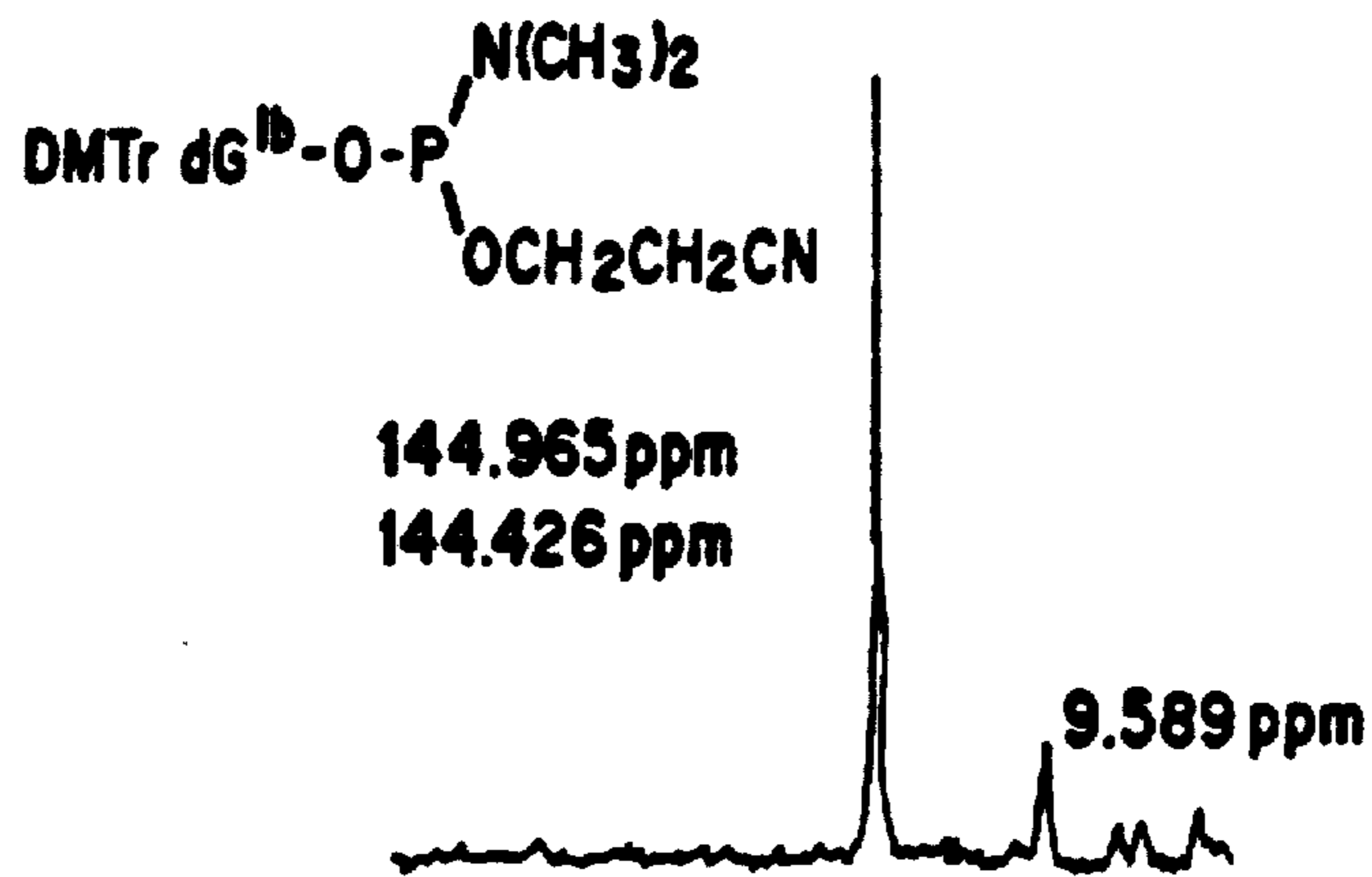


FIG. 2

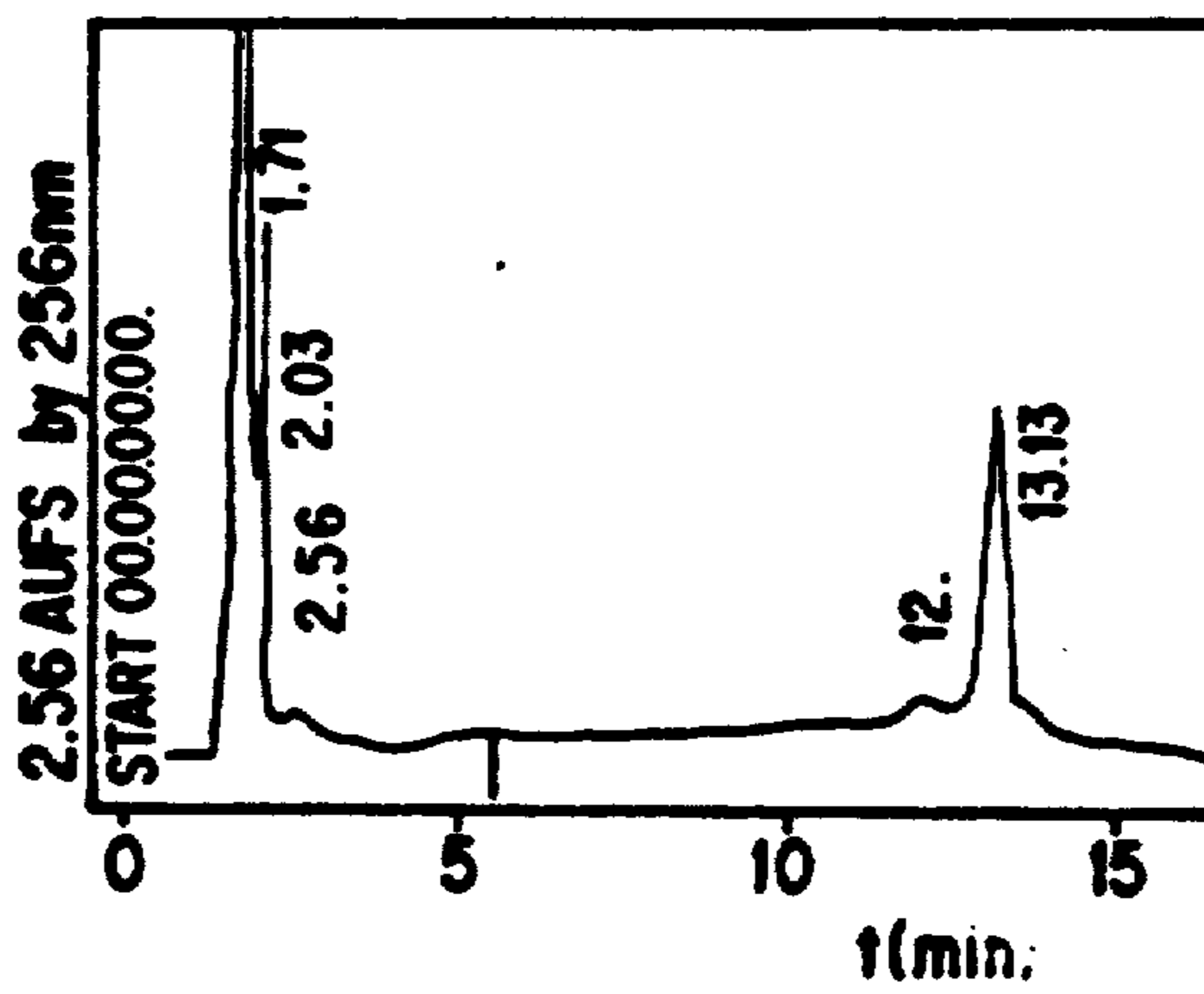


FIG. 3

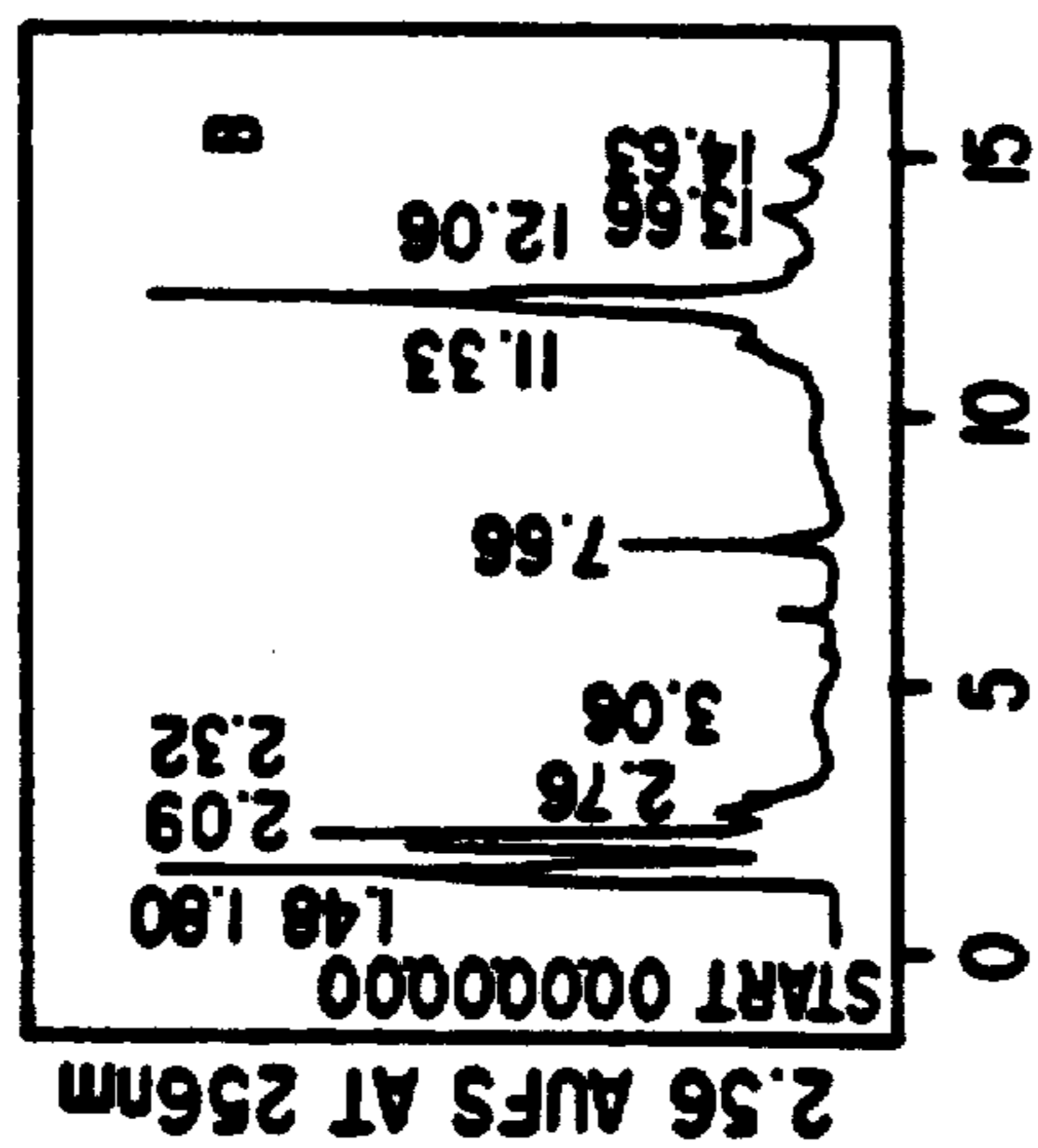


FIG. 6a



FIG. 6b

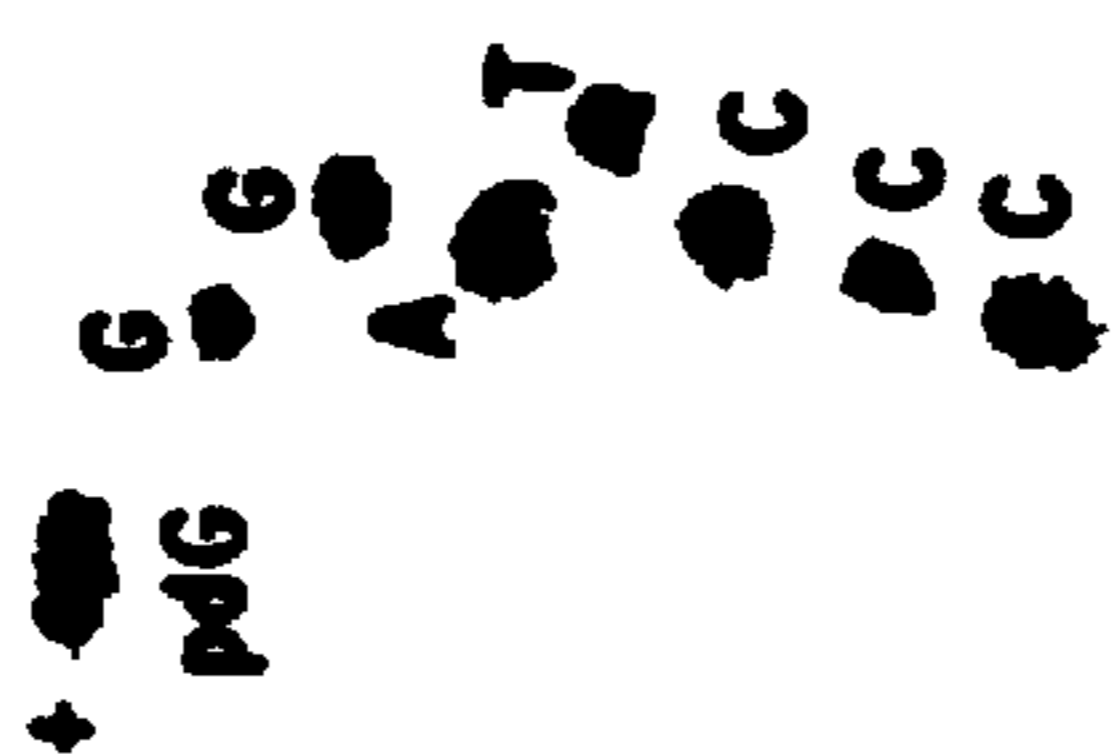


FIG. 6c

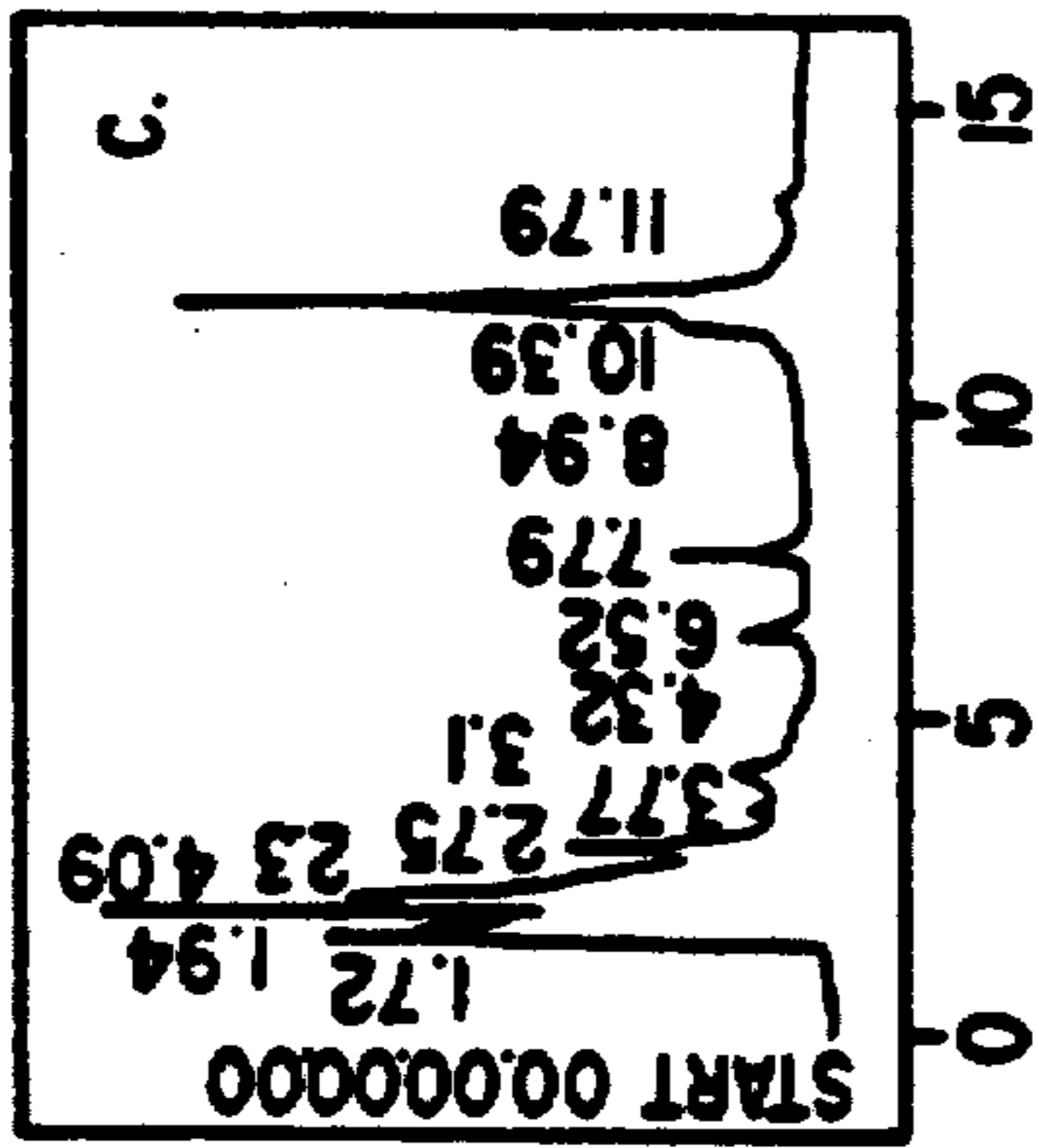


FIG. 7a



FIG. 7b

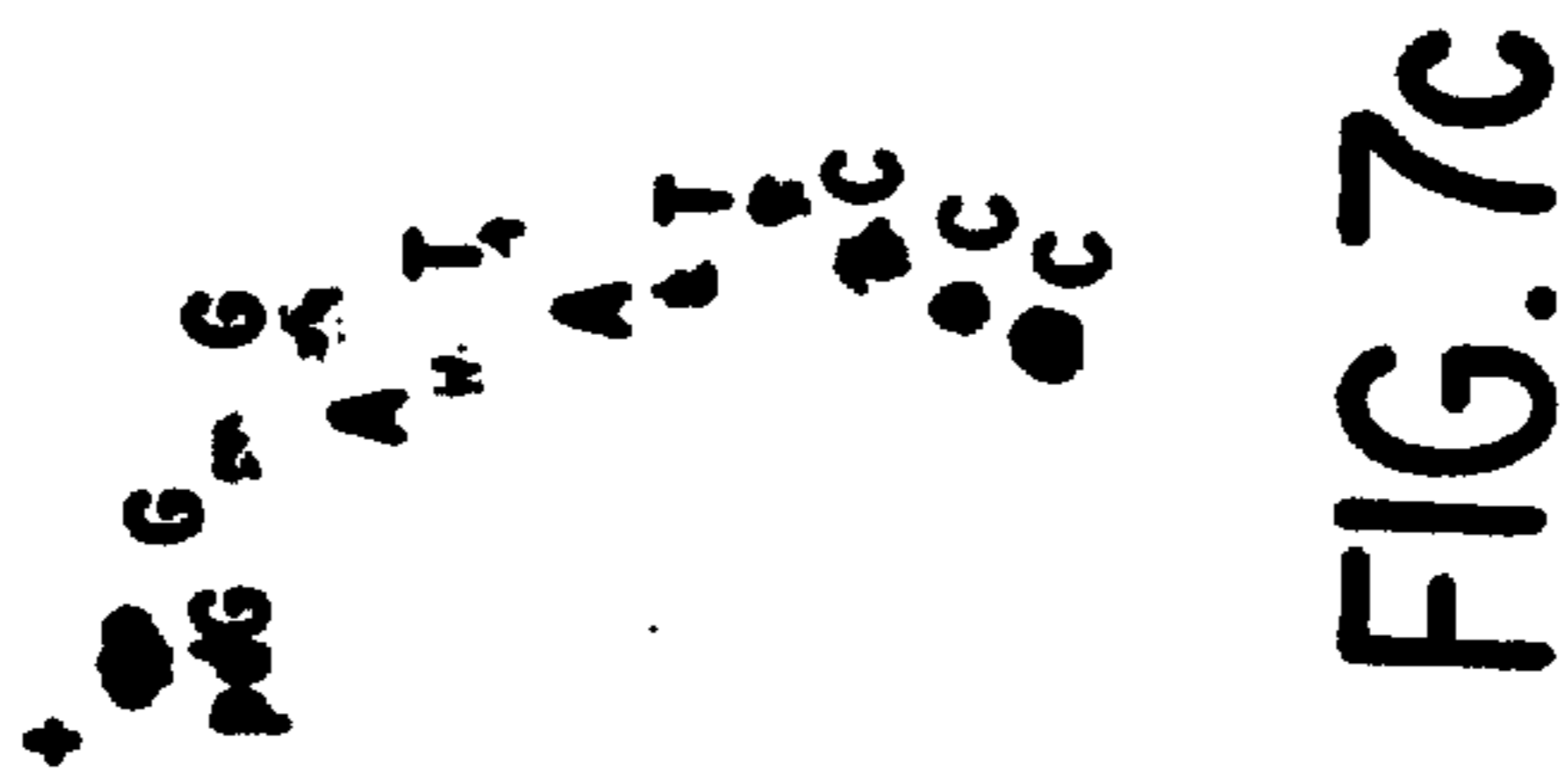


FIG. 7c

PROCESS FOR THE PREPARATION OF OLIGONUCLEOTIDES

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.

DESCRIPTION

The invention relates to a process for the preparation of oligonucleotides of the general formula I indicated in claim 1. The oligonucleotides prepared according to the invention have defined sequences and can be used as specific primers and probes and are of great importance for the synthesis of complete genes (Arzneimittelforschung 30, 3a, 548, (1980)).

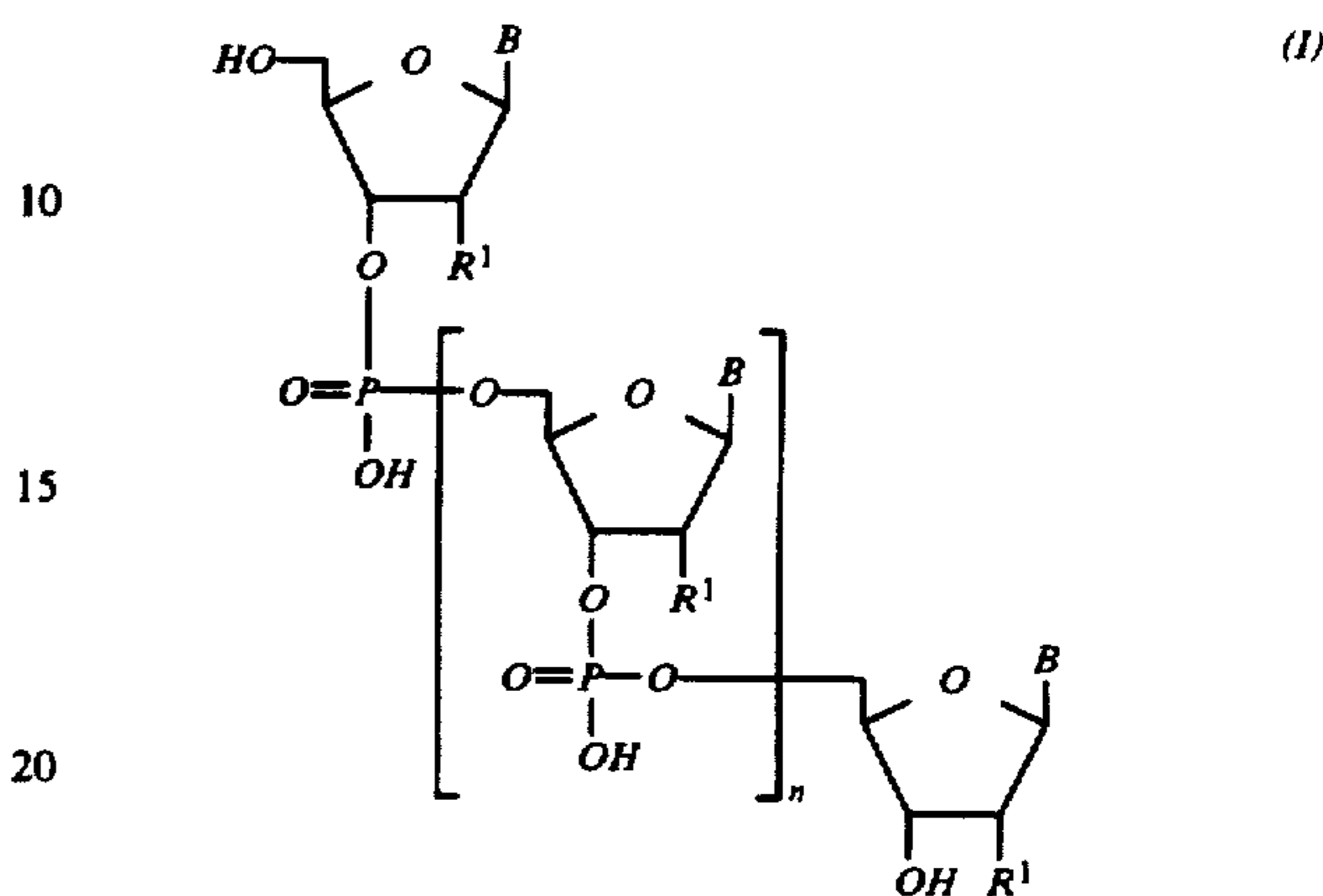
According to the most recent state of the art, oligonucleotides are prepared by either the phosphate or phosphite triester method using polymeric carriers (Nachr. Chem. Tech. Lab. 29, 230 (1981)). In order to be able to construct defined sequences, it is necessary for the individual units (nucleoside or nucleotides) to be provided with suitable protective groups. In this context, base-labile acyl groups are generally used for the protection of the exocyclic amino groups on the heterocyclic nucleobases, and a base-labile ester bond is used to attach the oligonucleotide chain to the polymeric carrier in a customary manner, and acid-labile trityl ether groups are used to protect the primary 5'-OH group. The phosphate protective group used in the phosphate triester method is customarily either the 2-chlorophenyl or the 4-chlorophenyl group, with an ester-type bond, which can only be removed by attack of a base or a nucleophile on the phosphorus atom. This type of step is inherently undesirable since it involves the risk of cleavage of the internucleotide phosphate ester bond. This risk has been greatly reduced by the use of oximate anions (Tetrahedron Lett. 19, 2727 (1978)), although these also attack the phosphorus atom in an undesired manner in the crucial step and, moreover, have the disadvantage that a relatively small amount of desired oligonucleotide is contaminated with every large amount of involatile salts which are difficult to extract. This not only makes the working up and subsequent purification of the synthesized oligonucleotide difficult but also leads to considerable material losses.

In the phosphite triester method, the methyl group with an ester-type bond is customarily used as the phosphate protective group which can be removed by attack of a nucleophile on the methyl C atom (J. Amer. Chem. Soc. 99, 3526 (1977)). Since attack on the P atom is avoided, there is likewise avoidance of the risk of cleavage of the internucleotide bond. The nucleophile customarily used is thiophenol/triethylamine, which are unpleasant to manipulate and, moreover, lead to involatile compounds which are difficult to extract and which, as mentioned above, both make work-up difficult and lead to considerable material losses.

Although the actual synthesis of oligonucleotides by the solid phase/phosphite or phosphate triester method takes place very efficiently and rapidly, the preparation of oligonucleotides of defined sequence remains very time-consuming. This is primarily due to the problems of the subsequent work-up and purification which take up a multiple of the actual synthesis time. The process

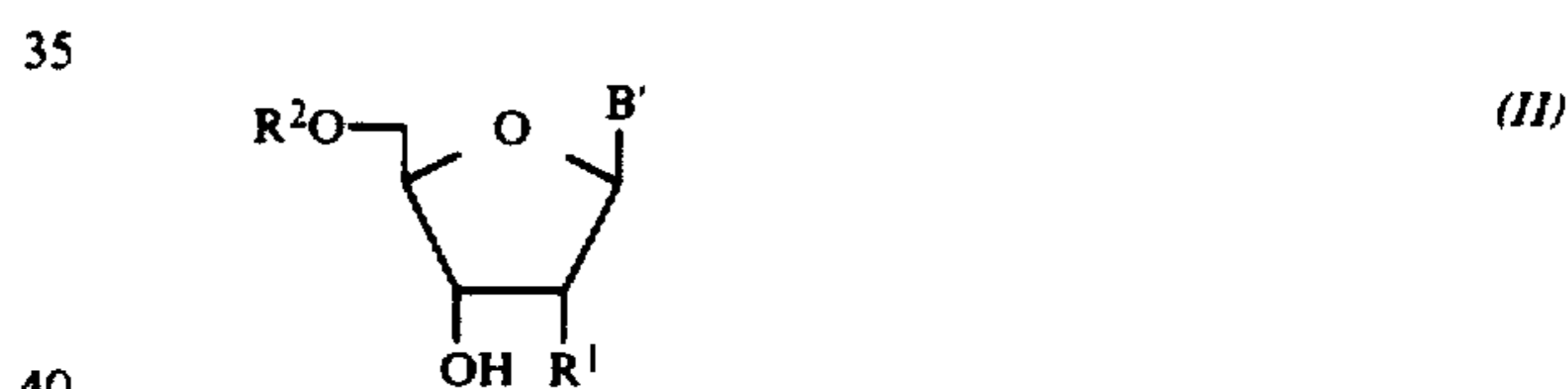
of the invention operates at this point and provides in this connection a crucial technical improvement.

In order to obtain compounds of the formula I indicated in claim 1,



in which B denotes a nucleobase, for example adenine (A), guanine (G), cytosine (C), thymine (T) or uracil (U) or their analogs, and R¹ denotes hydrogen, hydroxyl or hydroxyl which is protected by the protective groups customary in nucleotide chemistry, and n denotes an integer from 1 to 200, according to the invention a variety of defined reaction steps are carried out, as follows:

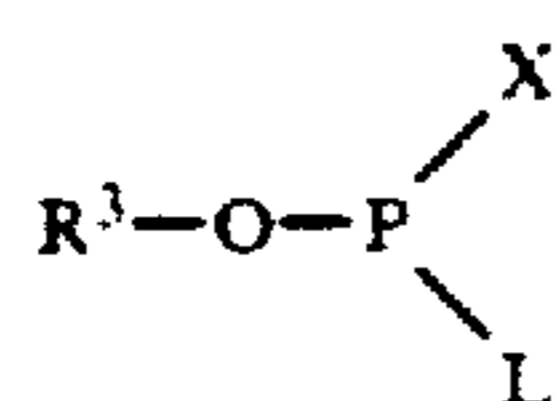
(a) Reaction of a nucleoside of the general formula II.



R¹ of the general formula II can be hydrogen; in this case the compounds of the formula I are oligodeoxynucleotides. The group R¹ can also be hydroxyl or hydroxyl which is, where appropriate, protected by the protective groups customary in nucleotide chemistry. Examples of protective groups of this type are trityl, monomethoxytrityl and dimethoxytrityl, acyl, for example acetyl, benzoyl; tetrahydropyranyl, methoxytetrahydropyranyl, o-nitrobenzyl and silyl ethers, such as, for example, t-butyl-diphenylsilyl ethers. A general review of the protective groups customary in nucleotide chemistry is to be found in, for example, Tetrahedron 1981, pages 363-369, Liebigs Ann. Chem. 1978, 839-850, and Nucleic Acids Research, Symposium Series No. 7, 1980, 39-59.

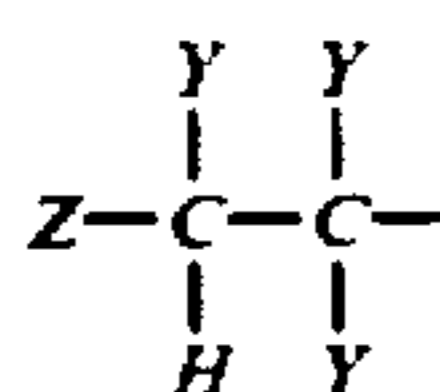
R² is likewise a protective group customary in nucleotide chemistry according to the above mentioned publications, preferably the acid-labile 4,4'-dimethoxytrityl or 4,4',4''-trimethoxytrityl group. B' can likewise have a protective group customary in nucleotide chemistry according to the above mentioned prior publications.

The nucleoside of the formula II is reacted according to the invention with a phosphine derivative of the general formula III according to claim 1.

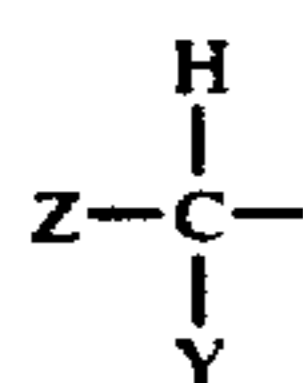


In the general formula, X denotes chlorine, bromine, CN or SCN; L denotes chlorine, bromine, CN, SCN or an amino radical of the formula—NR₂⁴ (formula VIII), where the groups R⁴ denote primary, or secondary or tertiary alkyl radicals having 1-10 carbon atoms, or together denote a cycloalkyl radical having 5-7 carbon atoms, optionally with alkyl branches, and/or can contain one or two nitrogen, oxygen and/or sulfur atoms as heteroatoms. The group L can also form a reactive heterocyclic radical, the imidazolyl, triazolyl, tetrazolyl, 3-nitro-1,2,4-triazolyl, thiazolyl, pyrrolyl, benzotriazolyl (optionally with substituents in the phenyl moiety) or benzohydroxytriazolyl (optionally with substituents in the phenyl ring) and the like.

R³ is the phosphine derivative of the general formula (III) is, according to the invention, a group of the general formula VII,

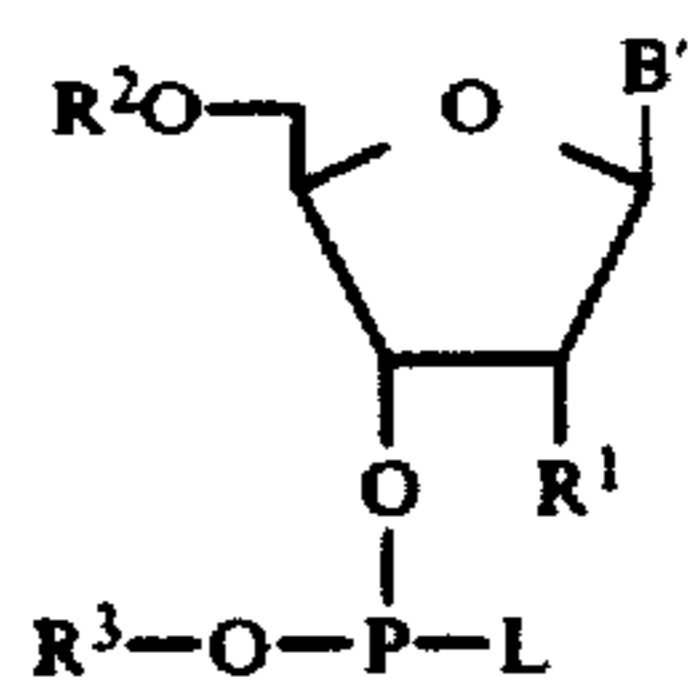


which can be removed with the aid of bases by β -elimination and in which Y denotes hydrogen, methyl or ethyl. Z represents an electron-attracting group, for example, halogen, such as fluorine, chlorine or bromine, CN or NO₂. Z can also denote phenyl, phenylthio, phenylsulfoxy or phenylsulfonyl, it being possible for the phenyl radicals to be substituted in the o,o'-position and/or p-position with halogen, CN or NO₂. It is also possible for one of the groups CF₃, CCl₃ or CBr₃ to replace the group



The reaction according to step a takes place in the presence of an organic base.

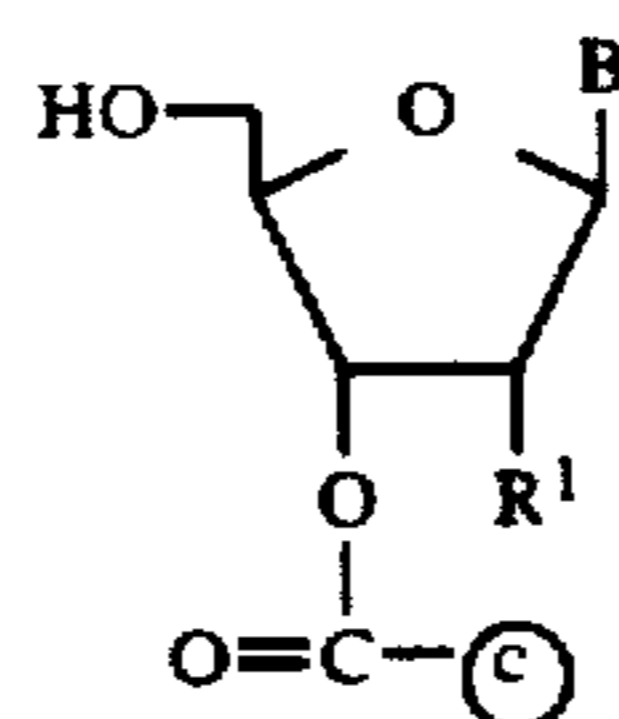
(b) Reaction of the nucleoside-[phosphorous] phosphorus acid derivative, of the formula IV, obtained in step a.



The reaction of the compound according to formula IV is carried out with a nucleoside of the general formula V according to claim 1, which is bound to a polymeric carrier.

III

5

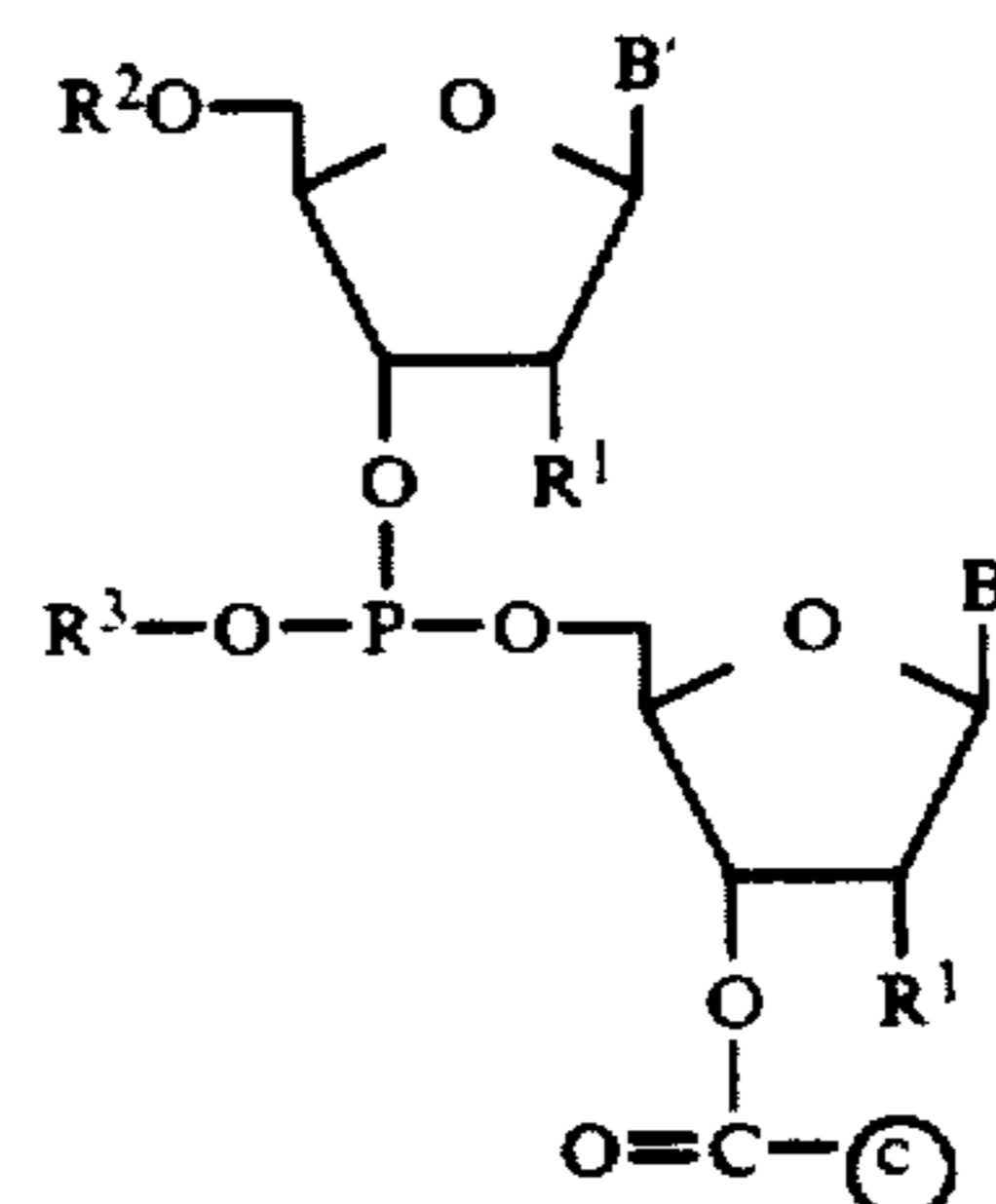


(V)

It is possible to use soluble or insoluble, that is to say crosslinked, polymeric carriers, for example modified silica gel, glass, especially "controlled pore glass", polyester, polyamide, polyvinyl alcohol, polysiloxane, polystyrene or the like. Ester bonds are suitable and preferred for the attachment between the carrier and the nucleoside, including those derived from the levulinyl or β -benzoylpropionyl radical; the latter ester bonds can be cleaved with hydrazine under neutral conditions. The acid-labile trityl ether bond, optionally with substituents in the phenyl rings, is also suitable as a method of attachment, compare Liebigs Ann. Chem. 1974, 959.

(c) Oxidation of the carrier-bond nucleotide-nucleoside, of the general formula VI, obtained in step b.

25



(VI)

30

35

40

45

50

55

60

65

Oxidation leads to a phosphate group; this can be carried out with, for example, iodine/H₂O, H₂O₂ or organic peracids or, in general, by oxidation by introduction of O, S or Se.

(d) Blocking of free primary 5'-OH groups which have not been reacted in the reaction according to step b (in the product of the formula V).

These free hydroxyl groups are blocked with a permanent protective group, for example by reaction with acetic anhydride.

(e) Elimination of the protective group(s) R².

The elimination is carried out using, for example, a protonic acid or Lewis acid, such as ZnBr₂ or dialkylaluminum chloride, when R² represents a trityl group or a methoxy derivative thereof.

(f) Introduction of further nucleoside phosphate or oligonucleoside phosphate units.

Steps a-e can be repeated to introduce at least one nucleoside phosphate moiety. Of course, when oligonucleoside phosphate units are employed, the chains are lengthened by more than one nucleoside phosphate unit.

(g) Elimination of all protective groups.

This elimination can be carried out in such a manner that, using aqueous ammonia, in one step the N-acyl groups on the heterocyclic bases, the ester bond between the oligonucleotide and the carrier (the latter can, where appropriate, also be cleaved with hydrazine under neutral conditions) and the phosphate protective group are eliminated by β -elimination in accordance

with the general scheme 1 at the end of the description. An oligonucleotide having only a 5'-terminal trityl protective group is then obtained, and this can be purified directly in a manner known per se, after removal of the volatile base (ammonia), by high-pressure liquid chromatography (HPLC) on reverse phase material.

The intermediates of the general formula IV according to claim 1 are new compounds. They are in the form of very stable compounds which can be prepared in the pure form and are easy to manipulate but nevertheless are very reactive in the sense of forming internucleotide bonds. The use of R³ as a protective group which can be removed by bases via β -elimination makes it possible for the first time to eliminate all the protective groups, apart from the 5'-trityl group, in one step where, in an advantageous manner, by the use of volatile bases the desired oligonucleotide is contaminated with foreign materials to only a very small extent and thus directly afterwards can be purified by reverse phase HPLC due to the hydrophobic 5'-trityl group which is still present.

A further advantage of the process of the invention results from the fact that, due to the removal of the protective group by β -elimination, no attack on the P-atom takes place and thus none of the newly formed internucleotide bonds can be cleaved during the deprotection. Thus, the process of the invention takes very much less time and leads to overall purer products than do the processes hitherto available.

The invention is illustrated in detail below by means of examples, the phosphine derivatives used being those in which R³ is a β -cyanoethyl group. Details of the reaction and physical characteristics of the compounds prepared can be seen in schemes 2 and 3, Table 1, and FIGS. 1-7 at the end of the description.

EXAMPLE 1

Preparation of phosphine derivatives of the general formula III:

β -Cyanoethyl phosphoramidochloridite:

A general summary of the reaction can be seen in scheme 2.

Apart from some improvements, dichloro- β -cyanoethoxyphosphine (1) is prepared as in Can. J. Chem. 58, 2686 (1980):

300 ml of ether and 79.0 g (1 mol) of pyridine are added through a dropping funnel to 137.5 g (1.0 mol) of PCl₃ in a three-neck flask; the mixture is cooled to -78° C. under argon. Then a solution of 71.0 g (1 mol) of β -cyanoethanol in 150 ml of dry ether is added dropwise over the course of 1 to 1.5 hours. The cooling bath is removed; stirring is continued at room temperature for a further 3 hours (where necessary, another 300 ml of ether are added in order to ensure better stirrability). The stirrer and dropping funnel are removed under argon; the mixture is stored at 0° C. overnight. The solid salts are removed under argon; the precipitate is washed twice with 75 ml of ether each time. The combined organic phases are concentrated in vacuo; the residue is finally distilled in vacuo; boiling point 70°-75° C./0.4 mm Hg.

β -Cyanoethyl phosphoramidochloridite (3):

A solution of 17.2 g (0.1 mol) of β -cyanoethyl phosphorodichloridite (1) in 60 ml of ether is added dropwise, over the course of 1 to 1.5 hours, to a solution of the N-trimethylsilylated secondary amine (0.1 mol) or secondary amine (0.2 mol) in 30 ml of ether at -20° C. under argon. After stirring at room temperature for 20

hours, the amine hydrochloride is removed; the remaining solution is concentrated. The residue is finally distilled in vacuo in a short-path distillation apparatus.

The physical properties of the compounds thus obtained are summarized in Table 1.

FIGS. 1a, 1b and 1c show ³¹P NMR spectra of three different β -cyanoethyl phosphoramidochloridites.

The N-morpholine derivative is too unstable to heat for distillation to be possible. Nevertheless, the preparation is so pure that the residue can be used directly for the preparation of the activated nucleoside derivatives. The purity is usually greater than 95% according to the ³¹P NMR spectra.

Nucleoside β -cyanoethyl phosphoramidites:

The preparation of the appropriately protected nucleoside β -cyanoethyl phosphoramidites can be seen in scheme 3.

The synthesis is analogy to Tetrahedron Lett. 22, 1859 (1981), with some improvements, provides good yields.

3.0 mmol of the N-protected 5'-dimethoxytritylated deoxynucleoside are dried azeotropically using THF/toluene, dissolved in 15 ml of dry THF, and 12.0 mmol of N,N,N-diisopropylethylamine are added. 6.0 mmol of the β -cyanoethyl phosphoramidochloridite are added dropwise to the solution under argon, with vigorous stirring, over the course of 2 minutes. After a short time (2 to 5 minutes), the amine hydrochloride precipitates out. The suspension is stirred for a further 30 to 40 minutes. The amine hydrochloride is filtered off under argon and thoroughly washed with dry THF (10 to 15 ml). The entire organic phase is concentrated and dissolved in argon-saturated ethyl acetate (100 ml). The organic phase is extracted twice with 50 ml each time of argon-saturated 10% aqueous sodium carbonate solution. The organic phases are dried with sodium sulfate and evaporated under reduced pressure to give a foam. The foam is dissolved in a little ethyl acetate or toluene and precipitated in n-hexane at -78° C. The activated nucleosides are stable for several months when stored at -20° C. under argon.

FIG. 2 shows the ³¹P NMR spectrum of one of the activated deoxynucleosides.

Synthesis of d(CGGTACCG)

100 mg of "controlled pore glass" (CPG) loaded with a total of 8 μ mol of N-isobutyryldeoxyguanine (compare Tetrahedron Lett. 24, 747 (1983)) are consecutively condensed with the 5'-dimethoxytritylated N-acylated β -cyanoethyl N,N-diisopropylphosphoramidites of the deoxynucleosides C, C, A, T, G, G and C, in each case 20 to 25 equivalents of the phosphoramidite in acetonitrile being activated with 75-80 equivalents of sublimed tetrazole. The condensations are complete within 30 minutes at the most; the coupling yield is greater than 94%. After each condensation, oxidation with I₂/H₂O and blocking of unreacted 5'-OH groups with acetic anhydride are carried out. Then the dimethoxytrityl group is eliminated either with 3% trichloroacetic acid in nitromethane/1% methanol or ZnBr₂/nitromethane/1% H₂O.

The overall yield of the protected octanucleotide at the end of all condensation steps is 55% based on carrier-bound deoxyguanosine.

Complete deprotection and cleavage off from the carrier is achieved in one step by reaction of the glass

beads with concentrated aqueous ammonia (3 ml) at 50° C. for 16 hours. The glass beads are then thoroughly washed with 50% aqueous methanol (3 times with 3 ml each time). The liquid phase is removed by evaporation (removal of the methanol) and freeze-drying. Then an aliquot is filtered through a millipore filter and purified by HPLC or RP 18 as can be seen in FIG. 3.

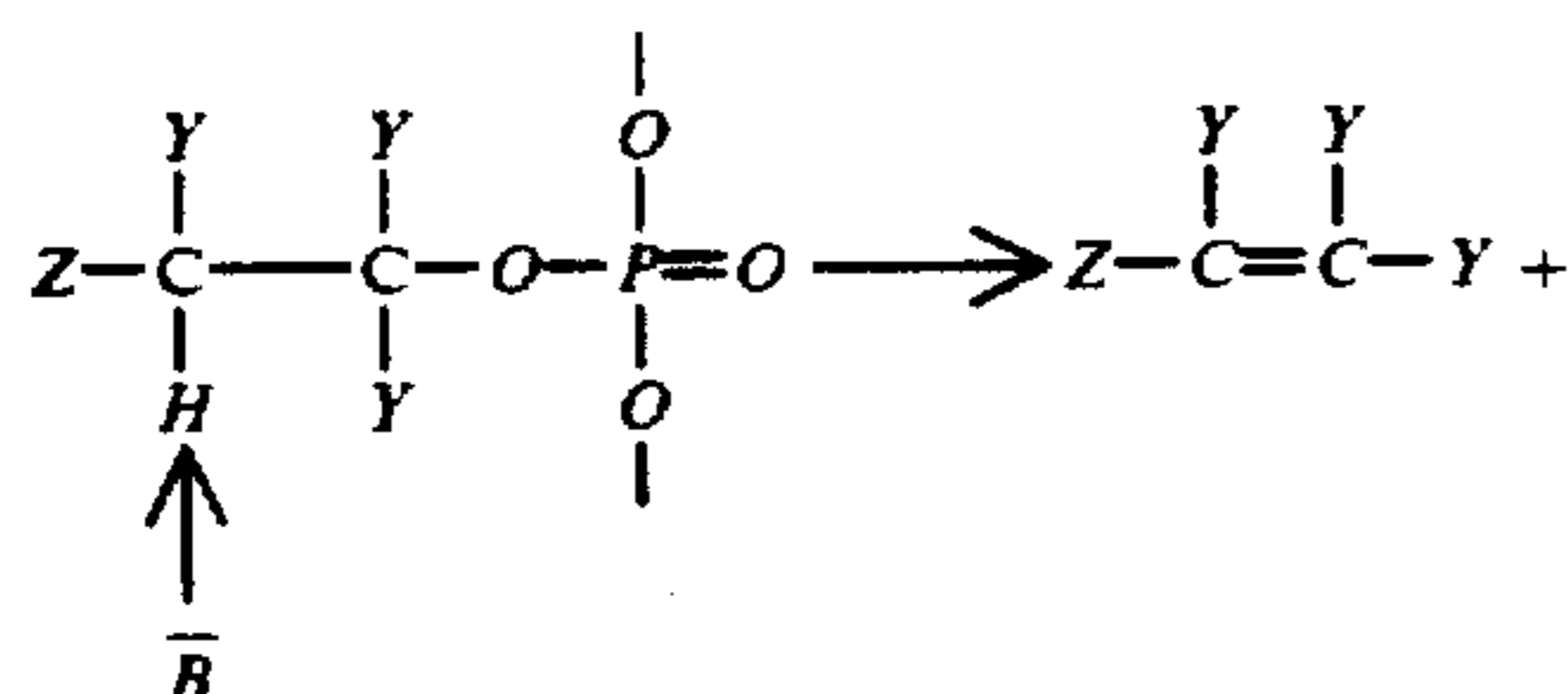
The fractions which contain the 5'-dimethoxytritylated oligonucleotide are collected; the volatile buffer is removed in a rotary evaporator in vacuo, 1 ml of 80% strength acetic acid is added to the residue. After 45 minutes at room temperature, the acetic acid is removed by freeze-drying.

The material thus obtained is phosphorylated in the customary manner (Liebig's Ann. Chem. 1978, 982) with T4-polynucleotide kinase and γ -³²P-ATP. The resulting product is characterized by polyacrylamide gel electrophoresis comparing with a homo-oligo-dT chain length standard (Nucleic Acids Res. 6, 2096 (1979), FIG. 4) and by sequencing according to FIG. 5 (Liebig's Ann. Chem. 1978, 982).

FIGS. 6a to 6c show the results (HPLC, gel electrophoresis, sequencing) of the synthesis of d(GGGATCCC) using the nucleoside β -cyanoethyl N,N-dimethylphosphoramidites, FIGS. 6a to 6c show the results (HPLC, g, electrophoresis, sequencing) of the synthesis of d(GGGATATCCC) using the nucleoside β -cyanoethyl N,N-morpholinophosphoramidites.

The results given in FIGS. 3, 6a and 7a were obtained by using a gradient from 10 to 25 vol. % CH₃CN, 5 min, and 25 to 29 vol. % CH₃CN, 30 min, in 0.1M triethylammonium acetate at pH 7.0.

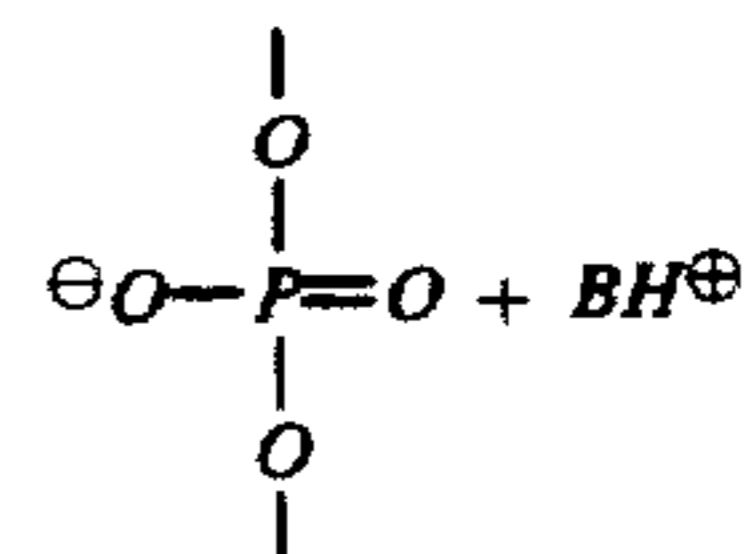
Scheme 1
Removal of the Group R by β -Elimination



-continued

Scheme 1

Removal of the Group R by β -Elimination



SCHEME 2

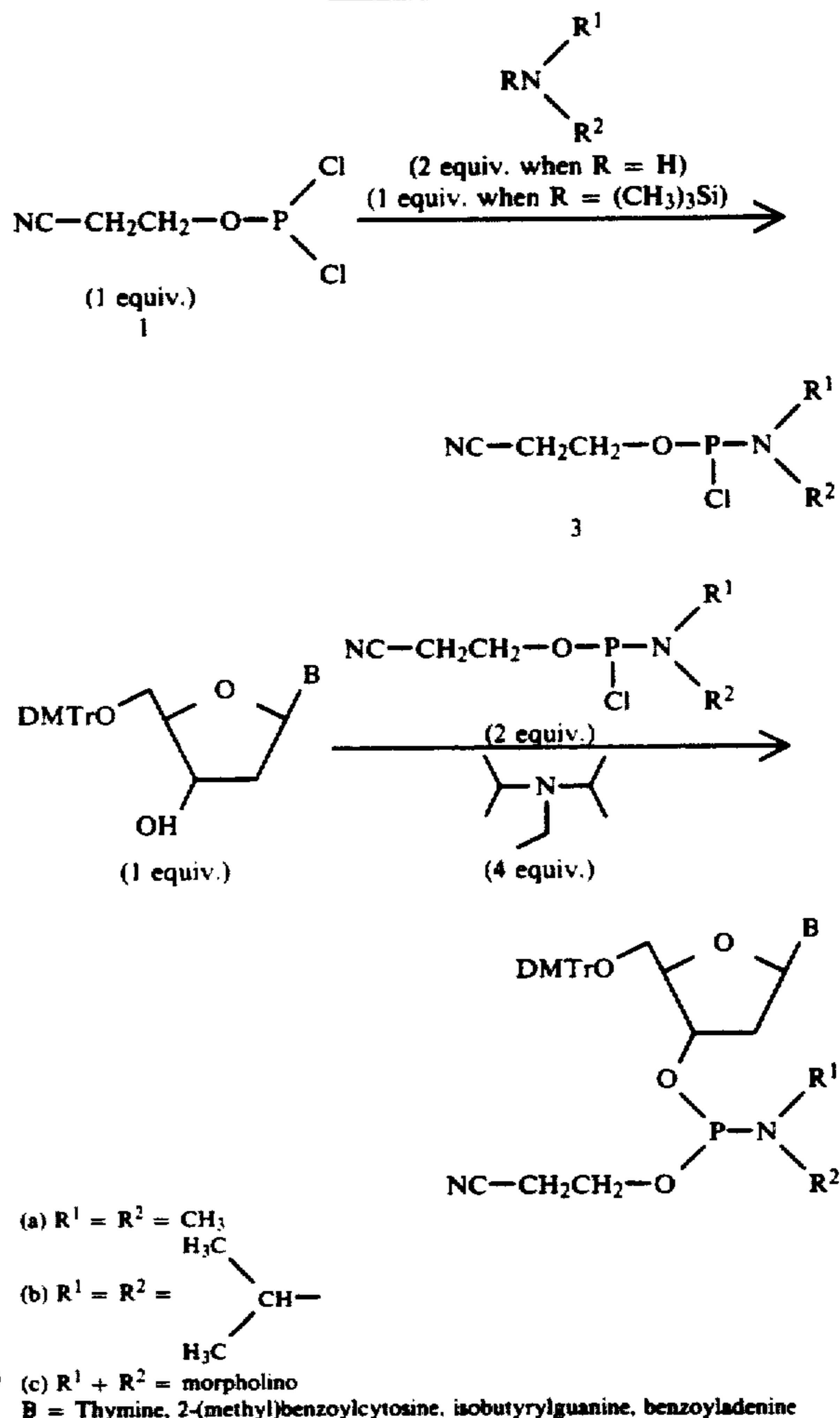


TABLE 1

| Compound | Physical data of β -cyanoethyl phosphoramidochloridites | | |
|--|---|--|---|
| | 3a L = N,N-dimethylamino | 3b L = N,N-diisopropylamino | 3c L = N-morpholino |
| Boiling point | 90-92°/0.6 mm | 103-5°/0.06 mm | — |
| Chemical shift ⁽²⁾ in ³¹ P NMR in CH ₃ CN | 175.97 ppm | 179.82 ppm | 168.22 ppm |
| Chemical shift in ¹ H NMR in ppm | 4.01, 4.17(2t, P—OCH ₂ , 2H) 2.71(t, —CH ₂ —CN, 2H) 2.7(d, N(CH ₃) ₂ , 6H) | 4.02, 4.2(2t, POCH ₂ , 2H) 3.8(m, N(CH ₂) ₂ , 2H) 2.77(t, —CH ₂ CH, 2H) 1.29(d, N—CH(CH ₃) ₂ , 12H) | 3.96, 4.1(2t, POCH ₂ , 2H) 3.67(t, O(CH ₂) ₂ , 4H) 3.17(m, P—N(CH ₂) ₂ , 4H) 2.74(t, CH ₂ —CN ₂ , 2H) |
| Mass spectrum | $\left(\frac{m}{e}\right)^+ = 180, 182(+2), 145$ | $\left(\frac{m}{e}\right)^+ = 236, 238(+2), 201$ | $\left(\frac{m}{e}\right)^+ = 222, 224(+2), 187$ |

TABLE 1-continued

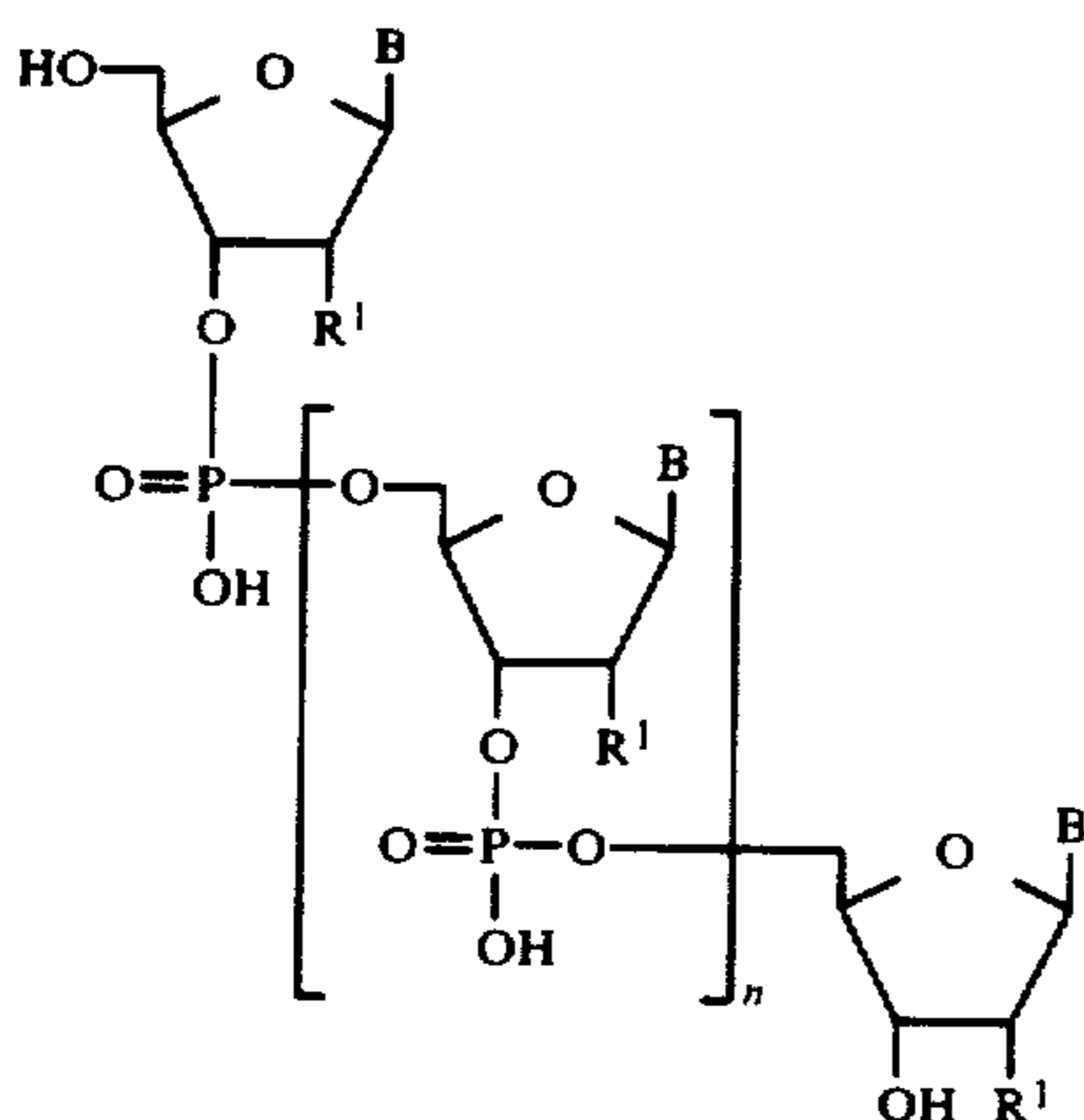
| Physical data of β -cyanoethyl phosphoramidochloridites | | | |
|---|--|--|---|
| Compound | 3a L = N,N-dimethylamino | 3b L = N,N-diisopropylamino | 3c L = N-morpholino |
| | (-Cl), 136(-C ₂ H ₆ N), 110 (-C ₃ H ₄ NO) | (-Cl), 166(-C ₃ H ₄ NO), 136(-C ₆ H ₁₄ N) | (-Cl), 152(-C ₃ H ₄ NO), 136(-C ₄ H ₈ O) |

(1) The crude product after removal of amine hydrochloride and compounds volatile under high vacuum at room temperature has a purity of 93-95% according to the ³¹P NMR spectrum

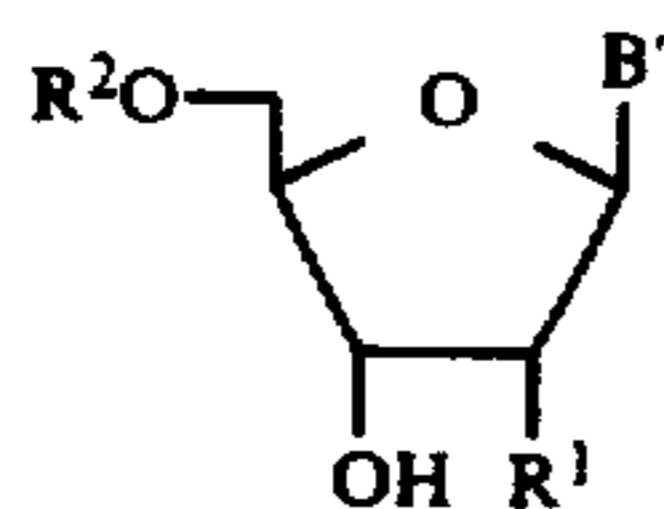
(2) The chemical shifts are determined in acetone-d₆ with 80% strength H₃PO₄ as the external standard.

We claim:

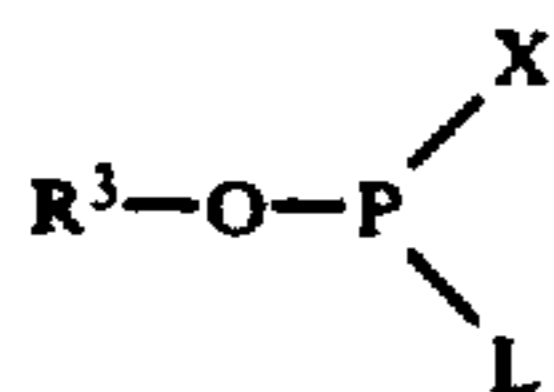
1. A process for the preparation of oligonucleotides of the [general] formula I



in which B denotes a nucleoside base, R¹ denotes hydrogen, hydroxyl or hydroxyl which is protected by, where appropriate, a removable protective group and n denotes an integer from 1 to 200, comprising the steps of
(a) reacting a nucleoside of the [general] formula II

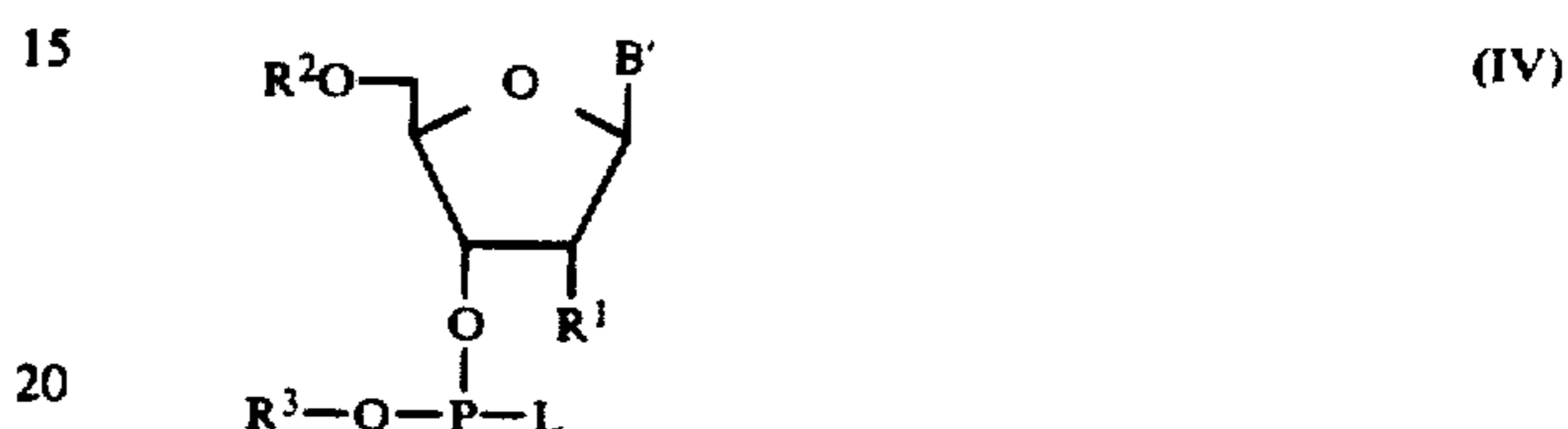


in which R¹ as defined as above, and R² denotes a removable protective group and B' denotes the nucleotide base B protected by the protective groups which can be eliminated, with a phosphine derivative of the [general] formula III

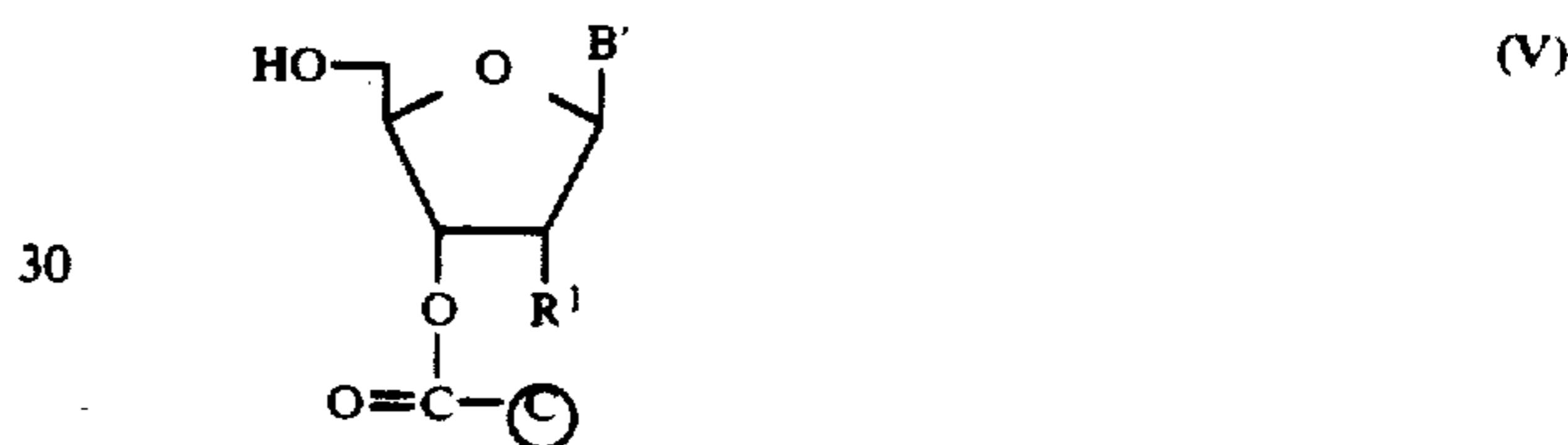


in which R³ is a protective group which can be eliminated, and X and L are groups which react with hydroxyl groups in the sugar moieties of the nucleotides or nucleosides, in the presence of a base to thereby form a nucleotide phosphite

(b) reacting the nucleotide phosphite obtained in step (a) and represented by the formula IV:

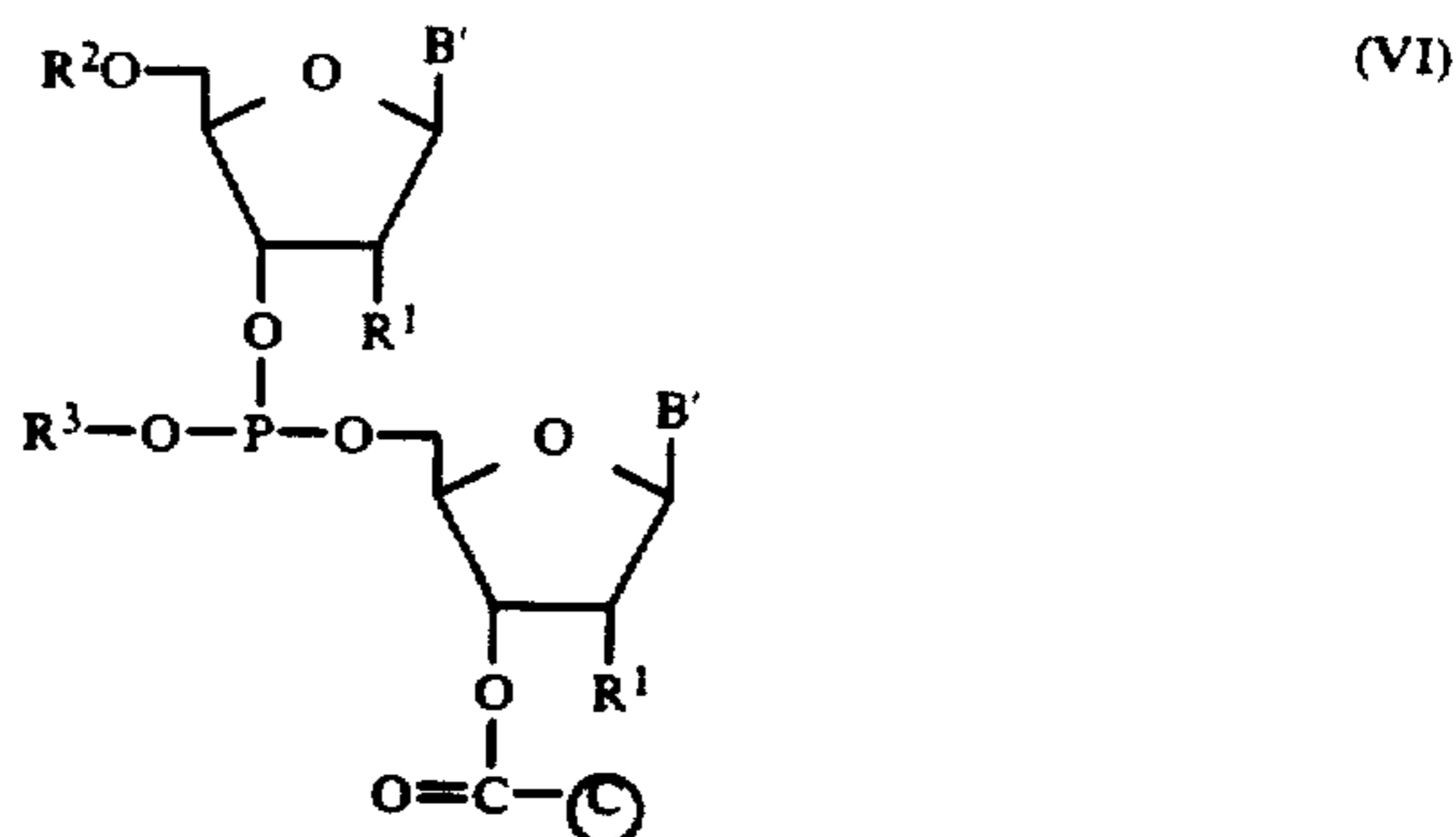


in which B', R¹, R², R³ and L are as defined above, with a nucleoside, of the [general] formula V, bound to a polymeric carrier



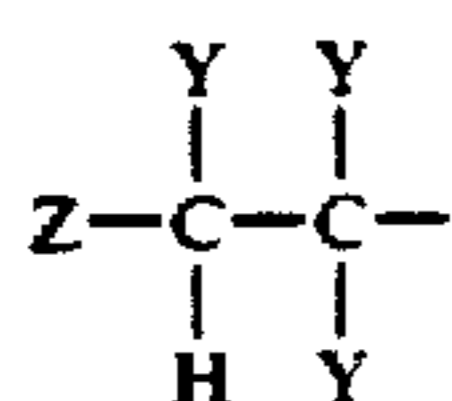
in which B' and R¹ are as defined above and C denotes the polymeric carrier;

(c) oxidizing the carrier-bound nucleoside-nucleotides obtained in step (b) and represented by the formula:



in which B', R¹, R², R³ and C are as defined above, with formation of phosphotriester groups,

(d) blocking free primary 5'-OH groups, which have not been reacted in the reaction according to step (b), with permanent protective groups;
(e) eliminating the protective group R²;
(f) optionally repeating steps (a) to (e) to introduce further nucleoside phosphate or oligonucleoside phosphate units; and
(g) cleaving the nucleoside carrier bond and optionally eliminating the protective groups present in the oligonucleoside phosphates, which process comprises using in step (a) as the phosphine derivative of the [general] formula III a compound in which R³ denotes a group of the formula VII



in which the groups Y, which can be identical or different, represent hydrogen, methyl, and/or ethyl and Z represents an electron-attracting group, where, in the phosphine derivative of the formula III, X is chlorine, bromine, CN or SCN and L is CN or SCN, a secondary amino radical of the formula (VIII)



where the groups R^4 are primary, secondary, or tertiary alkyl radicals having 1-10 carbon atoms, or together form a cycloalkyl radical having 5-7 carbon atoms, which can contain one or two nitrogen, oxygen, or sulfur atoms as heteroatoms, or are imidazole, triazole, tetrazole, 3-nitro-1,2,4-triazole, thiazole, pyrrole, benzotriazole, benzohydroxytriazole, imidazole substituted in the phenyl moiety, triazole substituted in the phenyl moiety, tetrazole substituted in the phenyl moiety, 3-nitro-1,2,4-triazole substituted in the phenyl moiety, thiazole substituted in the phenyl moiety, pyrrole substituted in the phenyl moiety, benzotriazole substituted in the phenyl moiety, or benzohydroxytriazole substituted in the phenyl moiety.

2. The process as claimed in claim 1, in which is used a phosphine derivative of the formula III in which X is chlorine or bromine, and L is a secondary amino radical of the formula (VIII)

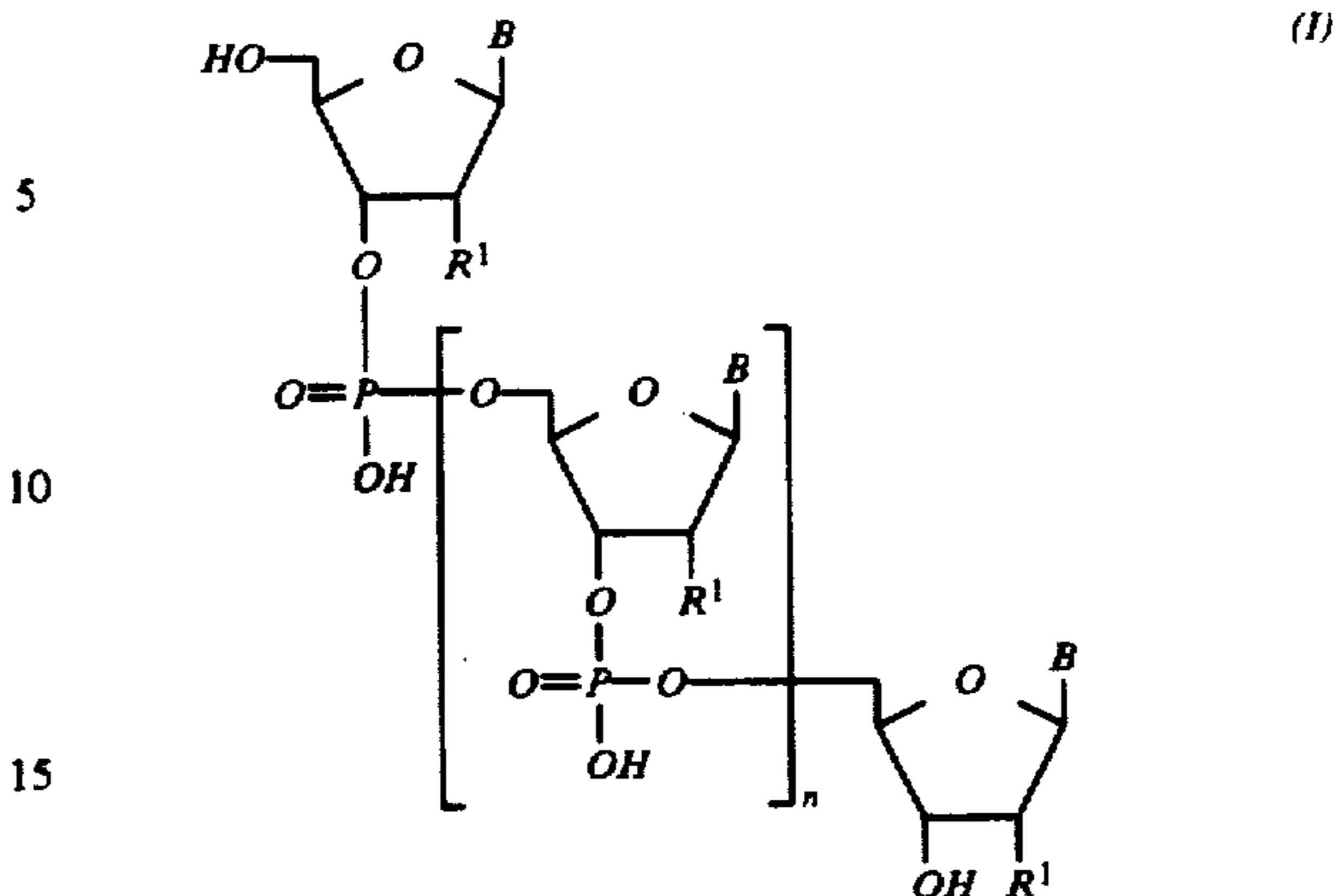


where the groups R^4 are primary, secondary or tertiary alkyl radicals having 1-10 carbon atoms, or together form a cycloalkyl radical having 5-7 carbon atoms, which can contain one or two nitrogen, oxygen or sulfur atoms as heteroatoms, or are imidazole, triazole, tetrazole, 3-nitro-1,2,4-triazole, thiazole, pyrrole, benzotriazole, benzohydroxytriazole, imidazole substituted in the phenyl moiety, triazole substituted in the phenyl moiety, tetrazole substituted in the phenyl moiety, 3-nitro-1,2,4-triazole substituted in the phenyl moiety, thiazole substituted in the phenyl moiety, pyrrole substituted in the phenyl moiety, benzotriazole substituted in the phenyl moiety, or benzohydroxytriazole substituted in the phenyl moiety.

3. The process as claimed in claim 1 or 2, in which is used a phosphine derivative of the formula (III) in which X is chlorine, L is an N,N-dimethylamino, diethylamino or -diisopropylamino group or N-morpholino group, and R^3 is a β -cyanoethyl group.

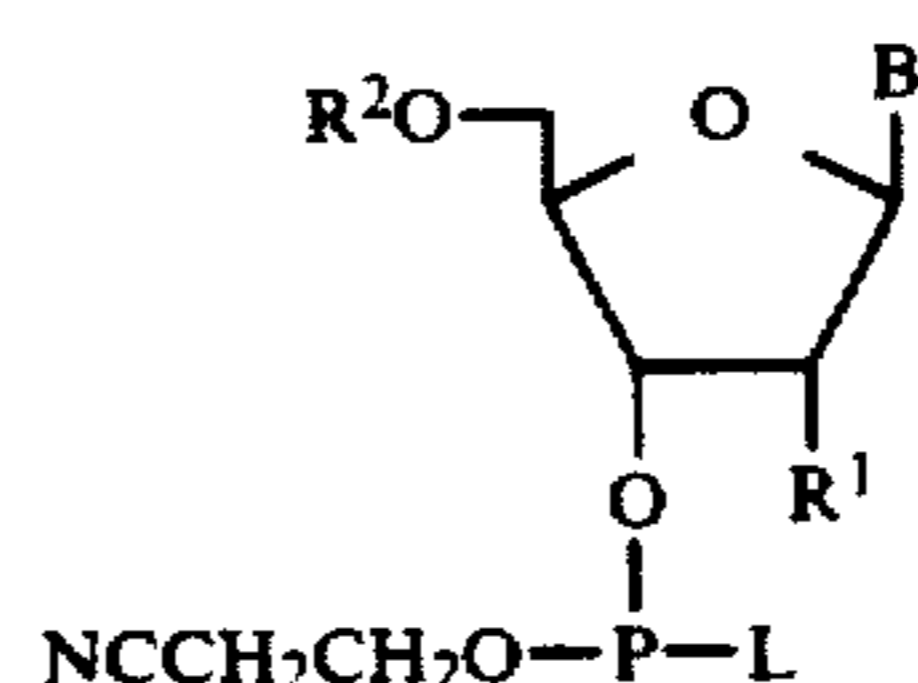
4. A method of preparing oligonucleotides of the [general] formula:

(VII)

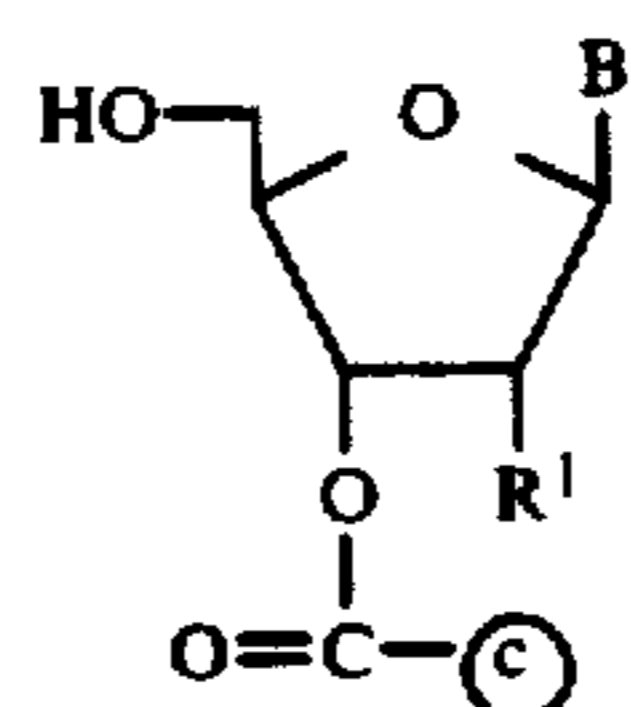


wherein B is a nucleoside base, R^1 is hydrogen, hydroxyl or hydroxyl which is protected by removable nucleoside protective groups, and n denotes an integer from 1 to 200, comprising the steps of:

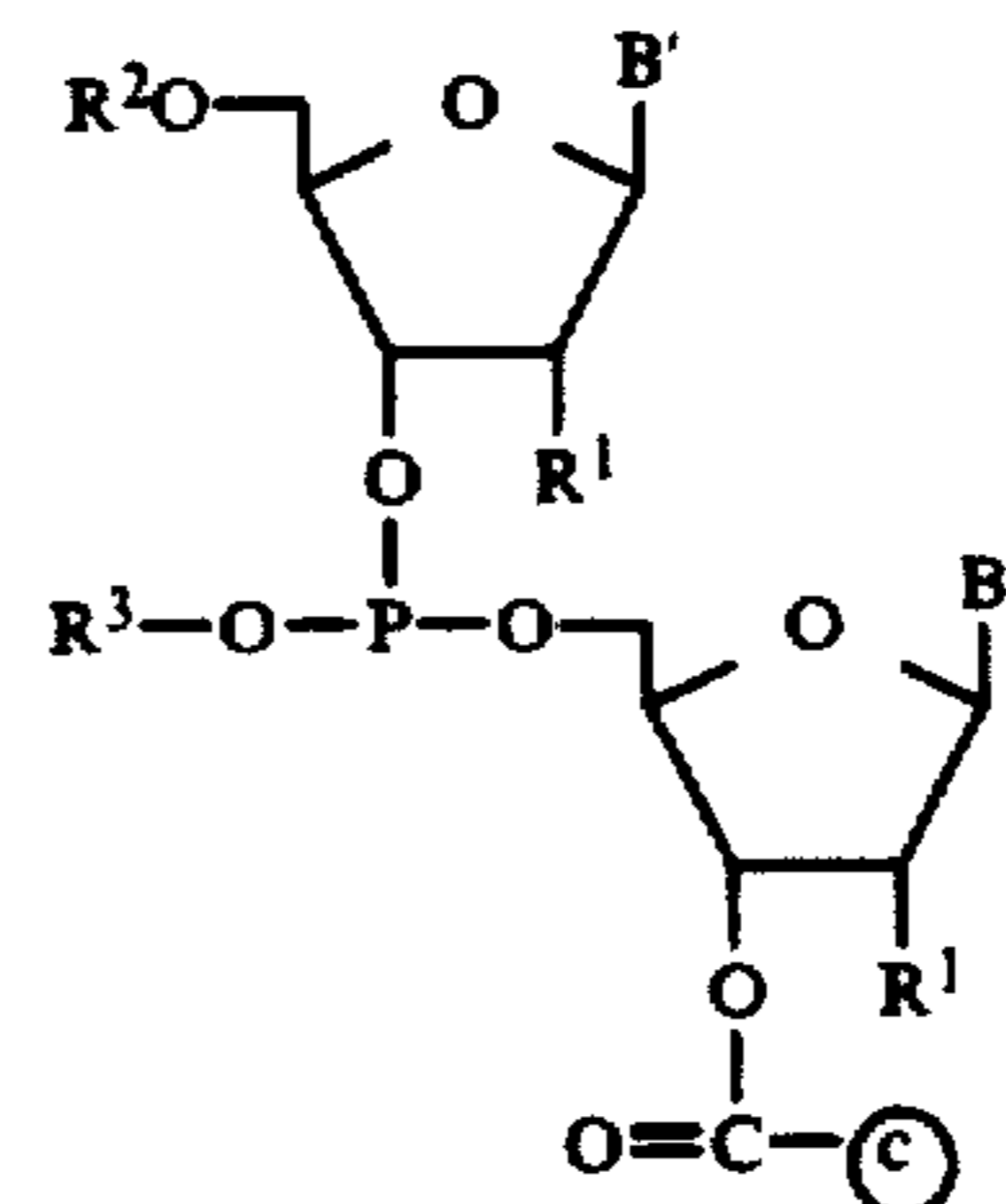
(a) reacting a nucleotide phosphite represented by the formula:



wherein B' is a nucleoside base B protected by, where appropriate, a base protective group which can be eliminated, R^1 is as defined above, R^2 is 4,4' dimethoxytrityl or 4,4',4'' trimethoxytrityl; and L is N,N-dimethylamino, N,N-diethylamino, N,N-diisopropylamino, or N-morpholino, with a nucleoside bound to a polymeric carrier, of the [general] formula:



wherein B' and R^1 are as defined above and C represents the polymeric carrier, to produce a carrier bound nucleoside-nucleotide of the formula:



wherein B' , R^1 , R^2 , R^3 and C are as defined above;
(b) oxidizing the carrier bound nucleoside-nucleotide;

13

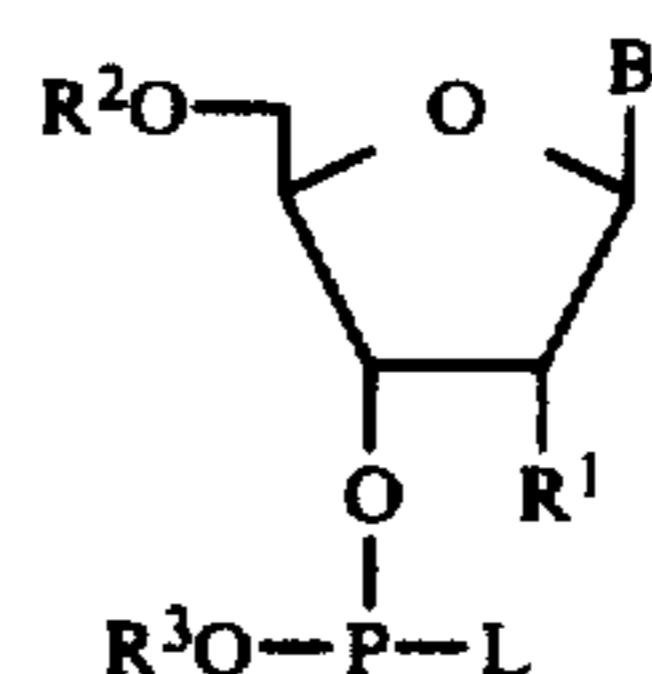
- (c) blocking free primary 5'—OH groups, which have not been reacted in the reaction of step (a), with permanent protective groups;
- (d) eliminating the protecting group R²;
- (f) repeating steps (a) to (d) to introduce further nucleoside phosphate units; and
- (g) cleaving the nucleoside-carrier bond and optionally eliminating protective groups present in the oligonucleoside phosphates.
5. A method of synthesizing oligonucleotides, comprising the steps of:
- coupling a nucleoside β-cyanoethyl-protected phosphoramidite to a carrier-bound nucleoside to produce a carrier bound nucleoside-nucleotide having a phosphite triester linkage;
 - oxidizing the phosphite triester to form a phosphate triester;
 - optionally coupling additional nucleoside β-cyanoethyl-protected phosphoramidites to the carrier bound nucleoside-nucleotide and, after each coupling step, oxidizing the resulting phosphite triester to form a phosphate triester, to form a carrier bound polynucleotide;
 - removing the β-cyanoethyl protecting groups; and
 - removing the polynucleotide from the carrier.
6. A method of claim 5, wherein the nucleoside β-cyanoethyl phosphoramidite is a nucleoside β-cyanoethyl N,N-dimethylphosphoramidite, N,N-diethylphosphoramidite, [N,N-dipropylphosphoramidite] N,N-diisopropylphosphoramidite or N,N-morpholino phosphoramidite.
7. A method of claim 6, wherein the carrier is controlled port glass.
8. A method of claim 7, wherein the β-cyanoethyl protecting group is removed with simultaneous removal of the polynucleotide from the carrier, by concentrated aqueous ammonia.
9. In a method of polynucleotide synthesis, comprising sequentially coupling nucleotide phosphoramidites to produce a polynucleotide, wherein the phosphorus atoms of the nucleotide phosphoramidites are protected by methyl groups, the improvement wherein the phosphorus protecting group are cyanoethyl groups.
10. A method of synthesizing oligonucleotides, comprising the steps of:
- coupling a nucleoside β-cyanoethyl-protected phosphoramidite to a nucleoside, the nucleoside being bound to a polymeric carrier via an ester bond to produce a carrier-bound nucleoside-nucleotide having a phosphite triester linkage;
 - oxidizing the phosphite triester to form a phosphate triester linkage;
 - sequentially coupling additional nucleoside β-cyanoethyl protected phosphoramidite to the carrier-bound nucleoside-nucleotide, and after each coupling step, oxidizing the resulting phosphite triester linkage to a phosphate triester to produce a carrier-bound polynucleotide;
 - treating the carrier bound polynucleotide with concentrated ammonia to remove the β-cyanoethyl phosphate protecting group and hydrolyzing the ester bond to the carrier to remove the polynucleotide from the carrier.
11. A method of claim 10, wherein the nucleoside β-cyanoethyl phosphoramidite is a nucleoside β-cyanoethyl N,N-dimethylphosphoramidite, N,N-diethylphosphoramidite; [N,N-dispropylphosphoramidite] N,N-

14

diisopropylphosphoramidite or N,N-morpholino phosphoramidite.

12. A method of claim 10, wherein the carrier is controlled pore glass.

13. A protected nucleotide having the formula:



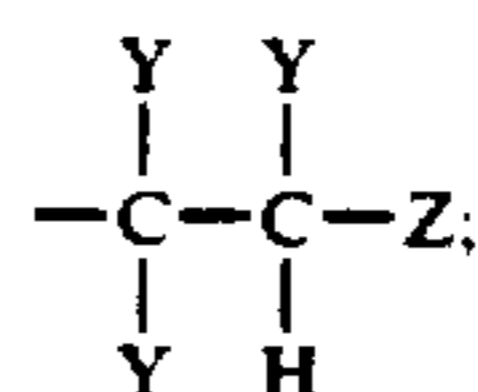
15 where,

B' is a nucleoside base protected, where appropriate, by a base protective group which can be eliminated;

R¹ is H, OH, or a hydroxyl group which is protected by a removable nucleoside protective group;

R² is a removable protective group;

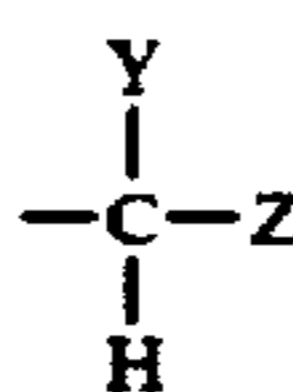
R³ is



30 L is CN, SCN, or NR₂⁴;

R⁴ is a primary, secondary or tertiary alkyl radical having 1-10 carbon atoms, or R₂⁴ is a cycloalkyl radical having 5-7 carbon atoms or a cycloalkyl radical having 5-7 atoms comprising atoms and one or two nitrogen, oxygen or sulfur atoms as heteroatoms;

Y is H, CH₃, or CH₂CH₃; and Z is a [halogen] halogen CN, NO₂, phenyl substituted in the o, o' or p positions with a halogen, CN or NO₂ radical, phenylthio, phenylsulfoxy, or phenylsulfonyl, where the phenyl radicals, may be substituted in the o, o' or p positions with a halogen, CN or NO₂ radical, or where the group



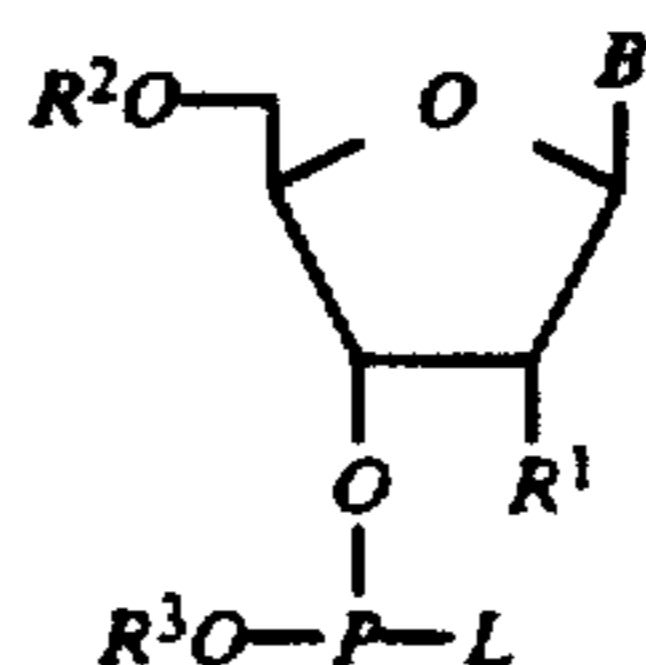
may be replaced by CF₃, CCl₃, or CBr₃.

14. A protected nucleotide as in claim 13, wherein Z is CN.

15. A protected nucleotide as in claim 14, wherein R₃ is CH₂—CH₂—CN.

16. A protected nucleotide as in claim 13, wherein R₂ is 4,4'-dimethoxytrityl or 4,4''-trimethoxytrityl.

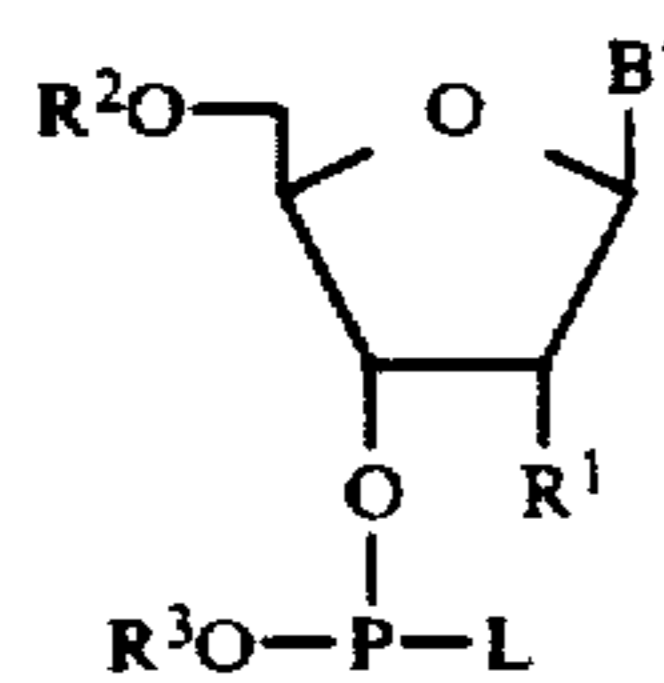
17. A protected nucleotide having the formula:



where,

15

R^1 is H, OH, or a hydroxyl group which is protected, where appropriate, by a removable nucleoside protective group;
 R^2 is 4,4'-dimethoxytrityl or 4,4',4''-trimethoxytrityl;
 B' is a nucleoside base protected by a base protective group which can be eliminated
 R^3 is CH_2-CH_2-CN ; and
 L is N,N-dimethylamino, N,N-diethylamino, N,N-diisopropylamino or N-morpholino.
 18. A protected nucleotide having the formula:



16

where,
 R^1 is H, OH, or a hydroxyl group which is protected by a protective group selected from the group consisting of trityl groups, acyl groups and silyl ether groups;
 R^2 is 4,4'-dimethoxytrityl or 4,4',4''-trimethoxytrityl;
 B' is a nucleoside base selected from the group consisting of adenine, guanine, cytosine, thymine, uracil and analogs thereof which are protected by acyl groups or Schiff bases;
 R^3 is CH_2-CH_2-CN ; and
 L is N,N-dimethylamino, N,N-diethylamino, N,N-diisopropylamino, or N-morpholino.
 19. A protected nucleotide of claim 18, wherein
 15 **[R₁]** R^1 is H and B' is adenine, guanine, cytosine, thymine or uracil wherein the adenine or **[guanine]** cytosine is protected by a benzoyl group and the guanine is protected by an **[isobutyl]** *isobutyryl* group.
 * * * * *

20

25

30

35

40

45

50

55

60

65

UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

PATENT NO. : Re. 34,069

Page 1 of 2

DATED : September 15, 1992

INVENTOR(S) : Hubert Koster and Nanda D. Sinha

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

column 6, line 19, change "is" to

---in---

In Claim 1, column 9, lines 39-40, delete ", where appropriate,".

In Claim 1, column 9, lines 52, 66 and 67; in Claim 4, column 12, line 23 and in Claim 9, column 13, lines 40 and 42, change "nucleotide" to
---nucleoside---

In Claim 1, column 9, line 53, before "by", insert ---where appropriate--- and before "protective", delete ---the---

In Claim 1, column 11, line 23, after "as", change "hereoatoms" to
---heteroatoms---

In Claim 11, column 14, line 1, change "diispropylphoshoramidite" to
---diisopropylphosphoramidite---

In Claim 13, column 14, line 34, after "comprising" insert ---carbon---

In Claim 13, column 14, line 38, change "halogen CN" to
--[halogen,] CN--

In Claim 13, column 14, line 42, change "halgen" to
---halogen---

In Claim 15, column 15, line 55, before "-CH₂", delete "1 is CH₂-CH₂-CN" and insert ---is CH₂-CH₂-CN---

In Claim 16, column 14, change "4,4"-trimethoxytrityl" to
---4,4',4"-trimethoxytrityl---

UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

Page 2 of 2

PATENT NO. : Re. 34,069

DATED : September 15, 1992

INVENTOR(S) : Hubert Koster and Nanda D. Sinha

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

In Claim 17, column 15, line 2, before "by", delete
---, where appropriate,---

In Claim 17, column 15, line 5, before "by", insert
---, where appropriate,---

Signed and Sealed this
Nineteenth Day of March, 1996

Attest:



BRUCE LEHMAN

Attesting Officer

Commissioner of Patents and Trademarks



US001034069B1

REEXAMINATION CERTIFICATE (2776th)

United States Patent [19]

Köster et al.

[11] **B1 Re. 34,069**[45] Certificate Issued **Jan. 23, 1996**

[54] PROCESS FOR THE PREPARATION OF OLIGONUCLEOTIDES

[75] Inventors: **Hubert Köster**, Concord, Mass.;
Nanda D. Sinha, San Rafael, Calif.[73] Assignee: **Millipore Corporation**, Bedford, Mass.

Reexamination Requests:

No. 90/003,113, Jul. 1, 1993
No. 90/003,309, Jan. 12, 1994
No. 90/003,459, Jun. 10, 1994

Reexamination Certificate for:

Patent No.: **1,034,069**
Issued: **Sep. 15, 1992**
Appl. No.: **481,572**
Filed: **Feb. 16, 1990**

Related U.S. Patent Documents

Reissue of:

[64] Patent No.: **4,725,677**
Issued: **Feb. 16, 1988**
Appl. No.: **752,178**
Filed: **Aug. 10, 1984**[51] Int. Cl.⁶ **C07H 15/12; C07H 17/00**
[52] U.S. Cl. **536/25.34; 536/26.5; 536/26.71;**
536/25.3; 987/189
[58] Field of Search **536/27, 28, 29,**
536/25.3, 25.34

[56] References Cited

U.S. PATENT DOCUMENTS

4,415,732 11/1983 Caruthers et al. 536/27

FOREIGN PATENT DOCUMENTS

0040099 11/1981 European Pat. Off. .

OTHER PUBLICATIONS

Adamiak et al., "Nucleoside 3'-phosphotriesters, as Key Intermediates for the Oligoribonucleotide Synthesis III.," *Nucl. Acids Res.* 3(12):3397 (1976).van Boom et al., "2,2,2-Trichloroethyl 2-chlorophenyl Phosphorochloridate, A Convenient Reagent for the Formation of Internucleotide Linkages", *Tet. Letters*, 11:869 (1976).Adams et al., "Hindered Dialkylamino Nucleoside Phosphate Reagents in the Synthesis of Two DNA 51-mers," *J. Am. Chem. Soc.* 105:661 (1983).Applied Biosystems, Users Manual: *PCR-MATE EP DNA Synthesizer* (Rev. A 1989) Appendix 1.Beaucage et al., "Deoxynucleoside Phosphoramidites—A New Class of Key Intermediates For Deoxypolynucleoside Synthesis," *Tet. Lett.* 22 (20):1859 (1981).Beld et al., "Bis-[2-(methylsulfonyl)ethyl] phosphochloridate, a new phosphorylating agent," *Recl. Trav. Chim. Pays-Bas*, 103:196 (1984).Caruthers, "Gene Synthesis Machines," *Science* 230:281 (1985) (Caruthers V) (Exhibit 15).Caruthers et al., "Chemical Synthesis of Deoxyoligonucleotides and Deoxyoligonucleotide Analogs," *Methods in Enzym.* 211:3 (1992).

Cherbuliez & Rabinowitz, "112 Recherches sur la formation

et la transformation des esters XXI", *Helv. Chem. Act.*, 43:863 (1960). (partial translation included).Dahl et al., "Mechanistic Studies on the Phosphoramidite Coupling Reaction In Oligonucleotide Synthesis," *Nucl. Acids Res.* 15(4):1729 (1987).Damha & Ogilvie, "Oligoribonucleotide Synthesis: The Silylphosphoramidite Method," *Methods in Molecular Biology* 20:81 (1993).Damha & Ogilvie, "Synthesis and Spectroscopic Analysis of Branched RNA Fragments," *J. Org. Chem.* 53:3710 (1988).Daub & van Tamelen, "Synthesis of Oligoribonucleotides Based on the Facile Cleavage of Methyl Phosphotriester Intermediates," *J. Am. Chem. Soc.* 99:3526 (1977).Fourrey & Varenne, "A new and General Procedure for the Preparation of Deoxynucleotide Phosphoramidites," *Tet. Lett.* 24(19):1983.Gasparutto et al., "Studies on the Formation of the Internucleotidic Bond in RNA Synthesis Using Dialkylamino Phosphoramidites," *Nucleosides & Nucleotides* 9(8):1087 (1990).Gilham & Tener, "A New Method of Phosphorylation," *Chem. & Indus.* 542 (1959).Ikehara et al., *Advances in Carbohydrate Chemistry & Biochemistry* 30,135 (1979) (review).Letsinger & Mahadevan, "Oligonucleotide Synthesis on a Polymer Support," *J. Am. Chem. Soc.* 87:3526 (1965).Letsinger & Ogilvie, "A Convenient Method for Stepwise Synthesis of Oligothymidate Derivative, in Large-Scale Quantities," *J. Am. Chem. Soc.* 89:4801 (1967).Letsinger & Ogilvie, "Synthesis of Oligothymidylates via Phosphotriester Intermediates," *J. Am. Chem. Soc.* 91(12):3350 (1969).

(List continued on next page.)

Primary Examiner—Ronald W. Griffin

[57] ABSTRACT

The invention relates to a process for the preparation of oligonucleotides by the following steps: reaction of a nucleoside with a phosphine derivative, reaction of the nucleotide derivative thus obtained with a nucleoside bonded to a polymeric carrier, oxidation of the carrier-bound nucleoside-nucleotide thus obtained with formation of phosphotriester groups, blocking of free primary 5'-OH groups, elimination of a protective group from the terminal 5'-OH group, where appropriate single or multiple repetition of the abovementioned steps to introduce further nucleoside phosphate or oligonucleoside phosphate units, and cleavage of the nucleoside-carrier bond and, where appropriate, elimination of all protective groups present in the oligonucleoside phosphates. The phosphine derivative used is a compound of the general formula III



in which X and L can react with OH groups of the sugar units in the oligonucleotides, and R³ is a protective group which can be liberated by β-elimination.

Letsinger et al., "Developments in Syntheses of Oligoribonucleotides and Their Organic Derivatives," *J. Am. Chem. Soc.* 91(12):3360 (1969).

OTHER PUBLICATIONS

- Letsinger et al., "Synthesis of Thymidine Oligonucleotides by Phosphite Triester Intermediates," *J. Am. Chem. Soc.* 98(12): 3655 (1975).
- Marugg et al., "Use of 2-Cyano-1,1-Dimethylethyl As A Protecting Group In The Synthesis of DNA Via Phosphite Intermediates," *Rec. Trav. Chim. Pays-Bas* 103:97 (1984).
- McBride & Caruthers, "An investigation of Several Deoxynucleotide Phosphoramidites Useful for Synthesizing Deoxyoligonucleotides," *Tet. Lett.* 24(3):245 (1983).
- Ogilvie et al., "The Chemical Synthesis of Oligoribonucleotides VIII," *Nucl. Acids Res.* 8(9):2105 (1980).
- Ogilvie & Nemer, "Silica gel as a Solid Support in the Synthesis of Oligoribonucleotides," *Tet. Lett.* 21:4159 (1980).
- Schaller et al., "Studies on Polynucleotides XXIV," *J. Am. Chem. Soc.* 85:3821 (1963).
- Tener, "2-cyanoethyl Phosphate and its Use in the Synthesis of Phosphate Esters," *J. Am. Chem. Soc.* 83:159 (1961).
- Usman et al., "Automated Chemical Synthesis of Long Oligoribonucleotides Using 2'-O-Silylated Ribonucleotide 3'-O-phosphoramidites of a 43-Nucleotide Sequence Similar to the 3'-Half Molecule of *Escherichia Coli* Formylmethionine tRNA," *J. Am. Chem. Soc.* 109:7845 (1987).
- Bender and Ogilvie, "Polynucleotide Synthesis", European Patent Application No. 0 40 099 (Filing date Dec. 5, 1981).
- Weimann & Khorana, "Studies on Polynucleotides XIII," *J. Am. Chem. Soc.* 84:419 (1962).
- Agarwal et al., "Studies on Polynucleotides CXLIII, a rapid and convenient method for the synthesis of deoxyribo-oligonucleotides carrying 5'-phosphate end groups using a new protecting group", *J. Am. Chem. Soc.*, 98:1065 (1976).
- Charubala et al., "Synthesis of Inosinate Trimer 12'p5'12'p5'1 and Tetramer 12'p5'12'p5'12'p5'1", *Tet. Letters*, 23:4789 (1982).
- Crea et al., "Synthesis of Oligonucleotides on Cellulose by a Phosphotriester Method", *Nucl. Acids Res.*, 8:2331 (1980).
- Dorper and Winnacker, "Improvements in the Phosphoramidite Procedure for the Synthesis of Oligodeoxyribonucleotides", *Nucl. Acids Res.*, 11:2575 (1983).
- Himmelsbach and Pfeleiderer, "Bis-(p-Nitrophenylethyl) Phosphomonochlorides, A New Versatile Phosphorylating Agent", *Tet. Letters*, 23:4793 (1982).
- Horn et al., "Synthesis of Oligonucleotides on cellulose, Part II: design and synthetic strategy to the synthesis of 22 oligodeoxynucleotides coding for Gastric Inhibitory Polypeptide (GIP)", *Nucl. Acids Res.*, Sym series 7:225 (1980).
- Ichiba and Pfeleiderer, "Chemical Synthesis of the Ribo-hexamer CpApApCpCpA", *Nucl. Acids Res. Sym. ser.*, 9:169 (1981).
- Ikehara et al., *Advances in Carbohydrate Chemistry & Biochemistry*, 36:135 (1979).
- Kabachnik and Medved, "Some Properties of Amides of β -Chloroethyl Phosphonates, β -chloroethylphosphonic acid, and vinylphosphonic acids" (Inst. Hetero-org Compds., Moscow), *Izv. Akad. Nauk SSR, Ser. Khim* (8):1365-70 (1966) (Russ) (CA 66, 1967, 54802u).
- Khorana et al., "Studies on Polynucleotides XIII, Stepwise Synthesis of Deoxyribo-oligonucleotides, An Alternative Approach and the Synthesis of Thymidine Di-, Tri- and Tetranucleotides Bearing 3'-Phosphomonoester End Groups", *J. Am. Chem. Soc.*, 84:419 (1962).
- Kohli et al., "Deoxypolynucleotides: Part I-Synthesis of Thymidylyl-(3'-5')-thymidylyl-(3'-5')-thymidine & a Comparison of Phosphodiester & Phosphotriester Approaches", *Indian. Journ. of Chemistry*, 17B:253 (1979).
- Kohli et al., "Synthesis of Deoxyribonucleotides Corresponding to Codons of Amino Acids by Phosphotriester Approach", *Indian. Journ. of Chemistry*, 17B:257 (1979).
- Letsinger et al., "Stepwise Synthesis of Oligodeoxyribonucleotides On An Insoluble Polymer Support", *J. Am. Chem. Soc.*, 88:5319 (1966).
- Letsinger et al., "Synthesis of Thymidine Oligonucleotides by Phosphite Triester Intermediates", *J. Am. Chem. Soc.*, 98:3655 (1976).
- McBride & Caruthers, "An Investigation of Several Deoxynucleotide Phosphoramidites Useful for Synthesizing Deoxyoligonucleotides", *Tet. Letters*, 24(3):245-248.
- Ogilvie & Nemer, "Silica Gel as Solid Support in the Synthesis of Oligoribonucleotides", *Tet. Letters*, 21:4159-4162.
- Pfleiderer and Beiter et al., "Solution Synthesis of Protected Di-2'-Deoxynucleoside Phosphotriesters via The Phosphoramidite Approach", *Tet. Letters*, 25:1975 (1984).
- Schaller et al., "The Stepwise Synthesis of Specific Deoxypolynucleotides(4), Protected Derivatives of Deoxyribonucleosides and New Synthesis of Deoxyribonucleosides-3'-Phosphates", *J. Am. Chem. Soc.*, 85:3821 (1963).
- Schaller and Khorana, "Studies on Polynucleotides XXVII, The Stepwise Synthesis of Deoxyribopolynucleotides (7), The Synthesis of Polynucleotides Containing Deoxycytidine and Deoxyguanosine in Specific Sequences and of Homologous Deoxycytidine Polynucleotides Terminating in Thymidine", *J. Am. Chem. Soc.*, 85:3841 (1963).
- Smith et al., "The Methyl Group as Phosphate Protecting Group in Nucleotide Syntheses", *Tet. Letters*, 21:861-864 (1980).
- Srivastava et al., "Studies on Oligonucleotide triester Synthesis, The Effect of Internucleotide Protecting Groups", *J. Carbohydrates, Nucleosides, Nucleotides*, 8(6):495(1981).
- Usman et al., "Total Chemical Synthesis of a 77-nucleotide-long RNA Sequence having Methionine-acceptance activity", *Proc. Natl. Acad. Sci. USA*, 85:5764 (1988).
- Weimann, Schaller and Khorana, "Studies on Polynucleotides XXVI, The Stepwise Synthesis of Specific Deoxyribopolynucleotides (6), The Synthesis of Thymidylyl-(3'-5')-deoxyadenylyl-(3'-5')-thymidylyl-(3'-5')-thymidylyl-(3'-5')-thymidine and of Polynucleotides Containing Thymidine and Deoxyadenosine in Alternating Sequence", *J. Am. Chem. Soc.*, 85:3835-3841 (1963).
- Pon et al., *Nucleic Acids Research* 13:6447 (1985).
- Letsinger et al., "Oligonucleotide Synthesis Utilizing β -benzoyl propionyl, a Blocking Group with a Trigger for Selective Cleavage", *J. Am. Chem. Soc.* 89:7146 (1967).
- Narang, "DNA Synthesis", *Tetrahedron* 39:3 (1983).
- Caruthers et al., "Total Synthesis of the Structural Gene for an Transfer Ribonucleic Acid from Yeast", *J. Mol. Biol.* 72:375 (1972).
- J. Smrt, "Oligonucleotidic Compounds, XL, Aspects of the Triester Synthesis in the Ribo Series," *Collection Czechoslov. Chem. Commun.*, vol. 37, pp. 1870-1877 (1972).

B1 Re. 34,069

1

**REEXAMINATION CERTIFICATE
ISSUED UNDER 35 U.S.C. 307**

NO AMENDMENTS HAVE BEEN MADE TO
THE PATENT

2

AS A RESULT OF REEXAMINATION, IT HAS BEEN
DETERMINED THAT:

The patentability of claims **1-19** is confirmed.

* * * * *