

DESCRIPTION

CROSSLINKED ARTIFICIAL NUCLEIC ACID ALNA

5 TECHNICAL FIELD

[0001]

The present invention relates to a monomer of amino LNA, and an oligomer containing the same.

10 BACKGROUND ART

[0002]

A therapeutic method for diseases with nucleic acid medicine include an antisense method, an antigen method, a method with aptamer, and a method with siRNA, and so on. Among them, the antisense method is a method in which a disease is treated or prevented by introducing with an oligonucleotide (antisense strand) complementary to disease-related mRNA or non-coding RNA from the outside to form a double strain, and accordingly, by adjusting the function of RNA involved in the disease to treat or prevent the disease.

[0003]

Though various artificial nucleic acids have been developed as materials for the nucleic acid medicine, problems such as avoiding toxicity in organs where nucleic acid medicines are likely to accumulate, such as the liver

and kidneys after systemic administration, and developing sufficient medicinal efficacy remain, there is still no molecule that should be a trump card. A 2'-amino LNA which was developed by Wengel et al., in 1998 (hereinafter, described as "ALNA") (Patent document 1, and Non-Patent Document 1) can synthesize various artificial nucleic acids by modifying the substituent at 2' position and so on, and many ALNA derivatives have been synthesized so far, mainly including alkyl or acyl derivatives (Patent documents 2 to 4, and Non-Patent Documents 2 to 11), and a characteristic tissue distribution after systemic administration has been reported (Non-Patent Document 12), however, it is insufficient in terms of pharmacological activity and accordingly, the derivatives have not been applied to pharmaceuticals so far.

[0004]

We apply the efficient synthesis method of GuNA previously found (Patent Document 5 and Non-Patent Document 13) to synthesize a wide variety of novel ALNA derivatives, and perform screening specialized in pharmacological activity evaluation, and accordingly, have found out a novel artificial nucleic acid having excellent in vitro and in vivo pharmacological activity as compared with the existing ALNA, and completed the invention of the present application.

CITATION LIST

PATENT DOCUMENT

[0005]

- 5 Patent Document 1: WO 99/014226 pamphlet
 Patent Document 2: WO 2013/013068 pamphlet
 Patent Document 3: WO 2014/124952 pamphlet
 Patent Document 4: WO 2016/128583 pamphlet
 Patent Document 5: WO 2017/047816 pamphlet

10 NON-PATENT DOCUMENT

[0006]

- Non-Patent Document 1: S.K. Singh et al., J. Org. Chem.
1998, 63, 6078-6079
- Non-Patent Document 2: M. D. Sorensen et al., Chem.
15 Commun. 2003, 2130
- Non-Patent Document 3: B. R. Babu et al., Chem. Commun.
2005, 13, 1705-1707
- Non-Patent Document 4: T. Bryld et al., Nucleosides,
Nucleotides & Nucleic Acids 2007, 26, 1645-1647
- 20 Non-Patent Document 5: M. Kalek et al., J. Am. Chem.
Soc. 2007, 129, 9392-9400
- Non-Patent Document 6: T. Umemoto et al., Org. Biomol.
Chem. 2009, 7, 1793-1797
- Non-Patent Document 7: M. W. Johannsen et al., Org.
25 Biomol. Chem. 2011, 9, 243-252

Non-Patent Document 8: A. S. Jørgensen et al., Chem. Commun. 2013, 49, 10751-10753

Non-Patent Document 9: I. K. Astakhova et al., Acc. Chem. Res. 2014, 47, 1768-1777

5 Non-Patent Document 10: C. Lou et al., Chem. Commun. 2015, 51, 4024-4027

Non-Patent Document 11: A. Ries et al., J. Org. Chem. 2016, 81, 10845-10856

10 Non-Patent Document 12: K. Fluiter et al., Chem. Bio. Chem. 2005, 6, 1104-1109

Non-Patent Document 13: H. Sawamoto et al., Org. Lett. 2018, 20, 1928-1931

SUMMARY OF INVENTION

15 (PROBLEMS TO BE SOLVED BY INVENTION)

[0007]

The present invention relates to a novel bridged artificial nucleic acid, as well as monomer containing the same as a monomer.

20

(MEANS TO SOLVE PROBLEMS)

[0008]

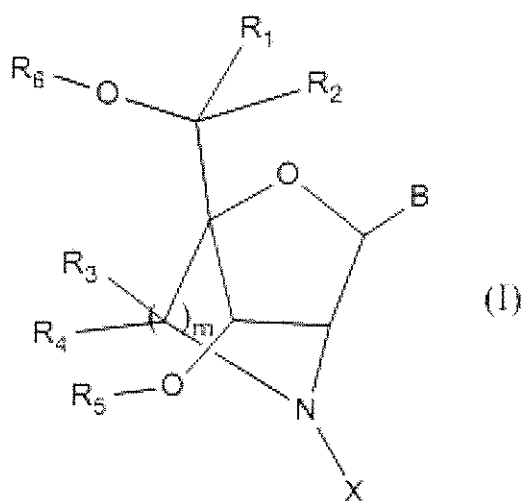
In order to solve the above-mentioned problems to be solved, the inventors have intensively studied to find out
25 a novel 2'-amino LNA and salts thereof, as well as oligomer

containing the same as a monomer, thereby completed the present invention.

[0009]

That is, the present invention includes the following
5 Items [1] to [33], which should not be limited thereto.

Item [1] A compound represented by general formula I:



[wherein

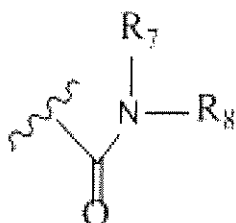
B represents a base moiety of nucleic acid wherein the
10 base moiety may be optionally substituted with one or more
substituents;

R₁, R₂, R₃ and R₄ represent independently of each other
a hydrogen atom, or a C₁₋₆ alkyl group which may be optionally
substituted with one or more substituents;

15 R₅ and R₆ represent independently of each other a
hydrogen atom, a protecting group for hydroxy group, or a
phosphate group which may be optionally substituted with
substituents;

m is an integer of 1 or 2;

X represents a group represented by the following formula (II-1), (II-2) or (II-3):



(II-1)



(II-2)



(II-3)

;

the symbols:



which is described in the formula (II-1), (II-2) or (II-3) represents a binding point to 2'-amino group;

R₇ and R₈ represent independently of each other a hydrogen atom, a C₁₋₆ alkyl group which may be optionally substituted with one or more substituents, a C₂₋₆ alkenyl group which may be optionally substituted with one or more substituents, a C₂₋₆ alkynyl group which may be optionally substituted with one or more substituents, or an aromatic group which may be optionally substituted with one or more substituents;

A represents an aromatic group;

M represents a sulfonyl group which is substituted with one substituent selected from the group consisting of a C₁₋₆ alkyl group which may be optionally substituted with one or

more substituents and an aromatic group which may be optionally substituted with one or more substituents.]

or salts thereof

(hereinafter, the compound represented by formula (I) or salts thereof is referred to as "Present compound" or "Compound of the present invention").

[2] The compound according to [1] or salt thereof wherein B represents an adeninyl group which may be optionally substituted with one or more substituents, a guaninyl group which may be optionally substituted with one or more substituents, a cytosinyl group which may be optionally substituted with one or more substituents, a 5-methylcytosinyl group which may be optionally substituted with one or more substituents, or an uracilyl group which may be optionally substituted with one or more substituents.

[3] The compound according to [1] or [2] or salts thereof wherein R_1 , R_2 , R_3 and R_4 represent independently of each other a hydrogen atom; and

m is an integer of 1.

[4] The compound according to any one of [1] to [3] or salts thereof wherein

X represents a group represented by formula (II-1); and

R_7 and R_8 represent independently of each other a hydrogen atom, a C_{1-6} alkyl group which may be optionally substituted with one or more substituents, or an aromatic

group which may be optionally substituted with one or more substituents.

[5] The compound according to [4] or salts thereof wherein R_7 and R_8 represent independently of each other a hydrogen atom, or a C_{1-3} alkyl group which may be optionally substituted with one or more substituents.

[6] The compound according to [5] or salts thereof wherein one of R_7 and R_8 represents a hydrogen atom, and the other thereof represents a methyl group.

[7] The compound according to [5] or salts thereof wherein one of R_7 and R_8 represents a hydrogen atom, and the other thereof represents an isopropyl group.

[8] The compound according to any one of [1] to [3] or salts thereof wherein

X represents a group represented by formula (II-2);

A represents a five or six membered heteroaryl group containing one or more heteroatoms selected from the group consisting of nitrogen atom, oxygen atom and sulfur atom, which may be optionally substituted with one or more substituents.

[9] The compound according to [8] or salts thereof wherein A represents a five or six membered heteroaryl group containing two or three heteroatoms selected from the group consisting of nitrogen atom, oxygen atom and sulfur atom, provided that at least two nitrogen atoms are included, which

may be optionally substituted with one or more substituents, and the substituents is selected independently of each other a group consisting of a C₁₋₃ alkyl group which may be optionally substituted with one or more halogen atoms, a
5 halogen atom, and an amino group which may be optionally substituted with one or more C₁₋₃ alkyl group.

[10] The compound according to [9] or salts thereof wherein the five or six membered heteroaryl group is selected from a group consisting of a triazolyl group, an oxadiazolyl group,
10 a thiadiazolyl group, a pyrimidinyl group, and a pyrazinyl group, each may be optionally substituted with one or more substituents.

[11] The compound according to [10] or salts thereof wherein the triazolyl group which may be optionally substituted with
15 one or more substituents represents 1,5-dimethyl-1,2,4-triazol-3-yl group.

[12] The compound according to [10] or salts thereof wherein the oxadiazolyl group which may be optionally substituted with one or more substituents represents 5-methyl-1,2,4-
20 oxadiazol-3-yl group.

[13] The compound according to [10] or salts thereof wherein the thiadiazol group which may be optionally substituted with one or more substituents represents 3-methyl-1,2,4-thiadiazol-5-yl group.

25 [14] The compound according to any one of [1] to [3] or salts

thereof wherein

X represents a group represented by formula (II-3); and

M represents a sulfonyl group which is substituted with one or more substituents selected from the group consisting of a C₁₋₆ alkyl group which may be optionally substituted with one or more substituents, and an aryl group which may be optionally substituted with one or more substituents.

[15] The compound according to [14] or salts thereof wherein M represents a sulfonyl group which is substituted with one or more substituents selected from the group consisting of a methyl group and a phenyl group.

[16] The compound according to according to any one of claims [1] to [15] or salts thereof wherein R₆ represents a hydrogen atom or DMTr group, and R₅ represents a hydrogen atom or -P(O(CH₂)₂CN)(N(iPr)₂).

[17] A compound selected from the group consisting of

3-[[[(1R,4R,6R,7S)-4-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-6-(5-methyl-2,4-dioxo-pyrimidin-1-yl)-2-methylsulfonyl-5-oxa-2-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 1);

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-

yl]benzamide (Present compound 2);

[9-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-2-(2-methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate (Present compound 3);

N-[9-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-purin-6-yl]benzamide (Present compound 4);

3-[[(1R, 3R, 4R, 7S)-5-(benzenesulfonyl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 5);

N-[1-[(1R, 3R, 4R, 7S)-5-(benzenesulfonyl)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 6);

(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-methyl-3-(5-methyl-2,4-

dioxo-pyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 7);

(1R,3R,4R,7S)-3-(4-benzamide-5-methyl-2-oxo-pyrimidin-1-yl)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-methyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 8);

[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(methylcarbamoyl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-2-(2-methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate (Present compound 9);

(1R,3R,4R,7S)-3-(6-benzamidepurin-9-yl)-1-[[bis(4-methoxyphenyl)phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-methyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 10);

(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-isopropyl-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 11);

(1R,3R,4R,7S)-3-(4-benzamide-5-methyl-2-oxopyrimidin-1-yl)-1-[[bis(4-methoxyphenyl)phenyl-methoxy]methyl]-7-[2-

cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-isopropyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 12);

5 [9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(isopropylcarbamoyl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-2-(2-methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate (Present compound 13);

10 (1R,3R,4R,7S)-3-(6-benzamidepurin-9-yl)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-isopropyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 14);

15 (1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N,N-dimethyl-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 15);

20 (1R,3R,4R,7S)-3-(4-benzamide-5-methyl-2-oxo-pyrimidin-1-yl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N,N-dimethyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 16);

25 3-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-

methoxy)methyl]-3-(5-methyl-2,4-dioxo-pyrimidin-1-yl)-5-pyrimidin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 17);

5 1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-pyrimidin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 18);

10 N'-[9-(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-pyrimidin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-N,N-dimethyl-6-formamidine (Present compound 19);

15 N'-[9-(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-pyrimidin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]purin-6-yl]benzamide (Present compound 20);

20 3-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-3-(5-methyl-2,4-dioxopyrimidin-1-yl)-5-pyrimidin-4-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 21);

25 N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-

methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-pyrimidin-4-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 22);

5 3-[[(1R, 3R, 4R, 7S) -1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-3-(5-methyl-2,4-dioxo-pyrimidin-1-yl)-5-[4-(trifluoromethyl)pyrimidin-2-yl]-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present
10 compound 23);

 N-[1-[(1R, 3R, 4R, 7S) -1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-[4-(trifluoromethyl)pyrimidin-2-yl]-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 24);

 3-[[(1R, 3R, 4R, 7S) -1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-5-(5-chloropyrimidin-2-yl)-3-(5-methyl-2,4-dioxo-pyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile
20 (Present compound 25);

 N-[1-[(1R, 3R, 4R, 7S) -1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-5-(5-chloropyrimidin-2-yl)-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-4-

yl]benzamide (Present compound 26);

3-[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy]methyl]-5-[6-(dimethylamino)pyrimidin-4-yl]-3-(5-
methyl-2,4-dioxo-pyrimidin-1-yl)-2-oxa-5-
5 azabicyclo[2.2.1]heptan-7-yl]oxy-
(diisopropylamine)phosphanyl]oxypropanenitrile (Present
compound 27);

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy]methyl]-7-[2-
10 cyanoethoxy(diisopropylamino)phosphanyl]oxy-5-[6-
dimethylamino)pyrimidin-4-yl]-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-
yl]benzamide (Present compound 28);

3-[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
15 methoxy]methyl]-5-[2-(dimethylamino)pyrimidin-4-yl]-3-(5-
methyl-2,4-dioxo-pyrimidin-1-yl)-2-oxa-5-
azabicyclo[2.2.1]heptan-7-yl]oxy-
(diisopropylamino)oxypropanenitrile (Present compound 29);

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
20 methoxy]methyl]-7-[2-cyanoethoxy-
(diisopropylamino)phosphanyl]oxy-5-[2-
(dimethylamino)pyrimidin-4-yl]-7-hydroxy-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-
yl]benzamide (Present compound 30);

25 3-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-

methoxy)methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-pyrazin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 31);

5 N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-pyrazine-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 32);

10 3-[[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present
15 compound 33);

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-
20 methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 34);

N'-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(1,5-dimethyl-1,2,4-
25 triazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-

1H-purin-2-yl]-N,N-dimethylformamidine (Present compound 35);

N-[9-(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]purin-6-yl]benzamide (Present compound 36);

3-[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-(5-methyl-1,2,4-oxadiazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 37);

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(5-methyl-1,2,4-oxadiazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 38);

3-[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-(3-methyl-1,2,4-thiadiazol-5-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 39);

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(3-methyl-1,2,4-thiadiazol-5-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 40);

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]dihydropyrimidin-2-one (Present compound 41);

N'-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy(diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-N,N-dimethylformamidine (Present compound 42);

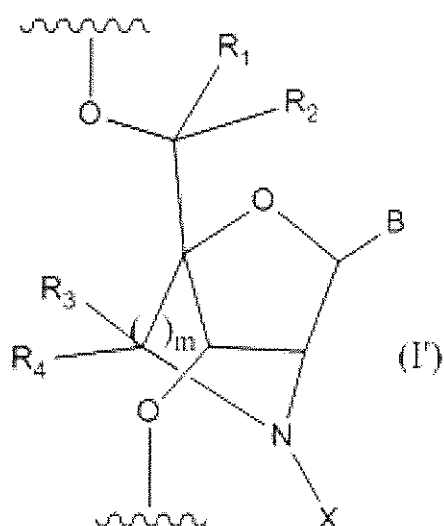
N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-2-methylpropane amide (Present compound 43); and

N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-2-methylpropane amide (Present compound 44).

[0010]

(Oligonucleotide)

[18] An oligonucleotide compound comprising one or more nucleosides represented by general formula (I'):



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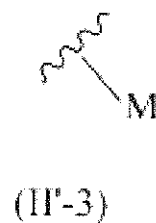
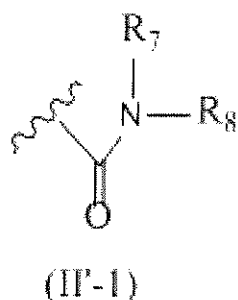
[wherein,

B represents a base moiety of nucleic acid wherein the base moiety may be optionally substituted with one or more substituents;

10 R_1 , R_2 , R_3 and R_4 represent independently of each other a hydrogen atom, or a C1-6 alkyl group which may be optionally substituted with one or more substituents;

m is an integer of 1 or 2;

15 X represents a group represented by the following formula (II'-1), (II'-2), or (II'-3):



the symbols:



which is described in the formula (II'-1), (II'-2) or (II'-
 5 3) represents a binding point to 2'-amino group;

A represents an aromatic group containing one or more heteroatoms selected from the group consisting of nitrogen atom, oxygen atom and sulfur atom, which may be optionally substituted with one or more substituents;

10 M represents a sulfonyl group which is substituted with one or more substituents selected from the group consisting of a C₁₋₆ alkyl group which may be optionally substituted with one or more substituents, and an aromatic group which may be optionally substituted with one or more substituents.]

15 or salts thereof

(hereinafter, the compound represented by formula (I') or salts thereof is referred to as "Present oligonucleotide compound" or "Oligonucleotide compound of the present invention").

20 [19] The compound according to [18] or salts thereof wherein

B represents an adeninyl group which may be optionally substituted with one or more substituents, a guaninyl group which may be optionally substituted with one or more substituents, a cytosinyl group which may be optionally substituted with one or more substituents, a 5-methylcytosinyl group which may be optionally substituted with one or more substituents or an uracilyl group which may be optionally substituted with one or more substituents.

[20] The oligonucleotide compound according to [18] or [19] or salts thereof wherein

R_1 , R_2 , R_3 and R_4 represent independently of each other a hydrogen atom; and

m is an integer of 1.

[21] (Urea compound) The oligonucleotide compound according to any one of [18] to [20] or salts thereof, wherein

X represents a group represented by formula (II'-1); and

R_7 and R_8 represent independently of each other a hydrogen atom, a C1-6 alkyl group which may be optionally substituted with one or more substituents, or an aromatic group which may be optionally substituted with one or more substituents.

[22] The oligonucleotide compound according to [21] or salts thereof wherein R_7 and R_8 represent independently of each

other a hydrogen atom, a C1-3 alkyl group which may be optionally substituted with one or more substituents.

[23] The oligonucleotide compound according to [22] or salts thereof wherein one of R₇ and R₈ represents a hydrogen atom,
5 and the other thereof represents a methyl group.

[24] The oligonucleotide compound according to [23] or salts thereof wherein one of R₇ and R₈ represents a hydrogen atom, and the other thereof represents an isopropyl group.

[25] (Aryl type compound) The oligonucleotide compound
10 according to any one of claims 18 to 20 or salts thereof wherein

X represents a group represented by formula (II'-2);
and

A represents a five or six membered heteroaryl group
15 containing one or more heteroatoms selected from the group consisting of nitrogen atom, oxygen atom and sulfur atom, which may be optionally substituted with one or more substituents.

[26] The oligonucleotide compound according to [25] or salts
20 thereof wherein

A represents a five or six membered heteroaryl group containing two or three heteroatoms selected from nitrogen atom, oxygen atom, and sulfur atom, provided that at least two nitrogen atoms are included, which may be optionally
25 substituted with one or more substituents, and the

substituent is selected independently of each other from a group consisting of a C1-3 alkyl group which may be optionally substituted with one or more halogen atoms, a halogen atom, and an amino group which may be optionally substituted with one or more C1-3 alkyl groups.

[27] The oligonucleotide compound according to [26] or salts thereof wherein the five or six membered heteroaryl group is selected from the group consisting of a triazolyl group, an oxadiazolyl group, a thiadiazolyl group, a pyrimidinyl group, and a pyrazinyl group, each may be optionally substituted with one or more substituents.

[28] The oligonucleotide compound according to [27] or salts thereof wherein the triazolyl group which may be optionally substituted with one or more substituents represents 1,5-dimethyl-1,2,4-triazol-3-yl group.

[29] The oligonucleotide compound according to [27] or salts thereof wherein the oxadiazolyl group which may be optionally substituted with one or more substituents represents 5-methyl-1,2,4-oxadiazol-3-yl group.

[30] The oligonucleotide compound according to [27] or salts thereof wherein the thiadiazolyl group which may be optionally substituted with one or more substituents represents 3-methyl-1,2,4-thiadiazol-5-yl group.

[31] (Sulfonamide type compound) The oligonucleotide compound according to any one of [18] to [20] or salts

thereof, wherein

X represents a group represented by formula (II'-3);
and

5 M represents a sulfonyl group which is substituted with
one or more substituents selected from the group consisting
of a C1-6 alkyl group which may be optionally substituted
with one or more substituents, and an aryl group which may
be optionally substituted with one or more substituents.

10 [32] The compound according to any one of [18] to [31] or
salts thereof wherein M represents a sulfonyl group which is
substituted with one or more substituents selected from the
group consisting of a methyl group and a phenyl group.

[33] The oligonucleotide compound according to any one of
[1] to [32] or salts thereof wherein one or more phosphate
15 bonds between nucleotides represent a phosphorothioate bond.

(EFFECT OF INVENTION)

[0011]

According to the present invention, novel 2'-amino LNA
(hereinafter, is abbreviated as "ALNA") and oligomer
20 comprising these ALNA as a monomer (hereinafter, is
abbreviated as "ALNA oligomer") can be prepared. The ALNA
oligomer of the present invention has a functional inhibition
of a target microRNA in vitro, or a potential target gene
knockdown activity, and when administered to a living body,
25 has a potential target gene knockdown activity in many organs

(such as muscle), which accordingly can be expected to use as novel nucleic acid medicine.

MODE FOR CARRYING OUT THE INVENTION

5 [0012]

Hereinafter, the present invention is further explained in details. All of the publications cited herein are incorporated herein by reference.

(Definition)

10 First, the definitions of the terms as used herein are described.

[0013]

The term of "C₁₋₆ alkyl which may be optionally substituted with one or more substituents" as used herein
15 encompasses any straight chain alkyl group having 1 to 6 carbon atoms (C₁₋₆), preferably 1 to 4 carbon atoms (C₁₋₄), more preferably 1 to 3 carbon atoms (C₁₋₃), any branched chain alkyl group having 3 to 6 carbon atoms in which the identical or different branched chains are contained, any cyclic alkyl
20 group having 3 to 6 carbon atoms, and any combinations thereof which have 4 to 6 carbon atoms. Specific examples of the straight chain alkyl group having 1 to 6 carbon atoms include methyl, ethyl, normal (which is abbreviated as "n")-propyl, iso (which is abbreviated as "i")-propyl, n-butyl,
25 n-pentyl, and n-hexyl. Specific examples of the branched

chain alkyl group having 3 to 6 carbon atoms in which the identical or different branched chains are contained include i-propyl, i-butyl, tert (which is abbreviated as "t")- butyl, sec (which is abbreviated as "s")- butyl, neopentyl, isopentyl, and the others, and also examples of the cyclic alkyl group having 3 to 6 carbon atoms include preferably a 3 to 6 membered monocyclic cycloalkyl group, and specific examples thereof include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, and the others, which are not limited thereto. Examples of the substituents include one or more (or preferably one to three) identical or different groups selected from the group consisting of a hydroxy group, a halogen atom, a nitro group, a cyano group, a C₁₋₆ alkyl group, a C₂₋₆ alkenyl group, a C₂₋₆ alkynyl group, a C₁₋₆ alkoxy group, an aryl group, an aryloxy group, an amino group which may be optionally substituted with C₁₋₃ alkyl group, an oxo group, a thioxo group, and a C₁₋₆ alkyl group which may be optionally substituted with one or more halogen atoms, and any groups not being affected under a reaction condition for oligomerization is preferably included.

[0014]

The term of "C₁₋₃ alkyl group which may be optionally substituted with one or more substituents" as used herein represents methyl, ethyl, n-propyl, or i-propyl, each may be substituted with one or more halogens, as defined in the

above term of "C₁₋₃ alkyl group which may be optionally substituted with one or more substituents". Specific examples thereof include perfluoroalkyl (such as trifluoromethyl, pentafluoroethyl, or heptafluoropropyl) and
5 the others.

[0015]

The term of "C₂₋₆ alkenyl which may be optionally substituted with one or more substituents" as used herein encompasses any straight chain alkenyl group having 2 to 6
10 carbon atoms (C₂₋₆), preferably having 2 to 4 carbon atoms (C₂₋₄), more preferably having 2 to 3 carbon atoms (C₂₋₃), any branched chain alkenyl group having 3 to 6 carbon atoms in which the identical or different branched chains are contained, any cyclic alkenyl group having 3 to 6 carbon
15 atoms, and any combinations thereof that have 4 to 6 carbon atoms. Specific examples of the straight chain alkenyl group having 2 to 6 carbon atoms include ethenyl, 1-propenyl, 2-propenyl, 1-butenyl, 2-butenyl, 1-pentenyl, 2-pentenyl, 3-pentenyl, 4-pentenyl, 1-hexenyl, and the others. Specific
20 examples the branched chain alkenyl group having 3 to 6 carbon atoms in which the identical or different branched chains are contained include isopropenyl, 1-methyl-1-propenyl, 1-methyl-2-propenyl, 2-methyl-1-propenyl, 2-methyl-2-propenyl, 1-methyl-2-butenyl and the others, and
25 also examples of the cyclic alkenyl group having 3 to 6

carbon atoms include a three to six membered monocyclic cycloalkenyl group, and specific examples thereof include cyclopropenyl, cyclobutenyl, cyclopentenyl, cyclohexenyl, and the others. Examples of the substituents include one or more (or preferably one to three) identical or different groups selected from the group consisting of a hydroxy group, a halogen atom, a nitro group, a cyano group, a C₁₋₆ alkyl group, a C₂₋₆ alkenyl group, a C₂₋₆ alkynyl group, a C₁₋₆ alkoxy group, an aryl group, an aryloxy group, an amino group which may be optionally substituted with C₁₋₃ alkyl group, an oxo group, a thioxo group, and a C₁₋₆ alkyl group which may be optionally substituted with one or more halogen atoms, and any groups not being affected under a reaction condition for oligomerization is preferably included.

[0016]

The term of "a C₂₋₆ alkynyl which may be optionally substituted with one or more substituents" as used herein encompasses any straight chain alkynyl group having 2 to 6 carbon atom (C₂₋₆), preferably any straight chain alkynyl group having 2 to 4 carbon atoms (C₂₋₄), more preferably any straight chain alkynyl group having 2 to 3 carbon atoms (C₂₋₃), any branched chain alkynyl group having 3 to 6 carbon atoms which have the identical or different branched chains, any cyclic alkynyl group having 3 to 6 carbon atoms, and any combinations thereof which have 4 to 6 carbon atoms.

Specific examples of the straight chain alkynyl group having 2 to 6 carbon atoms include ethynyl, 1-propynyl, 2-propynyl, 1-butynyl, 2-butynyl, 1-pentynyl, 2-pentynyl, 3-pentynyl, 4-pentynyl, 1-hexynyl and the others. Specific examples of

5 the branched chain alkynyl group having 3 to 6 carbon atoms in which the identical or different branched chains are contained include isopropynyl, 1-methyl-1-propynyl, 1-methyl-2-propynyl, 2-methyl-1-propynyl, 2-methyl-2-propynyl, 1-methyl-2-butynyl, and the others, and also examples of the

10 cyclic alkynyl group having 3 to 6 carbon atoms include preferably a three to six membered monocyclic cycloalkynyl group, and specific examples thereof include cyclobutynyl, cyclopentynyl, cyclohexynyl and the others, which are not limited thereto. Examples of the substituents include one

15 or more (or preferably one to three) identical or different groups selected from the group consisting of a hydroxy group, a halogen atom, a nitro group, a cyano group, a C₁₋₆ alkyl group, a C₂₋₆ alkenyl group, a C₂₋₆ alkynyl group, a C₁₋₆ alkoxy group, an aryl group, an aryloxy group, an amino group which
20 may be optionally substituted with C₁₋₃ alkyl group, an oxo group, a thioxo group, and a C₁₋₆ alkyl group which may be optionally substituted with one or more halogen atoms, and any groups not being affected under a reaction condition for oligomerization is preferably included.

25 [0017]

The term of "an aromatic group which may be optionally substituted with one or more substituents" as used herein means to encompass both of an aryl group and a heteroaryl group. The aryl group and the heteroaryl group may be
5 independently of each other substituted with one or more substituents. Examples of the substituents include one or more (or preferably one to three) identical or different groups selected from the group consisting of a hydroxy group, a halogen atom, a nitro group, a cyano group, a C₁₋₆ alkyl
10 group which may be optionally substituted with one or more substituents, a C₂₋₆ alkenyl group which may be optionally substituted with one or more substituents, a C₂₋₆ alkynyl group which may be optionally substituted with one or more substituents, a hydroxy group, a C₁₋₆ alkoxy group, an aryloxy
15 group, an amino group which may be optionally substituted with one or more C₁₋₃ alkyl groups, an C₁₋₆ alkyl groups which may be optionally substituted with one or more halogen atoms, and an aryl group, preferably a C₁₋₆ alkyl group which may be optionally substituted with one or more substituents, an
20 amino group which may be optionally substituted with one or more C₁₋₃ alkyl groups, and a halogen atom, and more preferably a trifluoromethyl group, a dimethylamino group, a chloro atom and the others, and any groups not being affected under a reaction condition for oligomerization is preferably
25 included.

[0018]

The term of "aryl (group)" as used herein represents a functional group or a substituent group which is derivatized from aromatic hydrocarbon group and encompasses a group composed of a plurality of cycles, and specifically, represents a monovalent group having 6 to 14 carbon atoms in which one hydrogen atom is excluded from aromatic hydrocarbon atom, consisting of one or more five membered and/or six membered ring, and include, for example, phenyl, indenyl, naphthyl, phenanthrenyl, anthracenyl and the others. Also examples of the substituents for the aryl group include one or more (or preferably one to three) identical or different groups selected from the group consisting of a hydroxy group, a halogen atom, a nitro group, a cyano group, a C₁₋₆ alkyl group which may be optionally substituted with one or more substituents, a C₂₋₆ alkenyl group which may be optionally substituted with one or more substituents, a C₂₋₆ alkynyl group which may be optionally substituted with one or more substituents, a C₁₋₆ alkoxy group, an aryloxy group, an amino group, an amino group which may be optionally substituted with one or more C₁₋₃ alkyl groups, and an aryl group, preferably a C₁₋₆ alkyl group which may be optionally substituted with one or more substituents, an amino group which may be optionally substituted with one or more C₁₋₃ alkyl groups, and a halogen atom, and more preferably a

trifluoromethyl group, a dimethylamino group, a chloro atom and the others, and any groups not being affected under a reaction condition for oligomerization is preferably included.

Specific examples of the aryl group which may be optionally substituted with one or more substituents include 2-methylphenyl, 3-methylphenyl, 4-methylphenyl, 2,6-dimethylphenyl, 2,4-dimethylphenyl, 2-chlorophenyl, 3-chlorophenyl, 4-chlorophenyl, 2,4-dichlorophenyl, 2,5-dichlorophenyl, 2,6-dichlorophenyl, 2-bromophenyl, 4-methoxyphenyl, 4-chlorophenyl, 2-nitrophenyl, 4-nitrophenyl, 2-nitrophenyl, 2,6-dinitrophenyl, 2-trifluoromethylphenyl, 3-trifluoromethylphenyl, 4-trifluoromethylphenyl, 2-dimethylaminophenyl, 3-dimethylaminophenyl, 4-dimethylaminophenyl, and biphenyl and the others. Preferred examples of the aryl group include a phenyl group which is substituted with halogen atom, C₁₋₆ alkoxy group or nitro group, or an unsubstituted phenyl group.

[0019]

The term of "heteroaryl (group)" as used herein represents a monovalent group in which one hydrogen atom is excluded from optional heteroaromatic compound containing one or more heteroatoms (such as nitrogen atom, oxygen atom, and/or sulfur atom) in a ring structure and having 5 to 12 carbon atoms, consisting of one or more five membered and/or

six membered ring, and preferably five membered cyclic or six membered cyclic heteroaryl group. Specific examples of the heteroaryl group include a five membered cyclic heteroaryl group selected from pyrrolyl, furyl, thienyl, pyrazolyl, imidazolyl, triazolyl, tetrazolyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, oxadiazolyl, and thiadiazolyl; a six membered cyclic heteroaryl group selected from pyridyl, pyridazinyl, pyrimidinyl, and pyrazinyl, and preferably triazolyl, oxadiazolyl, thiadiazolyl group, pyrimidinyl, or pyrazinyl. Also examples of the substituents for the heteroaryl group include one or more (or preferably one to three) identical or different groups selected from the group consisting of a hydroxy group, a halogen atom, a nitro group, a cyano group, a C₁₋₆ alkyl group which may be optionally substituted with one or more substituents, a C₂₋₆ alkenyl group which may be optionally substituted with one or more substituents, a C₂₋₆ alkynyl group which may be optionally substituted with one or more substituents, a C₁₋₆ alkoxy group, an aryloxy group, an amino group, an amino group which may be optionally substituted with one or more C₁₋₃ alkyl groups, and an aryl group, preferably a C₁₋₆ alkyl group which may be optionally substituted with one or more substituents, an amino group which may be optionally substituted with one or more C₁₋₃ alkyl groups, and a halogen atom, and more preferably a

trifluoromethyl group, a dimethylamino group, a chloro atom and the others, and any groups not being affected under a reaction condition for oligomerization is preferably included.

Specific examples of the heteroaryl group which may be optionally substituted with one or more substituents include 1,2,4-triazolyl, 1,5-dimethyl-1,2,4-triazolyl, 1,2,4-oxadiazolyl, 5-methyl-1,2,4-oxadiazol-3-yl, 1,2,4-thiadiazolyl, 3-methyl-1,2,4-thiadiazol-5-yl, 1,3-pyrimidinyl, 1,5-pyrimidinyl, 4-trifluoromethyl, 1,5-pyrimidinyl, 4-dimethylamino-2,4-pyrimidinyl, 3-dimethylamino-2,4-pyrimidinyl, 3-chloro-1,5-pyrimidinyl, and 1,4-pyridazinyl, and preferably 1,5-dimethyl-1,2,4-triazol-3-yl, 1,2,4-oxadiazolyl, 3-methyl-1,2,4-thiadiazol-5-yl, 1,5-pyrimidinyl, 4-trifluoromethyl-1,5-pyrimidinyl, 4-dimethylamino-2,4-pyrimidinyl, 3-dimethylamino-2,4-pyrimidinyl, 3-chloro-1,5-pyrimidinyl, and 1,4-pyridazinyl, and more preferably 1,5-dimethyl-1,2,4-triazol-3-yl, 1,2,4-oxadiazol-3-yl, and 3-methyl-1,2,4-thiadiazol-5-yl.

[0020]

Examples of the term of "five or six membered heteroaryl group containing one or more heteroatoms selected from the group consisting of nitrogen atom, oxygen atom and sulfur atom, which may be optionally substituted with one or more substituents" as used herein include a five membered cyclic

or six membered cyclic heteroaryl group which may optionally substituted with one or more substituents, which is defined in the above-mentioned term of "heteroaryl group".

[0021]

5 The term of "five or six membered heteroaryl group containing two or three heteroatoms selected from nitrogen atom, oxygen atom, and sulfur atom, provided that at least two nitrogen atoms are included, which may be optionally substituted with one or more substituents" as used herein
10 represents triazolyl, oxadiazolyl, thiadiazolyl, pyrimidinyl, and pyridazinyl, each may be substituted with one or more substituents, which is defined in the above-mentioned term of "heteroaryl group.

[0022]

15 The term of "sulfonyl group which is substituted with one substituent selected from the group consisting of a C₁-6 alkyl group which may be optionally substituted with one or more substituents and an aromatic group which may be optionally substituted with one or more substituents"
20 represents a sulfonyl (S(O)₂) group which is substituted with one substituents thereof. The C₁₋₆ alkyl group which may be optionally substituted with one or more substituents, or the aromatic group which may be optionally substituted with one or more substituents are as defined above, and specific
25 examples of the substituents include methyl, trifluoromethyl,

phenyl, 4-methylphenyl, and the others, preferably methyl and phenyl.

[0023]

The term of "C₁₋₆ alkoxy group" as used herein represents
5 a monovalent group in which the C₁₋₆ alkyl binds to an oxygen atom, and represents a C₁₋₆ alkyl-O group. Specific examples thereof include methoxy, ethoxy, n-propoxy, i-propoxy, n-butoxy, t-butoxy, s-butoxy, 3-methylbutoxy, and the others, which are not limited thereto.

10 [0024]

The term of "aryloxy (group)" as used herein represents a monovalent group in which the aryl group is bonded to an oxygen atom. Specific examples thereof include phenoxy, p-tolyl and the others, which are not limited thereto.

15 [0025]

The term of "amino group which may be optionally substituted with one or more C₁₋₃ alkyl groups" as used herein represents an amino group which may be optionally substituted with one or more hydrogen atoms may be optionally substituted
20 with a C₁₋₃ alkyl group. Specific examples thereof include methylamino, ethylamino, isopropylamino, dimethylamino, diethylamino, methylethylamino, and the others, which are not limited thereto.

[0026]

25 Examples of the term of "halogen (atom)" as used herein

include fluorine atom (fluoro), chlorine atom (chloro), bromine atom (bromo), and iodine atom (iodo), and fluorine atom or chloro atom is preferred.

[0027]

5 Herein when the compound of the present invention, an intermediate compound, or a starting material and the others has a functional group (such as, a hydroxy group, an amino group, a carboxyl group and the others), the functional group may be protected with a protecting group that is usually
10 used in an organic synthetic chemistry according to the method described in Theodora W. Greene, Peter G. M. Wuts, "Protective Groups in Organic Synthesis" 4th. ed., John Wiley & Sons, Inc. 1999, and after the reaction, the protecting group may be then removed to obtain the desirable compound.
15 Examples of the protecting group include the protecting group that is usually used in an organic synthetic chemistry as described in the same document, and is also usually used in an organic synthetic chemistry, and examples of each of the protecting group depending on the functional group are
20 described below.

[0028]

 The term of "protecting group" described in terms of "a protecting group for hydroxy group", "a protecting group for amino group", "a protecting group for phosphate group", or
25 "a protecting group for mercapto group" as used herein is

not particularly limited to any group as long as it can protect amino group, hydroxy group, phosphate group or mercapto group stably during a nucleic acid synthesis. Specific examples thereof include a protecting group that is

5 stable under acidic or neutral condition and also can be cleaved by a chemical method such as a hydrogenolysis, a hydrolysis, an electrolysis, and a photolysis. Examples of the protecting group includes an alkyl group having 1 to 6 carbon atoms; an alkenyl group having 2 to 6 carbon atoms;

10 an alkynyl group having 2 to 6 carbon atoms; an acyl group; a tetrahydropyranyl group or a tetrahydrothiopyranyl group; a tetrahydrofuranyl group or a tetrahydrothiofuranyl group; a silyl group; a methyl group substituted with alkoxy group having 1 to 6 carbon atoms; a methyl group substituted with

15 alkoxy group having 1 to 6 carbon atoms, said alkoxy group being substituted with alkoxy group having 1 to 6 carbon atoms; a methyl group substituted with alkoxy group having 1 to 6 carbon atoms said alkoxy group being substituted with halogen atom; an ethyl group substituted with alkoxy group

20 having 1 to 6 carbon atoms; an ethyl group substituted with halogen atom; a methyl group substituted with one to three aryl groups; a methyl group substituted with one to three aryl groups, said aryl groups being substituted with alkyl group having 1 to 6 carbon atoms, alkenyl group having 2 to

25 6 carbon atoms, alkynyl group having 2 to 6 carbon atoms,

alkoxy group having 1 to 6 carbon atoms, halogen atom, and/or cyano group; a carbonyl group substituted with alkoxy group having 1 to 6 carbon atoms; an aryl group substituted with halogen atom, alkoxy group having 1 to 6 carbon atoms and/or
5 nitro group; a carbonyl group substituted with alkoxy group having 1 to 6 carbon atoms, said alkoxy group being substituted with halogen atom and/or silyl group substituted with alkyl group having 1 to 6 carbon atoms; an alkenyloxycarbonyl group; an aralkyloxycarbonyl group which
10 may be optionally substituted with alkoxy group having 1 to 6 carbon atoms and/or aryl group substituted with nitro group; and the others.

[0029]

The term of "protecting group for hydroxy group"
15 represents a protecting group that is usually used in an organic synthetic chemistry (in particular, nucleic acid synthesis), and includes, for example, an aliphatic acyl group; an aromatic acyl group; an aminocarbonyl group which may be substituted; an alkoxycarbonyl group which may be
20 substituted; an aliphatic sulfonyl group; an aromatic sulfonyl group; a methyl group substituted with one to three aryl groups; a methyl group substituted with one to three aryl groups, said aryl group being substituted with alkyl group having 1 to 6 carbon atoms, alkoxy group having 1 to 6
25 carbon atoms, halogen atom and/or cyano group; or a silyl

group. Specific examples thereof include benzyl (Bn), 4,4'-dimethoxytrityl (DMTr), 4-methoxytrityl, triphenylmethyl, 2-naphthylmethyl, diphenylaminocarbonyl (DPC), cyanoethoxycarbonyl, tetrahydropyranyl, trimethylsilyl, triethylsilyl, triisopropylsilyl, t-butyldimethylsilyl, t-butyldiphenylsilyl, triphenylsilyl, 4-methoxybenzyl (p-methoxybenzyl), 3,4-dimethoxybenzyl, 2,6-dimethoxybenzyl, p-phenylbenzyl, methanesulfonyl, trifluoromethanesulfonyl, methoxymethyl, benzoyl (Br), phenoxyacetyl, and acetyl and the others, which should not be limited thereto. Benzyl (Bn), 4,4'-dimethoxytrityl (DMTr), t-butyldimethylsilyl, t-butyldiphenylsilyl, trimethylsilyl (TMS), diphenylaminocarbonyl (DPC), methanesulfonyl, and trifluoromethanesulfonyl are preferably included, and 4,4'-dimethoxytrityl (DMTr) is more preferably included.

[0030]

The term of "phosphate group which may be optionally substituted" as used herein represents a phosphate, a phosphite, or a hydrophosphite, each of which may have optionally substituents (which encompasses protecting group). The term encompasses a phosphate group represented by a formula of $-P(R^{P1})R^{P2}$, wherein R^{P1} and R^{P2} represent independently of each other a hydroxy group, a hydroxy group that is protected with a protecting group for nucleic acid synthesis, a mercapto group, a mercapto that is protected

with a protecting group for nucleic acid synthesis, an amino group, an alkoxy group having 1 to 5 carbon atoms, an alkylthio group having 1 to 6 carbon atoms, a cyanoalkoxy group having 1 to 6 carbon atoms, or an amino group that is substituted with alkyl group having 1 to 6 carbon atoms. Here the groups of the above-mentioned formula wherein R^{P1} represents OR^{P1a} and R^{P2} represents NR^{P2a} is referred to as "phosphoramidite group", which is a preferred example. The R^{P1a} represents an alkyl group having 1 to 5 carbon atoms or a cyanoalkyl group having 1 to 6 carbon atoms, and the R^{P2a} represents an alkyl group having 1 to 6 carbon atoms. Specific examples of "phosphoramidite group" include a group represented by a formula of $-P(O(CH_2)_2CN)(N(iPr)_2)$, or a group represented by a formula of $-P(OCH_3)(N(iPr)_2)$ and the others, which is not limited thereto. The group represented by $-P(O(CH_2)_2CN)(N(iPr)_2)$ is preferably included. Herein "iPr" represents an isopropyl group.

[0031]

The term of "phosphate group which may be optionally substituted with substituents" as used herein may form a chiral auxiliary containing a phosphorus atom. Specific examples of the chiral auxiliary containing a phosphorus atom include an optical active bicyclic oxazaphospholidine, which is described in non-patent document, N. Oka et al., J. AM. CHEM. SOC. 2008, 130, 16031, and an optical active 2-

thio-1,3,2-oxa-thiaphospharane containing five valent phosphorus atom, which is described in non-patent document, K. W. Knouse et al., Science 10. 1126/science. aau3369 (2018), which is not limited thereto.

5 [0032]

The term of "protecting group for amino group" represents a protecting group that is usually used in an organic synthetic chemistry (in particular, nucleic acid synthesis), and include, for example, an aliphatic acyl
10 group; an aromatic acyl group; an alkoxycarbonyl group which may be optionally substituted; a methyl group substituted with one to three aryl groups; and a methyl group substituted with aryl groups, said aryl group being substituted with halogen atom and/or cyano group. Specific examples thereof
15 include acetyl (Ac), phenoxyacetyl (Pac), t-butylphenoxyacetyl group (Tac), p-isopropyl phenoxyacetyl group (iPc-Pac), trifluoroacetyl, propionyl group, isobutyryl, benzoyl (Bz), methoxycarbonyl, ethoxycarbonyl, t-butoxycarbonyl (Boc), trimethylsilylethoxycarbonyl (Teoc),
20 cyanoethoxycarbonyl (Ceoc), benzyloxycarbonyl (Cbz), allyloxycarbonyl, 9-fluorenylmethoxycarbonyl (Fmoc), dimethylaminomethylenyl, 2,2,2-trichloroethoxycarbonyl, t-amylloxycarbonyl, 4-methoxybenzyl, triphenylmethyl, 2-nitrobenzenesulfonyl, 2,4-dinitrobenzenesulfonyl or 2-
25 (trimethylsilyl)ethoxymethyl and the others, which are not

limited thereto. Preferably, isobutyryl, benzoyl (Bz), t-butoxycarbonyl (Boc), and trimethylsilylethoxycarbonyl (Teoc) are included.

[0033]

5 The term of "protecting group for a phosphate group" represents a protecting group that is usually used in an organic synthetic chemistry (in particularly, nucleic acid synthesis), and examples thereof include an alkyl group having 1 to 6 carbon atoms and/or an alkyl group having 1 to
10 6 carbon atoms substituted with cyano group; an aralkyl group; an aralkyl group substituted with aryl group, said aryl group being substituted with nitro group and/or halogen atom; an aryl group substituted with alkyl group having 1 to 6 carbon atoms, halogen atom, or nitro group; 2-cyanoethyl
15 group; 2,2,2-trichloroethyl group; benzyl group; 2-chlorophenyl group; and 4-chlorophenyl group, and the others, which are not limited thereto.

[0034]

 The term of "protecting group for mercapto group"
20 represents a protecting group that is usually used in an organic synthetic chemistry (in particularly, nucleic acid synthesis), and includes, for example, an aliphatic acyl group, an aromatic acyl group, a benzoyl group (Bz), and the others, which are not limited thereto.

25 [0035]

The term of "leaving group" as used herein represents a partial substrate having electron pairs that is created when the substrate is cleaved by a cleavage of heterolysis during reaction(s), and encompasses a halogen atom (such as fluorine atom, chlorine atom, bromine atom, iodine atom), and "a leaving group for hydroxy". Examples of the leaving group for hydroxyl include a sulfonyloxy group (such as paratolunesulfonyloxy, mesyloxy, and trifluoromethanesulfonyloxy), an acyloxy group (preferably, saturated or unsaturated acyloxy group having 1 to 8 carbon atoms, for example, aryl group represented by $R^L-C(=O)-O-$ wherein the R^L represents an aryl group which may be optionally substituted with alkyl group (the total number of carbon atom is preferably 6 to 8, for example, phenyl, or p-tolyl), an aryloxy group which may be optionally substituted with alkyl group (the total number of carbon atom is preferably 6 to 8, for example, phenoxy, or p-tolyloxy), an aralkyl group (the total number of carbon atom is preferably 7 to 9, for example, benzyl), an arylalkenyl group (the total number of carbon atom is preferably 8 or 9, for example, cinnamyl), an aralkyloxy group (the total number of carbon atom is 7 to 15, for example, benzyloxy, or 9-fluorenylmethyloxy), an alkoxy group (a straight chain or branched chain alkoxy group, for example, methoxy, ethoxy, t-butoxy), and specific examples of the leaving group include

iodo, bromo, chloro, fluoro, mesyloxy, methanesulfonyloxy,
 trifluoromethanesulfonyloxy, ethanesulfonyloxy, 2,2,2-
 trifluoroethanesulfonyloxy, propanesulfonyloxy, iso-
 propanesulfonyloxy, butanesulfonyloxy,
 5 nonafluorobutanesulfonyloxy, heptafluoropropan-1-
 sulfonyloxy, pentanesulfonyloxy,
 pentafluoroethanesulfonyloxy, pentanesulfonyloxy,
 cyclopentanesulfonyloxy, hexanesulfonyloxy,
 cyclohexanesulfonyloxy, o-toluenesulfonyloxy, m-
 10 toluenesulfonyloxy, p-toluenesulfonyloxy,
 benzenesulfonyloxy, o-bromobenzenesulfonyloxy, m-
 bromobenzenesulfonyloxy, p-bromobenzenesulfonyloxy, o-
 nitrobenzenesulfonyloxy, m-nitrobenzenesulfonyloxy, and p-
 nitrobenzenesulfonyloxy, and the others, which are not
 15 limited thereto. Preferred examples of the leaving group
 include methanesulfonyloxy (mesyloxy; Ms-O-),
 trifluoromethanesulfonyloxy, and p-toluenesulfonyloxy.

[0036]

Examples of the term of "acyl group" as used herein
 20 include an aliphatic acyl group and aromatic acyl group.
 Specific examples of the aliphatic acyl group include an
 alkylcarbonyl group, such as formyl, acetyl, propionyl,
 butyryl, isobutyryl, pentanoyl, pivaloyl, valeryl,
 isovaleryl, octanoyl, nonanoyl, decanoyl, 3-methylnonanoyl,
 25 8-methylnonanoyl, 3-ethyloctanoyl, 3,7-dimethyloctanoyl,

undecanoyl, dodecanoyl, tridecanoyl, tetradecanoyl, pentdecenoyl, hexadecanoyl, 1-methylpentdecenoyl, 14-methylpentadecanoyl, 13,13-dimethyltetradecanoyl, heptadecanoyl, 15-methylhexadecanoyl, octadecanoyl, 1-methylheptadecanoyl, nonadecanoyl, eicosanoyl, and henaicosanoyl; an aryloxyalkylcarbonyl group such as phenoxyacetyl (pac); a carboxylated alkylcarbonyl group such as succinoyl, glutaroyl, and adipoyl; a carbonyl group that is substituted with alkyl group having 1 to 6 carbon atoms, said alkyl group being substituted with halogen atom, such as chloroacetyl group, dichloroacetyl, trichloroacetyl, trifluoroacetyl; an alkoxyalkylcarbonyl group having 1 to 6 carbon atoms such as methoxyacetyl; and an unsaturated alkylcarbonyl group such as (E)-2-methyl-2-butenoyl. Also, examples of the aromatic acyl group include an arylcarbonyl group such as benzoyl, α -naphthoyl, and β -naphthoyl; a halogenoarylcarbonyl group such as 2-bromobenzoyl and 4-chlorobenzoyl; an arylcarbonyl group that is substituted with alkyl group having 1 to 6 carbon atoms, such as 2,4,6-trimethylbenzoyl, and 4-toluoyl; an arylcarbonyl group that is substituted with alkoxy group having 1 to 6 carbon atoms, such as 4-anisoyl; a carboxylated arylcarbonyl group such as 2-carboxybenzoyl, 3-carboxybenzoyl, and 4-carboxybenzoyl; a nitroated arylcarbonyl group such as 4-nitrobenzoyl and 2-nitrobenzoyl; a carbonylated arylcarbonyl group that is

substituted with alkoxy group having 1 to 6 carbon atoms, such as 2-(methoxycarbonyl)benzoyl; and an arylated arylcarbonyl group such as 4-phenylbenzoyl, which are not limited thereto.

5 [0037]

The term of "aralkyl group" as used herein represents an alkyl group of 1 to 6 carbon atoms, preferably 1 to 4 carbon atoms, more preferably 1 to 3 carbon atoms, said alkyl group being substituted with aromatic hydrocarbon group (such as, 6 to 14 membered monocyclic, bicyclic, or tricyclic aromatic hydrocarbon group). Specific examples thereof include benzyl, phenethyl, 1-naphthylmethyl, 2-naphthylmethyl, and the others, but which are not limited thereto.

15 [0038]

Examples of term of "silyl group" as used herein include a silyl group that is substituted with alkyl group having 1 to 6 carbon atoms, such as trimethylsilyl, triethylsilyl, isopropyldimethylsilyl, t-butyldimethylsilyl, methyl diisopropylsilyl, methyl di-t-butylsilyl, triisopropylsilyl; a silyl group that is substituted with one or two aryl group and also three alkyl groups having 1 to 6 carbon atoms, such as t-butyldiphenylsilyl, diphenylmethylsilyl, butyldiphenylbutylsilyl, diphenylisopropylsilyl, and phenyldiisopropylsilyl; and

triphenylsilyl, but which are not limited thereto.

[0039]

The term of " β form" as used herein represents the a compound having a stereochemistry in which the direction of substituting the substituted base moiety of nucleic acid at 1' position of a ribose of nucleic acid and the direction of substituting the substituted 5'-position side chain at 4' position of a ribose of nucleic acid are the same direction of the substitution. For a bridged artificial nucleic acid 2',4'-LNA, the β form represents the compound having the stereo configuration in which the direction of substituting the substituted base moiety of nucleic acid at 1' position of a ribose of nucleic acid and the direction of substituting the substituted 5'-position side chain that is not used for bridging substituted at 4' position of ribose of nucleic acid are the same direction of the substitution.

[0040]

The term of " β selective" as used herein represents that a β form can be obtained.

[0041]

The "a base moiety of nucleic acid" in the term of "a base moiety of nucleic acid which may be optionally substituted with one or more substituents" as used herein includes, for example, a base moiety of natural nucleic acid, and a base moiety of unnatural nucleic acid, which contains

an aromatic heterocycle, and encompasses a monocyclic ring, a bicyclic ring, and a tricyclic ring. Here it should be clearly understood that base moieties of various nucleic acid that has been now considered "unnatural" to a skilled person in the art, could be found in nature hereafter. Thus "nucleic acid base moiety" includes not only any heterocycle contained in publicly known purine and pyrimidine, but also their heterocyclic analogues and tautomers. Specific examples of the base moiety of nucleic acid include adenine, guanine, thymine, cytosine, uracil, purine, xanthine, diaminopurine, 8-oxo-N⁶-methyladenine, 7-deazaxanthine, 7-deazaguanine, N⁴,N⁴-ethanocytosine, N⁶,N⁶-ethano-2,6-diaminopurine, 5-methylcytosine, 5-(C³-C⁶)-alkynylcytosine, 5-fluorocytosine, 5-bromouracil, pseudoisocytosine, 2-hydroxy-5-methyl-4-triazolopyridine, isocytosine, isoguanine, inosine, N⁶-arylpurine, N⁶-acylpurine, N⁶-benzylpurine, N⁶-halopurine, N⁶-vinylpurine, N⁶-acetylenic purine, N⁶-acylpurine, N⁶-hydroxyalkylpurine, N⁶-thioalkylpurine, N²-alkylpurine, N⁶-alkylpyrimidine, N⁴-acylpyrimidine, N⁴-benzylpurine, N⁴-halopyrimidine, N⁴-vinylpyrimidine, N⁴-acetylenic pyrimidine, N⁴-acetylpyrimidine, N⁴-hydroxyalkylpyrimidine, N⁶-thioalkylpyrimidine, 6-azapyrimidine, 6-azacytosine, 2-and/or 4-mercaptopyrimidine, uracil, C⁵-alkylpyrimidine, C⁵-benzylpyrimidine, C⁵-halopyrimidine, C⁵-vinylpyrimidine,

C⁵-acetylenicpyrimidine, C⁵-acylpyrimidine, C⁵-hydroxyalkylpurine, C⁵-amidopyrimidine, C⁵-cyanopyrimidinyl, C⁵-nitropyrimidine, C⁵-aminopyrimidine, N²-alkylpurine, N²-alkyl-6-thiopurine, 5-cytidinyl, 5-azauracilyl, 5 trazolopyridinyl, imidazolopyridinyl, pyrrolopyrimidinyl, pyrazolopyrimidinyl, and the others, which are not limited thereto. Preferred examples of the base moiety of nucleic acid include adenine, guanine, 2,6-diaminopurine, thymine, 2-thiothymine, cytosine, 5-methylcytosine, uracil, 5-10 fluorocytosine, xanthine, 6-aminopurine, 2-aminopurine, 6-chloro-2-aminopurine, and 6-chloropurine, and particular preferred examples of the base moiety of nucleic acid include adenine, guanine, cytosine, 5-methylcytosine, thymine, and uracil. The base moiety of the nucleic acid may be further 15 substituted with one or more substituents, and examples of the substituents include hydroxy group, C₁₋₆ alkoxy group, mercapto group, C₁₋₆ alkylthio group, amino group, amino group substituted with C₁₋₆ alkyl group, C₁₋₆ alkyl group, C₁₋₆ alkynyl group, oxo group, thioxo group, and halogen atom. 20 If necessary or if desirable, the functional oxygen atom, sulfur atom and nitrogen atom on the base moiety may be protected and/or deprotected. The appropriate protecting group is well known to a skilled person in the art, and encompasses, for example, the above-mentioned protecting 25 group for hydroxy group and protecting group for amino group,

and includes diphenylaminocarbonyl group, silyl group (such as trimethylsilyl group, dimethylhexylsilyl group, t-butyltrimethylsilyl group, and t-butylphenylsilyl group), trityl group, alkyl group, acyl group (such as acetyl group, propionyl group, isobutyryl group, benzoyl group (Bz), phenoxyacetyl group (Pac)), alkoxycarbonyl group (such as t-butoxycarbonyl group (Boc), benzyloxycarbonyl group (Cbz), diphenylaminocarbonyl group (DPC), cyanoethoxycarbonyl group (Ceoc)), sulfonyl group (such as methanesulfonyl group, and p-toluenesulfonyl group), dimethylaminomethylenyl group, and the others, which are not limited thereto.

[0042]

The term of "artificial nucleic acid" as used herein encompasses artificial nucleoside, artificial nucleotide (herein, one nucleoside or nucleotide is sometimes referred to as a monomer) or artificial oligonucleotide. These artificial nucleic acids are not natural nucleic acids, but nucleic acids that can be produced only artificially. Examples of these artificial nucleic acids include those in which the nucleic acid base moiety contains an unnatural base, those having a sugar in which the sugar moiety is modified, and/or those containing an unnatural phosphate group as a phosphate moiety, and the artificial nucleic acids as used herein represents those in which the sugar moiety contains an unnatural sugar, in particular, those containing

a (deoxy) ribose in which the carbon atoms at 2' position and 4' position are bridged.

[0043]

The term of "artificial oligonucleotide" as used herein
5 represents a substance in which two or more of the identical
or different "artificial nucleotides" are bonded with each
other via a phosphodiester bond or a thiophosphodiester bond
and the like, and includes a substance in which preferably
2 up to 100, more preferably 5 up to 50, most preferably 10
10 up to 30 artificial nucleotides are bonded, or a substance
in which the nucleotides together with these complementary
strands forms a double strand. Herein, an oligonucleotide
in which two or more nucleotides are bonded with each other
is sometimes referred to as oligomer.

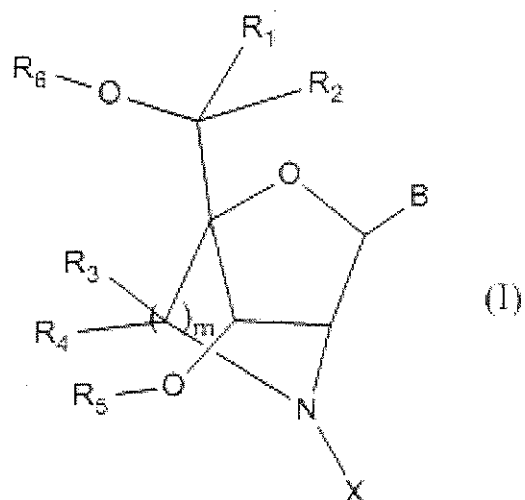
15 [0044]

Embodiments of the compounds of the present invention
are described.

[0045]

Compound represented by general formula I

20 According to one embodiment of the present application,
a compound represented by general formula I:



[wherein

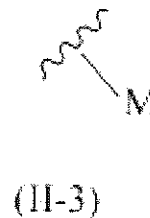
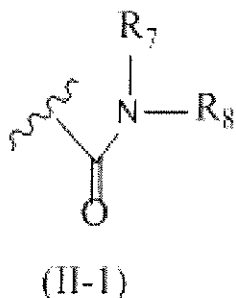
B represents a base moiety of nucleic acid wherein the base moiety may be optionally substituted with one or more substituents;

R_1 , R_2 , R_3 and R_4 represent independently of each other a hydrogen atom, or a C_{1-6} alkyl group which may be optionally substituted with one or more substituents;

R_5 and R_6 represent independently of each other a hydrogen atom, a protecting group for hydroxy group, or a phosphate group which may be optionally substituted with substituents;

m is an integer of 1 or 2;

X represents a group represented by the following formula (II-1), (II-2) or (II-3):



the symbols:



which is described in the formula (II-1), (II-2) or (II-3)

5 represents a binding point to 2'-amino group;

R_7 and R_8 represent independently of each other a hydrogen atom, a C_{1-6} alkyl group which may be optionally substituted with one or more substituents, a C_{2-6} alkenyl group which may be optionally substituted with one or more substituents, a C_{2-6} alkynyl group which may be optionally substituted with one or more substituents, or an aromatic group which may be optionally substituted with one or more substituents;

A represents an aromatic group;

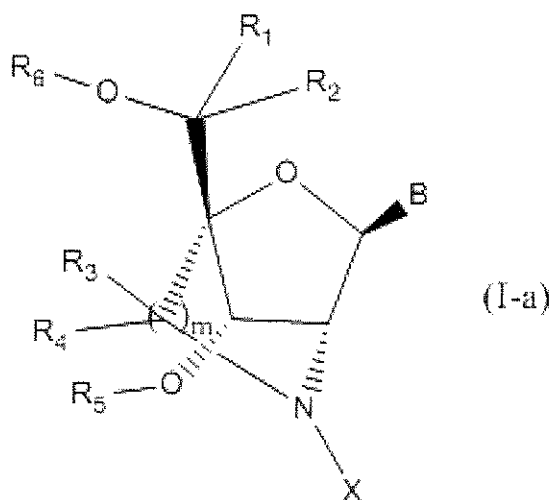
15 M represents a sulfonyl group which is substituted with one substituent selected from the group consisting of a C_{1-6} alkyl group which may be optionally substituted with one or more substituents and an aromatic group which may be optionally substituted with one or more substituents.]

20 or salts thereof is provided.

[0046]

According to one embodiment of the present application, the compound represented by general formula (I) or salts thereof represents a compound represented by general formula

5 I-a:



or salts thereof is preferably included (wherein R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , B , X and m are the same as defined in a general formula (I)).

10 [0047]

According to one embodiment of the present application, the compound represented by formula (I) or salts thereof wherein a ring B in the formula (I) represents an adeninyl group which may be optionally substituted with one or more substituents, a guaninyl group which may be optionally substituted with one or more substituents, a cytosinyl group which may be optionally substituted with one or more substituents, a 5-methylcytosinyl group which may be

15

optionally substituted with one or more substituents or an uracilyl group which may be optionally substituted with one or more substituents is provided.

[0048]

5 According to one embodiment of the present application, the compound represented by formula (I) or salts thereof wherein in the formula (I), a ring B represents a 5-methylcytosinyl group which may be optionally substituted with one or more substituents or a 5-uracilyl group which
10 may be optionally substituted with one or more substituents is provided.

[0049]

 According to one embodiment of the present application, the compound represented by formula (I) or salts thereof
15 wherein in the formula (I), R_1 , R_2 , R_3 and R_4 represent independently of each other a hydrogen atom, and m is an integer of 1 is provided.

[0050]

 According to one embodiment of the present application,
20 the compound represented by formula (I) or salts thereof wherein in the formula (I), R_6 represents a hydrogen atom or DMTr group, and R_5 represents a hydrogen atom or -
 $P(O(CH_2)_2CN)(N(i\text{pr})_2)$.

[0051]

25 According to one embodiment of the present application,

the compound represented by formula (I) or salts thereof wherein in the formula (I), R_6 represents DMTr group, and R_5 represents a hydrogen atom or $-P(O(CH_2)_2CN)(N(ipr)_2)$.

[0052]

5 According to one embodiment of the present application, the compound represented by formula (I) is roughly divided into the formula (II-1), (II-2) or (II-3).

First, the compound wherein in the formula (I), X represents a group represented by formula (II-1) (hereinafter, referred to as "Urea type compound") is included.

[0053]

15 According to one embodiment of the present application, a compound represented by general formula (I) or salts thereof wherein

X represents a group represented by formula (II-1); and

20 R_7 and R_8 represent independently of each other a hydrogen atom, a C1-6 alkyl group which may be optionally substituted with one or more substituents, or an aromatic group which may be optionally substituted with one or more substituents is provided.

[0054]

25 According to one embodiment of the present application, a compound represented by general formula (I) or salts

thereof wherein

X represents a group represented by formula (II-1); and

R₇ and R₈ represent independently of each other a
hydrogen atom, a C1-3 alkyl group which may be optionally
5 substituted with one or more substituents, or an aromatic
group which may be optionally substituted with one or more
substituents
is provided.

[0055]

10 According to one embodiment of the present application,
a compound represented by general formula (I) or salts
thereof wherein

X represents a group represented by formula (II-1); and

R₇ and R₈ represent independently of each other a
15 hydrogen atom, a methyl group, an isopropyl group, or a
phenyl group
is provided.

[0056]

20 According to one embodiment of the present application,
a compound represented by general formula (I) or salts
thereof wherein

X represents a group represented by formula (II-1); and

a combination of R₇ and R₈ represents a combination
wherein both of R₇ and R₈ represent a hydrogen atom; a
25 combination wherein both of R₇ and R₈ represent a methyl

group; a combination wherein R₇ represents a hydrogen atom and R₈ represents a methyl group; a combination wherein R₇ represents a hydrogen atom and R₈ represents an isopropyl group; or a combination wherein R₇ represents a hydrogen atom and R₈ represents a phenyl group is provided.

[0057]

According to one embodiment of the present application, a compound represented by formula (I) or salts thereof wherein

10 X represents a group represented by formula (II-1); and a combination of R₇ and R₈ represents a combination wherein both of R₇ and R₈ represent a hydrogen atom; a combination wherein both of R₇ and R₈ represent a methyl group; a combination wherein one of R₇ and R₈ represents a hydrogen atom and the other thereof represents a methyl group; a combination wherein one of R₇ and R₈ represents a hydrogen atom and the other thereof represents an isopropyl group; or a combination wherein one of R₇ and R₈ represents a hydrogen atom and the other thereof represents a phenyl group

20 group is provided.

[0058]

According to one embodiment of the present application, a compound represented by formula (I) or salts thereof

25 wherein

X represents a group represented by formula (II-1); and
one of R₇ and R₈ represents a hydrogen atom, and the
other thereof represents a methyl group,
is provided.

5 [0059]

According to one embodiment of the present application,
a compound represented by formula (I) or salts thereof
wherein

X represents a group represented by formula (II-1); and
10 one of R₇ and R₈ represents a hydrogen atom, and the
other thereof represents an isopropyl group,
is provided.

[0060]

Also the compound wherein in the formula (I), X
15 represents a group represented by formula (II-2)
(hereinafter, referred to as "heteroaryl type compound") may
be included.

[0061]

According to one embodiment of the present application,
20 a compound represented by formula (I) or salts thereof
wherein

X represents a group represented by formula (II-2); and

A represents a five or six membered heteroaryl group
containing one or more heteroatoms selected from the group
25 consisting of nitrogen atom, oxygen atom and sulfur atom,

which may be optionally substituted with one or more substituents,
is provided.

[0062]

5 According to one embodiment of the present application,
a compound represented by formula (I) or salts thereof
wherein

X represents a group represented by formula (II-2); and

A represents a five or six membered heteroaryl group
10 containing two or three heteroatoms selected from nitrogen
atom, oxygen atom, and sulfur atom, provided that at least
two nitrogen atoms are included, which may be optionally
substituted with one or more substituents,
is provided.

15 [0063]

According to one embodiment of the present application,
a compound represented by formula (I) or salts thereof
wherein

X represents a group represented by formula (II-2); and

20 A represents a five or six membered heteroaryl group
containing one or more heteroatoms selected from the group
consisting of nitrogen atom, oxygen atom and sulfur atom,
provided that at least two nitrogen atoms are included, which
may be optionally substituted with one or more substituents,
25 and the substituent is selected independently of each other

from a group consisting of a C₁₋₃ alkyl group which may be optionally substituted with one or more halogen atoms, a halogen atom, and an amino group which may be optionally substituted with one or more C₁₋₃ alkyl group,

5 is provided.

[0064]

According to one embodiment of the present application, a compound represented by formula (I) or salts thereof wherein

10 X represents a group represented by formula (II-2); and

A represents a five or six membered heteroaryl group containing one or more heteroatoms selected from the group consisting of nitrogen atom, oxygen atom and sulfur atom, provided that at least two nitrogen atoms are included, which
15 may be optionally substituted with one or more substituents, and the substituent is selected independently of each other from a group consisting of a C₁₋₃ alkyl group which may be optionally substituted with one or more halogen atoms, a halogen atom, and an amino group which may be optionally
20 substituted with one or more C₁₋₃ alkyl group,
is provided.

[0065]

According to one embodiment of the present application, a compound represented by formula (I) or salts thereof
25 wherein

X represents a group represented by formula (II-2); and
the five or six membered heteroaryl group as A group is
selected from the group consisting of a triazolyl group, an
oxadiazolyl group, a thiadiazolyl group, a pyrimidinyl group,
5 and a pyrazinyl group, each may be optionally substituted
with one or more substituents,
is provided.

[0066]

According to one embodiment of the present application,
10 a compound represented by formula (I) or salts thereof
wherein

X represents a group represented by formula (II-2); and
the triazolyl group which may be optionally substituted
with one or more substituents represents 1,5-dimethyl-1,2,4-
15 triazol-3-yl group,
is provided.

[0067]

According to one embodiment of the present application,
a compound represented by formula (I) or salts thereof
20 wherein

X represents a group represented by formula (II-2); and
the oxadiazolyl group which may be optionally
substituted with one or more substituents represents 5-
methyl-1,2,4-oxadiazol-3-yl group,
25 is provided.

[0068]

According to one embodiment of the present application,

a compound represented by formula (I) or salts thereof wherein

X represents a group represented by formula (II-2); and
the thiadiazolyl group which may be optionally
5 substituted with one or more substituents represents 3-
methyl-1,2,4-thiadiazol-5-yl group,
is provided.

[0069]

According to one embodiment of the present application,
10 a compound represented by formula (I) or salts thereof
wherein

X represents a group represented by formula (II-3); and
M represents a sulfonyl group which is substituted with
one substituent selected from a C1-6 alkyl group which may be
15 optionally substituted with one or more substituents, and an
aryl group which may be optionally substituted with one or
more substituents,
is provided.

[0070]

20 According to one embodiment of the present application,
a compound represented by formula (I) or salts thereof
wherein

X represents a group represented by formula (II-3); and
M represents a sulfonyl group which is substituted with
25 one substituent selected from the group consisting of a
methyl group which may be optionally substituted with one or
more substituents, and a phenyl group which may be optionally

substituted with one or more substituents,
is provided.

[0071]

According to one embodiment of the present application,
5 a compound represented by formula (I) or salts thereof
wherein

X represents a group represented by formula (II-3); and

M represents a sulfonyl group which is substituted with
one or more substituents selected from the group consisting
10 of a methyl group and a phenyl group,
is provided.

[0072]

According to one embodiment of the present application,
as specific examples of the compound represented by formula
15 (I) or salts thereof, one or more compounds selected from the
following compounds or salts thereof are provided.

The compounds selected from the group consisting of the
following compounds:

3-[[(1R, 4R, 6R, 7S) -4-[[bis (4-methoxyphenyl) -phenyl-
20 methoxy]methyl]-6-(5-methyl-2,4-dioxo-pyrimidin-1-yl)-2-
methylsulfonyl-5-oxa-2-azabicyclo[2.2.1]heptan-7-yl]oxy-
(diisopropylamino)phosphanyl]oxypropanenitrile (Present
compound 1);

N-[1-[(1R, 3R, 4R, 7S) -1-[[bis (4-methoxyphenyl) -
25 phenylmethoxy]methyl]-7-[2-cyanoethoxy-
(diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-

yl]benzamide (Present compound 2);

[9-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-2-(2-methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate (Present compound 3);

N-[9-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-purin-6-yl]benzamide (Present compound 4);

3-[[(1R, 3R, 4R, 7S) -5-(benzenesulfonyl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 5);

N-[1-[(1R, 3R, 4R, 7S)-5-(benzenesulfonyl)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 6);

(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-methyl-3-(5-methyl-2,4-

dioxo-pyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 7);

(1R,3R,4R,7S)-3-(4-benzamide-5-methyl-2-oxo-pyrimidin-1-yl)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-methyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 8);

[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(methylcarbamoyl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-2-(2-methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate (Present compound 9);

(1R,3R,4R,7S)-3-(6-benzamidepurin-9-yl)-1-[[bis(4-methoxyphenyl)phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-methyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 10);

(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-isopropyl-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 11);

(1R,3R,4R,7S)-3-(4-benzamide-5-methyl-2-oxopyrimidin-1-yl)-1-[[bis(4-methoxyphenyl)phenyl-methoxy]methyl]-7-[2-

cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-isopropyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 12);

5 [9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(isopropylcarbamoyl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-2-(2-methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate (Present compound 13);

10 (1R,3R,4R,7S)-3-(6-benzamidepurin-9-yl)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-isopropyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 14);

15 (1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N,N-dimethyl-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 15);

20 (1R,3R,4R,7S)-3-(4-benzamide-5-methyl-2-oxo-pyrimidin-1-yl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N,N-dimethyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 16);

25 3-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-

methoxy)methyl]-3-(5-methyl-2,4-dioxo-pyrimidin-1-yl)-5-pyrimidin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 17);

5 1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-pyrimidin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 18);

10 N'-[9-(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-pyrimidin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-N,N-dimethyl-6-formamidine (Present compound 19);

15 N'-[9-(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-pyrimidin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]purin-6-yl]benzamide (Present compound 20);

20 3-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-3-(5-methyl-2,4-dioxopyrimidin-1-yl)-5-pyrimidin-4-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 21);

25 N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-

methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-pyrimidin-4-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 22);

5 3-[[(1R, 3R, 4R, 7S) -1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-3-(5-methyl-2,4-dioxo-pyrimidin-1-yl)-5-[4-(trifluoromethyl)pyrimidin-2-yl]-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present
10 compound 23);

 N-[1-[(1R, 3R, 4R, 7S) -1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-[4-(trifluoromethyl)pyrimidin-2-yl]-2-oxa-5-
15 azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 24);

 3-[[(1R, 3R, 4R, 7S) -1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-5-(5-chloropyrimidin-2-yl)-3-(5-methyl-2,4-dioxo-pyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-
20 yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 25);

 N-[1-[(1R, 3R, 4R, 7S) -1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-5-(5-chloropyrimidin-2-yl)-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-2-oxa-5-
25 azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-4-

yl]benzamide (Present compound 26);

3-[[(1R, 3R, 4R, 7S) -1-[[bis (4-methoxyphenyl) -phenyl-
methoxy]methyl]-5-[6-(dimethylamino)pyrimidin-4-yl]-3-(5-
methyl-2,4-dioxo-pyrimidin-1-yl)-2-oxa-5-
5 azabicyclo[2.2.1]heptan-7-yl]oxy-
(diisopropylamine)phosphanyl]oxypropanenitrile (Present
compound 27);

N-[1-[(1R, 3R, 4R, 7S) -1-[[bis (4-methoxyphenyl) -phenyl-
methoxy]methyl]-7-[2-
10 cyanoethoxy(diisopropylamino)phosphanyl]oxy-5-[6-
dimethylamino)pyrimidin-4-yl]-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-
yl]benzamide (Present compound 28);

3-[[(1R, 3R, 4R, 7S) -1-[[bis (4-methoxyphenyl) -phenyl-
15 methoxy]methyl]-5-[2-(dimethylamino)pyrimidin-4-yl]-3-(5-
methyl-2,4-dioxo-pyrimidin-1-yl)-2-oxa-5-
azabicyclo[2.2.1]heptan-7-yl]oxy-
(diisopropylamino)oxypropanenitrile (Present compound 29);

N-[1-[(1R, 3R, 4R, 7S) -1-[[bis (4-methoxyphenyl) -phenyl-
20 methoxy]methyl]-7-[2-cyanoethoxy-
(diisopropylamino)phosphanyl]oxy-5-[2-
(dimethylamino)pyrimidin-4-yl]-7-hydroxy-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-
yl]benzamide (Present compound 30);

25 3-[1-[(1R, 3R, 4R, 7S) -1-[[bis (4-methoxyphenyl) -phenyl-

methoxy)methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-pyrazin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 31);

5 N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-pyrazine-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 32);

10 3-[[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present
15 compound 33);

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound
20 34);

N'-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-

1H-purin-2-yl]-N,N-dimethylformamidine (Present compound 35);

N-[9-(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-
5 (diisopropylamino)phosphanyl]oxy-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]purin-6-yl]benzamide (Present compound 36);

3-[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-(5-
10 methyl-1,2,4-oxadiazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 37);

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-
15 (diisopropylamino)phosphanyl]oxy-5-(5-methyl-1,2,4-oxadiazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 38);

20 3-[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-(3-methyl-1,2,4-thiadiazol-5-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present
25 compound 39);

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(3-methyl-1,2,4-thiadiazol-5-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 40);

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]dihydropyrimidin-2-one (Present compound 41);

N'-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy(diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-N,N-dimethylformamidine (Present compound 42);

N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-2-methylpropane amide (Present compound 43); and

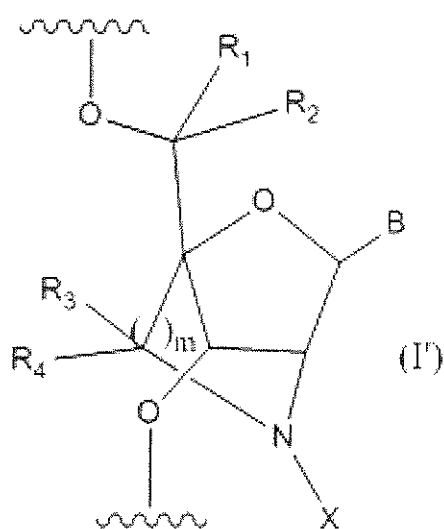
N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-2-methylpropane amide (Present compound 44).

[0073]

Oligonucleotide

Compound represented by general formula (I')

According to one embodiment of the present application,
 5 an oligonucleotide compound comprising one or more
 nucleosides represented by general formula (I'):



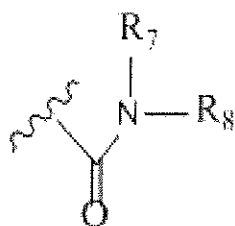
[wherein,

B represents a base moiety of nucleic acid wherein the
 10 base moiety may be optionally substituted with one or more
 substituents;

R₁, R₂, R₃ and R₄ represent independently of each other
 a hydrogen atom, or a C1-6 alkyl group which may be
 optionally substituted with one or more substituents;

15 m is an integer of 1 or 2;

X represents a group represented by the following
 formula (II'-1), (II'-2), or (II'-3):



(II'-1)



(II'-2)



(II'-3)

;

the symbols:



which is described in the formula (II'-1), (II'-2) or (II'-
 5 3) represents a binding point to 2'-amino group;

A represents an aromatic group containing one or more heteroatoms selected from the group consisting of nitrogen atom, oxygen atom and sulfur atom, which may be optionally substituted with one or more substituents;

10 M represents a sulfonyl group which is substituted with one or more substituents selected from the group consisting of a C₁₋₆ alkyl group which may be optionally substituted with one or more substituents, and an aromatic group which may be optionally substituted with one or more substituents.]

15 or salts thereof,

is provided.

[0074]

According to one embodiment of the present application, an oligonucleotide compound represented by formula (I') or
 20 salts thereof wherein

in the formula (I'), the ring B represents an adeninyl group which may be optionally substituted with one or more substituents, a guaninyl group which may be optionally substituted with one or more substituents, a cytosinyl group which may be optionally substituted with one or more substituents, a 5-methylcytosinyl group which may be optionally substituted with one or more substituents or an uracil group which may be optionally substituted with one or more substituents,

is provided.

[0075]

According to one embodiment of the present application, wherein

in the formula (I'), the ring B represents a 5-methylcytosinyl group which may be optionally substituted with one or more substituents or a 5- uracil group which may have one or more protecting groups,

is provided.

[0076]

According to one embodiment of the present application, an oligonucleotide compound represented by formula (I') or salts thereof wherein

in the formula (I'), R_1 , R_2 , R_3 and R_4 represent independently of each other a hydrogen atom, and m is an integer of 1,

is provided.

[0077]

According to one embodiment of the present application,
an oligonucleotide compound represented by formula (I') or
5 salts thereof wherein

in the formula (I'), one or more phosphate bonds between
nucleotides represent a phosphorothioate bond, and is/are
sulfurized,

is provided.

10 [0078]

The compound wherein in the formula (I'), X represents
a group represented by formula (II'-1) (hereinafter,
referred to as "Urea type compound" may be included.

[0079]

15 According to one embodiment of the present application,
an oligonucleotide compound represented by formula (I') or
salts thereof wherein

X represents a group represented by formula (II'-1);
and

20 R₇ and R₈ represent independently of each other a
hydrogen atom, a C₁₋₆ alkyl group which may be optionally
substituted with one or more substituents, or an aromatic
group which may be optionally substituted with one or more
substituents,

25 is provided.

[0080]

According to one embodiment of the present application,
an oligonucleotide compound represented by formula (I') or
salts thereof wherein

5 X represents a group represented by formula (II'-1);
and

 R₇ and R₈ represent independently of each other a
hydrogen atom, a C1-6 alkyl group which may be optionally
substituted with one or more substituents, or an aromatic
10 group which may be optionally substituted with one or more
substituents,
is provided.

[0081]

 According to one embodiment of the present application,
15 an oligonucleotide compound represented by formula (I') or
salts thereof wherein

 X represents a group represented by formula (II'-1);
and

 R₇ and R₈ represent independently of each other a
20 hydrogen atom, a C1-3 alkyl group which may be optionally
substituted with one or more substituents, or an aromatic
group which may be optionally substituted with one or more
substituents,
is provided.

25 [0082]

According to one embodiment of the present application,
an oligonucleotide compound represented by formula (I') or
salts thereof wherein

X represents a group represented by formula (II'-1);
5 and

R₇ and R₈ represent independently of each other a
hydrogen atom, a methyl group, an isopropyl group, or a
phenyl group,
is provided.

10 [0083]

According to one embodiment of the present application,
an oligonucleotide compound represented by formula (I') or
salts thereof wherein

X represents a group represented by formula (II'-1);
15 and

a combination of R₇ and R₈ represents a combination
wherein both of R₇ and R₈ represent a hydrogen atom; a
combination wherein both of R₇ and R₈ represent a methyl
group; a combination wherein R₇ represents a hydrogen atom
20 and R₈ represents a methyl group; a combination wherein R₇
represents a hydrogen atom and R₈ represents an isopropyl
group; or a combination wherein R₇ represents a hydrogen atom
and R₈ represents a phenyl group,
is provided.

25 [0084]

According to one embodiment of the present application,
an oligonucleotide compound represented by formula (I') or
salts thereof wherein

X represents a group represented by formula (II'-1);
5 and

a combination of R₇ and R₈ represents a combination
wherein both of R₇ and R₈ represent a hydrogen atom; a
combination wherein both of R₇ and R₈ represent a methyl
group; a combination wherein one of R₇ and R₈ represents a
10 hydrogen atom and the other thereof represents an isopropyl
group; a combination wherein one of R₇ and R₈ represents a
hydrogen atom and the other thereof represents an isopropyl
group; or a combination wherein one of R₇ and R₈ represents
a hydrogen atom and the other thereof represents a phenyl
15 group,
is provided.

[0085]

According to one embodiment of the present application,
an oligonucleotide compound represented by formula (I') or
20 salts thereof wherein

X represents a group represented by formula (II'-1);
and

one of R₇ and R₈ represents a hydrogen atom and the
other thereof represents a methyl group,
25 is provided.

[0086]

According to one embodiment of the present application, an oligonucleotide compound represented by formula (I') or salts thereof wherein

5 X represents a group represented by formula (II'-1);
and

one of R₇ and R₈ represents a hydrogen atom and the other thereof represents an isopropyl group,
is provided.

10 [0087]

Also a compound wherein in the formula (I'), X represents a group represented by formula (II'-2) (hereinafter, referred to as "heteroaryl type compound") may be included.

15 [0088]

According to one embodiment of the present application, an oligonucleotide compound represented by formula (I') or salts thereof wherein

X represents a group represented by formula (II'-2);
20 and

A represents a five or six membered heteroaryl group containing one or more heteroatoms selected from the group consisting of nitrogen atom, oxygen atom and sulfur atom, which may be optionally substituted with one or more
25 substituents,

is provided.

[0089]

According to one embodiment of the present application,
an oligonucleotide compound represented by formula (I') or
5 salts thereof wherein

X represents a group represented by formula (II'-2);
and

A represents a five or six membered heteroaryl group
containing two or three heteroatoms selected from nitrogen
10 atom, oxygen atom, and sulfur atom, provided that at least
two nitrogen atoms are included, which may be optionally
substituted with one or more substituents,
is provided.

[0090]

15 According to one embodiment of the present application,
an oligonucleotide compound represented by formula (I') or
salts thereof wherein

X represents a group represented by formula (II'-2);
and

20 A represents a five or six membered heteroaryl group
containing one or more heteroatoms selected from nitrogen
atom, oxygen atom, and sulfur atom, provided that at least
two nitrogen atoms are included, which may be optionally
substituted with one or more substituents, and the
25 substituent is selected independently of each other from a

C1-3 alkyl group which may be optionally substituted with one or more halogen atoms, a halogen atom, and an amino group which may be optionally substituted with one or more C1-3 alkyl groups,

5 is provided.

[0091]

According to one embodiment of the present application, an oligonucleotide compound represented by formula (I') or salts thereof wherein

10 X represents a group represented by formula (II'-2);
and

A represents a five or six membered heteroaryl group containing one or more heteroatoms selected from nitrogen atom, oxygen atom, and sulfur atom, provided that at least
15 two nitrogen atoms are included, which may be optionally substituted with one or more substituents, and the substituent is selected independently of each other from a C1-3 alkyl group which may be optionally substituted with one or more halogen atoms, a halogen atom, and an amino group
20 which may be optionally substituted with one or more C1-3 alkyl groups,
is provided.

[0092]

According to one embodiment of the present application,
25 an oligonucleotide compound represented by formula (I') or

salts thereof wherein

X represents a group represented by formula (II'-2);
and

the five or six membered heteroaryl group as A group is
5 selected from the group consisting of a triazolyl group, an
oxadiazolyl group, a thiadiazolyl group, a pyrimidinyl group,
and a pyrazinyl group, each may be optionally substituted
with one or more substituents,
is provided.

10 [0093]

According to one embodiment of the present application,
an oligonucleotide compound represented by formula (I') or
salts thereof wherein

X represents a group represented by formula (II'-2);
15 and

the triazole group which may be optionally substituted
with one or more substituents represents 1,5-dimethyl-1,2,4-
triazol-3-yl group,
is provided.

20 [0094]

According to one embodiment of the present application,
an oligonucleotide compound represented by formula (I') or
salts thereof wherein

X represents a group represented by formula (II'-2);
25 and

the oxadiazolyl group which may be optionally substituted with one or more substituents represents 5-methyl-1,2,4-oxadiazol-3-yl group, is provided.

5 [0095]

According to one embodiment of the present application, an oligonucleotide compound represented by formula (I') or salts thereof wherein

X represents a group represented by formula (II'-2);
10 and

the thiadiazolyl group which may be optionally substituted with one or more substituents represents 3-methyl-1,2,4-thiadiazol-5-yl group, is provided.

15 [0096]

According to one embodiment of the present application, an oligonucleotide compound represented by formula (I') or salts thereof wherein

X represents a group represented by formula (II'-3);
20 and

M represents a sulfonyl group which is substituted with one substituent selected from the group consisting of a C1-6 alkyl group which may be optionally substituted with one or more substituents and an aryl group which may be
25 optionally substituted with one or more substituents,

is provided.

[0097]

According to one embodiment of the present application,
an oligonucleotide compound represented by formula (I') or
5 salts thereof wherein

X represents a group represented by formula (II'-3);
and

M represents a sulfonyl group which is substituted with
one substituent selected from the group consisting of a
10 methyl group which may be optionally substituted with one or
more substituents, and a phenyl group which may be optionally
substituted with one or more substituents,
is provided.

[0098]

15 According to one embodiment of the present application,
an oligonucleotide compound represented by formula (I') or
salts thereof wherein

X represents a group represented by formula (II'-3);
and

20 M represents a sulfonyl group which is substituted with
one or more substituents selected from the group consisting
of a methyl group and a phenyl group,
is provided.

[0099]

25 According to one embodiment of the present application,

an oligonucleotide compound represented by formula (I') or salts thereof wherein

X represents a group represented by formula (II'-3);
and

5 M represents a sulfonyl group which is substituted with one or more substituents selected from the group consisting of a methyl group and a phenyl group,
is provided.

[0100]

10 According to one embodiment of the present application, the oligonucleotide or salts thereof is made by binding two or more the nucleotides, and preferably contains 2 to 100 nucleotides, more preferably 5 to 50 nucleotides, and particularly preferably those in which 10 to 30 artificial
15 nucleotides are bonded or are combined together with these complementary strands forms a double strand.

[0101]

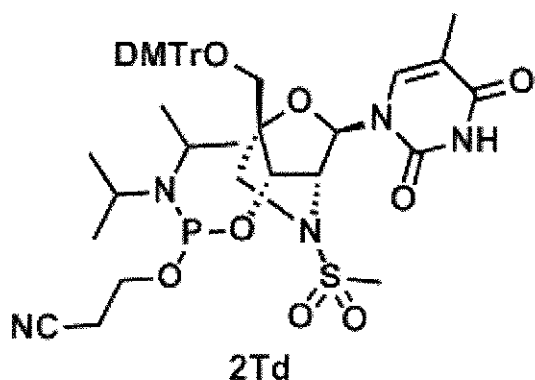
The term of "salt" in the expression of "salts of compound" used herein alkali metal salts such as sodium salt,
20 potassium salt, lithium salt; alkaline earth metal salts such as calcium salt, and magnesium salts; metal salts such as aluminium salt, iron salt, zinc salt, copper salt, nickel salt, and cobalt salt; inorganic salts such as ammonium salt; amine salts, for example, organic salts such as t-octylamine
25 salt, dibenzylamine salt, morpholine salt, glucosamine salt,

phenylglycine alkyl ester salt, ethylenediamine salt, N-methylglucamine salt, guanidine salt, diethylamine salt, triethylamine salt, hexylamine salt, dicyclohexylamine salt, N,N'-dibenzylethylenediamine salt, chlorprocaine salt, 5 procaine salt, diethanolamine salt, N-benzyl phenethylamine salt, piperazine salt, tetramethylammonium salt, and tris(hydroxymethyl)aminomethane salt; inorganic acid salts, for example, hydrohalic acids (such as hydrofluoric acid, hydrochloric acid, hydrobromic acid, and hydroiodic acid), 10 nitrate salt, perchloric acid salt, sulfate salt, and phosphate salt; salts of alkanesulfonic acid having one to six carbon atoms, such as methanesulfonate salt, trifluoromethanesulfonate salt, ethanesulfonate salt; arylsulfonic acid salts such as benzene sulfonate salt, and p-toluene sulfonate salt; organic acid salts such as acetate salt, 15 malate salt, fumarate salt, succinate salt, citrate salt, tartrate salt, oxalate salt, and maleate salt; and amino acid salts such as glycine salt, lysine salt, arginine salt, ornithine salt, glutamic acid salt, and aspartic acid salt, 20 which should not be limited thereto. The salts are encompassed by pharmaceutically acceptable salts.

[0102]

Each compound represented by general formula (I) described herein encompasses enantiomer form, diastereomer 25 form, or mixtures thereof. For example, a stereo

configuration of sugar moiety in the structure of each compound encompasses α form and β form, however, β form is preferred. When the above-mentioned compounds represented by the above formula (I) is obtained in the form of diastereomer or enantiomer, they can be separated by well-known methods in organic synthesis (for example, sugar synthesis), such as chromatography method or fractional crystallization method. For example, the compound 2Td represented by the following formula:



has β form as a stereo configuration of sugar moiety.

[0103]

Each compound represented by general formula I described herein compasses the compounds labeled with isotopes (such as ^3H , ^{13}C , ^{14}C , ^{15}N , ^{18}F , ^{32}P , ^{35}S or ^{125}I) and deuterated transformers.

[0104]

The oligonucleotide or analogues thereof described herein may be mixed with auxiliary agents that are usually used in the formulation of pharmaceuticals, such as

excipients, binders, preservatives, antioxidants, disintegrants, lubricants, and flavors, to make parenterally-administered formulation or liposome formulations. Also, the oligonucleotide or analogues thereof is mixed with pharmaceutical carriers that are usually used in the pharmaceutical arts to formulate into topical formulations such as solutions, creams and ointments. [0105]

(Preparation method)

Hereinafter, the preparation method of the compound of the present invention is described.

Optically active substances having absolute configuration in each compound represented by the below-mentioned starting materials and intermediate compounds may be prepared by using as starting material optically active substance(s) or separating the isomers that created in the intermediate stages for synthesis. Also, the below-mentioned replacement reaction of nucleic acid base in which nucleic acid base is replaced may proceed a β -selective transglycosylation effectively, and the obtained compound represented by general formula I may be obtained as a desired β former selectively.

[0106]

The compound of the present invention or pharmaceutically acceptable salts thereof may be prepared

according to the following method, but which should not be limited thereto.

For starting materials, unless specific processes therefor are stated, they may be used as commercially available ones, or can be prepared according to the publicly known method or a similar method thereto.

[0107]

The abbreviated symbols as used herein mean the following meanings respectively.

- 10 THF: tetrahydrofuran
 DMSO: dimethyl sulfoxide
 MsCl: methanesulfonyl chloride
 DMAP: N,N-dimethyl-4-aminopyridine
 $i\text{Pr}_2\text{NEt}$, DIPEA: N,N-diisopropylethylamine
 15 BSA: N,O-bis(trimethylsilyl)acetamide
 TMSOTf: trimethylsilyl trifluoro-methanesulfonate
 TBSOTf: t-butyldimethylsilyl trifluoro
 methanesulfonate
 Bn: benzyl
 20 TMS: trimethylsilyl
 TBDPS: tert-butyldiphenylsilyl
 Ac: acetyl
 MOE: methoxyethyl
 CH_2Cl_2 : dichloromethane
 25 MeCN: acetonitrile

DMTr: 4,4'-dimethoxytrityl

[0108]

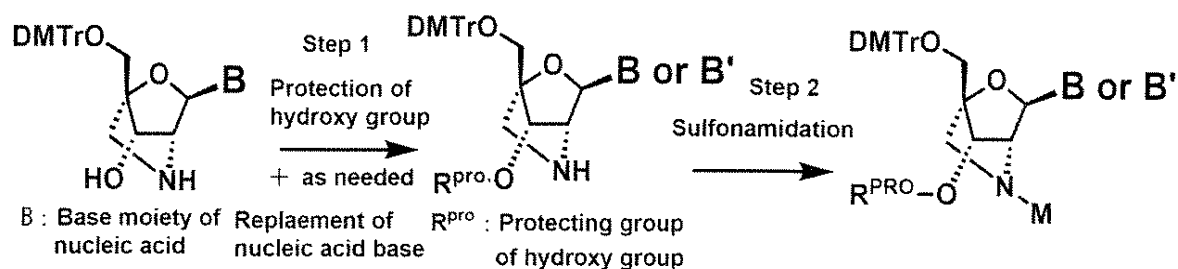
First, the summary of a preparation method for the compound of the present invention (monomer) is described.

5 [0109]

(Sulfonamide type compound)

The preparation method of the compound of the present invention represented by general formula I wherein X represents a group represented by formula (II-3), so-called
10 sulfonamide type compound is described. The typical example of the preparation scheme is described, and is not limited thereto.

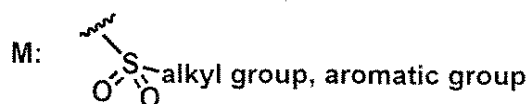
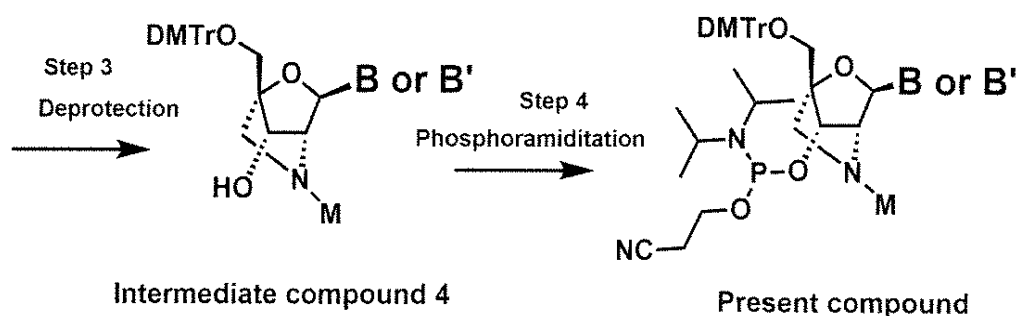
Reaction scheme 1



Starting compound 1

Intermediate compound 2

Intermediate compound 3



[0110]

(Step 1)

A protecting group for hydroxy group is introduced as
 5 R_s group in the general formula (I) by using a starting
 material 1 (for example, which can be prepared according to
 the process described in WO 2017/047816 A1) which is a sugar
 compound, according to a publicly known method to prepare an
 intermediate compound 2. Alternatively, a basic moiety of
 10 nucleic acid in the starting material 1 (abbreviated as "B")
 is subjected to a nucleic acid base exchange reaction with
 other basic moiety of nucleic acid which may be optionally

protected (abbreviated as "B'") and a protecting group for hydroxy group is also introduced at the same time as the nucleic acid base exchange reaction to prepare an intermediate compound 2.

5 This introduction of the protecting group for hydroxy group can be conducted under a reaction condition which is known in organic synthetic chemistry (for example, nucleic acid synthesis) depending on a kind of protecting group. For example, when a protecting group for hydroxy group is a
10 silyl type protecting group (such as TMS), a silyl agent (such as BSA, hexamethyl disilazane, TMS chloride) can be used. In the reaction, Lewis acid (such as TMSOTf, TBSOTf, and tin chloride) may be added.

 The silylating agent may be used in about 1 to about 20
15 molar equivalent(s), and the Lewis acid may be used in a catalytic amount (about 0.05 molar equivalents) to about 2 molar equivalents as opposed to 1 mole of the reaction substrates.

 The solvents may be any ones unless they affect the
20 reaction adversely, and the reaction may be carried out in appropriate solvents (for example, ethers such as THF; halogenated hydrocarbons such as dichloromethane, dichloroethane, and chloroform; hydrocarbons such as benzene and toluene; acetonitrile; water; or mixed solvents thereof).
25 The reaction may proceed preferably at 0 °C to high

temperature, and particularly preferably at room temperature to about 60 °C.

[0111]

Also, according to a nucleic acid exchange reaction
5 (transglycosylation reaction), a pyrimidine base moiety of nucleic acid base such as thymine group (T) and uracil group (U) can be replaced with other nucleic acid base moiety (for example, adenine (A), guanine (G), uracil (U), thymine (T), cytosine (C), or 5-methylcytosine (^{Me}C) to obtain an
10 intermediate compound 2 wherein the nucleic acid base moiety is replaced. The base exchange reaction can be carried out, for example, in the presence of Lewis acid (such as TMSOTf and TBSOTf), and the reaction can be facilitated by reacting with the silylating agent.

15 [0112]

(Step 2)

Next, using the intermediate compound 2 as a starting material, an alkyl sulfonyl halide or an aromatic sulfonyl halide can be reacted with the intermediate compound 2
20 according to a publicly known method to undergo a sulfonamidation on an amino nitrogen atom, resulting in preparing an intermediate compound 3.

Examples of the sulfone amidation reagent used in a sulfone amidation reaction include an alkyl sulfonyl halide
25 reagent (such as methane sulfonyl chloride (MsCl)), an alkyl

sulfonic anhydride reagent (such as methane sulfonic anhydride), an aromatic sulfonyl halide reagent (such as benzene sulfonyl chloride), or an aromatic sulfonic anhydride (such as benzene sulfonic anhydride), and these
5 reagents can be used in the presence of a base (such as triethyl amine, and DIPEA).

The sulfonamidation reagent may be used in approximately molar equivalents to somewhat excess molar equivalents (for example, about 1.0 molar equivalent) as
10 opposed to 1 mole of the reaction substrates.

The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in appropriate solvents (for example, ethers such as THF; halogenated hydrocarbons such as dichloromethane,
15 dichloroethane, and chloroform; hydrocarbons such as benzene and toluene; acetonitrile; water; or mixed solvents thereof). The reaction can proceed preferably at -25 °C to room temperature, and particularly preferably at 0 °C to room temperature.

20 [0113]

(Step 3)

Next, the protecting groups for hydroxy group which is introduced in the step 1 is deprotected by using the intermediate compound 3 as a starting material according to
25 a publicly known method to prepare an intermediate compound

4.

The present deprotection reaction can be carried out in appropriate reaction condition (for example, reagents to be used) depending on the kinds of the protecting group. For example, when a protecting group for hydroxy group (abbreviated as "R^{PRO}") is a silyl type protecting group (such as TMS), the reaction can be carried out by hydrolysis under acidic condition (such as acetic acid-THF-water) or by treating with a fluoride ion-donating reagent (such as TBAF).

The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in an appropriate solvents (for example, ethers such as THF, halogenated hydrocarbon atoms such as dichloromethane, water, or mixed solvents thereof).

The reaction may proceed preferably at -25 °C to 100 °C, particularly preferably at 0 °C to 50 °C as a reaction temperature.

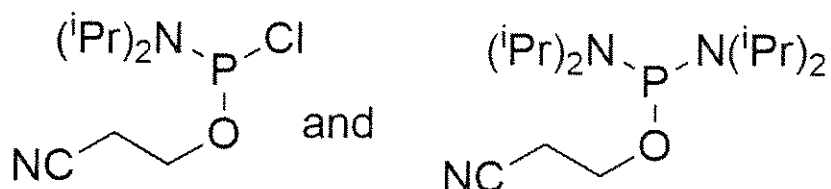
[0114]

(Step 4)

Further, the intermediate compound 4 as a starting material is subjected to a phosphoramidation reaction under reaction conditions that have been generally known in organic synthesis (in particular, nucleic acid synthesis) (for example, in terms of reagent(s)) to prepare a desired compound of the present invention as a sulfone amide type

compound.

Examples of the phosphoramiditation reagent include agents by the following formula:



5 , which should not be limited thereto.

The phosphoramiditation reaction may be conducted in the presence of appropriate bases (such as DIPEA) or appropriate acids (such as diisopropyl ammonium tetrazolide, and 4,5-dicyanoimidazole).

10 The phosphoramiditation reagent may be used in 1 to about 10 molar equivalent(s), preferably 1 to about 5 molar equivalent(s) (such as 3 molar equivalents) as opposed to 1 mole of the reaction substrates.

15 The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in appropriate solvents (for example, ethers such as THF; halogenated hydrocarbons such as dichloromethane, dichloroethane, and chloroform; acetonitrile; water; or mixed solvents thereof). The reaction can proceed preferably
20 at -25 °C to 60 °C, and particularly preferably at 0 °C to room temperature.

[0115]

Here in any stages from the timing of before the above step 1 to the timing of after the above step 4, a substituent on the ring of the nucleic acid base may be appropriately modified (for example, a benzoylation reaction or a
5 acetylation reaction on an amino substituent on a cytosine ring).

[0116]

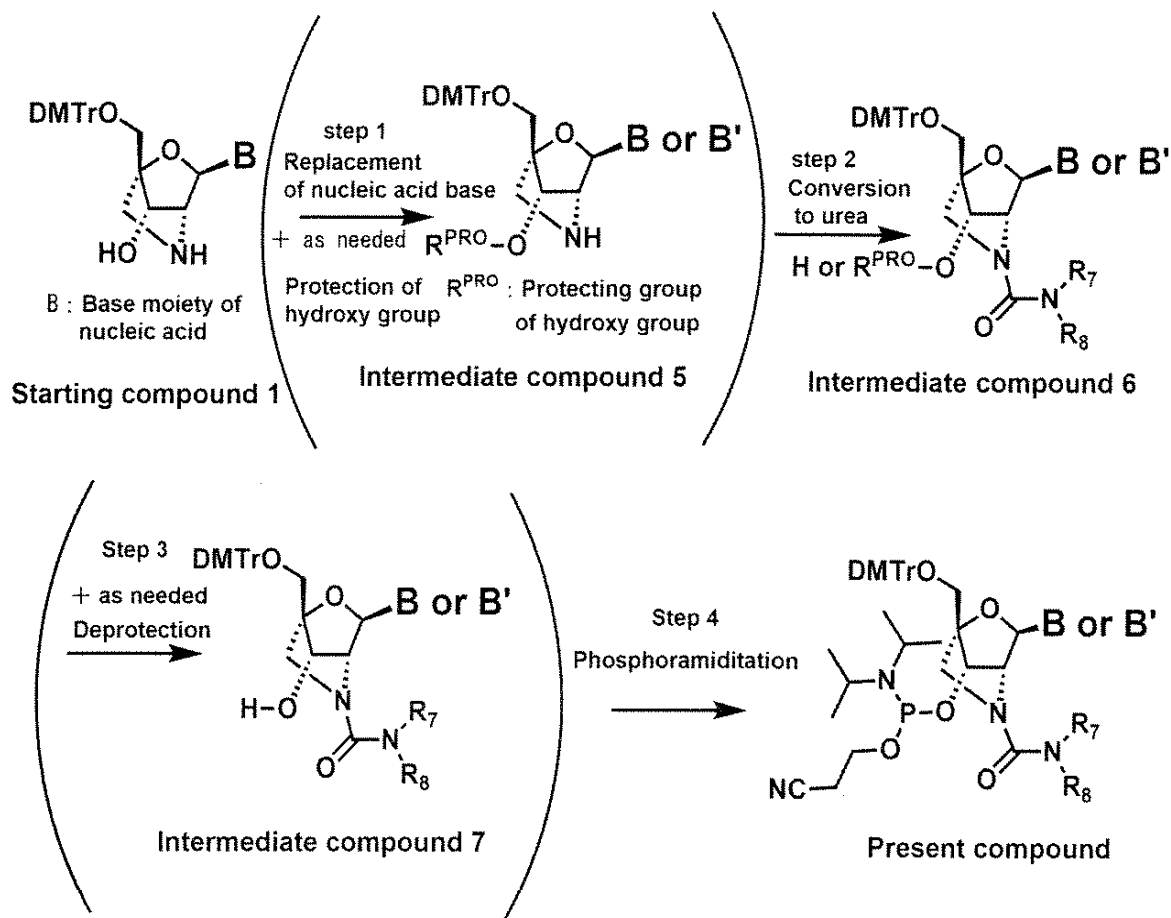
Also, among the above step (1) to step (4), a plural steps (for example, step (1) and step (2), and step (2) and
10 step (3)) may be carried out successively.

[0117]

(Urea type compound)

A method for preparing a compound of the present invention represented by general formula I wherein X
15 represents a group represented by formula (II-1), which is so-called urea type compound, is described. Typical examples of the preparation scheme is indicated in the following reaction scheme 2, which should not be limited thereto.

Reaction scheme 2



[0118]

(Step 1)

Before undergoing a reaction for converting into an urea according to a publicly known method, similarly to the above reaction scheme 1 using the above starting compound 1 as a starting material, a basic moiety of nucleic acid in the starting compound 1 (abbreviated as "B") is subjected to a nucleic acid base exchange reaction with other basic moiety of nucleic acid which may be optionally protected (abbreviated as "B'"), and thereafter, the reaction for converting into an urea may be carried out. Also, similarly

to the above reaction scheme, after the nucleic acid base exchange reaction is carried out, a protecting group for hydroxy group may be introduced appropriately.

In a nucleic acid exchange reaction (transglycosylation reaction), according to a publicly known method, a pyrimidine base moiety of nucleic acid base such as thymine group (T) and uracil group (U) can be replaced with other nucleic acid base moiety (for example, adenine (A), guanine (G), uracil (U), thymine (T), cytosine (C), or 5-methylcytosine (^{Me}C) to obtain an intermediate compound 5 wherein the nucleic acid base moiety is replaced. The base exchange reaction can be carried out, for example, in the presence of Lewis acid (such as TMSOTf and TBSOTf), and the reaction can be facilitated by reacting with the silylating agent (such as BSA).

This introduction of the protecting group for hydroxy group can be conducted under a reaction condition which is known in organic synthetic chemistry (for example, nucleic acid synthesis) depending on a kind of protecting group. For example, when a protecting group for hydroxy group is a silyl type protecting group (such as TMS), a silyl agent (such as BSA, hexamethyl disilazane, TMS chloride) can be used. In the reaction, Lewis acid (such as TMSOTf, TBSOTf, and tin chloride) may be added.

The silylating agent may be used in about 1 to about 20 molar equivalent(s), and the Lewis acid may be used in a

catalytic amount (about 0.05 molar equivalents) to about 2 molar equivalents as opposed to 1 mole of the reaction substrates.

5 The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in appropriate solvents (for example, ethers such as THF; halogenated hydrocarbons such as dichloromethane, dichloroethane, and chloroform; hydrocarbons such as benzene and toluene; acetonitrile; water; or mixed solvents thereof).

10 The reaction may proceed preferably at 0 °C to high temperature, and particularly preferably at room temperature to about 60 °C.

[0119]

(Step 2)

15 Next, using the starting compound 1 or the intermediate compound 5 as a starting material, according to a publicly known method, the reagent for converting into urea is reacted with the starting compound 1 or the intermediate compound 5 to undergo a conversion reaction to a urea on the amino

20 nitrogen atom, results in preparing an intermediate compound 6.

Examples of the reagent for converting into urea used in the reaction for converting into urea include isocyanate, alkyl carbamoyl halide reagent (such as N-methylcarbamoyl

25 chloride) or N,N-dialkyl carbamoyl halide reagent (such as

N,N-dialkyl carbamoyl chloride), and these reagents can be used in the presence or absence of a base (such as DIPEA).

The reagent for converting into urea may be used in approximately molar equivalents to somewhat excess molar equivalents (for example, about 1.0 molar equivalent) as
5 opposed to 1 mole of the reaction substrates.

The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in appropriate solvents (for example, ethers such as THF;
10 halogenated hydrocarbons such as dichloromethane, dichloroethane, and chloroform; or mixed solvents thereof). The reaction may proceed preferably at -25 °C to room temperature, and particularly preferably at 0 °C to room temperature.

15 [0120]

(Step 3)

Next, as needed, when a protecting group for hydroxy group is introduced in the above-mentioned step 1, using the intermediate compound 6 as a starting material, according to
20 a publicly known method, the introduced protecting group for hydroxy group is deprotected to prepare an intermediate compound 7.

The present deprotection reaction can be carried out in appropriate reaction condition (for example, reagents to be
25 used) depending on the kinds of the protecting group. For

example, when a protecting group for hydroxy group (abbreviated as "R^{PRO}") is a silyl type protecting group (such as TMS), the reaction can be carried out by hydrolysis under acidic condition (such as acetic acid-THF-water) or by
5 treating with a fluoride ion-donating reagent (such as TBAF).

The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in an appropriate solvents (for example, ethers such as THF, halogenated hydrocarbon atoms such as dichloromethane, water,
10 or mixed solvents thereof).

The reaction may proceed preferably at -25 °C to 100°C, particularly preferably at 0 °C to 50 °C as a reaction temperature.

[0121]

15 (Step 4)

Further, similarly to the step 4 of the above-mentioned reaction scheme 1 ,the intermediate compound 6 or the intermediate compound 7 as a starting material is subjected to a phosphoramiditation reaction according to a publicly
20 known method to prepare a desired urea type (for example, methyl urea type, and isopropyl urea type) of the present compound.

Examples of the phosphoramiditation reagent include 2-cyanoethyl-N,N-diisopropylchlorophosphoramidite, and the
25 reagent can be used in the presence of a base (such as

triethylamine, DIPEA).

The phosphoramiditation reagent may be used in 1 to about 10 molar equivalent(s), preferably 1 to about 5 molar equivalent(s) (such as 3 molar equivalents) as opposed to 1
5 mole of the reaction substrates.

The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in appropriate solvents (for example, ethers such as THF; halogenated hydrocarbons such as dichloromethane,
10 dichloroethane, and chloroform; acetonitrile; water; or mixed solvents thereof). The reaction can proceed preferably at -25 °C to 60 °C, and particularly preferably at 0 °C to room temperature.

[0122]

15 Also, among the above step (1) to step (4), a plural steps (for example, step (1) and step (2), step (2) and step (3), and step (1) to step (3)) may be carried out successively.

[0123]

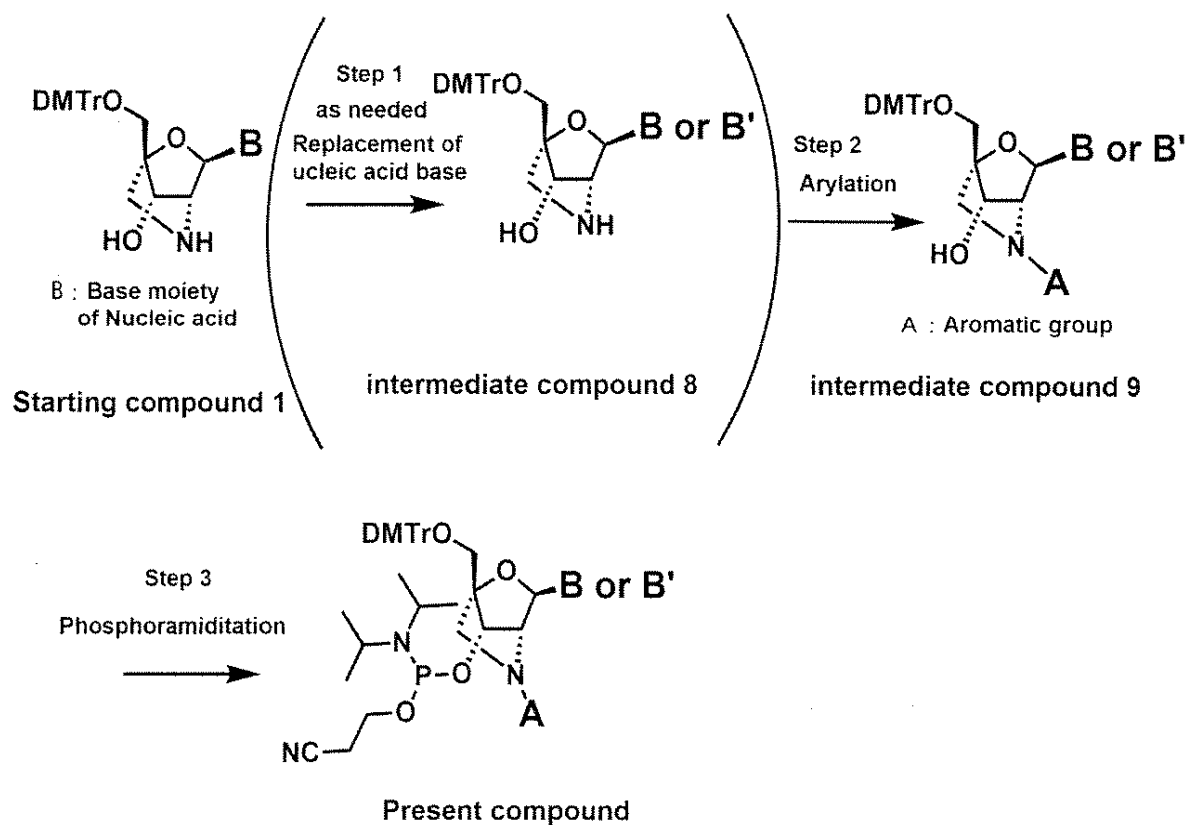
20 Here in any stages from the timing of before the above step 1 to the timing of after the above step 4, a substituent on the ring of the nucleic acid base may be appropriately modified (for example, a benzylation reaction or a acetylation reaction on an amino substituent on a cytosine
25 ring).

[0124]

(Aryl type compound)

A preparation method of the compound represented by formula I wherein X represents formula (II-2), so-called an aryl type of the present compound is described. Examples of the preparation scheme is shown in the following reaction scheme 3.

Reaction scheme 3



10 [0125]

(Step 1)

Before undergoing an arylation reaction according to a publicly known method, similarly to the above reaction scheme

1 using the above starting compound 1 as a starting material,
a basic moiety of nucleic acid in the starting compound 1
(abbreviated as "B") is subjected to a nucleic acid base
exchange reaction with other basic moiety of nucleic acid
5 which may be optionally protected (abbreviated as "B'"), and
thereafter, the arylation reaction may be carried out.

In a nucleic acid exchange reaction (transglycosylation
reaction), according to a publicly known method, a pyrimidine
base moiety of nucleic acid base such as thymine group (T)
10 and uracil group (U) can be replaced with other nucleic acid
base moiety (for example, adenine (A), guanine (G), uracil
(U), thymine (T), cytosine (C), or 5-methylcytosine (^{Me}C) to
obtain an intermediate compound 8 wherein the nucleic acid
base moiety is replaced. The base exchange reaction can be
15 carried out, for example, in the presence of Lewis acid (such
as TMSOTf and TBSOTf), and the reaction can be facilitated
by reacting with the silylating agent (such as BSA).

This introduction of the protecting group for hydroxy
group can be conducted under a reaction condition which is
20 known in organic synthetic chemistry (for example, nucleic
acid synthesis) depending on a kind of protecting group.
For example, when a protecting group for hydroxy group is a
silyl type protecting group (such as TMS), a silyl agent
(such as BSA, hexamethyl disilazane, TMS chloride) can be
25 used. In the reaction, Lewis acid (such as TMSOTf, TBSOTf,

and tin chloride) may be added.

The silylating agent may be used in about 1 to about 20 molar equivalent(s), and the Lewis acid may be used in a catalytic amount (about 0.05 molar equivalents) to about 2
5 molar equivalents as opposed to 1 mole of the reaction substrates.

The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in appropriate solvents (for example, ethers such as THF;
10 halogenated hydrocarbons such as dichloromethane, dichloroethane, and chloroform; hydrocarbons such as benzene and toluene; acetonitrile; water; or mixed solvents thereof). The reaction may proceed preferably at 0 °C to high temperature, and particularly preferably at room temperature
15 to about 60 °C.

[0126]

(Step 2)

Next, using the starting compound 1 or the intermediate compound 8 as a starting material, according to a publicly
20 known method, the arylation reagent is reacted with the starting compound 1 or the intermediate compound 8 to undergo an aromatization reaction on the amino nitrogen atom, results in preparing an intermediate compound 9.

Examples of the arylation reagent used in the arylation
25 reaction include a desired aryl halide compound (such as

chloropyrimidine, dichloropyrimidine, fluoropyrazine, chlorotriazole, chlorooxadiazole, or chlorothiadiazole), and these reagents can be used in the presence of a base (such as DIPEA).

5 The arylation reagent can be used in about one to about five molar equivalent(s) (for example, three molar equivalents) as opposed to 1 mole of the reaction substrates.

10 The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in appropriate solvents (for example, ethers such as THF; halogenated hydrocarbons such as dichloromethane, dichloroethane, and chloroform; aprotic polar solvents such as DMSO; or mixed solvents thereof). The reaction may proceed preferably at room temperature to high temperature, and particularly preferably at high temperature (such as 120 to 130 °C).

[0127]

(Step 3)

20 Further, similarly to the step 4 of the above-mentioned reaction scheme 1, the intermediate compound 9 or the intermediate compound 8 as a starting material is subjected to a phosphoramiditation reaction according to a publicly known method to prepare a desired aryl type (for example, aromatic type or heteroaryl type) of the present compound.

25 Examples of the phosphoramiditation reagent include 2-

cyanoethyl-N,N-diisopropylchlorophosphoramidite, and the reagents can be used in the presence of a base (such as triethylamine, DIPEA).

5 The phosphoramiditation reagent may be used in about 1 to about 10 molar equivalent(s), and preferably 1 to about 5 molar equivalent(s) (such as 1 to 3 molar equivalent(s)) as opposed to 1 mole of the reaction substrates.

10 The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in appropriate solvents (for example, ethers such as THF; halogenated hydrocarbons such as dichloromethane, dichloroethane, and chloroform; acetonitrile; or mixed solvents thereof). The reaction may proceed preferably at -25 °C to 60 °C, and particularly preferably at 0 °C to room
15 temperature.

[0128]

Also, among the above step (1) to step (3), a plural steps (for example, step (1) and step (2)) may be carried out successively.

20 [0123]

Here in any stages from the timing of before the above step 1 to the timing of after the above step 3, a substituent on the ring of the nucleic acid base may be appropriately modified (for example, a benzylation reaction or a
25 acetylation reaction on an amino substituent on a cytosine

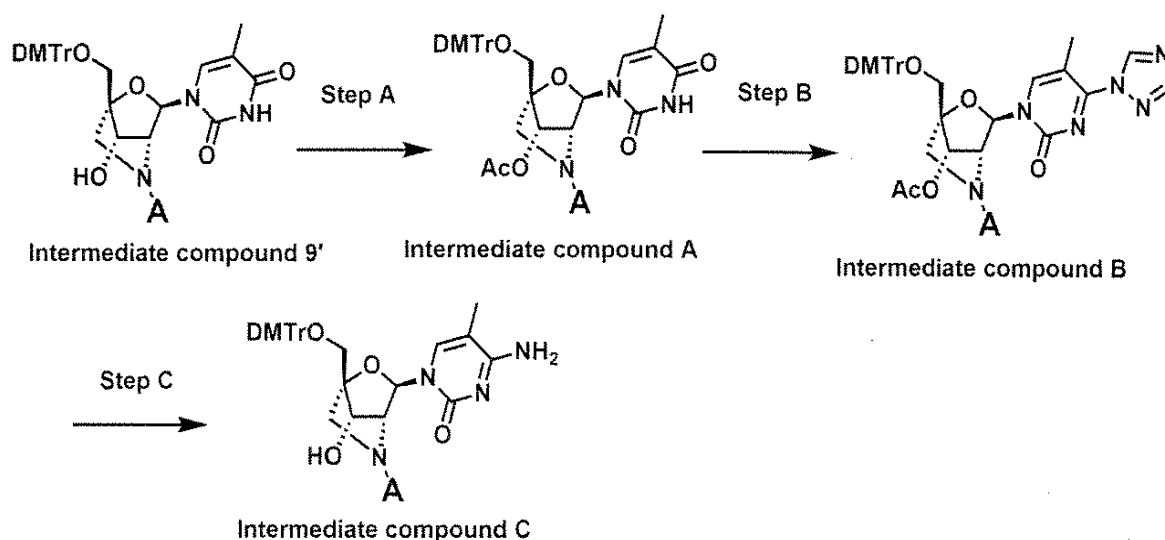
ring), or the substituents on the aromatic ring of the introduced aryl group may be modified (for example, a transformation of the chloro group into an amino group), or both of these substituents may be modified.

5 [0130]

(Alternative preparation method of nucleic acid base exchange reaction)

10 Instead of the nucleic acid base exchange reaction described in step 1, the partial transformation of nucleic acid base may be carried out by using a series of reaction as shown in the following reaction scheme 4. The series of reaction can be carried out according to the method described in the literature (for example, C.H.Kim. et al., J. Med. Chem. 1987, 330, 862).

15 Reaction scheme 4



[0131]

(Step A)

The intermediate compound 9' wherein the nucleic acid base moiety represents a thymine (such as the intermediate compound 2 obtained in the above step 2) as a starting material is reacted with an acetylation reagent (such as acetic anhydride) in the presence of a base (such as pyridine) to prepare an intermediate compound A wherein a hydroxy group is acylated. As needed, a catalytic amount of DMAP may be added as a reaction activating reagent.

The acetylation reagent can be used in about 1 to 3 molar equivalent(s) (such as about 1.5 molar equivalents) as opposed to 1 mole of the reaction substrates.

The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in appropriate solvents (for example, ethers such as THF; or pyridine used as a base). The reaction can proceed preferably at 0 °C to high temperature (such as room temperature).

[0132]

(Step B)

Next, after the thymine moiety is activated with a chlorinating reagent such as phosphoryl chloride, the intermediate compound A as a starting material is reacted with 1,2,4-triazole in the presence of a base (such as DIPEA) to prepare an intermediate compound B wherein the nucleic acid base moiety is substituted with 1,2,4-triazole group.

The 1,2,4-triazole can be used in excess molar equivalents (for example, about five to about 20 molar equivalents, including about 9 to 10 molar equivalents).

5 The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in appropriate solvents (for example, ethers such as THF; halogenated hydrocarbons such as dichloromethane, dichloroethane, and chloroform; acetonitrile; or mixed solvents thereof). The reaction can proceed preferably at 10 -25 °C to room temperature, and particularly preferably at 0 °C to room temperature.

[0133]

(Step C)

Also the intermediate compound B as a starting material 15 is reacted with a base (such as ammonia water) to remove the acetyl group which is introduced in the above step A, and the 1,2,4-triazole group which is modified in the step B, to prepare an intermediate compound C wherein the nucleic acid base moiety is 5-methylcytosine.

20 The ammonia water (such as 28% ammonia water) can be used in excess molar equivalents as opposed to 1 mole of the reaction substrates.

The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in 25 appropriate solvents (for example, ethers such as THF;

acetonitrile; or mixed solvents thereof). The reaction can proceed preferably at 0 °C to room temperature, and particularly preferably at room temperature.

[0134]

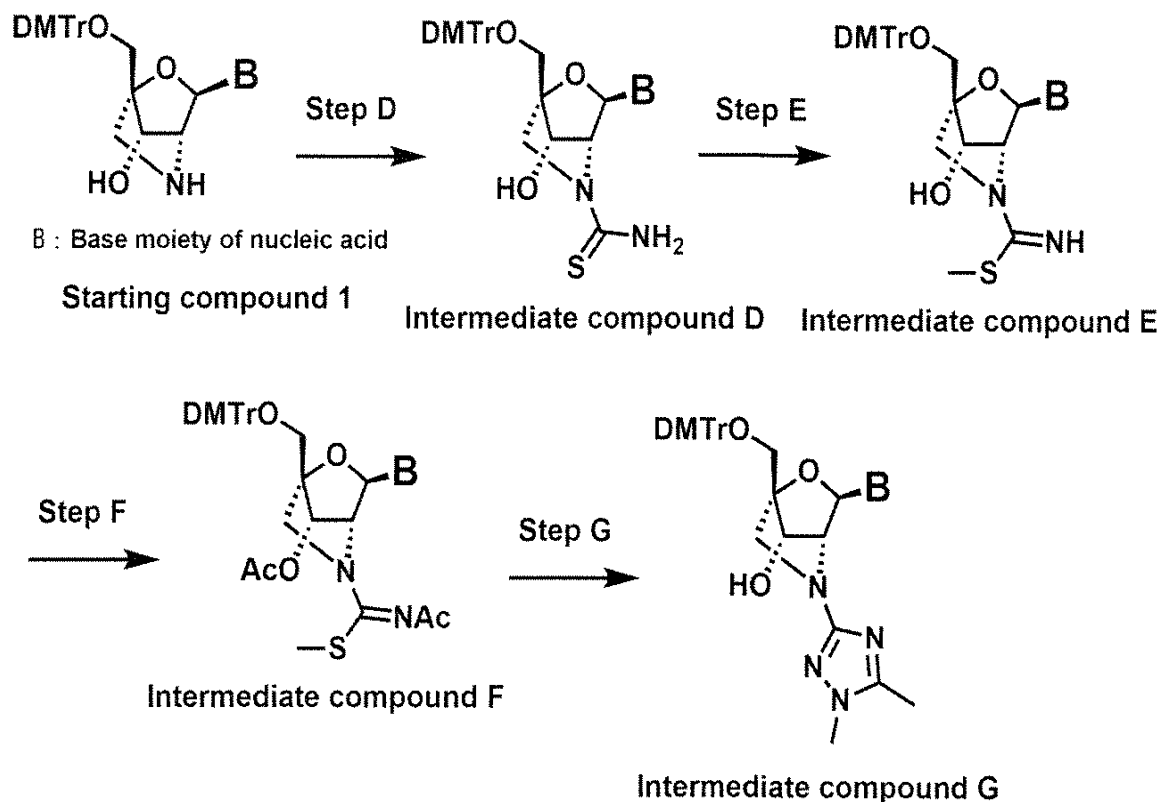
5 Using the obtained intermediate compound C as a starting material, as needed, the substituent on the ring of nucleic acid base is modified (for example, benzoylation or acetylation of amino substituent of the cytosine ring), or the substituent on the aromatic ring of the introduced aryl
10 group is modified (for example, a transformation of a chloro group into an amino group), and thereafter, the substituent is subjected to a phosphoramidation reaction to prepare a desired aryl type (for example, aromatic type, and heteroaryl type) of the present compound.

15 [0135]

(Alternative preparation method 1 for aromatization)

 Further, instead of an aromatization reaction with an aryl reagent as described in step 2, a series of reaction as shown in the following reaction step 5 can be used for an
20 aromatization reaction.

Reaction scheme 5



[0136]

(Step D)

The above starting material 1 is reacted with a reagent for converting into thiourea (such as 1,1-thiocarbonyl imidazole), followed by treating with ammonia water, to prepare an intermediate compound D wherein a group on the amino nitrogen atom is converted into a thiourea.

The reagent for converting into thiourea can be used in about 1 to 5 molar equivalent(s), particularly about 2 molar equivalents, as opposed to 1 mole of the reaction substrates.

The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in appropriate solvents (for example, ethers such as THF). The

reaction can proceed preferably at 0 °C to room temperature (for example, room temperature).

[0137]

(Step E)

5 Next, the intermediate compound D as a starting material is reacted with a methylation reagent (such as methyl iodide) to prepare an intermediate compound having S-methylated and carboimidated group or moiety.

10 The methylation reagent can be used in about 1 to 5 molar equivalent(s), particularly, 3 molar equivalents, as opposed to 1 mole of the reaction substrates.

15 The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in appropriate solvents (for example, ethers such as THF). The reaction can proceed preferably at 0 °C to room temperature, (for example, room temperature).

[0138]

(Step F)

20 Also the intermediate compound E as a starting material is reacted with an acetylation reagent (such as acetic anhydride) in the presence of a base (such as pyridine) to prepare an intermediate compound F having acylation on a carbodiimide group and a hydroxy group. As needed, a catalytic amount of DMAP may be added as a reaction
25 activating reagent.

The acetylation reagent can be use in about 1 to 5 molar equivalent(s) (for example, about 3 molar equivalent(s)) as opposed to 1 mole of the reaction substrates.

5 The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in appropriate solvents (for example, ethers such as THF, or pyridine used as a base). The reaction can proceed preferably at 0 °C to room temperature, (for example, room temperature).

10 [0139]

(Step G)

Also the intermediate compound F as a starting material is subjected to a ring closure reaction with methyl hydrazine to prepare an intermediate compound G wherein a 1,2,4-
15 triazole ring is formed.

The methyl hydrazine can be used in excess molar equivalent (for example, about 5 to 20 molar equivalents (such as about 10 molar equivalents)) as opposed to 1 mole of the reaction substrate.

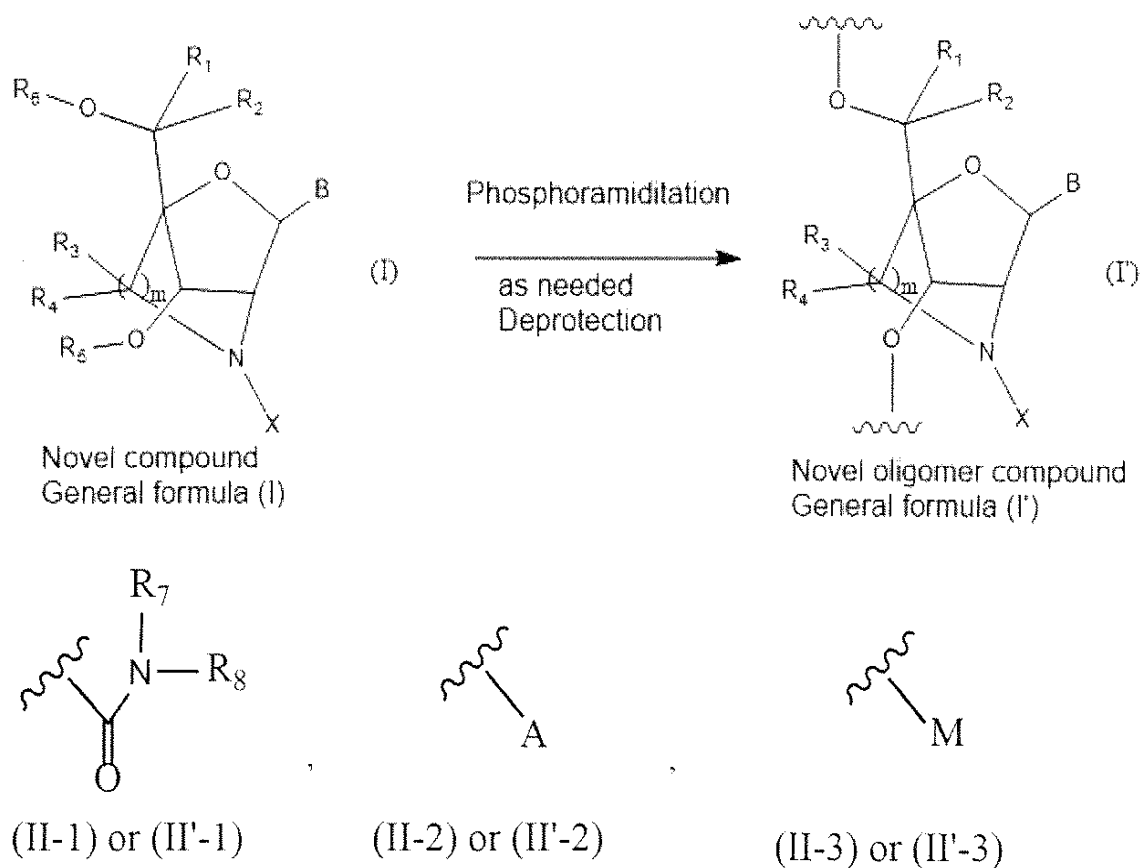
20 The solvents may be any ones unless they affect the reaction adversely, and the reaction may be carried out in appropriate solvents (for example, alcohols such as ethanol). The reaction can proceed preferably at 0 °C to high temperature (for example, room temperature).

25 [0140]

Using the obtained intermediate compound G as a starting material, as needed, the substituent on the ring of nucleic acid base is modified (for example, benzylation or the other ones of amino substituent of the cytosine ring), or the substituent on the aromatic ring of the introduced aryl group is modified (for example, a transformation of a chloro group into an amino group), and thereafter, the substituent is subjected to a phosphoramidation reaction to prepare a desired aryl type (for example, aromatic type, and heteroaryl type) of the present compound.

[0141]

(Preparation method of oligomer)



[0142]

The present compound or salts thereof can be used as a monomer starting material for preparation of oligonucleotide. The oligonucleotide can be synthesized by subjecting to
5 oligomerization reaction of the compound represented by general formula (I) or salts thereof, as needed followed by deprotecting the protecting group for amino group and the protecting group for hydroxy group.

For example, the oligomerization is not limited to any
10 method as long as it is the method that has been generally known in the synthetic chemistry (in particular, nucleic acid synthesis), and it can be used by an phosphoramidite method. The phosphoramidite method can be conducted, for example, according to the method described in WO 2014/046212
15 A1.

Using this oligomerization, oligonucleotide containing one or more nucleotides represented by general formula (I) can be prepared.

[0143]

20 That is, the method for preparing oligonucleotide represented by general formula (I') comprises the following step.

a) a step of oligomerization of the compound represented by general formula (I) or salts thereof, if necessary,
25 followed by deprotecting a protecting group for amino group,

and a protecting group for hydroxy group.

[0144]

Also, the method for preparing the compound represented by general formula (I) or salts thereof may comprise at least one of the following steps among the steps described herein.

a step of preparing the present compound or salts thereof;

a step of phosphoramiditation of the above intermediate compound 4, 7 or 9 or salts thereof to prepare a compound represented by general formula (I') or salts thereof; and

a step of deprotecting a protecting group for hydroxy group in the intermediate compound 3 or 6 to prepare the above intermediate compound 4, 7 or 9.

[0145]

(Preparation method of phosphorothioate oligomer)

In the synthesis of oligonucleotide by phosphoramiditation, an oxidizing agent is used in a conversion reaction of a phosphite group into a phosphate group. DDTT (((dimethylaminomethylidene)amino)-3H-1,2,4-dithiazaoline-3-one) or Beaucage reagent (3H-1,2-benzodithiol-3-one 1,1-dioxide) is used instead of the oxidizing agent used in the reaction to obtain a phosphorothioate oligomer having a protecting group wherein an oxygen atom at P=O of phosphate group is replaced with a

sulfur atom like P=S to make a thiophosphate group.

Examples

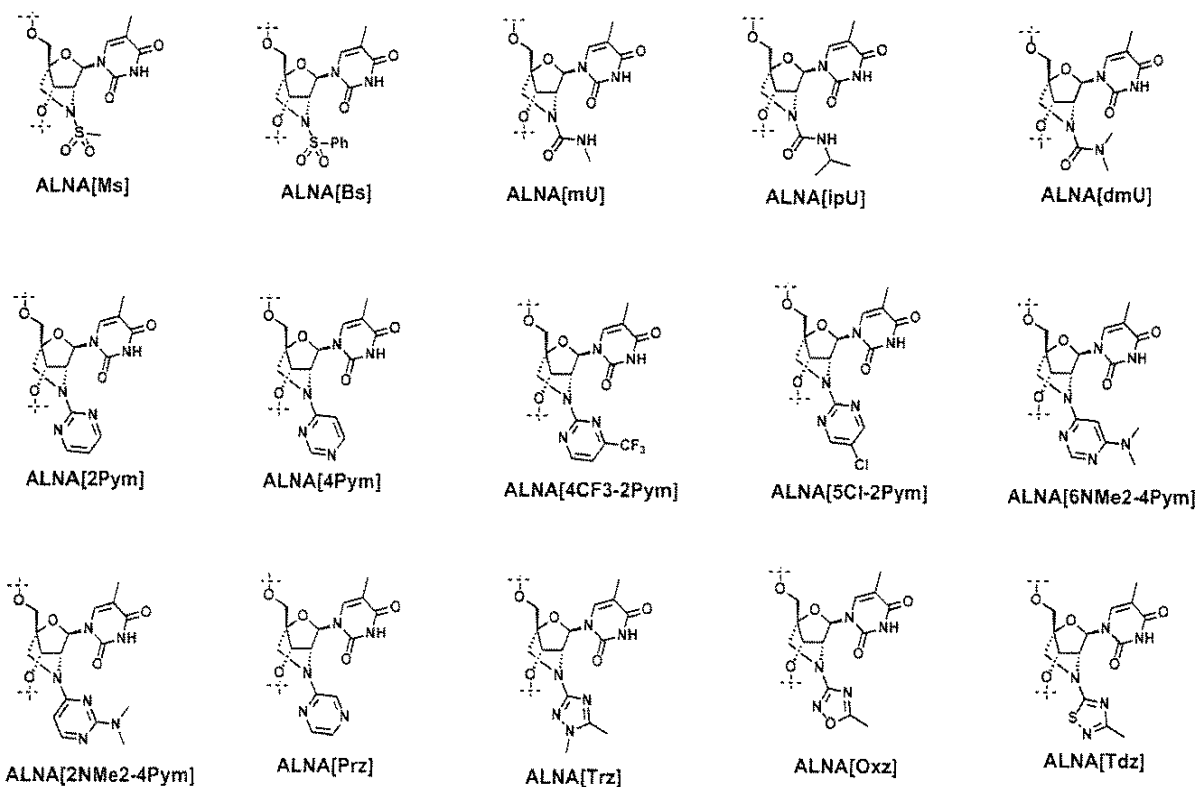
[0146]

5 Hereinafter, the present invention is further specifically described by Examples, however, the present invention is not limited thereto. The identification of compound was carried out by Mass spectrum, High performance liquid chromatography mass spectrometer; LCMS, NMR spectrum, 10 High-performance liquid chromatography (HPLC) and the like. In NMR spectrum, for proton nuclear magnetic resonance (^1H -NMR), an apparatus with 400 MHz of the resonance frequency was used, and for phosphorus nuclear magnetic resonance (^{31}P -NMR), an apparatus with 161.8 MHz of the resonance frequency 15 was used. As symbols used in NMR spectrum, "s" is "singlet", "d" is "doublet", "dd" is "double of doublet", "t" is "triplet", "td" is "double of triplet", "q" is "quartet", "quin" is "quintet", "sept" is "septet", "m" is "multiplet", "br" is "broad", "brs" is "broad singlet", "brd" is "broad 20 doublet", "brt" is "broad triplet", and "J" is "coupling constant".

[0147]

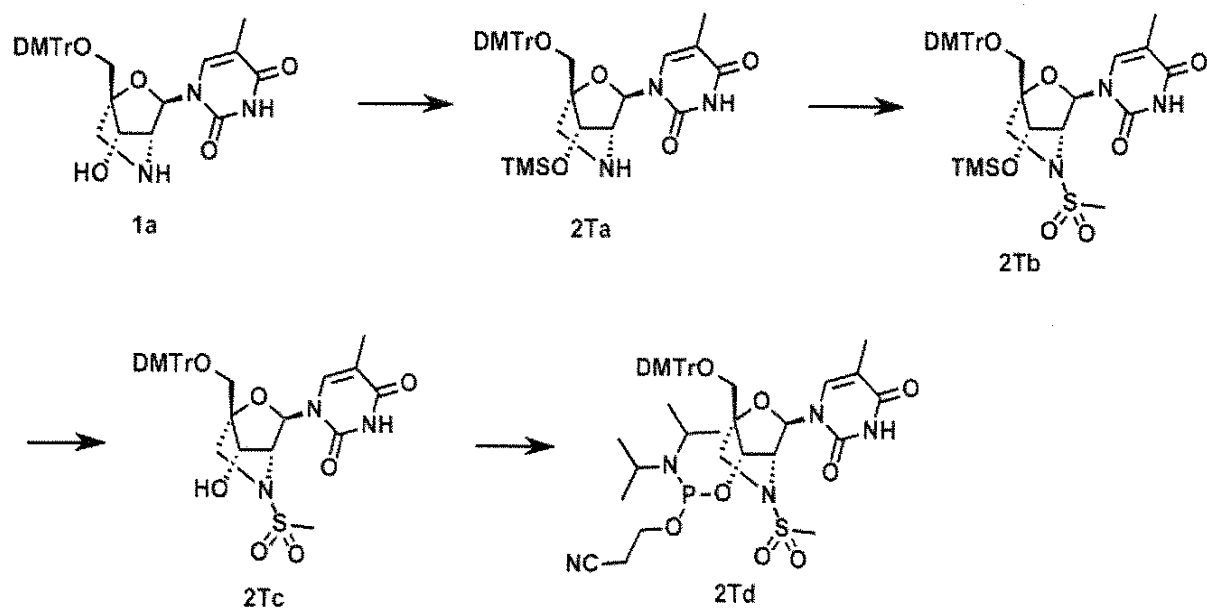
25 A structure and abbreviation of each artificial nucleic acid (hereinafter, as needed, described by abbreviation) is shown in the following structural formula.

Structure and Abbreviation of each artificial nucleic acid



[0148]

Synthesis of ALNA[Ms]-T



[0149]

Example 1 Synthesis of Compound 2Ta

1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy)methyl]-7-trimethylsilyloxy-2-oxa-5-
5 azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2,4-dione

A mixed solution of 1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2,4-dione (1a) (which was prepared according to the method described
10 in WO 2017/047816 A1) (12.35 g, 21.61 mmol) in dichloromethane was added BSA (13 mL, 53.17 mmol), and the mixture was stirred at room temperature for 8 hours. Thereto were added BSA (5.2 mL, 21 mmol) and TMSOTf (200 μ L, 1.0 mmol), and the mixture was stirred again at room temperature
15 for 15 hours. To the reaction solution was added 5% sodium bicarbonate water (50 mL), and the mixture was stirred, and extracted with dichloromethane. The aqueous layer was mixed with dichloromethane (61 mL), and extracted, and the dichloromethane organic layer was combined. After the
20 solvent was evaporated from the organic layer, the resulting residue was purified by silica gel chromatography (chloroform/methanol, 100/0 to 98/2) to obtain a compound 2Ta (10.49 g, yield 75%).

MS(ESI): m/z = 644 (M+H)⁺

[0150]

Example 2 Synthesis of Compound 2Tb

1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy)methyl]-5-methylsulfonyl-7-trimethylsilyloxy-
2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-
5 2,4-dione

To a mixed solution of compound 2Ta (10.19 g, 15.83 mmol) in dichloromethane (50 mL) was added DIPEA (5.4 mL, 31 mmol), and thereto was added mesyl chloride (1.4 mL, 18 mmol) under ice bath, and the mixture was stirred in ice bath for
10 1.5 hours. To the reaction solution was added 5% sodium bicarbonate water (20 mL), and the mixture was stirred, and extracted with dichloromethane. The aqueous layer was mixed with dichloromethane (20 mL), and extracted, and combined and mixed with a dichloromethane organic layer. After the
15 solvent was evaporated from the organic layer, the resulting residue was purified by silica gel chromatography (chloroform/methanol, 100/0 to 98/2) to obtain a compound 2Tb (6.56 g, yield 57%).

MS(ESI): m/z = 720 (M-H)⁻

20 [0151]

Example 3 Synthesis of Compound 2Tc

1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy)methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2,4-dione

25 To a mixed solution of compound 2Tb (3.02 g, 4.13 mmol)

in tetrahydrofuran (15 mL) was added TBAF (1.35 mL, 4.58 mmol, 1M tetrahydrofuran solution) in ice bath, and the mixture was stirred for 30 minutes. To the reaction solution were added 10% aqueous ammonium chloride solution (20 mL) and dichloromethane (20 mL), and the mixture was stirred, and extracted with dichloromethane. The aqueous layer was mixed with dichloromethane (20 mL), extracted, and the dichloromethane organic layer was mixed. After the solvent was evaporated from the organic layer, the resulting residue was purified by silica gel chromatography (chloroform/methanol, 100/0 to 98/2) to obtain a compound 2tc (2.47 g, yield 92%).

MS(ESI): m/z = 648 (M-H)⁻

[0152]

Example 4 Synthesis of Compound 2Td

3-[[[(1R,4R,6R,7S)-4-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-6-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-methylsulfonyl-5-oxa-2-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 1)

To a solution of the compound 2Tc (2.47 g, 3.80 mmol) in dichloromethane (12.5 mL) were added DIPEA (2.00 mL, 12.0 mmol), and 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (2.10 mL, 9.40 mmol), and the mixture was stirred at room temperature for 5 hours.

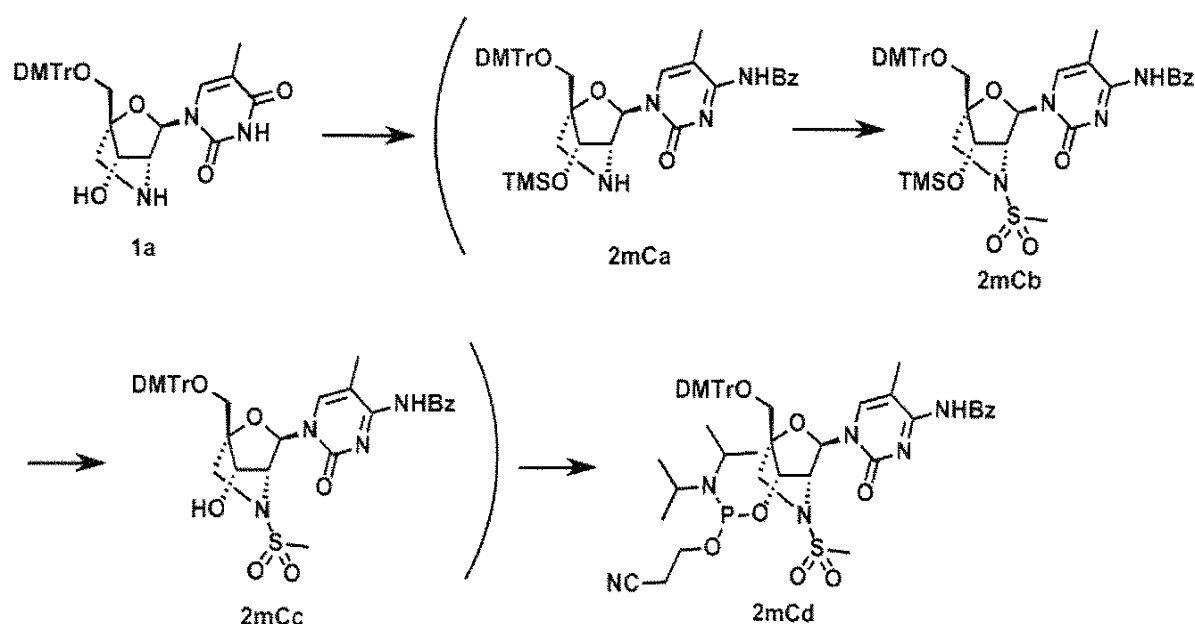
Then thereto was added 5% aqueous sodium hydrocarbonate solution (15 mL) and dichloromethane (10 mL) under ice-cooling, and the mixture was stirred at room temperature, and then extracted with dichloromethane. The aqueous layer
 5 was extracted with dichloromethane (10 mL), and combined and mixed with the dichloromethane organic layer. After the solvent was evaporated from the mixture under reduced pressure, the mixture was purified by silica gel chromatography (hexane/ethyl acetate, 60/40 to 40/60) to
 10 obtain a compound 2Td as the present compound 1 (2.13 g, yield 66%).

MS(ESI): $m/z = 850$ (M+H)⁺

³¹P-NMR (CDCl₃) δ : 150.30, 149.12

[0153]

15 Synthesis of ALNA[Ms]-mC



[0154]

Example 5 Synthesis of Compound 2mCc

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy]methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-
5 azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-
yl]benzamide

To a suspension solution of the compound 1a (10.33 g,
18.1 mmol) and N-(5-methyl-2-oxo-1H-pyrimidin-4-yl)
benzamide (12.42 g, 54.2 mmol) in toluene solution (180 ml)
10 was added BSA (44 mL, 180 mmol), and the mixture was stirred
at room temperature for 15 minutes. Thereto was added TMSOTf
(0.52 mL, 2.71 mmol), and the mixture was stirred at room
temperature for 2 hours. The reaction solution was added to
a mixed solution of IPE (100 mL), saturated sodium
15 bicarbonate water (50 mL) and water (50 mL), and the mixture
was stirred for 10 minutes, and insoluble materials were
removed through Celite (TM). The aqueous layer was removed
from the filtrates, and the organic layer was washed with
water (50 mL) and saturated brine (50 mL) successively. The
20 organic layer was passed through a phase separator, and the
solvent was evaporated. The residue was solubilized with
dichloromethane (90 mL), and thereto were added
triethylamine (5.0 mL, 36.0 mmol) and methane sulfonic
anhydride (3463 mg, 19.9 mmol), and the mixture was stirred
25 at room temperature for 40 minutes. Thereto was added

methane sulfonic anhydride (944 mg, 5.42 mmol), and the mixture was stirred at room temperature for 30 minutes. Thereto was added further methane sulfonic anhydride (944 mg, 5.42 mmol) and triethylamine (1.3 mL, 9.06 mmol), and
5 the mixture was stirred at room temperature for 20 minutes. The mixture was diluted with ethyl acetate (80 mL), and thereto were added water (40 mL) and saturated sodium bicarbonate water (40 mL), and the mixture was stirred. The aqueous layer was removed, and the organic layer was washed
10 with saturated brine (50 mL), and then passed through a phase separator, and a solvent was evaporated, and the resulting mixture was dried under vacuum. A residue was added by THF (90 mL) to make a solution, and thereto was added TBAF (0.1 M, THF solution, 21.7 mL, 21.7 mmol) under ice-cooling, and
15 the mixture was stirred under ice-cooling for 15 minutes. The mixture was diluted with ethyl acetate (90 mL), and thereto were added aqueous ammonium chloride solution (40 mL) and water (40 mL), and the mixture was stirred. The aqueous layer was removed, and the organic layer was passed
20 through a phase separator, and the solvent was evaporated. The resulting residue was purified by silica gel chromatography (chloroform/methanol, 100/0 to 95/5) to obtain a compound 2mCc (5702 mg, yields over three steps 38%).

25 MS(ESI): $m/z = 753$ (M+H)⁺

[0155]

Example 6 Synthesis of Compound 2mCd

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy]methyl]-7-[2-cyanoethoxy-

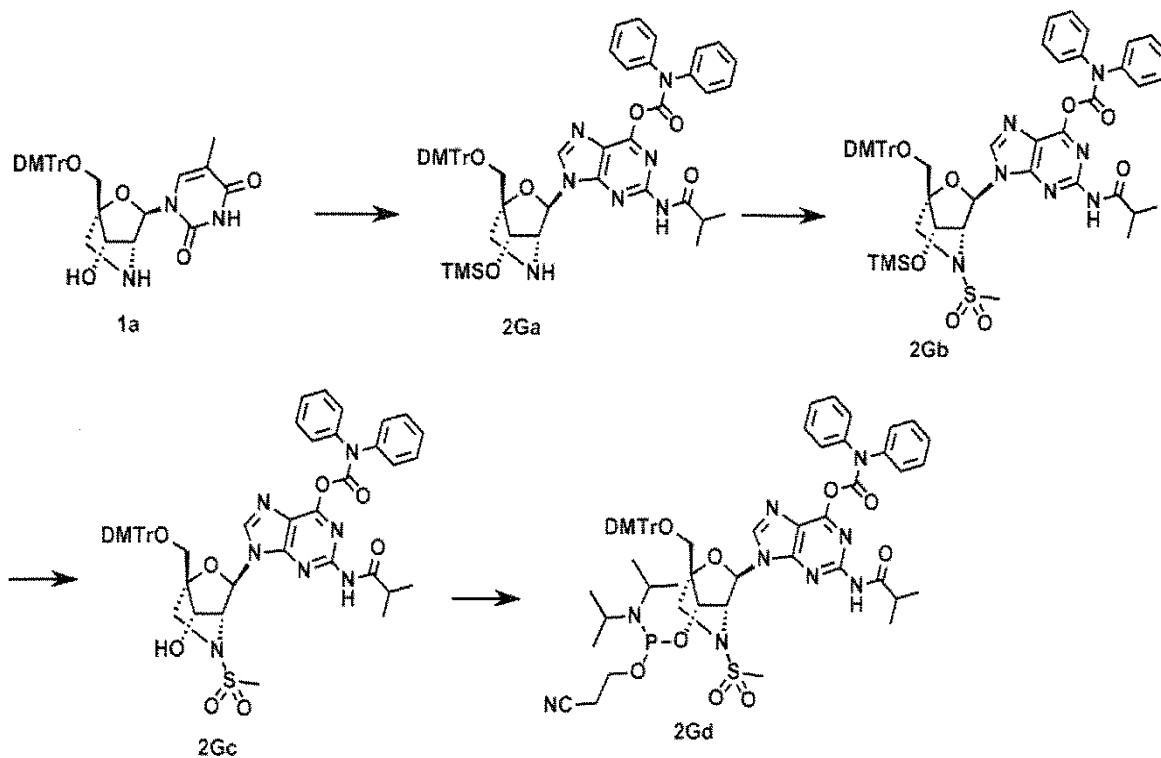
5 (diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-
yl]benzamide (Present compound 2)

The compound 2mCd as the present compound 2 (6809 mg,
yield 72%) was obtained from the compound 2mCc (8099 mg,
10 9.89 mmol) similarly to the Example 4.

MS(APCI): $m/z = 954$ (M+H)⁺

[0156]

Synthesis of ALNA[Ms]-G



[0157]

Example 7 Synthesis of Compound 2Ga

[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy)methyl]-7-trimethylsilyloxy-2-oxa-5-
5 azabicyclo[2.2.1]heptan-3-yl]-2-(2-
methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate

To a solution of the compound 1a (5040 mg, 8.817 mmol)
and a solution of [2-(2-methylpropanoylamino)-9H-purin-6-
yl]N,N-diphenyl carbamate (5400 mg, 12.97 mmol) in 1,2-
10 dichloroethane (24.7 mL) was added BSA (14.8 mL, 60.46 mmol),
and the mixture was stirred at 64 °C for 1 hour. Successively,
thereto was added TMSOTf (0.167 mL, 0.864 mmol) was added
gradually while maintaining the inner temperature at 60 °C,
and the mixture was stirred for 1 hour. After allowed to
15 stand to cool, the reaction solution was added gradually to
a mixed solution of chloroform (40 mL) and saturated aqueous
sodium hydrogen carbonate solution (40 mL), which was
prepared separately, and the mixture was stirred. After
suction filtering insoluble materials off, the filtrates
20 were extracted with ethyl acetate (50 mL). The organic layer
was washed with saturated brine (50 mL), and then dried over
sodium sulfate. Successively, after the filtrates were
evaporated under reduced pressure, the resulting residue was
purified by silica gel chromatography (chloroform/methanol,
25 100/0 to 97/3) to obtain a compound 2Ga (6350 mg, yield 79%).

MS(ESI): $m/z = 935$ (M+H)⁺

[0158]

Example 8 Synthesis of Compound 2Gb

[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
5 phenylmethoxy)methyl]-5-methylsulfonyl-7-trimethylsilyloxy-
2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-2-(2-
methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate

To a solution of the compound 2Ga (2370 mg, 2.537 mmol)
in dichloromethane (24.9 mL) was added triethylamine (0.692
10 mL, 4.984 mmol), and the mixture was stirred at room
temperature for 5 minutes. Successively, thereto was added
methanesulfonyl chloride (0.193 mL, 2.488 mmol) under ice-
cooling, and the mixture was stirred for 1 hour. To the
reaction solution were added methylene chloride (30 mL) and
15 saturated aqueous sodium hydrogen carbonate solution (30 mL)
under ice-cooling, and the mixture was separated with a
separatory funnel. The organic layer was washed with
saturated brine (30 mL), and then dried over sodium sulfate.
Successively, after the filtrates were evaporated under
20 reduced pressure, the resulting mixture was purified by
silica gel chromatography (hexane/ethyl acetate, 67/33 to
50/50) to obtain a compound 2Gb (2210 mg, yield 88%).

MS(ESI): $m/z = 1013$ (M+H)⁺

[0159]

25 Example 9 Synthesis of Compound 2Gc

[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-2-(2-methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate

5 To a solution of the compound 2Gb (2200 mg, 2.173 mmol) in THF (11 mL) was added TBAF (2.4 mL, 3.407 mmol) under ice-cooling, and the mixture was stirred at room temperature for 5 hours. To the reaction solution were added ethyl acetate (30 mL) and aqueous ammonium chloride solution (30
10 mL) under ice-cooling, and the mixture was filtered and separated with a separatory funnel. The organic layer was washed with saturated brine (30 mL), and then dried over sodium sulfate. Successively, after the filtrates were evaporated under reduced pressure, the resulting residue was
15 purified by silica gel chromatography (hexane/ethyl acetate, 60/40 to 50/50). In order to remove raw materials, the mixture was purified again by silica gel chromatography (chloroform/methanol, 100/0 to 97/3) to obtain a compound 2Gc (1070 mg, yield 52%).

20 MS(ESI): $m/z = 941$ (M+H)⁺
[0160]

Example 10 Synthesis of Compound 2Gd

[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-
25 (diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-

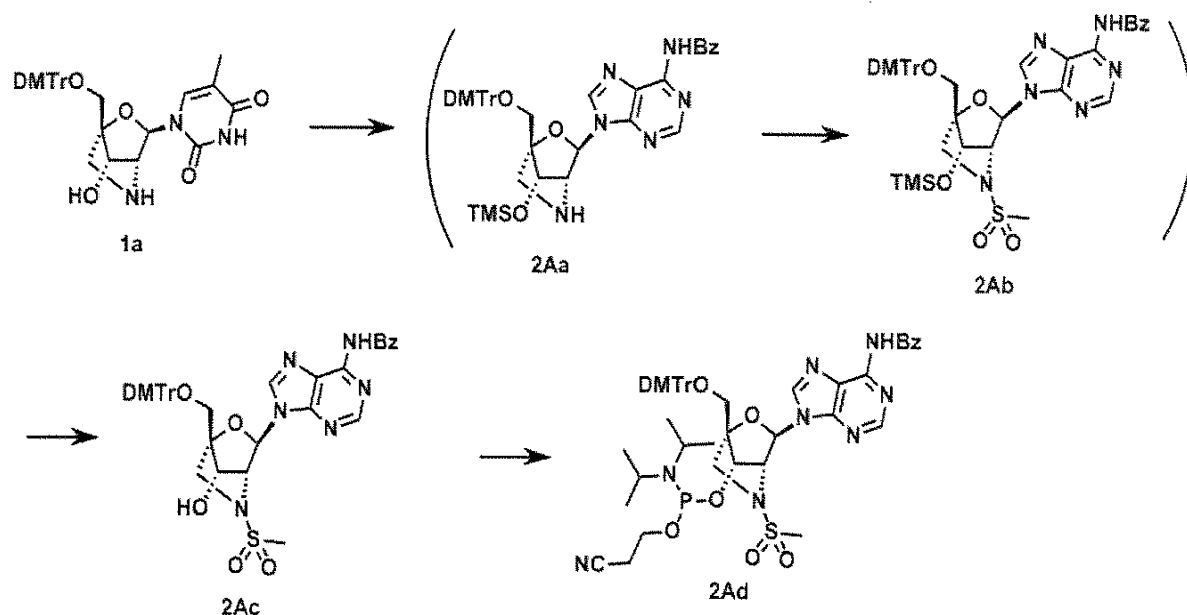
azabicyclo[2.2.1]heptan-3-yl]-2-(2-methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate
(Present compound 3)

To a solution of the compound 2Gc (1070 mg, 1.139 mmol)
5 in dichloromethane (5.690 mL) were added DIPEA (0.590 mg,
3.407 mmol) and 2-cyanoethyl N,N-diisopropylchloro
phosphoramidite (0.510 mL, 2.282 mmol) under ice-cooling,
and the mixture was stirred at room temperature for 22 hours.
To the reaction solution were added dichloromethane (20 mL)
10 and saturated sodium bicarbonate water (20 mL) under ice-
cooling, and the mixture was separated with a separatory
funnel. The aqueous layer was removed, and the organic layer
was passed through a phase separator. After the filtrates
were evaporated under reduced pressure, the resulting
15 residue was purified by silica gel chromatography
(hexane/ethyl acetate, 50/50 to 40/60) to obtain a compound
2Gd as the present compound 3 (548 mg, yield 42%).

MS(ESI): $m/z = 1141$ ($M+H$)⁺

[0161]

20 Synthesis of ALNA[Ms]-A



[0162]

Example 11 Synthesis of Compound 2Ac

N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
 5 phenylmethoxy)methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-
 azabicyclo[2.2.1]heptan-3-yl]-purin-6-yl]benzamide

To a suspension of the compound 1a (10.0 g, 17.6 mmol) and N-(9H-purin-6-yl)benzamide (5.04 g, 21.1 mmol) in toluene (176 mL) was added dropwise BSA (30 mL, 123 mmol), and the mixture was warmed to 60 °C, and stirred for 30 minutes. Thereto was added TMSOTf (1.03 mL, 5.34 mmol) and the mixture was stirred at 60 °C for 15 minutes. The reaction solution was added to the mixture solution of IPE (100 mL) and saturated sodium bicarbonate water (100 mL) while stirring under ice-cooling, and the mixture was stirred for 20 minutes, and insoluble materials were filtered off. After

the filtered residue was washed with Et₂O (20 mL) twice, the filtrates were separated with a separatory funnel, and washed with saturated brine (50 mL). After the mixture was passed through a phase separator, the mixture was concentrated under reduced pressure. The residue was solubilized with dichloromethane (24.9 mL), and thereto were added pyridine (4.57 mL, 56.5 mmol) and methane sulfonic anhydride (3614 mg, 20.75 mmol), and the mixture was stirred at room temperature for 30 minutes. Thereto were added saturated aqueous sodium hydrogen carbonate solution (50 mL) and water (50 mL), and the mixture was stirred, and separated with a separatory funnel. After the organic layer was washed with saturated brine (50 mL), the mixture was passed through a phase separator, and the solvent was evaporated under reduced pressure. The residue was added by THF (88 mL) to make a solution, and thereto was added TBAF (1M THF solution, 21.1 mL, 21.1 mmol) under ice-cooling, and the mixture was stirred under ice-cooling for 30 minutes. The mixture was diluted with ethyl acetate (80 mL), and thereto were added aqueous ammonium chloride solution (50 mL) and water (50 mL), and the mixture was stirred at room temperature. The mixture was separated with a separatory funnel, and the organic layer was washed with saturated brine (50 mL), and then passed through a phase separator, and the solvent was evaporated. The resulting residue was purified by silica gel

chromatography (chloroform/methanol, 100/0 to 95/5) to a compound 2Ac (10.6 g, yields over three steps 75%).

MS (ESI): $m/z = 763$ (M+H)⁺

[0163]

5 Example 12 Synthesis of Compound 2Ad

N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-purin-6-yl]benzamide (Present
10 compound 4)

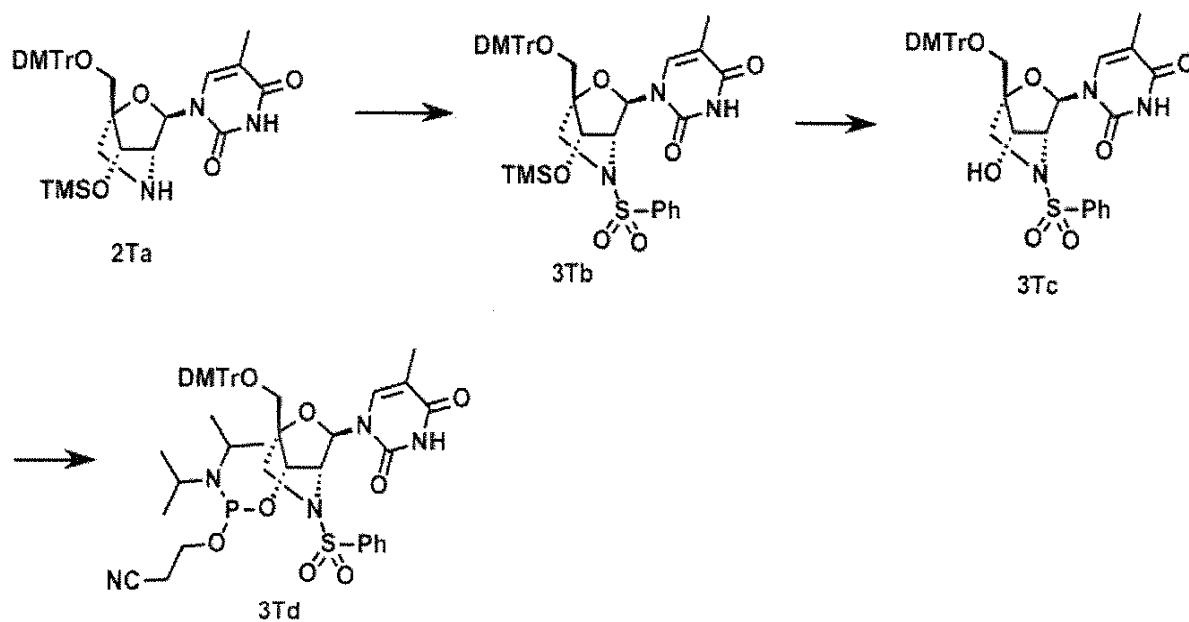
To a solution of the compound 2Ac (5.60 g, 7.01 mmol) in dichloromethane (40 mL) were added DIPEA (4.8 mL, 28.0 mmol) and 2-cyanoethyl N,N-diisopropylchloro phosphoramidite (4.7 mL, 21.0 mmol) in ice-cooling, and the mixture was stirred at
15 room temperature for 4 hours. Thereto was added ethanol (1.2 mL, 21.0 mmol) under ice-cooling, and the mixture was stirred at room temperature for 5 minutes. The mixture was placed again under ice-cooling, and thereto were added saturated sodium bicarbonate water (20 mL), and saturated brine (20 mL),
20 and the mixture was stirred, and the organic layer was passed through a phase separator. After the filtrates were evaporated under reduced pressure, the resulting residue was purified by silica gel chromatography (hexane/ethyl acetate, 67/33 to 10/90) to obtain a compound 2Ad as the present
25 compound 4 (5.44 g,

yield 77%).

MS(ESI): $m/z = 964$ (M+H)⁺

[0164]

Synthesis of ALNA[Bs]-T



5

[0165]

Example 13 Synthesis of Compound 3Tb

1-[(1R,3R,4R,7S)-5-(benzenesulfonyl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-trimethylsilyloxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-pyrimidin-2,4-dione

A compound 2Ta (1.00 g, 1.56 mmol) was azeotroped with toluene. To the residue were added dichloromethane (5 mL) and triethylamine (0.43 mL, 3.1 mmol), and the mixture was solubilized completely. The mixture solution was cooled

15

under ice-cooling, and thereto was added benzenesulfonyl chloride (0.21 mL, 1.6 mmol), and the mixture was stirred under ice-cooling for 4 hours. To the reaction solution was added 5% potassium carbonate solution (20 mL), and the mixture was stirred, and extracted with dichloromethane (5 mL). The aqueous layer was mixed with dichloromethane (5 mL), extracted, and the extracts were combined and mixed with the dichloromethane organic layer. After the solvent was evaporated from the organic layer, the resulting residue was purified by silica gel chromatography (hexane/ethyl acetate, 40/60 to 20/80) to obtain a compound 3Tb (0.78 g, yield 64%). MS(ESI): $m/z = 782$ (M-H)⁻

[0166]

Example 14 Synthesis of Compound 3Tc

1-[(1R,3R,4R,7S)-5-(benzenesulfonyl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2,4-dione

A compound 3Tc (0.59 g, yield 84%) was synthesized from the compound 3Tb (0.78 g, 0.99 mmol) similarly to the Example 3.

MS(ESI): $m/z = 710$ (M-H)⁻

[0167]

Example 15 Synthesis of Compound 3Td

3-[(1R,3R,4R,7S)-5-(benzenesulfonyl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-3-(5-methyl-2,4-dioxo-

pyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-
 (diisopropylamino)phosphanyl]oxypropanenitrile (Present
 compound 5)

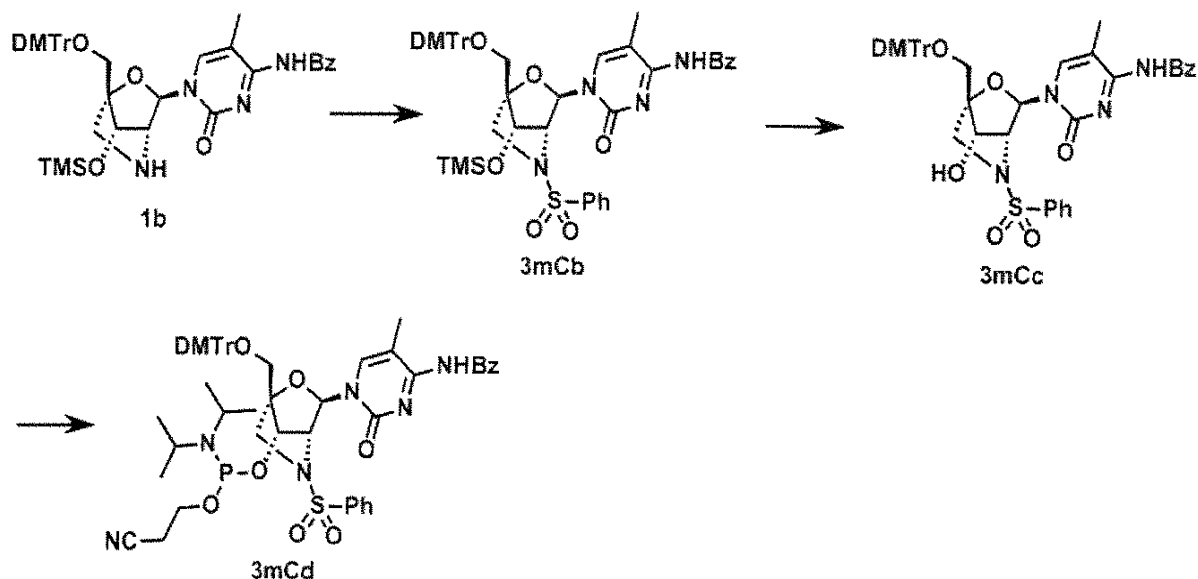
To a solution of the compound 3Tc (599 mg, 0.765 mmol)
 5 in acetonitrile (3 mL) were added diisopropylammonium
 tetrazolide (171 mg, 0.996 mmol), 2-Cyanoethyl N,N,N',N'-
 Tetraisopropylphosphordiamidite (0.36 mL, 1.1 mmol), and the
 mixture was stirred at room temperature for 19 hours. To the
 reaction solution were added ethyl acetate (10 mL) and water
 10 (5 mL), and after the mixture was stirred, the aqueous layer
 was separated. To the aqueous layer was added ethyl acetate
 (10 mL), and after the mixture was mixed, the organic layer
 was separated. The organic layer was combined and mixed with
 above organic layer, and the mixture was concentrated at 40
 15 °C under reduced pressure. The resulting residue was
 purified by silica gel chromatography (hexane/ethyl acetate,
 67/33 to 10/90) to obtain a compound 3Td as the present
 compound 5 (0.65 g, yield 86%).

MS(ESI): $m/z = 912$ (M+H)⁺

20 ³¹P-NMR (CDCl₃) δ : 149.68, 149.22

[0168]

Synthesis of ALNA[Bs]-mC



[0169]

Example 16 Synthesis of Compound 3Cb

5 N-[1-[(1R,3R,4R,7S)-5-(benzenesulfonyl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-trimethylsilyloxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide

10 A compound 3mCb (0.85 g, yield 71%) was synthesized from the compound 1b (1.00 g, 1.34 mmol) which was synthesized according to the method described in WO 2017/047816 A1 similarly to the Example 13.

MS(ESI): $m/z = 887$ ($M+H$)⁺

[0170]

Example 17 Synthesis of Compound 3mCb

15 N-[1-[(1R,3R,4R,7S)-5-(benzenesulfonyl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-

yl]benzamide

A compound 3mCc (0.78 g, yield 99%) was synthesized from the compound 3mCb (0.85 g, 0.96 mmol) similarly to the Example 14.

5 MS(ESI): $m/z = 813$ (M+H)⁺

[0171]

Example 18 Synthesis of Compound 3mCd

N-[1-[(1R,3R,4R,7S)-5-(benzenesulfonyl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 6)

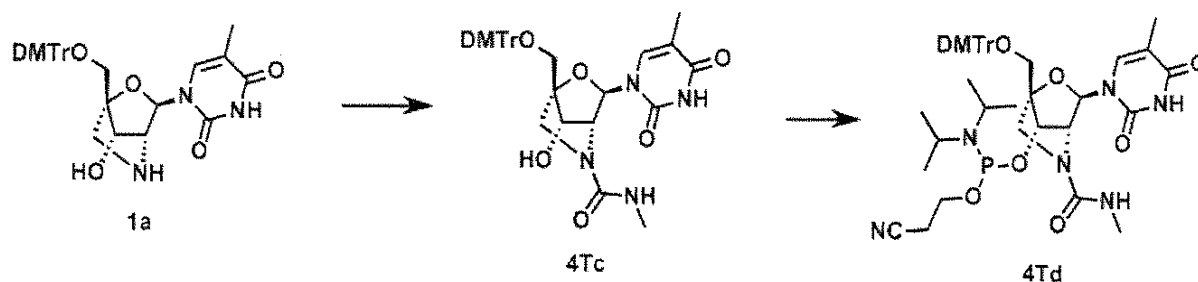
A compound 3mCd (0.53 g, yield 55%) as the present compound 6 was synthesized from the compound 3mCc (0.78 g, 0.89 mmol) similarly to the Example 15.

MS(ESI): $m/z = 1015$ (M+H)⁺

³¹P-NMR (CDCl₃) δ : 149.94, 149.34

[0172]

Synthesis of ALNA[mU]-T



[0173]

Example 19 Synthesis of Compound 4Tc

(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy)methyl]-7-hydroxy-N-methyl-3-(5-methyl-2,4-
5 dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-5-
carboxamide

A compound 1a (1.20 g, 2.10 mmol) was azeotroped with
toluene, and to the mixture solution of the residue in
tetrahydrofuran (6 mL) was added N,N-diisopropylethylamine
10 (0.61 mL, 3.5 mmol), and thereto was added a solution of N-
methylcarbamoyl chloride (0.20 g, 2.2 mmol) in
tetrahydrofuran (1 mL) in ice bath, and the mixture was
stirred for 3 hours in ice bath. To the reaction solution
were added 5% sodium bicarbonate water (20 mL) and ethyl
15 acetate (10 mL), and the mixture was stirred, and the mixture
was extracted with ethyl acetate. The aqueous layer (10 mL)
was extracted with ethyl acetate (10 mL) twice, and the
extracts were combined and mixed with the organic layer.
After the solvent was evaporated from the organic layer, the
20 resulting residue was purified by silica gel chromatography
(chloroform/methanol, 100/0 to 97/3) to obtain a compound
4Tc (1.28 g, yield 97%).

MS(ESI): m/z = 627 (M-H)⁻

[0174]

25 Example 20 Synthesis of Compound 4Td

(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy)methyl]-7-[2-cyanoethoxy-
(diisopropylamino)phosphanyl]oxy-N-methyl-3-(5-methyl-2,4-
dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-5-
5 carboxamide (Present compound 7)

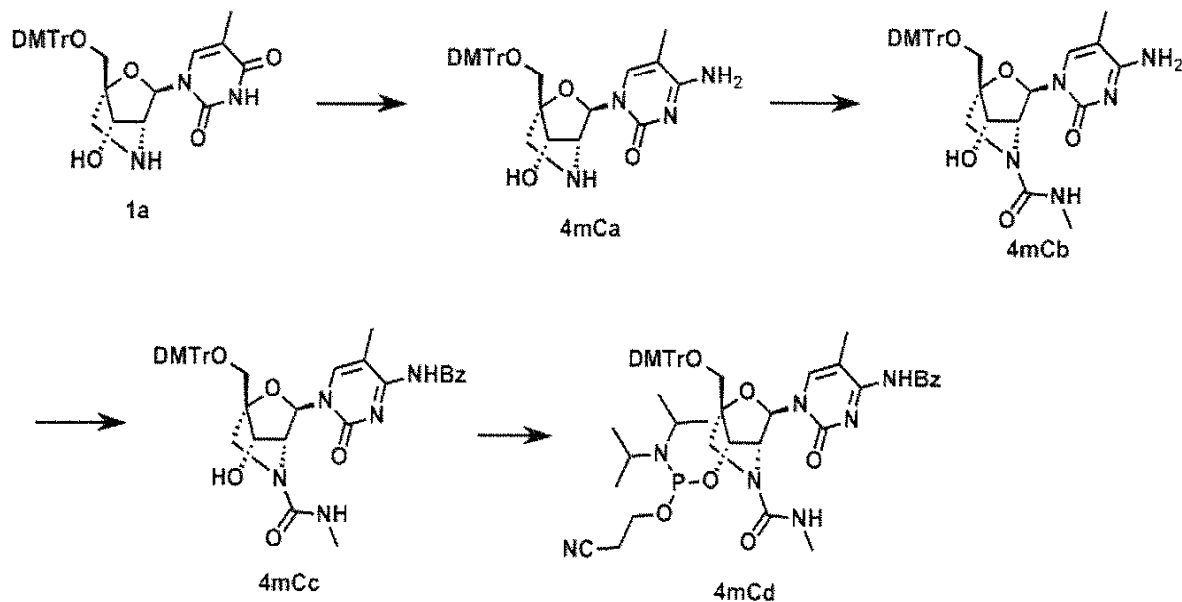
A compound 4Td (0.56 g, yield 37%) as the present
compound 7 was synthesized from the compound 4Tc (1.16 g,
1.8 mmol) similarly to the Example 15.

MS(ESI): $m/z = 828 (M-H)^-$

^{31}P -NMR (CDCl_3) δ : 149.31, 147.44

[0175]

Synthesis of ALNA[mU]-mC



[0176]

Example 21 Synthesis of Compound 4mCa

4-amino-1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-

phenylmethoxy)methyl]-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-pyrimidin-2-one

To a mixed solution of the compound 1a (105.0 g, 184 mmol), 5-methylcytosine (69.0 g, 551 mmol) and 1,2-dichloroethane (918 mL) was added dropwise BSA (584 mL, 2390 mmol) over 18 minutes. The mixture was warmed to 60 °C, and stirred for 20 minutes. After an inner temperature was cooled to 19.5°C under ice-cooling, thereto was added TMSOTf (10 mL, 55.2 mmol), and the mixture was warmed to 60 °C. After the mixture was stirred for 40 minutes, the reaction solution was cooled under ice-cooling, and thereto was added dropwise 5% aqueous sodium hydrocarbonate solution (525 mL). Next, thereto were added chloroform (1050 mL) and aqueous 15% sodium chloride solution (525 mL). After the resulting slurry was filtered through Celite, the filtered residue were washed with chloroform (525 mL) to obtain a residue 1 and a filtrate 1. To the residue 1 were added ethyl acetate (2100 mL) and water (2100 mL), and the mixture was stirred for 1 hour. After the mixture was filtered through Celite, the residue was washed with ethyl acetate (1050 mL) to obtain the filtrates 2. The filtrates 2 were separated with a separatory funnel, and the organic layer washed with water (525 mL) twice. On the other hand, the filtrates 1 were separated with a separatory funnel, and the aqueous layer was extracted with chloroform (525 mL). The organic layers

were combined, and washed with water 1050 mL twice. The resulting organic layer was concentrated under reduced pressure to obtain pale yellow solids (125.6 g). Thereto was added THF (550 mL), and the mixture was stirred, and thereto was added 1M TBAF/THF solution (276 mL, 276 mmol). After the mixture was stirred at room temperature for 3 hours, the reaction solution was quenched with water (550 mL), and thereto was added chloroform (550 mL), and the mixture was separated with a separatory funnel. The aqueous layer was extracted with chloroform (275 mL) again, and the organic layers were combined. The organic layer was washed with water, and then concentrated under reduced pressure. The resulting residue was purified by silica gel chromatography (chloroform/methanol, 20/1 to 4/1) to obtain a compound 4mCa (69.62 g, yield 62%).

MS(ESI): $m/z = 593 (M+Na)^+$

[0177]

Example 22 Synthesis of Compound 4mCb

(1R,3R,4R,7S)-3-(4-amino-5-methyl-2-oxo-pyrimidin-1-yl)-1-
[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-hydroxy-N-
methyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide

The compound 4mCa (1.00 g, 1.75 mmol) was azeotroped with toluene, and to the mixture solution of the residue in tetrahydrofuran (5 mL) was added triethylamine (0.49 mL, 3.5 mmol), and thereto was added a solution of N-methylcarbamoyl

chloride (0.17 g, 1.8 mmol) in tetrahydrofuran (1 mL) under ice-cooling, and the mixture was stirred for 3 hours in ice bath. To the reaction solution were added tap water (15 mL), ethyl acetate (1 mL) and chloroform (15 mL), and the mixture
5 was stirred, and extracted with organic solvent. The aqueous layer was extracted with chloroform (5 mL), and the organic layers were combined and mixed. The organic layer was dried over magnesium sulfate, and the solids were filtered. The solvents were evaporated from the filtrates to obtain a
10 compound 4mCb (1.18 g, yield quant.). The residue was used to a next reaction without purification.

MS(ESI): $m/z = 627$ (M-H)⁻

[0178]

Example 23 Synthesis of Compound 4mCc

15 (1R,3R,4R,7S)-3-(4-benzamide-5-methyl-2-oxo-pyrimidin-1-yl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-hydroxy-N-methyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide

The compound 4mCb (1.18 g) was azeotroped with toluene,
20 and to a mixture solution of the residue in tetrahydrofuran (5 mL) were added benzoic anhydride (0.51 g, 2.27 mmol) and methanol (0.01 mL, 0.25 mmol) at room temperature, and the mixture was stirred at 70 °C as a temperature of outer bath for 4 hours. Thereto were added 5% sodium bicarbonate water
25 (10 mL) and ethyl acetate (10 mL), and the mixture was

stirred, and extracted with organic solvent(s). The aqueous layer was extracted with ethyl acetate (15 mL), and the organic layers were combined and mixed. After the solvent was evaporated from the organic layer, the resulting residue was purified by silica gel chromatography (chloroform/methanol, 100/0 to 97/3) to obtain a compound 4mCc (0.99 g, yield 77 %).

MS(ESI): $m/z = 731$ (M-H)⁻

[0179]

10 Example 24 Synthesis of Compound 4mCd

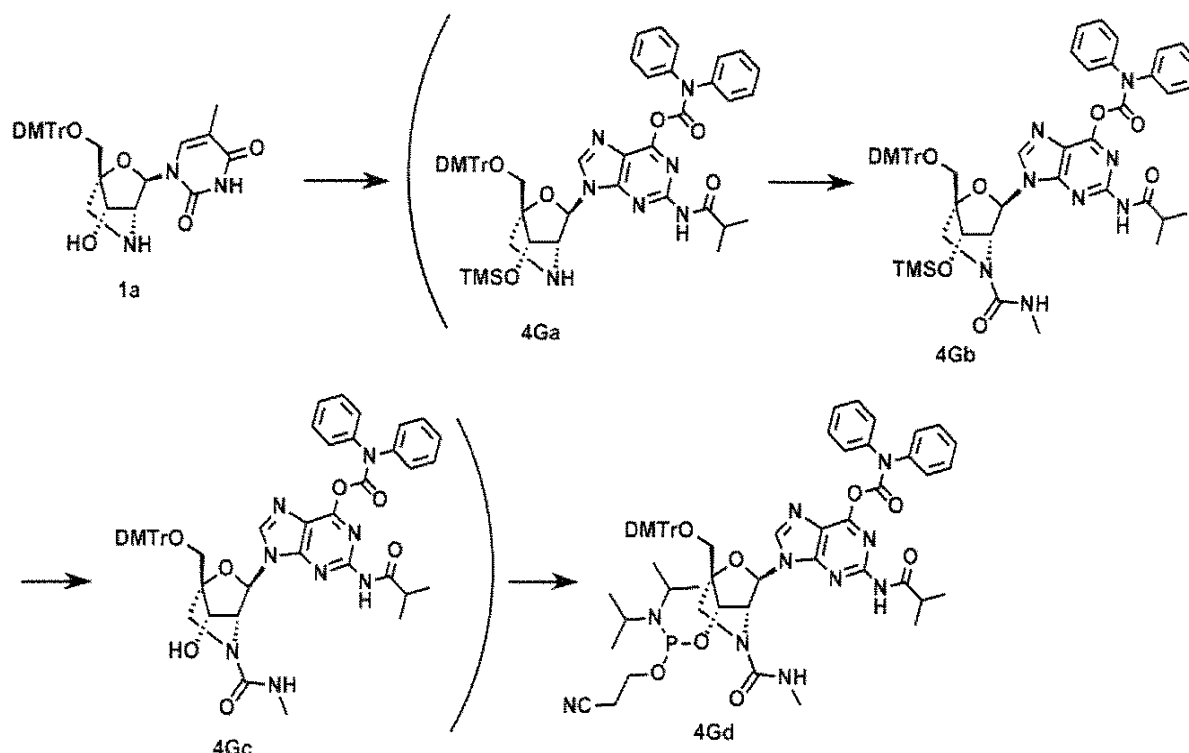
(1R,3R,4R,7S)-3-(4-benzamide-5-methyl-2-oxo-pyrimidin-1-yl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-methyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 8)

A compound 4mCd (0.96 g, yield 77 %) as the present compound 8 was synthesized from the compound 4mCc (0.98 g, 1.34 mmol) similarly to the Example 15.

MS(ESI): $m/z = 933$ (M+H)⁺

20 [0180]

Synthesis of ALNA[mU]-G



[0181]

Example 25 Synthesis of Compound 4Ga

[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
 5 phenylmethoxy)methyl]-7-trimethylsilyloxy-2-oxa-5-
 azabicyclo[2.2.1]heptan-3-yl]-2-(2-
 methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate

To a suspension of the compound 1a (30.5 g, 53.4 mmol) and [2-(2-methylpropanoylamino)-9H-purin-6-yl]N,N-diphenylcarbamate (26.7 g, 64.1 mmol) in toluene (267 mL) was added dropwise BSA (91 mL, 372 mmol) over about 20 minutes, and the mixture was warmed to 60 °C, and stirred for 5 minutes. Thereto was added further toluene (100 mL),

and the mixture was stirred at 60 °C for additional 10 minutes. Thereto was added TMSOTf (1.03 mL, 5.34 mmol) and the mixture was stirred at 60 °C for 20 minutes. The reaction solution was added to a mixture solution of IPE (100 mL),
5 saturated sodium bicarbonate water (50 mL) and water (50 mL) while stirring under ice-cooling, and the mixture was stirred for 2 hours, and insoluble materials were filtered off. After the residue was washed with Et₂O (20 mL) twice, the filtrate were separated with a separatory funnel, and the
10 organic layer was washed with water (50 mL) and saturated brine (50 mL) successively. The mixture was passed through a phase separator, concentrated under reduced pressure, and dried under vacuum to Compound 4Ga as crude product (48.67 g).

15 MS(ESI): m/z = 954 (M+H)⁺

[0182]

Example 26 Synthesis of Compound 4Gc

[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-N-methyl-2-oxa-5-
20 azabicyclo[2.2.1]heptan-3-yl]-2-(2-methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate

The compound 4Ga (17.4 g, 18.6 mmol) was solubilized by THF (100 mL), and thereto were added DIPEA (4.87 mL, 28.0 mmol), and N-methylcarbamoyl chloride (1830 mg, 19.6 mmol)
25 under ice-cooling, and the mixture was stirred for 30 minutes.

Thereto were added saturated sodium bicarbonate water (100 mL) and ethyl acetate (100 mL) successively, and the mixture was stirred at room temperature for 5 minutes. After the mixture was separated with a separatory funnel, the organic layer was washed with water (50 mL) and saturated brine (50 mL) successively, and the mixture was passed through a phase separator, and the solvent was evaporated, and dried under vacuum. The residue was added by THF (90 mL) to make a solution, and thereto was added TBAF (1M THF solution, 4.54 mL, 4.54 mmol) under ice-cooling, and the mixture was stirred under ice-cooling for 20 minutes. The mixture was diluted with ethyl acetate (100 mL), and thereto were added aqueous ammonium chloride solution (50 mL) and water (50 mL), and the mixture was stirred at room temperature for 5 minutes. The mixture was separated with a separatory funnel, and the organic layer was passed through a phase separator and the solvents were evaporated. The resulting residue was purified by silica gel chromatography (chloroform/methanol, 100/0 to 95/5) to obtain a compound 4Gc (9030 mg, yields over three steps 54%).

MS(ESI): $m/z = 919$ (M+H)⁺

[0183]

Example 27

Synthesis of Compound 4Gd

[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-

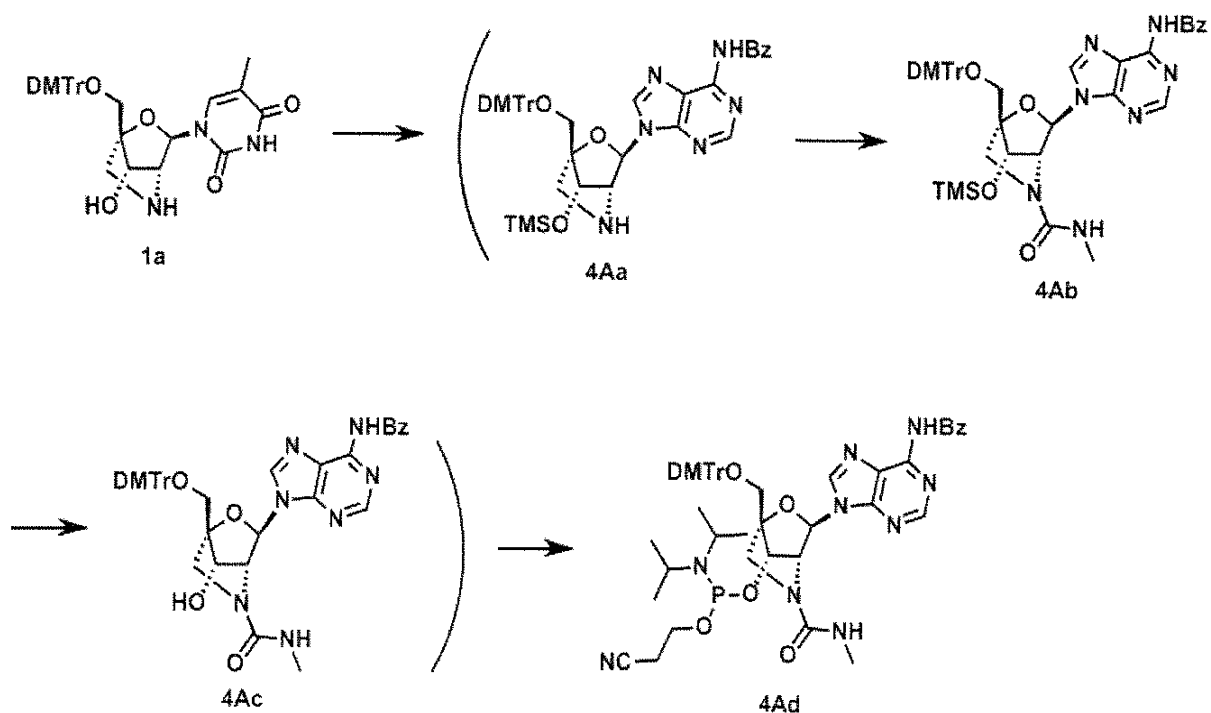
phenylmethoxy)methyl]-7-[2-cyanoethoxy-
 (diisopropylamino)phosphanyl]oxy-5-(methylcarbamoyl)-2-oxa-
 5-azabicyclo[2.2.1]heptan-3-yl]-2-(2-
 methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate
 5 (Present compound 9)

A compound 4Dc as the present compound 9 (8700 mg, yield
 79%) was synthesized from the compound 4Gc (9020 mg, 9.82
 mmol) similarly to the Example 12.

MS(ESI): $m/z = 1197$ ($M+H$)⁺

10 [0184]

Synthesis of ALNA[mU]-A



[0185]

Example 28 Synthesis of Compound 4Aa

N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-7-trimethylsilyloxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]purin-6-yl]benzamide

To a suspension of the compound 1a (30.0 g, 52.5 mmol) and N-(9H-purin-6-yl)benzamide (15.0 g, 62.7 mmol) in toluene (500 mL) was added dropwise BSA (90 mL, 368 mmol), and the mixture was warmed to 80 °C, and stirred for 20 minutes. Thereto was added TMSOTf (1.0 mL, 5.17 mmol) under ice-cooling, and the mixture was stirred at 60 °C for 20 minutes. To a mixed solution of IPE (200 mL) and saturated sodium bicarbonate water (200 mL) was added the reaction solution under ice-cooling while stirring, and the mixture was stirred for 20 minutes, and insoluble materials were filtered off. After the residues were washed with Et₂O (40 mL) twice, the filtrates were separated with a separatory funnel, and washed with saturated brine. After the organic layer was dried over sodium sulfate, insoluble materials were filtered off, and the filtrates were concentrated under reduced pressure, and dried under vacuum to obtain a compound 4Aa as a crude product (40.5 g).

MS(ESI): m/z = 757 (M+H)⁺

[0186]

Example 29 Synthesis of Compound 4Ab

(1R,3R,4R,7S)-3-(6-benzamidepurin-9-yl)-1[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-N-methyl-7-

trimethylsilyloxy-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide

The crude product 4Aa (8.20 g) was solubilized by THF (87 mL), and thereto were added DIPEA (3.00 mL, 17.2 mmol),
5 and N-methyl carbamoyl chloride (970 mg, 10.4 mmol) successively at room temperature, and the mixture was stirred for 30 minutes. Thereto were added ethyl acetate (100 mL), saturated sodium bicarbonate water (50 mL) and water (50 mL), and the mixture was separated with a separatory funnel, and
10 the organic layer was then washed with saturated brine (50 mL), and passed through a phase separator and the solvents were evaporated, and dried under vacuum to obtain a crude product 4Ab (8.70 g).

MS(ESI): $m/z = 813 (M-H)^-$

15 [0187]

Example 30 Synthesis of Compound 4Ac

(1R,3R,4R,7S)-3-(6-benzamidepurin-9-yl)-1[[bis(4-methoxyphenyl)phenylmethoxy]methyl]-7-hydroxy-N-methyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide

20 The crude product 4Ab (8.00 g) was added by THF (100 mL) to make a solution, and thereto was added TBAF (1M THF solution, 5.00 mL, 5.00 mmol) under ice-cooling, and the mixture was stirred under ice-cooling for 1 hour. The mixture was diluted with ethyl acetate (80 mL), and thereto
25 were added aqueous ammonium chloride solution (50 mL), and

water (50 mL), and the mixture was stirred at room temperature. The mixture was separated with a separatory funnel, the organic layer was washed with saturated brine (50 mL), and passed through a phase separatory, and the
5 solvents were evaporated. The resulting residue was purified by silica gel chromatography (chloroform/methanol, 100/0 to 97/3) to obtain a compound 4Ac (6100 mg, yields over three steps 84%).

MS(ESI): $m/z = 742$ (M+H)⁺

10 [0188]

Example 31 Synthesis of Compound 4Ad

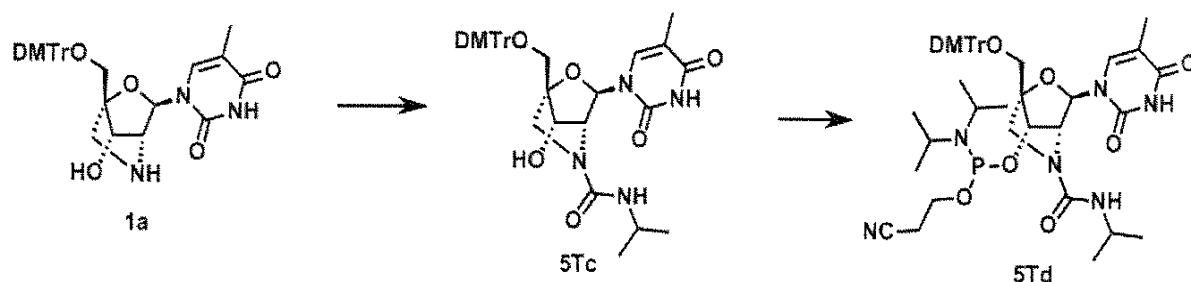
(1R,3R,4R,7S)-3-(6-benzamidepurin-9-yl)-1-[[bis(4-methoxyphenyl)phenylmethoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-methyl-2-oxa-5-
15 azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 10)

A compound 4Ad as the present compound 10 (3830 mg, yield 48%) was obtained from the compound 4Ac (6100 mg, 8.20 mmol) similarly to the Example 12.

20 MS(APCI): $m/z = 942$ (M+H)⁺

[0189]

Synthesis of ALNA[ipU]-T



[0190]

Example 32 Synthesis of Compound 5Tc

(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
 5 phenylmethoxy]methyl]-7-hydroxy-N-isopropyl-3-(5-methyl-
 2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-
 5-carboxamide

To a solution of the compound 1a (1500 mg, 2.624 mmol)
 in dichloromethane (26 mL) was added 2-isocyanato-propane
 10 (0.27 mL, 2.7 mmol) under ice-cooling, and the mixture was
 stirred at room temperature for 1 hour. Thereto were added
 a mixed solution of ethyl acetate (10 mL) and saturated
 bicarbonate solution (20 mL), and the mixture was separated
 with a separatory funnel, and the aqueous layer was then
 15 extracted with ethyl acetate 10 mL twice, and the combined
 organic layer was washed with saturated brine 10 mL. The
 organic layer was dried over anhydrous sodium sulfate, and
 filtered, and the solvents were then evaporated. The mixture
 was purified by silica gel chromatography (hexane/ethyl
 20 acetate, 50/50 to 0/100) to obtain a compound 5tc (1904 mg,

yield quant.).

MS(ESI): $m/z = 656(M-H)^-$

[0191]

Example 33 Synthesis of Compound 5Td

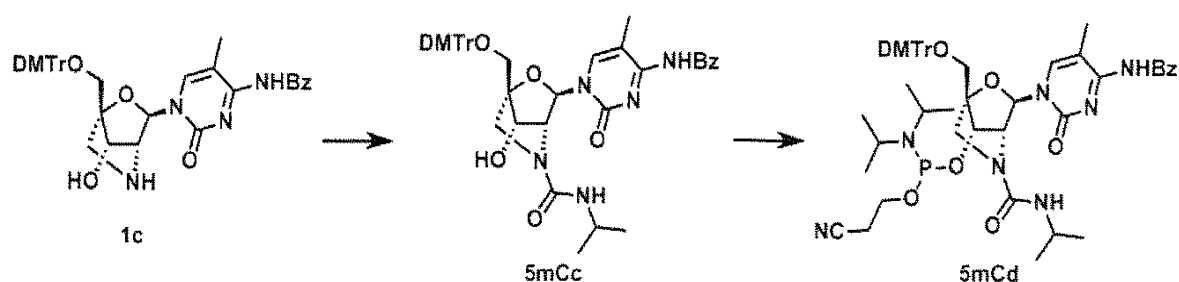
(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-isopropyl-3-(5-methyl-2,4-dioxo-pyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 11)

A compound 5td as the present compound 11 (1272.9 mg, yield 52%) was obtained from the compound 5Tc (2020 mg, 2.85 mmol) similarly to the Example 4.

MS(ESI): $m/z = 855(M-H)^-$

[0192]

15 Synthesis of ALNA[ipU]-mC



[0193]

Example 34 Synthesis of Compound 5mCc

(1R,3R,4R,7S)-3-(4-benzamide-5-methyl-2-oxo-pyrimidin-1-yl)-1-[[bis(4-methoxyphenyl)phenylmethoxy]methyl]-7-

20

hydroxy-N-isopropyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide

A compound 5mCc (1168 mg, yield 93%) was obtained from the compound 1c (1300 mg, 1.657 mmol) which was synthesized according to the method described in WO 2017/047816 A1.

MS(ESI): $m/z = 761$ (M+H)⁺

[0194]

Example 35 Synthesis of Compound 5mCd

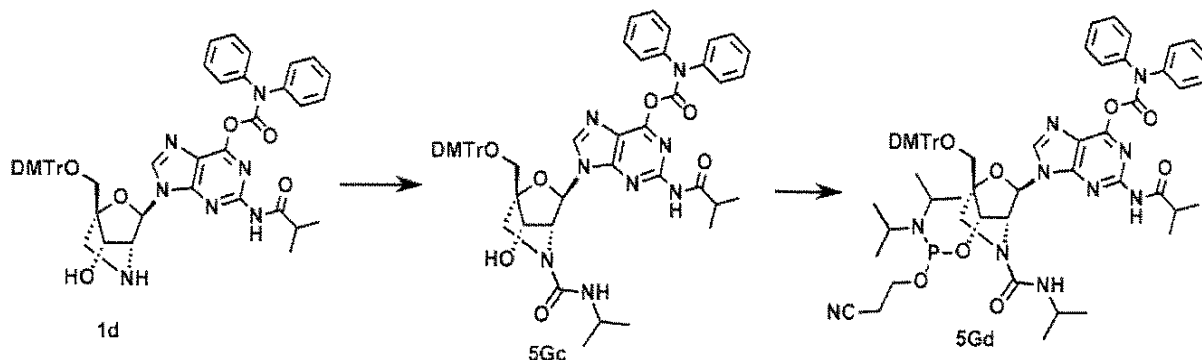
(1R,3R,4R,7S)-3-(4-benzamide-5-methyl-2-oxo-pyrimidin-1-yl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-isopropyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 12)

A compound 5mCd as the present compound 12 (1002.8 mg, yield 75%) was obtained from the compound 5mCc (1168 mg, 1.137 mmol) similarly to the Example 4.

MS(ESI): $m/z = 961$ (M+H)⁺

[0195]

Synthesis of ALNA[ipU]-G



[0196]

Example 36 Synthesis of Compound 5Gc

N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy]methyl]-7-hydroxy-5-(isopropylcarbamoyl)-2-
5 oxa-5-azabicyclo[2.2.1]heptan-3-yl]-2-(2-
methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate

To a solution of the compound 1d which was synthesized according to the method described in Working examples of WO 2017/047816 A1 (2.42 g, 2.81 mmol) in dichloromethane (28
10 mL) was added isopropyl isocyanate (0.31 mL, 3.09 mmol) under ice-cooling, and the mixture was stirred at room temperature for three hours. To the reaction solution was added saturated sodium bicarbonate water, and the mixture was stirred, passed through a phase separator, and the solvents
15 were then evaporated. The residue was purified by silica gel chromatography (hexane/ethyl acetate, 50/50 to 20/80) to obtain a compound 5Gc (1.41 g, yield 53%).

MS(ESI): m/z = 948 (M+H)⁺

[0197]

20 Example 37 Synthesis of Compound 5Gd

[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy]methyl]-7-[2-cyanoethoxy-
(diisopropylamino)phosphanyl]oxy-5-(isopropylcarbamoyl)-2-
oxa-5-azabicyclo[2.2.1]heptan-3-yl]-2-(2-
25 methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate

(Present compound 13)

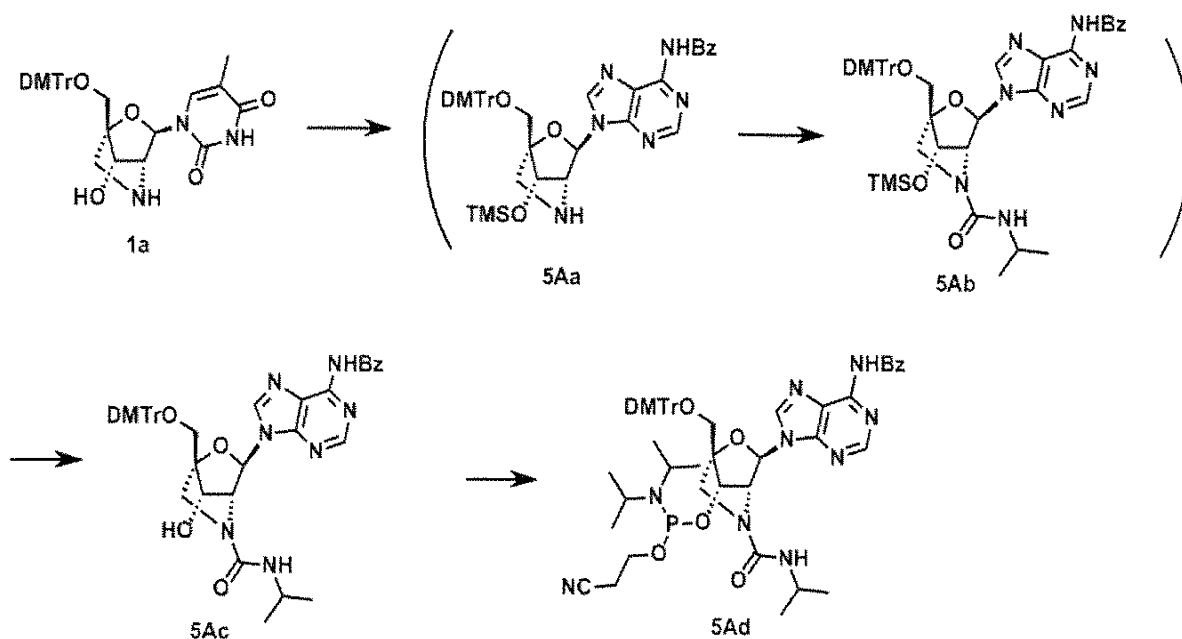
A compound 5Gd as the present compound 13 (1.20 g, yield 70%) was obtained from the compound 5Gc (1.41 g, 1.49 mmol) similarly to the Example 4.

5 MS(ESI): $m/z = 1148$ ($M+H$)⁺

³¹P-NMR (CDCl₃) δ : 149.42, 149.39

[0198]

Synthesis of ALNA[ipU]-A



10 [0199]

Example 38 Synthesis of Compound 5Ac

(1R,3R,4R,7S)-3-(6-benzamidepurin-9-yl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-hydroxy-N-isopropyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-carboxamide

15 To a suspension of the compound 1a (10.0 g, 17.5 mmol)

and N-(9H-purin-6-yl)benzamide (5.0 g, 21.0 mmol) in toluene (175 mL) was added dropwise BSA (30 mL, 122 mmol), and the mixture was warmed to 60 °C and stirred for 10 minutes, and thereto was added TMSOTf (0.34 mL, 1.8 mmol), and the mixture
5 was stirred at 60 °C for 15 minutes. The reaction solution was added to a mixed solution of the IPE (100 mL) and saturated sodium bicarbonate water (100 mL) under ice-cooling while stirring, and the mixture was stirred for 20 minutes, and insoluble materials were filtered off. After
10 the filtered residue was washed with Et₂O (10 mL), the filtrates were separated with a separatory funnel, and washed with saturated brine. The organic layer was dried over sodium sulfate, and insoluble materials were then filtered off, and concentrated under reduced pressure. The residue
15 was solubilized by dichloromethane (162 mL), and thereto was added isopropyl isocyanate (1.8 mL, 18 mmol) at room temperature, and the mixture was stirred for 60 minutes. Thereto were added ethyl acetate (100 mL), saturated sodium bicarbonate water (50 mL) and water (50 mL), and the mixture
20 was separated with a separatory funnel, and the organic layer was washed with saturated brine (50 mL), dried over sodium sulfate, filtered, and the solvents were then evaporated. The residue was added by THF (162 mL) to make a solution, and thereto was added TBAF (1M THF solution, 2.39 mL, 8.11
25 mmol) under ice-cooling, and the mixture was stirred at room

temperature for 24 hours. The mixture was diluted with ethyl acetate (80 mL), and thereto were added aqueous ammonium chloride solution (50 mL) and water (50 mL), and the mixture was stirred at room temperature. The mixture was separated with a separatory funnel, and the organic layer was washed with saturated brine (50 mL), and then dried over sodium sulfate, filtered, and solvents were evaporated. The resulting residue was purified by silica gel chromatography (hexane/ethyl acetate, 50/50 to 0/100) to obtain a compound 5Ac (10.1 g, yields over three steps 77%).

MS(ESI): $m/z = 770$ (M+H)⁺

[0200]

Example 39 Synthesis of Compound 5Ad

(1R,3R,4R,7S)-3-(6-benzamidepurin-9-yl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-isopropyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 14)

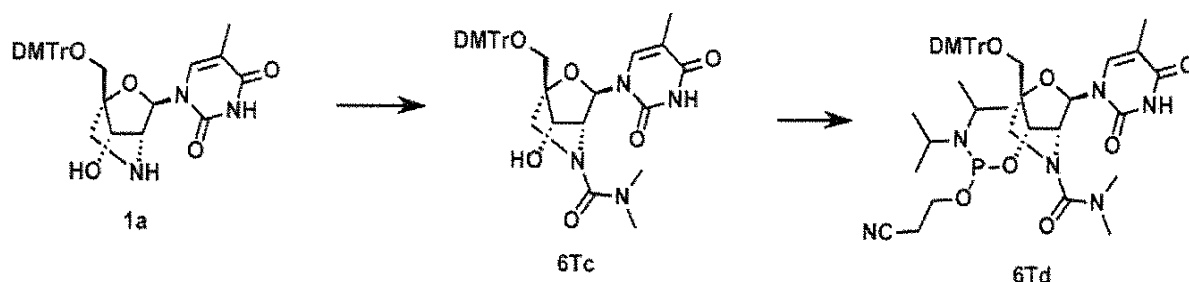
A compound 5d as the present compound 14 (4.6 g, yield 51%) was obtained from the compound 5c (7.0 g, 9.09 mmol) similarly to the Example 4.

MS(ESI): $m/z = 970$ (M+H)⁺

³¹P-NMR (CDCl₃) δ : 149.4

[0201]

Synthesis of ALNA[dmU]-T



[0202]

Example 40 Synthesis of Compound 6Tc

(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
 5 phenylmethoxy)methyl]-7-hydroxy-N,N-dimethyl-3-(5-methyl-
 2,4-dioxo-pyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-
 5-carboxamide

To a solution of the compound 1a (1512 mg, 2.645 mmol)
 in tetrahydrofuran (26 mL) was added N,N-
 10 diisopropylethylamine (0.483 mL, 2.773 mmol) and the mixture
 was cooled under ice-cooling, and thereto was added N,N-
 dimethyl carbamoyl chloride (0.255 mL, 2.77 mmol). The
 mixture was warmed to room temperature, and thereto was then
 added N,N-dimethyl formamide (1 mL, 12.9 mmol), and the
 15 mixture was stirred for 4 hours. Thereto was added a mixed
 solution of ethyl acetate (10 mL) and saturated bicarbonate
 water (20 mL), and the mixture was stirred, and the mixture
 was separated with a separatory funnel, and the aqueous layer
 was then extracted with ethyl acetate 10 mL twice, and the
 20 organic layers were combined, and the combined organic layers

were then washed with saturated brine 10 mL. The organic layer was dried over anhydrous sodium sulfate, filtered, and the solvents were evaporated. The mixture was purified by silica gel chromatography (hexane/ethyl acetate, 50/50 to 0/100) to obtain a compound 6Tc1 (1705 mg, yield quant.).

MS(ESI): $m/z = 642 (M-H)^-$

[0203]

Example 41 Synthesis of compound 6Td

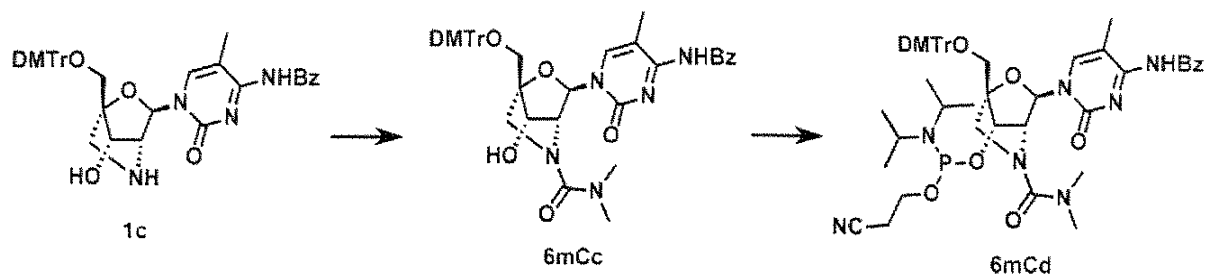
(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N,N-dimethyl-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 15)

A compound 6Td as the present compound 15 (1565 mg, yield 68%) was obtained from the compound 6Tc (1758 mg, 2.735 mmol) similarly to the Example 4.

MS(ESI): $m/z = 842 (M-H)^-$

[0204]

Synthesis of ALNA[dmU]-mC



[0205]

Exmample 42 Synthesis of Compound 6mCc

(1R,3R,4R,7S)-3-(4-benzamide-5-methyl-2-oxo-pyrimidin-1-yl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-hydroxy-N,N-dimethyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide

To a solution of the compound 1c (1300 mg, 1.657 mmol) in tetrahydrofuran (16.6 mL) was added N,N-diisopropyl ethyl amine (0.317 mL, 1.820 mmol), and the mixture was cooled under ice-cooling, and thereto was added N,N-dimethyl carbamoyl chloride (0.168 mL, 1.83 mmol), and the mixture was stirred at room temperature for 10 minutes. After the mixture was heated under reflux for 2 hours, thereto was added a mixed solution of ethyl acetate (10 mL) and saturated sodium bicarbonate water (20 mL) at room temperature, and the mixture was separated with a separatory funnel, the aqueous layer was extracted with ethyl acetate 10 mL twice, and the combined organic layers were washed with saturated brine 10 mL. The organic layer was dried over anhydrous sodium sulfate, and filtered, and the solvents were then evaporated. The mixture was purified by silica gel chromatography (hexane/ethyl acetate, 70/30 to 30/70) to obtain a compound 6mCc (1705 mg, yield 79%).

MS(ESI): $m/z = 747$ (M+H)⁺

[0206]

Example 43 Synthesis of Compound 6mCd

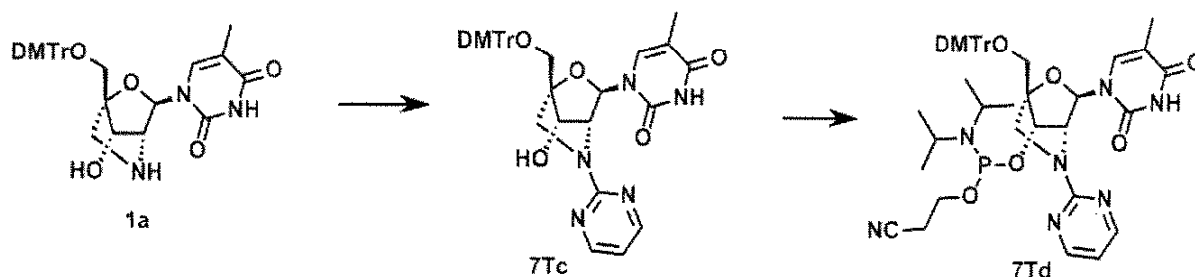
(1R,3R,4R,7S)-3-(4-benzamide-5-methyl-2-oxo-pyrimidin-1-yl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N,N-dimethyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide (Present compound 16)

A compound 6mCd as the present compound 16 (936.5 mg, yield 76%) was obtained from the compound 6mCc (1019 mg, 1.312 mmol) similarly to the Example 4.

MS(ESI): $m/z = 947$ (M+H)⁺

[0207]

Synthesis of ALNA[2Pym]-T



[0208]

Example 44 Synthesis of Compound 7Tc

1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-7-hydroxy-5-pyrimidin-2-yl]-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2,4-dione

A mixture of the compound 1a (2.00 g, 3.50 mmol), 2-chloropyrimidine (1.20 g, 10.5 mmol), DIPEA (3.05 mL, 17.5

mmol), and DMSO (17.5 mL) was stirred at 130 °C under nitrogen atmosphere for 6 hours. Ethyl acetate was added to the reaction solution, and the mixture was washed with water and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and insoluble materials were filtered off, and the filtrates were concentrated under reduced pressure. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 90/10) to obtain a compound 7Tc (1.74 g, yield 77%).

HRMS(MALDI): calculated value as C₃₆H₃₅N₅O₇ [M+Na]⁺: 672.2429, measured value: 672.2427

[0209]

Example 45 Synthesis of Compound 7Td

3-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-3-(5-methyl-2,4-dioxo-pyrimidin-1-yl)-5-pyrimidin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 17)

To a mixture of the compound 7Tc (1.73 g, 2.66 mmol), dichloromethane (13 mL), and DIPEA (1.39 mL, 7.98 mmol) was added 2-cyanoethyl N,N-diisopropyl chloro phosphoroamidite (1.19 mL, 5.32 mmol) under ice-cooling, and the mixture was stirred at room temperature under nitrogen atmosphere overnight. A saturated aqueous sodium hydrogen carbonate solution was added to the reaction solution, and the mixture

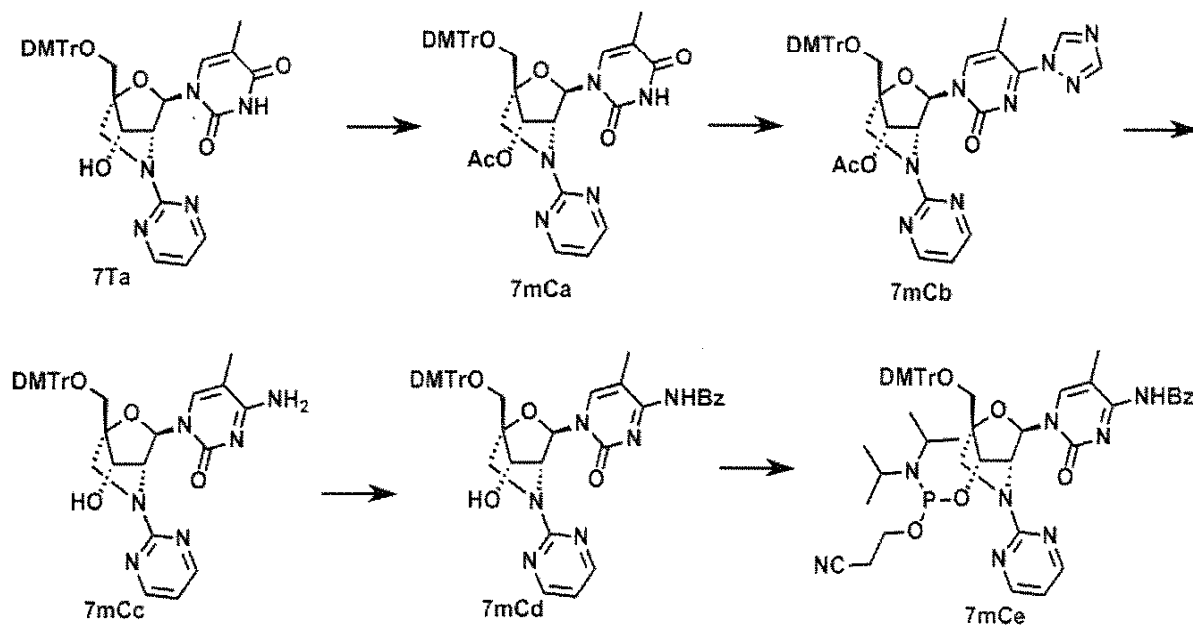
was extracted with chloroform. The organic layer was passed through a phase separator, and evaporated under reduced pressure. The residue was purified by column chromatography (SiO₂: hexane/ethyl acetate = 60/40 to 10/90, Diol: hexane/ethyl acetate = 70/30 to 20/80, NH: chloroform/methanol = 100/0 to 95/5) to obtain a compound 7Td as the present compound 17 (1.53 g, yield 68%).

HRMS(FAB): calculated value as C₄₅H₅₂N₇O₉P [M+H]⁺: 850.3693, measured value: 850.3699

³¹P-NMR (CDCl₃) δ: 148.72, 148.83

[0210]

Synthesis of ALNA[2Pym]-mC



[0211]

Example 46 Synthesis of Compound 7mCa

[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-3-(5-methyl-2,4-dioxo-pyrimidin-1-yl)-5-pyrimidin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]acetate

5 To a mixture of the compound 7Ta (3.68 g, 5.67 mmol), DMAP (69.3 mg, 0.567 mmol), and pyridine (28 mL) was added acetic anhydride (0.804 mL, 8.51 mmol), and the mixture was stirred at room temperature overnight. Ethyl acetate was added to the reaction solution, and the mixture was washed
10 with saturated aqueous sodium hydrogen carbonate solution, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and insoluble materials are filtered off, and concentrated under reduced pressure. Pyridine was azeotroped with ethyl acetate, and the mixture
15 was triturated with diisopropyl ether to obtain the compound 7mCa (3.49 g, yield 89%).

HRMS(FAB): calculated value as C₃₉H₃₈N₅O₈ [M+H]⁺: 692.2720, measured value: 692.2713

[0212]

20 Example 47 Synthesis of Compound 7mCb

[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-3-[5-methyl-2-oxo-4-(1,2,4-triazol-1-yl)pyrimidin-1-yl]-5-pyrimidin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]acetate

25 To a mixture of the compound 7mCa (3.46 g, 5.00 mmol),

1,2,4-triazole (3.11 g, 45.0 mmol), DIPEA (8.71 mL, 50.0 mmol) and acetonitrile (50 mL) was added phosphoryl chloride (0.792 mL, 8.50 mmol) under ice-cooling, and the mixture was stirred at room temperature under nitrogen atmosphere overnight. Ethyl acetate was added to the reaction solution, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and insoluble materials were filtered off, and the filtrates were concentrated under reduced pressure. The mixture was triturated with diisopropyl ether to obtain a compound 7mCb1 (3.40 g, yield 92%).

HRMS(FAB): calculated value as $C_{40}H_{38}N_8O_7$ $[M+H]^+$: 743.2942, measured value: 743.2945

[0213]

Example 48 Synthesis of Compound 7mCc

4-amino-1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-7-hydroxy-5-pyrimidin-2-yl]-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2-one

To a mixture of the compound 7mCb (3.40 g, 4.58 mmol) and acetonitrile (46 mL) was added 28% ammonia water (31 mL), and the mixture was stirred at room temperature for 3 days. Water was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was dried over anhydrous magnesium sulfate, and insoluble materials

were filtered off, and concentrated under reduced pressure. The mixture was triturated with diisopropyl ether and small amounts of ethyl acetate to obtain a compound 7mCc (2.65 g, yield 89%).

5 HRMS(FAB): calculated value as $C_{36}H_{36}N_6O_6$ $[M+H]^+$: 649.2775, measured value: 649.2766

[0214]

Example 49 Synthesis of Compound 7mCd

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
10 phenylmethoxy)methyl]-7-hydroxy-5-pyrimidin-2-yl]-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide

To a mixture of the compound 7mCc (2.62 g, 4.04 mmol), and DMF (20 mL) was added benzoic anhydride (959 mg, 4.24
15 mmol), and the mixture was stirred at room temperature under nitrogen atmosphere overnight. Ethyl acetate was added to the reaction solution, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, water, and saturated brine. The organic layer was dried over
20 anhydrous magnesium sulfate, insoluble materials were filtered off, and concentrated under reduced pressure. The residue was purified by column chromatography (SiO_2 : chloroform/methanol = 100/0 to 90/10) to obtain the compound 7mCd (1.42 g, yield 47%).

25 HRMS(FAB): calculated value as $C_{43}H_{40}N_6O_7$ $[M+H]^+$: 753.3037,

measured value: 753.3035

[0215]

Example 50 Synthesis of Compound 7mCe

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-

5 phenylmethoxy)methyl]-7-[2-cyanoethoxy-

(diisopropylamino)phosphanyl]oxy-5-pyrimidin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 18)

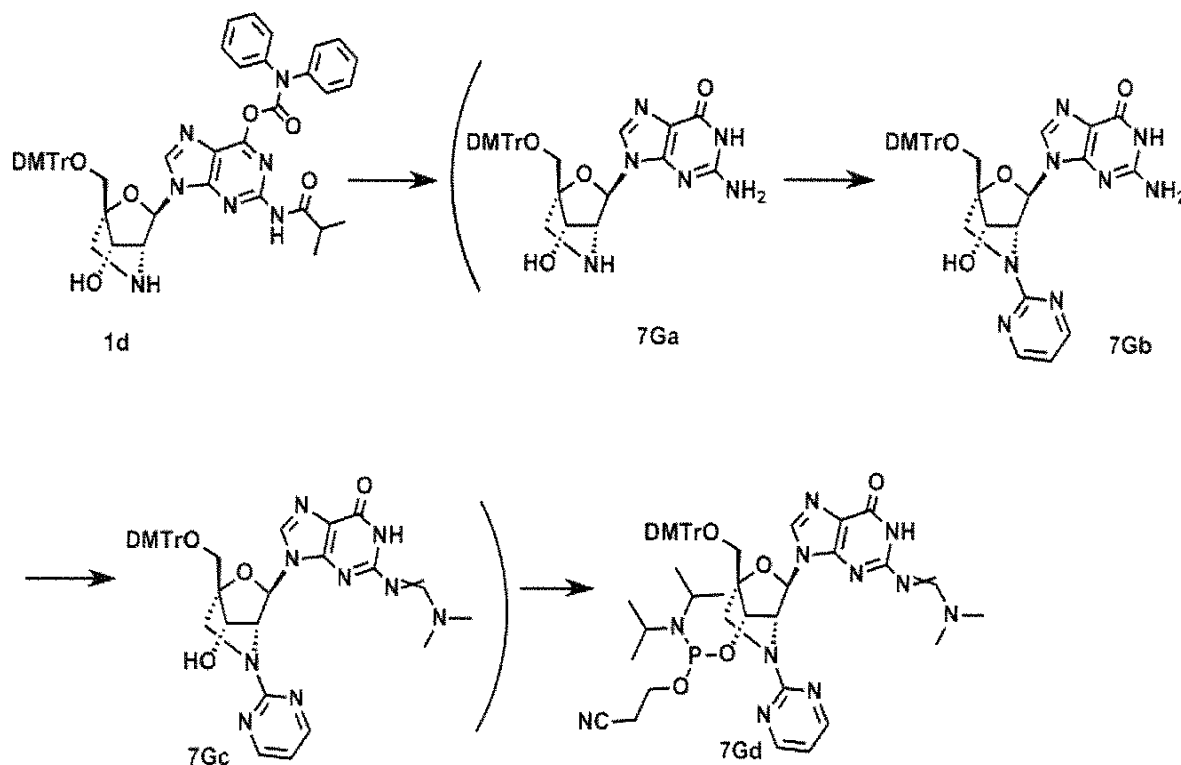
To a mixture of the compound 7mCd (1.40 g, 1.86 mmol),
10 dichloromethane (9 mL), and DIPEA (0.972 mL, 5.58 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (0.830 mL, 3.72 mmol), and the mixture was stirred at room temperature under nitrogen atmosphere overnight. To reaction solution was added saturated aqueous sodium
15 hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and the solvents were evaporated. The residue was purified by column chromatography (SiO₂: hexane/ethyl acetate = 80/20 to 30/70, NH: hexane/ethyl
20 acetate = 50/50 to 0/100) to obtain a compound 7mCe as the present compound 18 (1.01 g, yield 57%).

HRMS(FAB): calculated value as C₅₂H₅₇N₈O₈P [M+Na]⁺: 975.3935,
measured value: 975.3931

³¹P-NMR (CDCl₃) δ: 149.17, 149.23

25 [0216]

Synthesis of ALNA[2Pym]-G



[0217]

Example 51 Synthesis of Compound 7Gc

5 N'-[9-(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-7-hydroxy-5-pyrimidin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-N,N-dimethyl-formamidine

To a solution of the compound **1d** (3.00 g, 3.50 mmol) in
 10 methanol (20 mL) and THF (20 mL) was added 28% ammonia water (40 mL). Thereto were added methanol (20 mL) and THF (20 mL), and the mixture was allowed to stand at room temperature for 16 hours. Thereto was a solution of 40% methyl amine in

methanol (40 mL), and the mixture was allowed to stand at room temperature for 4 hours. After solvents were evaporated, the residue was azeotroped with toluene, and dried under reduced pressure. The residue was solubilized with DMSO (35 mL), and thereto were added DIPEA (4.8 mL, 28.0 mmol), and 2-fluoropyrimidine (0.66 mL, 10.5 mmol), and the mixture was stirred at 80 °C for 2 hours, and then stirred at 100 °C for additional 2 hours. Thereto was added 2-fluoropyrimidine (0.22 mL, 3.5 mmol) and DIPEA (1.6 mL, 9.3 mmol) additionally, and the mixture was stirred at 100 °C for 2 hours. To the reaction solution was N,N-dimethyl formamide dimethyl acetal (2.3 mL, 17.0 mmol) and the mixture was stirred at room temperature. After 1 hour, thereto were added ethyl acetate (200 mL), saturated sodium bicarbonate water (20 mL) and water (80 mL), and the mixture was separated with a separatory funnel. The organic layer was washed with water (50 mL) three times, and then washed with saturated brine. After the mixture was dried over sodium sulfate, insoluble materials were filtered off, and solvents were evaporated under reduced pressure. The aqueous layer at the separation with a separatory funnel was further extracted with chloroform (100 mL) three times, followed by washing with water (100 mL) and saturated brine. The chloroform layer was dried over sodium sulfate, and insoluble materials were then filtered off, and solvents were evaporated from the

filtrates under reduced pressure, and the residue was combined with the above extracts, and the mixture was purified by silica gel chromatography (chloroform/methanol, 100/0 to 92/8) to obtain a compound 7Gc (1.66 g, yields over
 5 three steps 65 %).

MS(ESI): $m/z = 731$ (M+H)⁺

[0218]

Example 52 Synthesis of Compound 7Gd

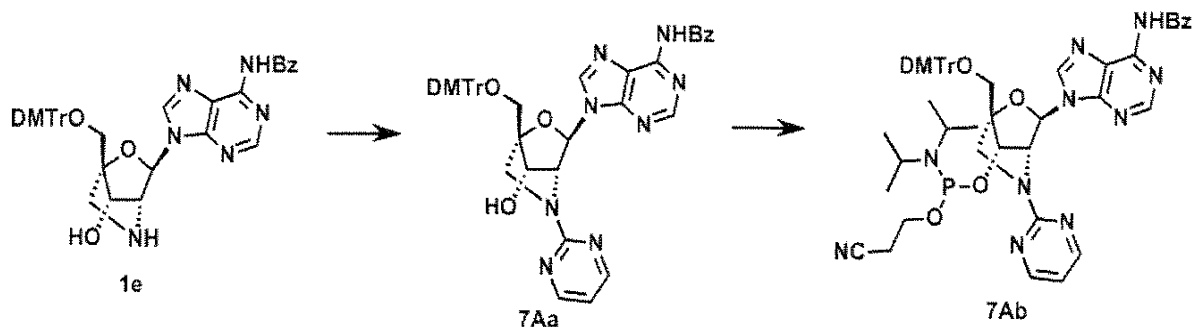
N'-[9-(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
 10 phenylmethoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-pyrimidin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-N,N-dimethyl-6-formamidine (Present compound 19)

A compound 7Gd as the present compound 19 (1.07 g, yield
 15 53%) was obtained from the compound 7Gc (1.58 g, 2.16 mmol) similarly to the Example 4.

MS(ESI): $m/z = 931$ (M+H)⁺

[0219]

Synthesis of ALNA[2Pym]-A



[0220]

Example 53 Synthesis of Compound 7Aa

N-[9-(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy)methyl]-7-hydroxy-5-pyrimidin-2-yl-2-oxa-5-
5 azabicyclo[2.2.1]heptan-3-yl]purin-6-yl]benzamide

A mixture of the compound 1e which was synthesized according to the method described in WO 2017/047816 A1 (500 mg, 0.730 mmol), 2-fluoropyrimidine (215 mg, 2.19 mmol), DIPEA (0.509 mL, 2.92 mmol) and DMSO (7 mL) was stirred at
10 130 °C under nitrogen atmosphere for 7 hours. Ethyl acetate was added to the reaction solution, and the mixture was washed with water and saturated brine. The organic layer was dried over anhydrous sodium sulfate, and insoluble materials were filtered off, and the filtrates were
15 concentrated by evaporating solvents under reduced pressure. The residue was purified by column chromatography (SiO₂: hexane/ethyl acetate = 50/50 to 0/100) to obtain a compound 7Aa (207 mg, yield 37 %).

HRMS(FAB): calculated value as C₄₃H₃₈N₈O₆ [M+Na]⁺: 785.2807,
20 measured value: 785.2807

[0221]

Example 54 Synthesis of Compound 7Ab

N'-[9-(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy)methyl]-7-[2-cyanoethoxy-
25 (diisopropylamino)phosphanyl]oxy-5-pyrimidin-2-yl-2-oxa-5-

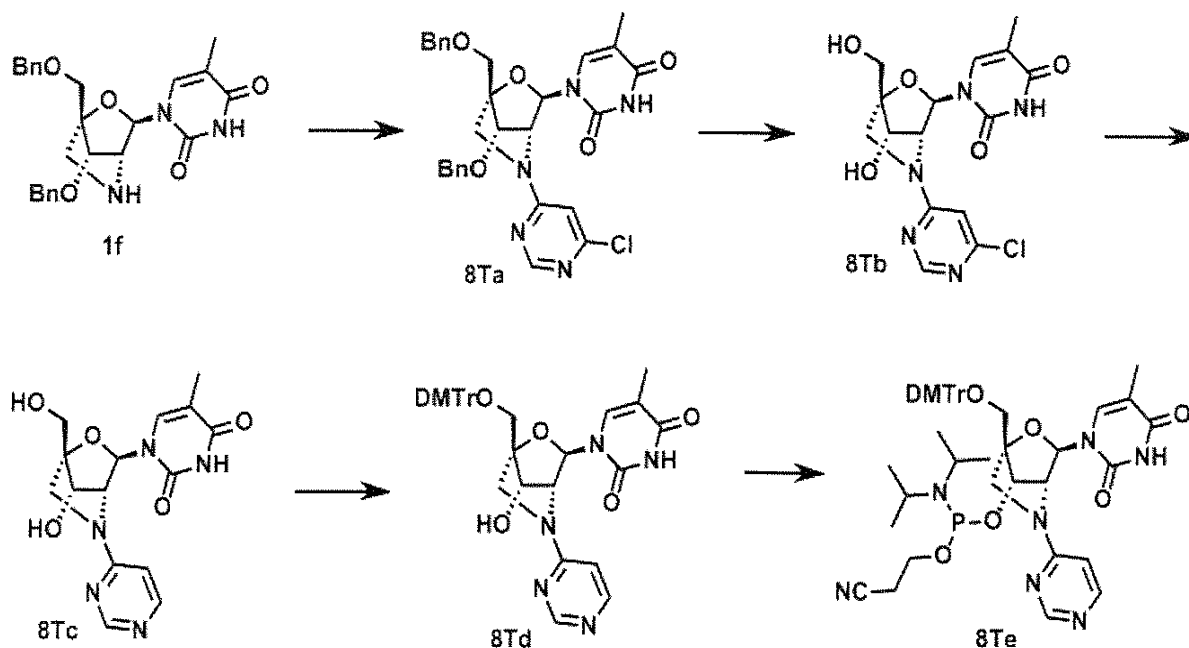
azabicyclo[2.2.1]heptan-3-yl]purin-6-yl]benzamide (Present compound 20)

To a mixture of the compound 7Aa (126 mg, 0.165 mmol), dichloromethane (3 mL) and DIPEA (86.2 μ L, 0.495 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (73.6 μ L, 0.330 mmol), and the mixture was stirred at room temperature under nitrogen atmosphere for 5 hours. Saturated aqueous sodium hydrogen carbonate solution was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and the solvent was evaporated. The residue was purified by column chromatography (NH: hexane/ethyl acetate = 50/50 to 0/100, SiO₂: hexane/ethyl acetate = 50/50 to 0/100, diol: hexane/ethyl acetate = 60/40 to 10/90) to obtain a compound 7Ab as the present compound 20 (110 mg, yield 69%). HRMS(FAB): calculated value as C₅₂H₅₅N₁₀O₇P [M+Na]⁺: 985.3885, measured value: 985.3878

³¹P-NMR (CDCl₃) δ : 149.33, 149.47

[0222]

Synthesis of ALNA[4Pym]-T



[0223]

Example 55 Synthesis of Compound 8Ta

1-[(1R,3R,4R,7S)-7-benzyloxy-1-(benzyloxy-methyl)-5-(6-chloropyrimidin-4-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-pyrimidin-2,4-dione

A mixture of the compound 1f which was synthesized according to the method described in WO 2017/047816 A1 (1.00 g, 2.22 mmol), 4,6-dichloropyrimidine (398 mg, 2.66 mmol), DIPEA (1.16 mL, 6.66 mmol) and EtOH (20 mL) (2 batches) was stirred at 120 °C under microwave irradiation for 5 hours for each batch. The batches were combined and insoluble materials of the reaction solutions were collected by filtration, washed with EtOH, and dried under reduced pressure to obtain a compound 8Ta (1.54 g, yield 62%).

HRMS(FAB): calculated value as $C_{29}H_{28}ClN_5O_5$ $[M+H]^+$: 562.1857,

measured value: 562.1859

[0224]

Example 56 Synthesis of Compound 8Tb

1-[(1R,3R,4R,7S)-5-(6-chloropyrimidin-4-yl)-7-hydroxy1-
5 (hydroxymethyl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-
methyl-pyrimidin-2,4-dione

To a mixture of the compound 8Ta (3.49 g, 6.21 mmol)
and dichloromethane (62 mL) was added trichloroborane (1.0
mol/L dichloromethane solution, 62 mL, 62.1 mmol) at -78 °C
10 under nitrogen atmosphere, and the mixture was warmed to
room temperature gradually overnight. MeOH (62 mL) was added
to the reaction solution slowly under ice-cooling, and the
mixture was stirred for a while. The reaction solution was
concentrated, and the residue was then triturated with ethyl
15 acetate to obtain a compound 8Tb (3.28 g) as a crude product.
HRMS(FAB): calculated value as C₁₅H₁₆ClN₅O₅ [M+H]⁺: 382.0918,
measured value: 382.0919

[0225]

Example 57 Synthesis of Compound 8Tc

20 1-[(1R,3R,4R,7S)-7-hydroxy-1-(hydroxymethyl)-5-pyrimidin-4-
yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-
pyrimidin-2,4-dione

To a mixture of the compound 8Tb (3.28 g, 6.21 mmol,
crude product) and MeOH (62 mL) was added 20% palladium
25 hydroxide (654 mg, 20 wt%), and the mixture was stirred at

room temperature under nitrogen atmosphere overnight. Water (31 mL) was added to the reaction solution to dissolve insoluble materials, and palladium was filtered through Celite. The filtrates were concentrated, and water was azeotroped with toluene. The residue was triturated with MeOH to obtain a compound 8Tc (1.75 g, yields over two steps 81%).

HRMS(FAB): calculated value as $C_{15}H_{17}N_5O_5$ $[M+H]^+$: 348.1302, measured value: 348.1308

[0226]

Example 58 Synthesis of Compound 8Td

1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-hydroxy-5-pyrimidin-4-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2,4-dione

To a mixture of the compound 8Tc (444 mg, 1.28 mmol) and pyridine (13 mL) was added 4,4'-dimethoxy trityl chloride (520 mg, 1.54 mmol), and the mixture was stirred at room temperature overnight. Saturated aqueous sodium hydrogen carbonate solution was added dropwise to the reaction solution under ice-cooling, and thereto was added ethyl acetate. The organic layer was washed with saturated aqueous sodium hydrogen carbonate solution, water and saturated brine. The organic layer was dried over anhydrous sodium sulfate, and concentrated. Pyridine was azeotroped with ethyl acetate, and the residue was triturated with

diisopropyl ether to obtain a compound 8Td (652 mg, yield 78%).

HRMS(FAB): calculated value as $C_{36}H_{35}N_5O_7$ $[M+H]^+$: 650.2609, measured value: 650.2600

5 [0227]

Example 59 Synthesis of Compound 8Te

3-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-pyrimidin-4-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 21)

10

To a mixture of the compound 8Td (300 mg, 0.462 mmol), dichloromethane (5 mL) and DIPEA (322 μ L, 1.85 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (309 μ L, 1.39 mmol) under ice-cooling, and the mixture was stirred at room temperature under nitrogen atmosphere overnight. Saturated aqueous sodium hydrogen carbonate solution was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and solvents were evaporated. The residue was purified by column chromatography (SiO_2 : chloroform/methanol = 100/0 to 90/10, diol: hexane/ethyl acetate= 50/50 to 0/100) to obtain a compound 8Te as the present compound 21 (263 mg, yield 67%).

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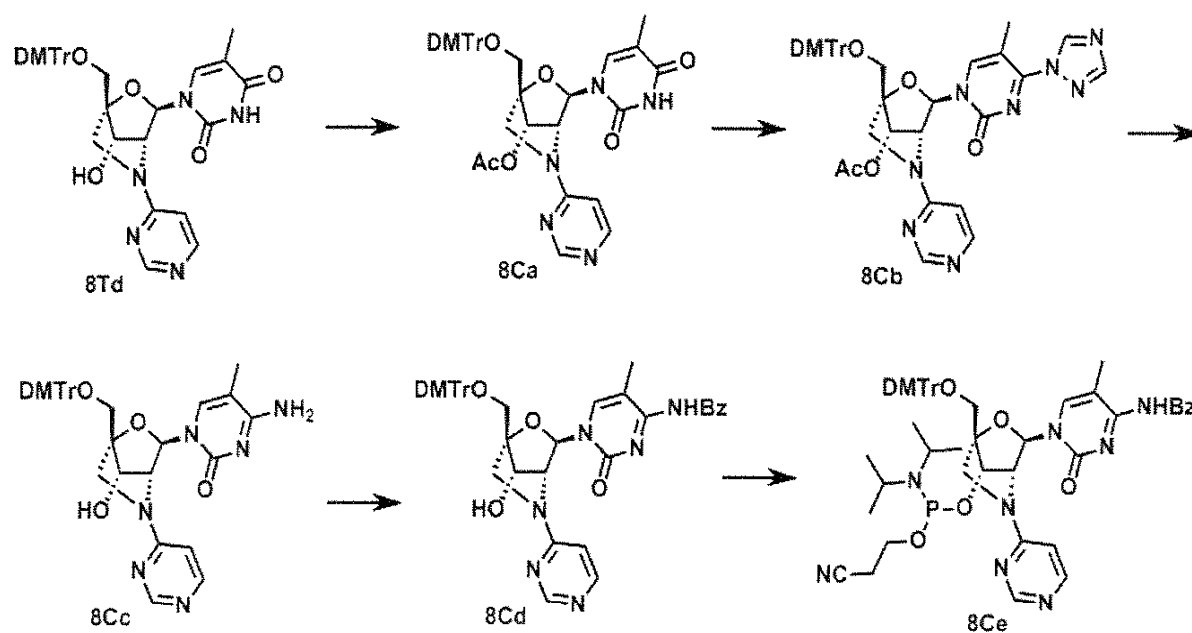
25 HRMS(FAB): calculated value as $C_{45}H_{52}N_7O_8$ $[M+H]^+$: 850.3693,

measured value: 850.3685

^{31}P -NMR (CDCl_3) δ : 148.93, 149.47

[0228]

Synthesis of ALNA[4Pym]-mC



5

[0229]

Example 60 Synthesis of Compound 8Ca

[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-pyrimidin-4-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]acetate

To a mixture of the compound 8Td (234 mg, 0.360 mmol), DMAP (4.4 mg, 0.0360 mmol) and pyridine (3.6 mL) was added acetic anhydride (51.1 μL , 0.540 mmol), and the mixture was stirred at room temperature overnight. Ethyl acetate was

15

added to the reaction solution, and the mixture was washed with 5% aqueous copper sulfate solution, water and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and concentrated. The residue was triturated with diisopropyl ether to obtain a compound 8Ca (219 mg, yield 88%).

HRMS(FAB): calculated value as $C_{38}H_{27}N_5O_8$ $[M+H]^+$: 692.2720, measured value: 692.2720

[0230]

10 Example 61 Synthesis of Compound 8Cb

[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-3-[5-methyl-2-oxo-4-(1,2,4-triazol-1-yl)pyrimidin-1-yl]-5-pyrimidin-4-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]acetate

15 To a mixture of the compound 8Ca (201 mg, 0.291 mmol), 1,2,4-triazole (181 mg, 2.62 mmol), DIPEA (507 μ L, 2.91 mmol) and acetonitrile (3 mL) was added phosphoryl chloride (46.1 μ L, 0.494 mmol) under ice-cooling, and the mixture was stirred at room temperature under nitrogen atmosphere overnight. Ethyl acetate was added to the reaction solution, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and concentrated. The mixture was triturated with diisopropyl ether to obtain a compound 8Cb (211 mg yield

20

25

98%).

HRMS(FAB): calculated value as $C_{40}H_{38}N_8O_7$ $[M+H]^+$: 743.2942,
measured value: 743.2939

[0231]

5 Example 62 Synthesis of Compound 8Cc

4-amino-1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy)methyl]-7-hydroxy-5-pyrimidin-4-yl]-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2-one

To a mixture of the compound 8Cb (613 mg, 0.825 mmol)
10 and acetonitrile (8.3 mL) was added 28% ammonia water (5.5
mL), and the mixture was stirred at room temperature
overnight. Acetonitrile of the reaction solution was
concentrated. Insoluble materials were collected by
filtration, and washed with water. The crude product was
15 triturated with ethyl acetate to obtain a compound 8Cc (380
mg, yield 71%).

HRMS(FAB): calculated value as $C_{36}H_{36}N_6O_6$ $[M+H]^+$: 649.2775,
measured value: 649.2773

[0232]

20 Example 63 Synthesis of Compound 8Cd

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy)methyl]-7-hydroxy-5-pyrimidin-4-yl]-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-
yl]benzamide

25 To a mixture of the compound 8Cc (378 mg, 0.583 mmol),

and pyridine (6 mL) was added benzoic anhydride (198 mg, 0.875 mmol), and the mixture was stirred at room temperature for 3 days. The reaction solution was concentrated, and the residue was purified by column chromatography (diol: 0.5% triethylaminechloroform/methanol = 100/0 to 90/10, SiO₂: 0.55% triethylaminechloroform/methanol = 100/0 to 90/10) to obtain a compound 8Cd (247 mg, yield 56%).

HRMS (FAB): calculated value as C₄₃H₄₀N₆O₇ [M+H]⁺: 753.3037, measured value: 753.3040

[0233]

Example 64 Synthesis of Compound 8Ce

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-pyrimidin-4-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 22)

To a mixture of the compound 8Cd (135 mg, 0.179 mmol), dichloromethane (2 mL), and DIPEA (125 µL, 0.716 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (120 µL, 0.537 mmol), and the mixture was stirred at room temperature under nitrogen atmosphere for 2 days. Saturated aqueous sodium hydrogen carbonate solution was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and the solvents were evaporated. The residue

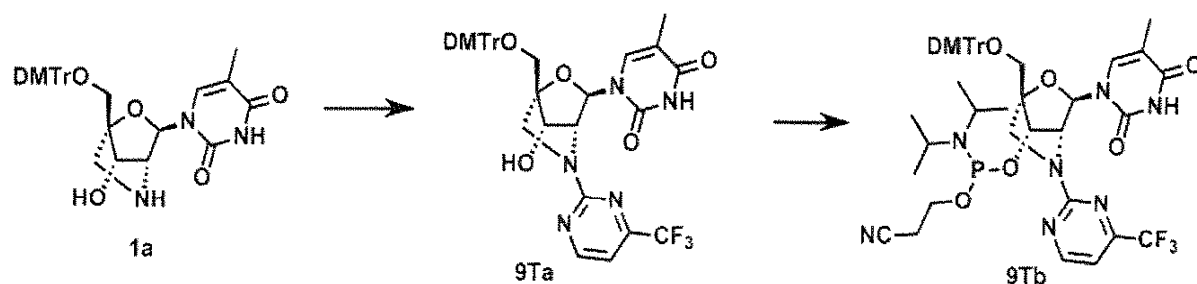
was purified by column chromatography (SiO₂: hexane/ethyl acetate= 50/50 to 0/100, diol: hexane/ethyl acetate= 60/40 to 10/90, NH: hexane/ethyl acetate = 50/50 to 0/100) to obtain a compound 8Ce as the present compound 22 (61.9 mg, yield 36%).

HRMS(FAB): calculated value as C₅₂H₅₇N₈O₈ [M+H]⁺: 953.4115, measured value: 953.4109

³¹P-NMR (CDCl₃) δ: 149.65, 150.24

[0234]

10 Synthesis of ALNA[4-CF₃-2Pym]-T



[0235]

Example 65 Synthesis of Compound 9Ta

1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-hydroxy-5-[4-(trifluoromethyl)pyrimidin-2-yl]-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2,4-dione

A mixture of the compound 1a (500 mg, 0.875 mmol), 2-chloro-4-(trifluoromethyl)pyrimidine (239 mg, 1.31 mmol), DIPEA (457 μL, 2.63 mmol) and EtOH (4.4 mL) was stirred at

120 °C under microwave irradiation for 30 minutes. 2-Chloro-4-(trifluoromethyl)pyrimidine (239 mg, 1.31 mmol) and DIPEA (457 µL, 2.63 mmol) were added additionally to the reaction solution, and the mixture was stirred at 120 °C under microwave irradiation for 1 hour. The reaction solution was concentrated, and thereto was added ethyl acetate, and the mixture was washed with water and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and concentrated. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 95/5) to obtain a compound 9Ta (545 mg, yield 87%).

HRMS(FAB): calculated value as C₃₇H₃₄F₃N₅O₇ [M+H]⁺: 718.2489, measured value: 718.2491

[0236]

Example 66 Synthesis of Compound 9Tb

3-[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-3-(5-methyl-2,4-dioxopyrimidin-1-yl)-5-[4-(trifluoromethyl)pyrimidin-2-yl]-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 23)

To a mixture of the compound 9Ta (527 mg, 0.734 mmol), dichloromethane (7 mL) and DIPEA (384 µL, 2.20 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (328 µL, 1.47 mmol) under ice-cooling, and the mixture was

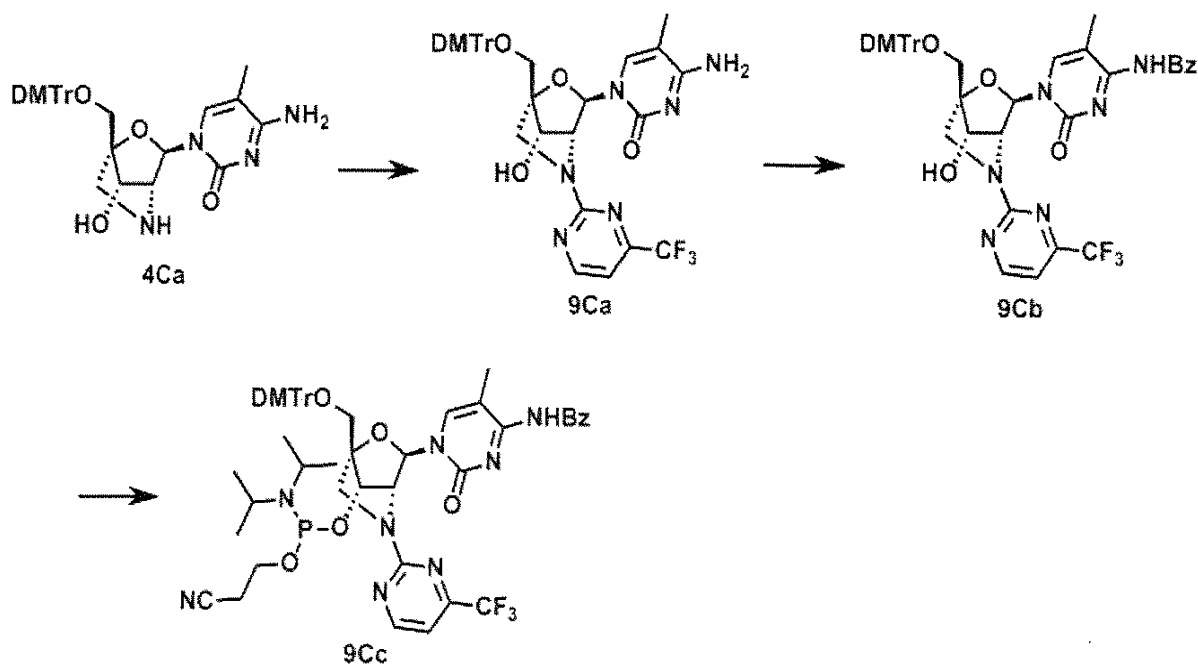
stirred at room temperature under nitrogen atmosphere overnight. Saturated aqueous sodium hydrogen carbonate solution was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was passed
 5 through a phase separator, and the solvents were evaporated. The residue was purified by column chromatography (SiO₂: hexane/ethyl acetate = 80/20 to 30/70, diol: hexane/ethyl acetate = 70/30 to 20/80) to obtain a compound 9Tb as the present compound 23 (530 mg, yield 79%).

10 HRMS(FAB): calculated value as C₄₆H₅₁F₃N₇O₈P [M+H]⁺: 918.3567,
 measured value: 918.3568

³¹P-NMR (CDCl₃) δ: 149.04, 149.12, 149.20, 149.47

[0237]

Synthesis of ALNA[4-CF₃-2Pym]-mC



[0238]

Example 67 Synthesis of Compound 9Ca

4-amino-1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy)methyl]-7-hydroxy-5-[4-
5 (trifluoromethyl)pyrimidin-2-yl]-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2-one

A mixture of the compound 4Ca (500 mg, 0.876 mmol), 2-chloro-4-(trifluoromethyl)pyrimidine (317 μ L, 2.63 mmol), DIPEA (610 μ L, 3.50 mmol), and EtOH (4.4 mL) was stirred at
10 150 °C under microwave irradiation for 1 hour. The reaction solution was concentrated, and thereto was added ethyl acetate, and the mixture was washed with water and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and concentrated. The residue was purified by
15 column chromatography (SiO₂: chloroform/methanol = 100/0 to 90/10) to obtain a compound 9Ca (186 mg, yield 30%).
HRMS(FAB): calculated value as C₃₇H₃₅F₃N₆O₆ [M+H]⁺: 717.2648, measured value: 717.2651

[0239]

20 Example 68 Synthesis of Compound 9Cb

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy)methyl]-7-hydroxy-5-[4-
(trifluoromethyl)pyrimidin-2-yl]-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-
25 yl]benzamide

To a mixture of the compound 9Ca (395 mg, 0.551 mmol), and DMF (5.5 mL) was added benzoic anhydride (131 mg, 0.579 mmol), and the mixture was stirred at room temperature overnight. Ethyl acetate was added to the reaction solution, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, water, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and concentrated. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 95/15) to obtain a compound 9Cb (335 mg, yield 74%).

HRMS(FAB): calculated value as C₄₄H₃₉F₃N₆O₇ [M+H]⁺: 821.2911, measured value: 821.2932

[0240]

Example 69 Synthesis of Compound 9Cc

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-[4-(trifluoromethyl)pyrimidin-2-yl]-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 24)

To a mixture of the compound 9Cb (316 mg, 0.385 mmol), dichloromethane (4 mL), and DIPEA (201 µL, 1.16 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (172 µL, 0.770 mmol), and the mixture was stirred at room temperature under nitrogen atmosphere overnight. Saturated

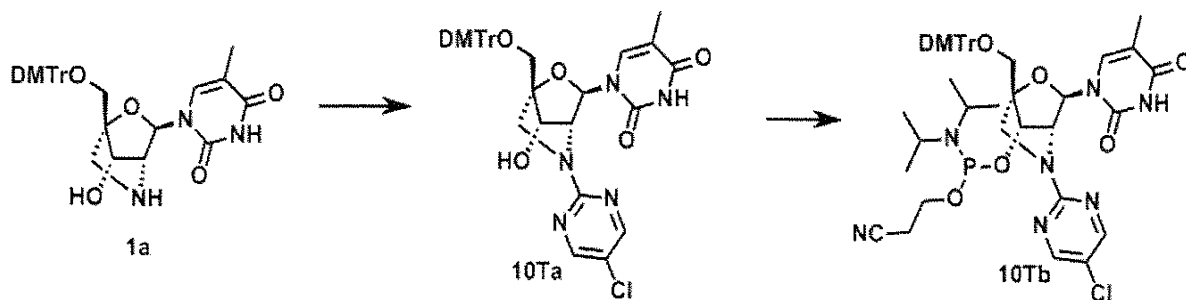
aqueous sodium hydrogen carbonate solution was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and solvents were evaporated. The residue was purified by column chromatography (SiO₂: hexane/ethyl acetate = 100/0 to 50/50, Diol: hexane/ethyl acetate = 100/0 to 50/50) to obtain a compound 9Cc as the present compound 24 (227 mg, yield 58%).

HRMS(FAB): calculated value as C₅₃H₅₆F₃N₈O₈P [M+H]⁺: 1021.3989, measured value: 1021.3993

³¹P-NMR (CDCl₃) δ: 149.15, 149.28, 149.36, 149.60

[0241]

Synthesis of ALNA[5-Cl-2Pym]-T



[0242]

Example 70 Synthesis of Compound 10Ta

1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-5-(5-chloropyrimidin-2-yl)-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2,4-dione

A mixture of the compound 1a (500 mg, 0.875 mmol, 2,5-dichloropyrimidine (196 mg, 1.31 mmol), DIPEA (457 μ L, 2.63 mmol) and EtOH (4.4 mL) was stirred at 150 $^{\circ}$ C under microwave irradiation for 1 hour. Insoluble materials of the reaction solutions were solubilized with chloroform, and the mixture was concentrated. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 90/10) to obtain a compound 10Ta (263 mg, yield 44%).

HRMS(FAB): calculated value as C₃₆H₃₄ClN₅O₇ [M+H]⁺: 684.2225, measured value: 684.2226

[0243]

Example 71 Synthesis of Compound 10Tb

3-[[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-5-(5-chloropyrimidin-2-yl)-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 25)

To a mixture of the compound 10Ta (240 mg, 0.351 mmol), dichloromethane (3.5 mL) and DIPEA (183 μ L, 1.05 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (157 μ L, 0.702 mmol) under ice-cooling, and the mixture was stirred at room temperature under nitrogen atmosphere for 4 days. Saturated aqueous sodium hydrogen carbonate solution was added to the reaction solutions, and the mixture was

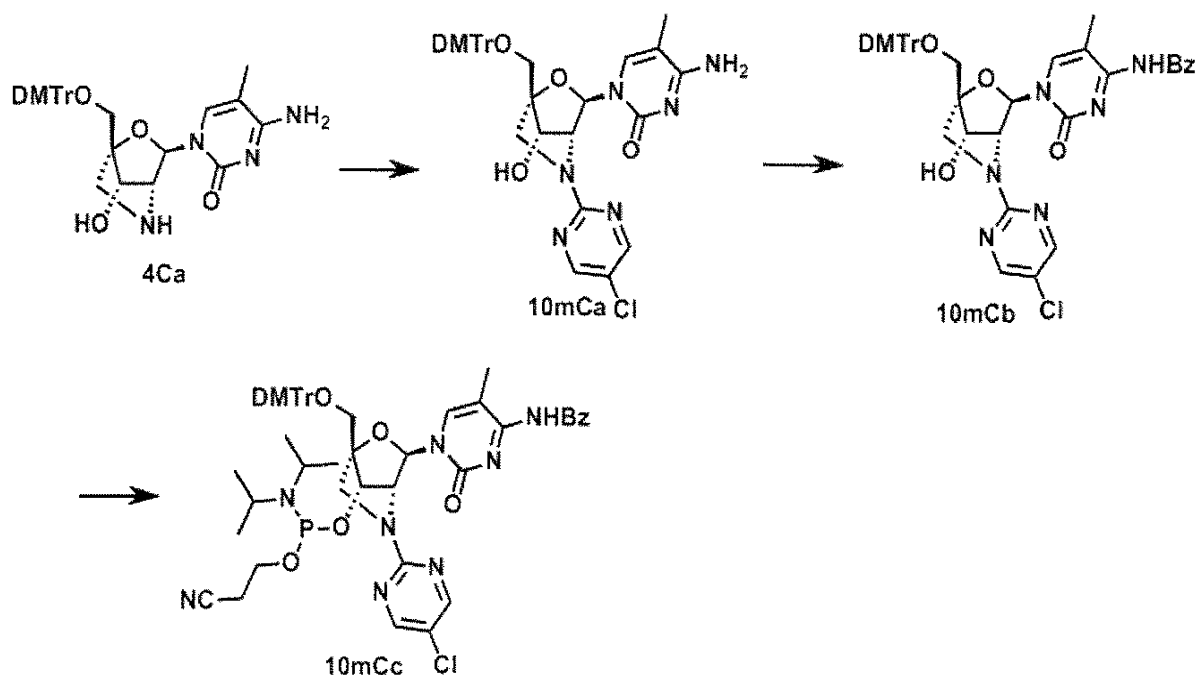
extracted with chloroform. The organic layer was passed through a phase separator, and the solvents were evaporated. The residue was purified by column chromatography (SiO₂: hexane/ethyl acetate = 70/30 to 20/80, diol: hexane/ethyl acetate = 70/30 to 20/80) to obtain a compound 10Tb (201 mg, yield 65%).

HRMS(FAB): calculated value as C₄₅H₅₁ClN₇O₈P [M+H]⁺: 884.3304, measured value: 884.3303

³¹P-NMR (CDCl₃) δ: 149.12

[0244]

Synthesis of ALNA[5-Cl-2Pym]-mC



[0245]

Example 72 Synthesis of Compound 10mCa

4-amino-1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-

phenylmethoxy)methyl]-5-(5-chloropyrimidin-2-yl)-7-hydroxy-
2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2-
one

A mixture of the compound 4Ca (500 mg, 0.876 mmol),
5 2,5-dichloropyrimidine (391 mg, 2.63 mmol), and DIPEA (610
μL, 3.50 mmol) and EtOH (4.4 mL) (2 batches) was stirred at
150 °C under microwave irradiation for 1 hour for each batch.
The reaction solution was concentrated, and water was added
thereto, and the mixture was extracted with chloroform. The
10 combined organic layer was passed through a phase separator,
and the solvents were evaporated. The residue was purified
by column chromatography (SiO₂: chloroform/methanol = 100/0
to 90/10) to obtain a compound 10mCa (381 mg, yield 32%).
HRMS (FAB): calculated value as C₃₆H₃₅ClN₆O₆ [M+H]⁺: 683.2385,
15 measured value: 683.2389

[0246]

Example 73 Synthesis of Compound 10mCb

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy)methyl]-5-(5-chloropyrimidin-2-yl)-7-hydroxy-
20 2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-
pyrimidin-4-yl]benzamide

To a mixture of the compound 10mCa (375 mg, 0.549 mmol)
and DMF (5.5 mL) was added benzoic anhydride (130 mg, 0.576
mmol), and the mixture was stirred at room temperature
25 overnight. To the reaction solution was added benzoic

anhydride (24.8 mg, 0.110 mmol) additionally, and the mixture was further stirred at room temperature overnight. Ethyl acetate was added to the reaction solution, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, water, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and concentrated. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 95/5) to obtain a compound 10mCb (422 mg, yield 98%).

HRMS (MALDI): calculated value as C₄₃H₃₉ClN₆O₇ [M+Na]⁺: 809.2461, measured value: 809.2463
[0247]

Example 74 Synthesis of Compound 10mCc

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy)methyl]-5-(5-chloropyrimidin-2-yl)-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-4-yl]benzamide (Present compound 26)

To a mixture of the compound 10mCb (409 mg, 0.520 mmol), dichloromethane (5 mL), and DIPEA (272 μ L, 1.56 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (232 μ L, 1.04 mmol), and the mixture was stirred at room temperature under nitrogen atmosphere overnight. Saturated aqueous sodium hydrogen carbonate solution was added to the reaction solution, and the mixture was extracted with

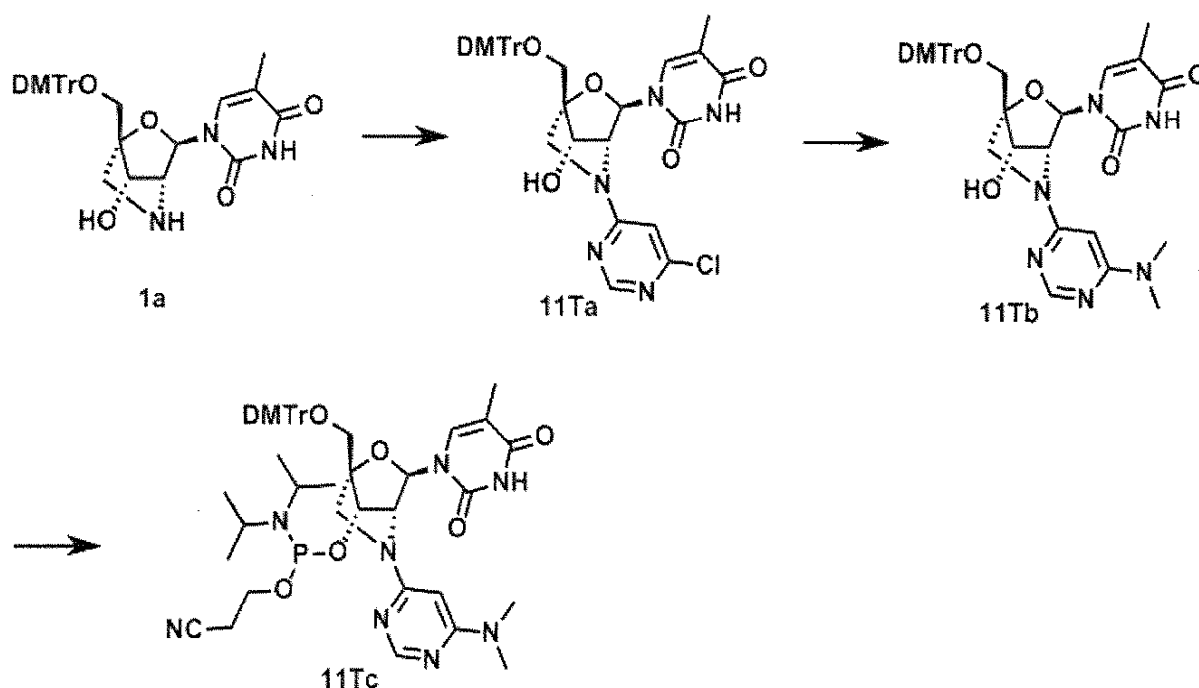
chloroform. The organic layer was passed through a phase separator, and the solvents were evaporated. The residue was purified by column chromatography (SiO₂: hexane/ethyl acetate= 100/0 to 50/50, diol: hexane/ethyl acetate = 100/0 to 50/50) to obtain a compound 10mCc as the present compound 26 (264 mg, yield 51%).

HRMS(MALDI): calculated value as C₅₂H₅₆ClN₈O₈P [M+Na]⁺: 1009.3540, measured value: 1009.3521

³¹P-NMR (CDCl₃) δ: 149.28

10 [0248]

Synthesis of ALNA[6-NMe₂-4Pym]-T



[0249]

Example 75 Synthesis of Compound 11Ta

1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-5-(6-chloropyrimidin-4-yl)-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2,4-dione

5 A mixture of the compound 1a (1.00 g, 1.75 mmol), 4,6-dichloropyrimidine (391 mg, 2.63 mmol), DIPEA (914 μ L, 5.25 mmol) and EtOH (18 mL) was stirred at 120 °C under microwave irradiation for 1 hour. The reaction was concentrated, the residue was purified by column chromatography (SiO₂:
10 chloroform/methanol = 100/0 to 95/5) to obtain a compound 11Ta (1.00 g, yield 84%).

HRMS(FAB): calculated value as C₃₆H₃₄ClN₅O₇ [M+H]⁺: 684.2225, measured value: 684.2214
[0250]

15 Example 76 Synthesis of Compound 11Tb

1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-5-[6-(dimethylamino)pyrimidin-4-yl]-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2,4-dione

20 To a mixture of the compound 11Ta (500 mg, 0.731 mmol), DIPEA (637 μ L, 3.66 mmol), and EtOH (5 mL) was added dimethylamine hydrochloride (179 mg, 2.19 mmol), and the mixture was stirred at 150 °C under microwave irradiation for 30 minutes. Insoluble materials were collected by filtration, and dried
25 under reduced pressure to obtain a compound 11Tb (386 mg,

yield 76%).

HRMS(FAB): calculated value as $C_{38}H_{40}N_6O_7$ $[M+H]^+$: 693.3037,
measured value: 693.3030

[0251]

5 Example 77 Synthesis of Compound 11Tc

3-[[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy]methyl]-5-[6-(dimethylamino)pyrimidin-4-yl]-
3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-
azabicyclo[2.2.1]heptan-7-yl]oxy-

10 (diisopropylamine)phosphanyl]oxypropanenitrile (Present
compound 27)

To a mixture of the compound 11Tb (335 mg, 0.484 mmol),
dichloromethane (5 mL), DIPEA (253 μ L, 1.45 mmol) was added
2-cyanoethyl N,N-diisopropylchlorophosphoramidite (216 μ L,
15 0.968 mmol) under ice-cooling, and the mixture was stirred
at room temperature under nitrogen atmosphere overnight. DMF
(2 mL) was added to the reaction solution, and the mixture
was stirred at room temperature under nitrogen atmosphere
overnight. To the reaction solution were added DIPEA (253
20 μ L, 1.45 mmol), and 2-cyanoethyl N,N-
diisopropylchlorophosphoramidite (216 μ L, 0.968 mmol)
additionally, and the mixture was stirred at room temperature
under nitrogen atmosphere for 2 days. Ethyl acetate was
added to the reaction solution, and the mixture was washed
25 with saturated aqueous sodium hydrogen carbonate solution,

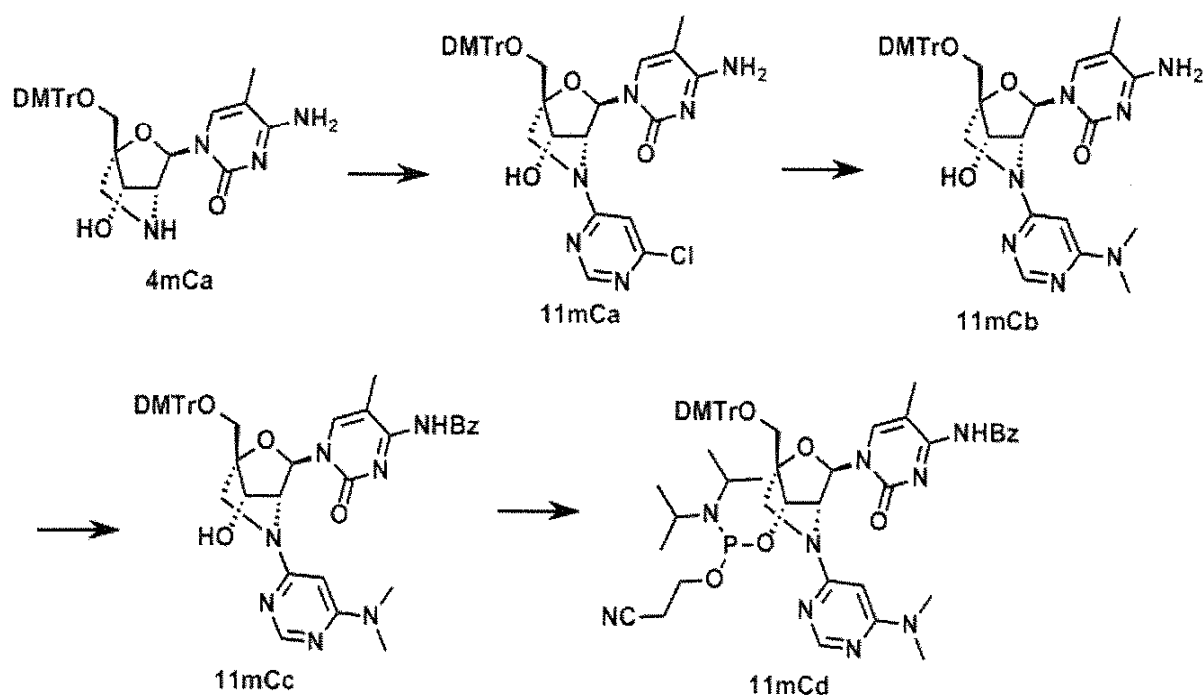
and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and concentrated. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 90/10, Diol: hexane/ethyl acetate = 50/50 to 0/100) to obtain a compound 11Tc as the present compound 27 (189 mg, yield 44%).

HRMS(FAB): calculated value as C₄₇H₅₇N₈O₈P [M+H]⁺: 893.4115, measured value: 893.4115

³¹P-NMR (CDCl₃) δ: 148.83, 148.99

[0252]

Synthesis of ALNA[6-NMe₂-4Pym]-mC



[0253]

Example 78 Synthesis of Compound 11mCa

4-amino-1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy)methyl]-5-(6-chloropyrimidin-4-yl)-7-hydroxy-
2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2-
one

5 A mixture of the compound 4mCa (2.00 g, 3.50 mmol),
4,6-dichloropyrimidine (626 mg, 4.20 mmol), DIPEA (1.83 mL,
10.5 mmol) and EtOH (17.5 mL) was stirred at 100 °C for 4
hours. The reaction solution was concentrated, and water
was added thereto, and the mixture was extracted with
10 chloroform. The organic layer was passed through a phase
separator, and the solvents were evaporated. The residue
was purified by column chromatography (SiO₂:
chloroform/methanol = 100/0 to 90/10) to obtain a compound
11mCa (1.34 g, yield 56%).
15 HRMS(FAB): calculated value as C₃₆H₃₅ClN₆O₆ [M+H]⁺: 683.2385,
measured value: 683.2384

[0254]

Example 79 Synthesis of Compound 11mCb

4-amino-1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-
20 phenylmethoxy)methyl]-5-[8-(dimethylamino)pyrimidin-4-yl]-
7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-
methylpyrimidin-2-one

A mixture of the compound 11mCa (1.12 g, 1.64 mmol),
dimethylamine (50% aqueous solution, 863 µL, 8.20 mmol), and
25 EtOH (1 mL) was stirred at 150 °C under microwave irradiation

for 30 minutes. The reaction was concentrated, and the residue was purified by column chromatography (NH: chloroform/methanol = 100/0 to 95/5) to obtain a compound 11mCb (1.09 g, yield 96%).

5 HRMS(MALDI): calculated value as $C_{38}H_{41}N_7O_6$ $[M+Na]^+$: 714.3011, measured value: 714.3003

[0255]

Example 80 Synthesis of Compound 11mCc

10 N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-5-[6-(dimethylamino)pyrimidin-4-yl]-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide

To a mixture of the compound 11mCb (1.08 g, 1.56 mmol) and DMF (8 mL) was added benzoic anhydride (389 mg, 1.72 mmol), and the mixture was stirred at room temperature overnight. Ethyl acetate was added to the reaction solutions, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, water, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and concentrated. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 95/5) to obtain a compound 11Cc (1.02 g, yield 82%).

20 HRMS(MALDI): calculated value as $C_{45}H_{45}N_7O_7$ $[M+H]^+$: 796.3453, measured value: 796.3466

25 [0256]

Example 81 Synthesis of Compound 11mCd

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy(diisopropylamino)phosphanyl]oxy-5-[(6-dimethylamino)pyrimidin-4-yl]-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 28)

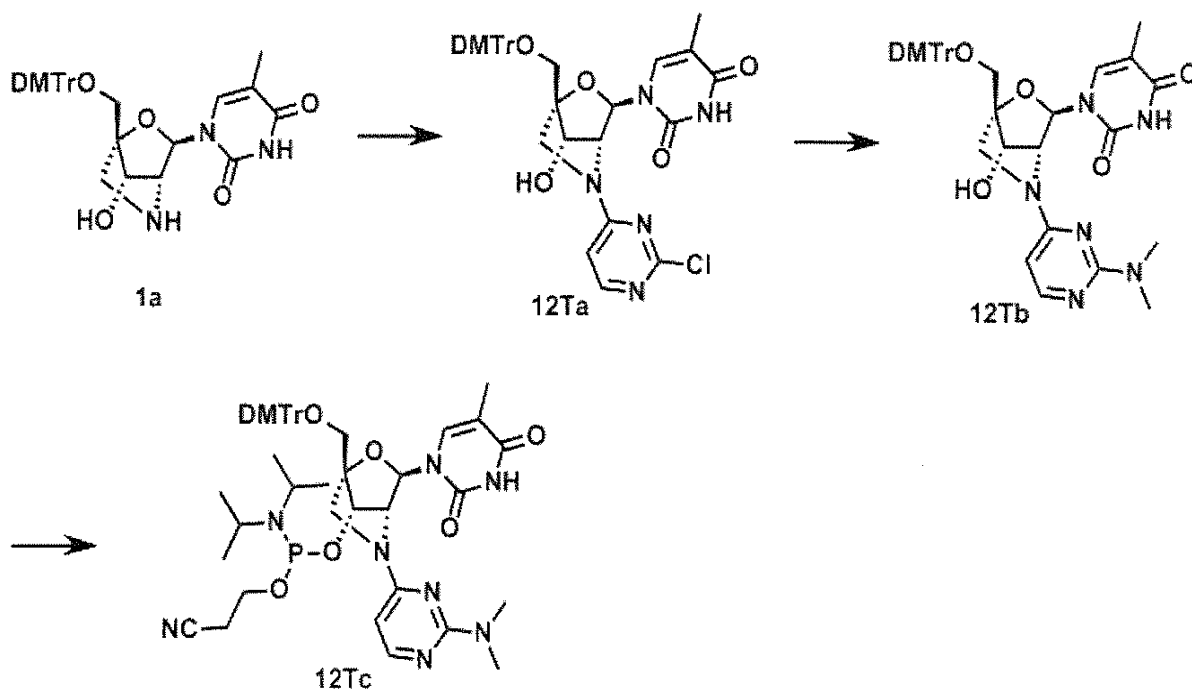
To a mixture of the compound 11mCc (972 mg, 1.22 mmol), dichloromethane (12 mL) and DIPEA (637 μ L, 3.66 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (545 μ L, 2.44 mmol), and the mixture was stirred at room temperature under nitrogen atmosphere overnight. Saturated aqueous sodium hydrogen carbonate solution was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and the solvents were evaporated. The residue was purified by column chromatography (SiO₂: hexane/ethyl acetate= 50/50 to 0/100, NH: hexane/ethyl acetate = 50/50 to 0/100) to obtain a compound 11mCd as the present compound 28 (788 mg, yield 65%).

HRMS(MALDI): calculated value as C₅₄H₆₂N₉O₈P [M+H]⁺: 996.4532, measured value: 996.4513

³¹P-NMR (CDCl₃) δ : 149.06, 149.17

[0257]

Synthesis of ALNA[2-NMe₂-4Pym]-T



[0258]

Example 82 Synthesis of Compound 12Ta

1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-5-(2-chloropyrimidin-4-yl)-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2,4-dione

A mixture of the compound 1a (2.00 g, 3.50 mmol), 2,4-dichloropyrimidine (782 mg, 5.25 mmol), DIPEA (1.83 mL, 10.5 mmol), and EtOH (17.5 mL) was stirred at 100 °C for 4.5 hours. The reaction solutions were concentrated, and water was added thereto, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and the solvents were evaporated. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 95/5) to obtain a compound

12Ta (1.86 g, yield 78%).

HRMS(FAB): calculated value as $C_{36}H_{34}ClN_5O_7$ $[M+H]^+$: 684.2225,
measured value: 684.2223

[0259]

5 Example 83 Synthesis of Compound 12Tb

1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy)methyl]-5-[2-(dimethylamino)pyrimidin-4-yl]-7-
hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-
methylpyrimidin-2,4-dione

10 To a mixture of the compound 12Ta (500 mg, 0.731 mmol),
DIPEA (637 μ L, 3.66 mmol), and EtOH (5 mL) was added dimethyl
amine hydrochloride (179 mg, 2.19 mmol), and the mixture was
stirred at 150 °C under microwave irradiation for 30 minutes.
The reaction solutions were concentrated, and the residue
15 was purified by column chromatography (SiO_2 :
chloroform/methanol = 100/0 to 95/5). The mixture was
trituated with diisopropyl ether to obtain a compound 12Tb
(648 mg) as a crude product.

HRMS(FAB): calculated value as $C_{39}H_{40}N_6O_7$ $[M+H]^+$: 693.3037,
20 measured value: 693.3041

[0260]

Example 84 Synthesis of Compound 12Tc

3-[[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy)methyl]-5-[2-(dimethylamino)pyrimidin-4-yl]-3-(5-
25 methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-

azabicyclo[2.2.1]heptan-7-yl]oxy-

(diisopropylamino)oxypropanenitrile (Present compound 29)

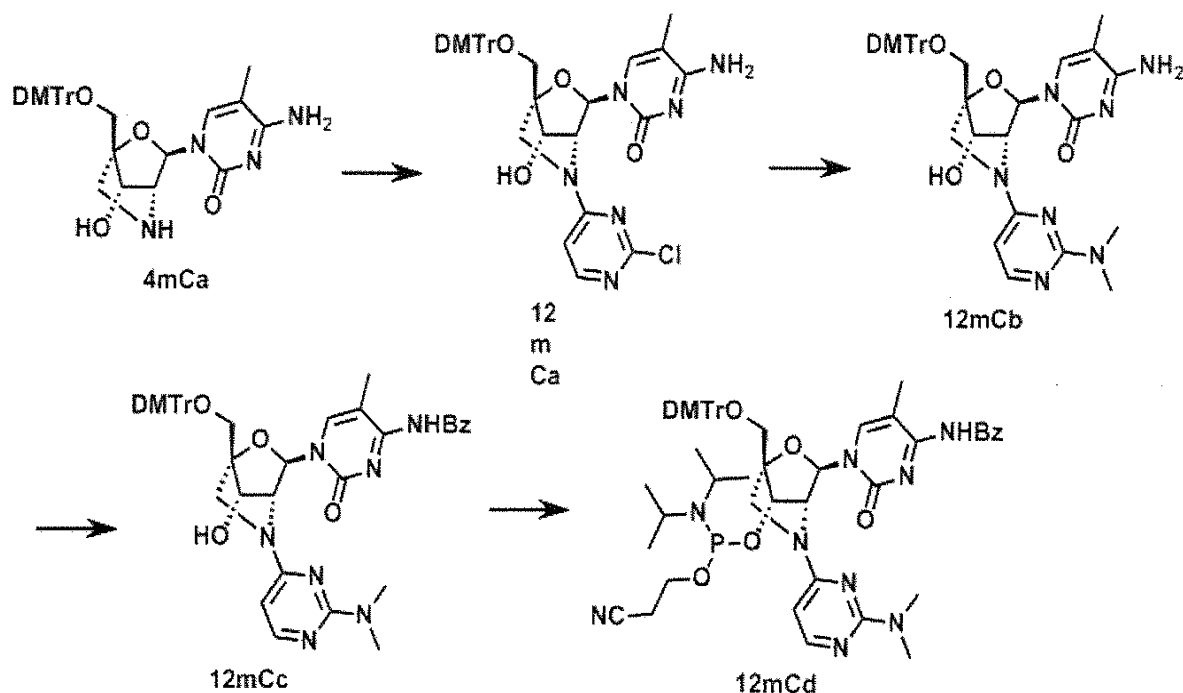
To a mixture of the compound 12Tb (638 mg, 0.731 mmol, crude product), dichloromethane (7 mL) and DIPEA (382 μ L, 2.19 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (326 μ L, 1.46 mmol), and the mixture was stirred at room temperature under nitrogen atmosphere for 5 days. Saturated aqueous sodium hydrogen carbonate solution was added to the reaction solutions, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and the solvents were evaporated. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 90/10, diol: hexane/ethyl acetate = 50/50 to 0/100) to obtain a compound 12Tc as the present compound 29 (304 mg, yields 47% over two steps).

HRMS(FAB): calculated value as C₄₇H₅₇N₈O₈P [M+H]⁺: 893.4115, measured value: 893.4128

³¹P-NMR (CDCl₃) δ : 148.91, 149.33

[0261]

Synthesis of ALNA[2-NMe₂-4Pym]-mC



[0262]

Example 85 Synthesis of Compound 12mCa

4-amino-1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-5-(2-chloropyrimidin-4-yl)-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2-one

A mixture of the compound 4mCa (2.00 g, 2.50 mmol), 2,4-dichloropyrimidine (627 mg, 4.20 mmol), DIPEA (1.83 mL, 10.5 mmol) and EtOH (17.5 mL) was stirred at 100 °C for 3 hours. The reaction solution was concentrated, and water was added thereto, and the mixture was extracted with chloroform. The organic layer was dried over anhydrous magnesium sulfate, and concentrated. The residue was purified by column chromatography (SiO₂: chloroform/methanol

= 100/0 to 90/10) to obtain a compound 12mCa (1.28 g, yield 54%).

HRMS(MALDI): calculated value as $C_{36}H_{35}ClN_6O_6$ $[M+Na]^+$: 705.2199, measured value: 705.2199

5 [0263]

Example 86 Synthesis of Compound 12mCb

4-amino-1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-5-[2-(dimethylamino)pyrimidin-4-yl]-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2-one

10

A mixture of the compound 12mCa (941 mg, 1.38 mmol), dimethyl amine (aqueous 50% solution, 725 μ L, 6.90 mmol), and EtOH (14 mL, 0.1 M) was stirred at 150 °C under microwave irradiation for 30 minutes. The reaction solution was concentrated, and the residue was purified by column chromatography (NH: chloroform/methanol = 100/0 to 90/10) to obtain a compound 12mCb (784 mg, yield 82%).

15

HRMS(MALDI): calculated value as $C_{38}H_{41}N_7O_6$ $[M+H]^+$: 692.3191, measured value: 692.3167

20 [0264]

Example 87 Synthesis of Compound 12mCc

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-5-[2-(dimethylamino)pyrimidin-4-yl]-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide

25

To a mixture of the compound 12mCb (754 mg, 1.09 mmol) and DMF (5.5 mL) was added benzoic anhydride (271 mg, 1.20 mmol), and the mixture was stirred at room temperature overnight. Ethyl acetate was added to the reaction solution, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, water, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and concentrated. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 95/5) to obtain a compound 12mCc (731 mg, yield 84%).

HRMS(MALDI): calculated value as C₄₅H₄₅N₇O₇ [M+H]⁺: 796.3453, measured value: 796.3440

[0265]

Example 88 Synthesis of Compound 12mCd

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-[2-(dimethylamino)pyrimidin-4-yl]-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 30)

To a mixture of the compound 12mCc (709 mg, 0.891 mmol), dichloromethane (9 mL) and DIPEA (466 µL, 2.67 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (397 µL, 1.78 mmol), and the mixture was stirred at room temperature under nitrogen atmosphere overnight. Saturated

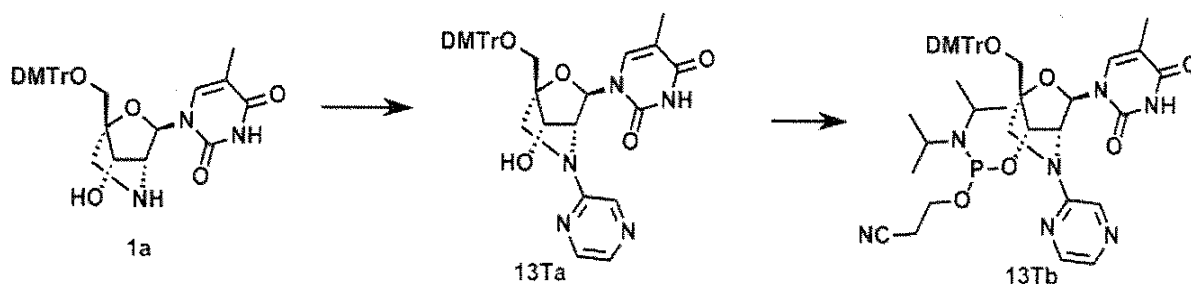
aqueous sodium hydrogen carbonate solution was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and the solvents were evaporated. The residue was purified by column chromatography (SiO₂: hexane/ethyl acetate = 50/50 to 0/100, NH: hexane/ethyl acetate = 50/50 to 0/100) to obtain a compound 12mCd as the present compound 30 (482 mg, yield 54%).

HRMS (MALDI): calculated value as C₅₄H₆₂N₉O₉P [M+H]⁺: 996.4532, measured value: 996.4526

³¹P-NMR (CDCl₃) δ: 149.19, 149.46, 149.65

[0266]

Synthesis of ALNA[Prz]-T



[0267]

Example 89 Synthesis of Compound 13Ta

[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-pyrazine-2-yl]-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-pyrimidin-2,4-dione

A mixture of the compound 1a (500 mg, 0.875 mmol), 2-

fluoropyrazine (201 μ L, 2.63 mmol), DIPEA (610 μ L, 3.50 mmol) and DMSO (8.8 mL) was stirred at 140 $^{\circ}$ C for 1.5 hours and at 150 $^{\circ}$ C for 4.5 hours under nitrogen atmosphere. Ethyl acetate was added to the reaction solutions, and the mixture was washed with water, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and concentrated. The residue was purified by column chromatography SiO₂: chloroform/methanol 100/0 to 90/10, SiO₂: hexane/ethyl acetate = 50/50 to 0/100) to obtain a compound 13Ta (227 mg, yield 40%).

HRMS (MALDI): calculated value as C₃₆H₃₅N₅O₇ [M+Na]⁺: 672.2429, measured value: 672.2409

[0268]

Example 90 Synthesis of Compound 13Tb

3-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-pyrazine-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 31)

To a mixture of the compound 13Ta (214 mg, 0.329 mmol), dichloromethane (6.6 mL), and DIPEA (172 μ L, 0.987 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (147 μ L, 0.658 mmol), and the mixture was stirred at room temperature under nitrogen atmosphere overnight. Saturated aqueous sodium hydrogen carbonate solution was added to the

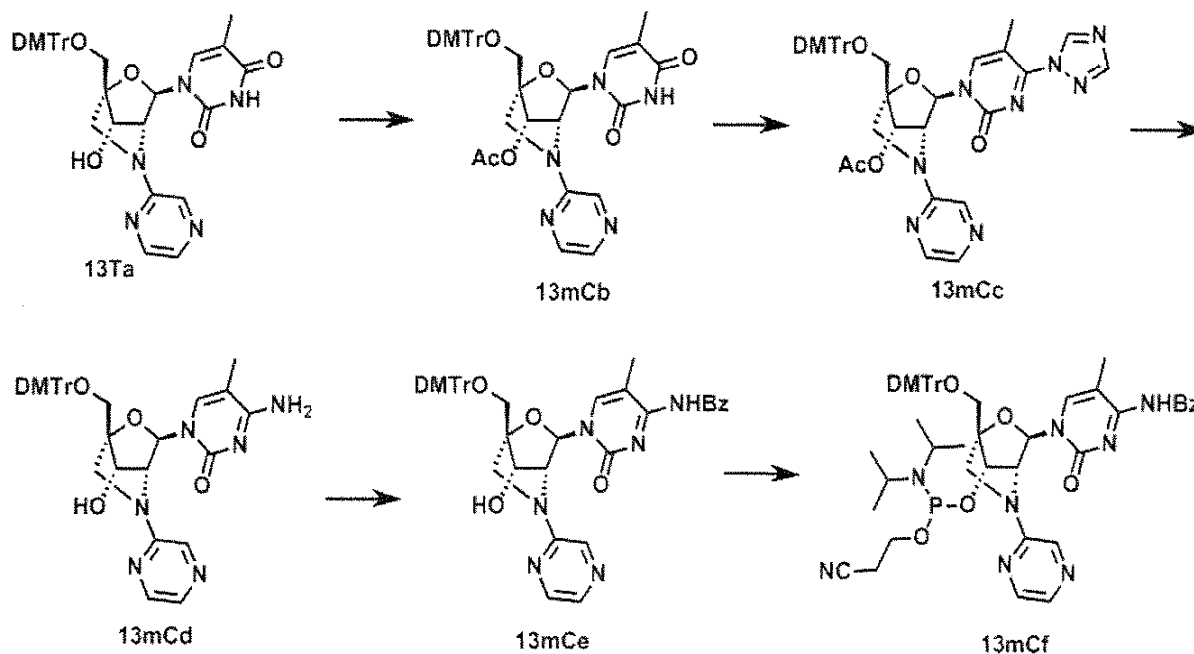
reaction solution, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and the solvents were evaporated. The residue was purified by column chromatography (SiO₂: hexane/ethyl acetate = 50/50 to 0/100, NH: ethyl acetate/methanol = 100/0 to 90/10) to obtain a compound 13Tb as the present compound 31 (204 mg, yield 73%).

HRMS(MALDI): calculated value as C₄₅H₅₂N₇O₈P [M+Na]⁺: 872.3507, measured value: 872.3507

³¹P-NMR (CDCl₃) δ: 149.12, 149.31

[0269]

Synthesis of ALNA[Prz]-mC



[0270]

Example 91 Synthesis of Compound 13mCb

[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-pyrazin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]acetate

To a mixture of the compound 13Ta (746 mg, 1.15 mmol),
 5 DMAP (14.0 mg, 0.115 mmol) and pyridine (5.8 mL) was added acetic anhydride (163 μ L, 1.73 mmol), and the mixture was stirred at room temperature overnight. Ethyl acetate was added to the reaction solution, and the mixture was washed
 10 with saturated aqueous sodium hydrogen carbonate solution, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and concentrated. Pyridine was azeotroped with ethyl acetate, and the residue was triturated with diisopropyl ether to obtain a compound 13MCb (712 mg, yield 90%).

15 HRMS(MALDI): calculated value as $C_{38}H_{37}N_5O_8$ $[M+Na]^+$: 714.2534, measured value: 714.2520
 [0271]

Example 92 Synthesis of Compound 13mCc

[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-3-(5-methyl-2-oxo-(4-1,2,4-triazol-1-yl)pyrimidin-1-yl)-5-pyrazin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]acetate
 20

To a mixture of the compound 13mCb (692 mg, 1.00 mmol),
 1,2,4-triazole (622 mg, 9.00 mmol), DIPEA (1.74 mL, 10.0
 25 mmol) and acetonitrile (10 mL) was added phosphoryl chloride

(158 μ L, 1.70 mmol) under ice-cooling, and the mixture was stirred at room temperature under nitrogen atmosphere for 1 hour. Ethyl acetate was added to the reaction solution, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and concentrated to obtain a compound 13mCc (753 mg) as a crude product.

HRMS(MALDI): calculated value as $C_{40}H_{38}N_8O_7$ $[M+Na]^+$: 765.2756, measured value: 765.2750

[0272]

Example 93 Synthesis of Compound 13mCd

4-amino-1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-pyrazin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2-one

To a mixture of the compound 13mCc (744 g, 1.00 mmol, crude product) and acetonitrile (10 mL) was added 28% ammonia water (6.7 mL), and the mixture was stirred at room temperature overnight. Water was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was dried over anhydrous magnesium sulfate, and concentrated to obtain a compound 13mCd (638 mg, yields over two steps 98%).

HRMS(MALDI): calculated value as $C_{36}H_{36}N_6O_6$ $[M+Na]^+$: 671.2589, measured value: 671.2591

[0273]

Example 94 Synthesis of Compound 13mCe

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-pyrazin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide

To a mixture of the compound 13mCd (625 mg, 0.963 mmol) and DMF (4.8 mL) was added benzoyl anhydride (327 mg, 1.44 mmol), and the mixture was stirred at room temperature overnight. Ethyl acetate was added to the reaction solution, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, water, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and concentrated. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 95/5) to obtain a compound 13mCe (245 mg, yield 34%).

HRMS(MALDI): calculated value as C₄₃H₄₀N₆O₇ [M+Na]⁺: 775.2851, measured value: 775.2845

[0274]

Example 95 Synthesis of Compound 13mCf

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-pyrazin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 32)

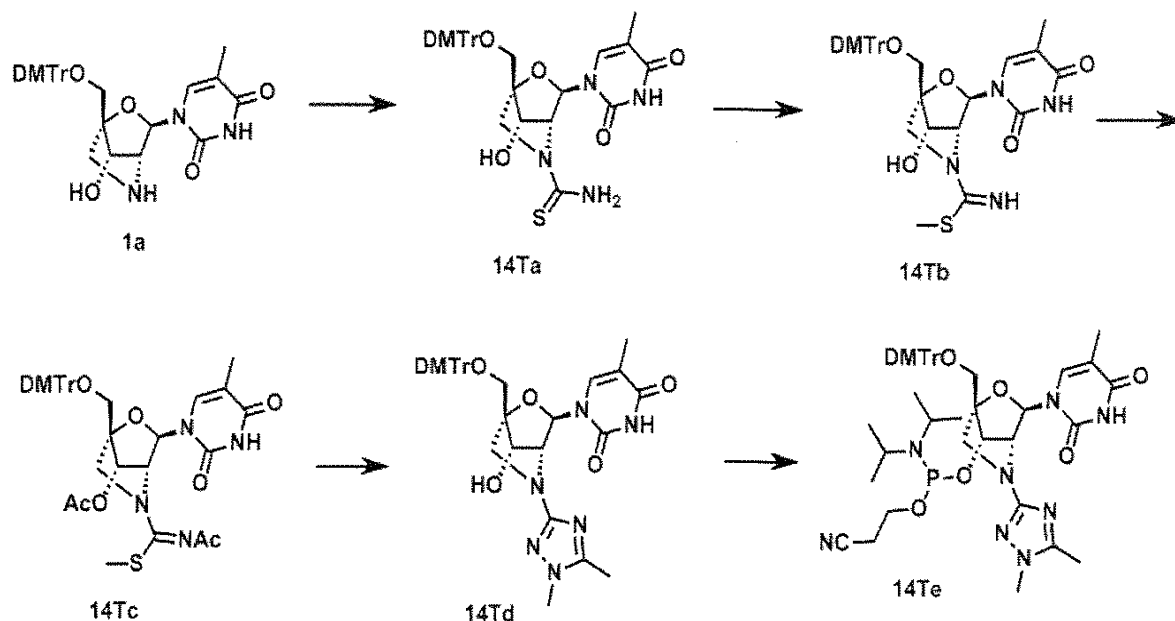
To a mixture of the compound 13mCe (565 mg, 0.751 mmol),

dichloromethane (7.5 mL) and DIPEA (392 μ L, 2.25 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (335 μ L, 1.50 mmol), and the mixture was stirred at room temperature under nitrogen atmosphere overnight. Saturated aqueous sodium hydrogen carbonate solution was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and the solvents were evaporated. The residue was purified by column chromatography (SiO₂: hexane/ethyl acetate = 70/30 to 20/80, NH: hexane/ethyl acetate = 50/50 to 0/100) to obtain a compound 13mCf as the present compound 32 (369 mg, yield 52%).

HRMS(MALDI): calculated value as C₅₀H₄₄N₆O₈ [M+Na]⁺: 975.3929, measured value: 975.3923

³¹P-NMR (CDCl₃) δ : 149.25, 149.49
[0275]

Synthesis of ALNA[Trz]-T



[0276]

Example 96 Synthesis of Compound 14Ta

(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-5-carbothioamide

To a mixture of the compound 1a (2.00 g, 3.50 mmol) and THF (17.5 mL) was added 1,1-thiocarbonyl diimidazole (1.25 g, 7.00 mmol), and the mixture was stirred at room temperature overnight. 28% Ammonia water (17.5 mL) was added to the reaction solution, and the mixture was stirred at room temperature overnight. The reaction solution was concentrated, and insoluble materials were filtered off. The insoluble materials were washed with water to obtain a compound 14Ta (2.35 g) as a crude product.

HRMS(MALDI): calculated value as $C_{33}H_{34}N_4O_7$ $[M+Na]^+$: 653.2040,

measured value: 653.2026

[0277]

Example 97 Synthesis of Compound 14Tb

methyl (1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
5 methoxy]methyl]-7-hydroxy-3-(5-methyl-2,4-dioxypyrimidin-1-
yl)-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxyimidothioate

To a mixture of the compound 14Ta (2.35 g, 3.50 mmol, crude product) and THF (17.5 mL), was added methyl iodide (654 μ L, 10.5 mmol), and the mixture was stirred at room
10 temperature for 2 days. Saturated aqueous sodium hydrogen carbonate solution was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was dried over anhydrous magnesium sulfate, and insoluble materials were filtered off, and concentrated under reduced
15 pressure. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 80/20) to obtain a compound 14Tb (1.88 g, yields over two steps 83%).

HRMS(MALDI): calculated value as C₃₄H₃₆N₄O₇ [M+H]⁺: 645.2377, measured value: 645.2374

20 [0278]

Example 98 Synthesis of Compound 14Tc

[(1R,3R,4R,7S)-5-[(Z)-N-acetyl-C-methylsulfanyl-carbamimidoyl]-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-
25 5-azabicyclo[2.2.1]heptan-7-yl]acetate

To a mixture of the compound 14Tb (1.87 g, 2.90 mmol ,
 DMAP (35.4 mg, 0.290 mmol) and pyridine (14.5 mL) was added
 acetic anhydride (822 μ L, 8.70 mmol), and the mixture was
 stirred at room temperature overnight. Ethyl acetate was
 5 added to the reaction solution, and the mixture was washed
 with 5% aqueous copper sulfate solution, water, and saturated
 brine. The organic layer was dried over anhydrous magnesium
 sulfate, and insoluble materials were filtered off, and the
 filtrates were then concentrated under reduced pressure to
 10 obtain a compound 14Tc (2.11 g, yield 100%).

HRMS(MALDI): calculated value as $C_{38}H_{40}N_4O_9S$ $[M+Na]^+$: 751.2408,
 measured value: 751.2398

[0279]

Example 99 Synthesis of Compound 14Td

15 1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
 methoxy)methyl]-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-7-
 hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-
 methylpyrimidin-2,4-dione

To a mixture of the compound 14Tc (1.66 g, 2.28 mmol)
 20 and EtOH (23 mL) was added methyl hydrazine (1.20 mL, 22.8
 mmol), and the mixture was stirred at room temperature
 overnight. The reaction solution was concentrated, and the
 residue was purified by column chromatography (SiO_2 :
 chloroform/methanol = 100/0 to 90/10) to obtain a compound
 25 14td (985 mg, yield 65%).

HRMS(MALDI): calculated value as $C_{36}H_{38}N_6O_7$ $[M+Na]^+$: 689.2694,
measured value: 689.2684

[0280]

Example 100 Synthesis of Compound 14Te

5 3-[[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy]methyl]-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-3-(5-
methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-
azabicyclo[2.2.1]heptan-7-yl]oxy-
(diisopropylamino)phosphanyl]oxypropanenitrile (Present
10 compound 33)

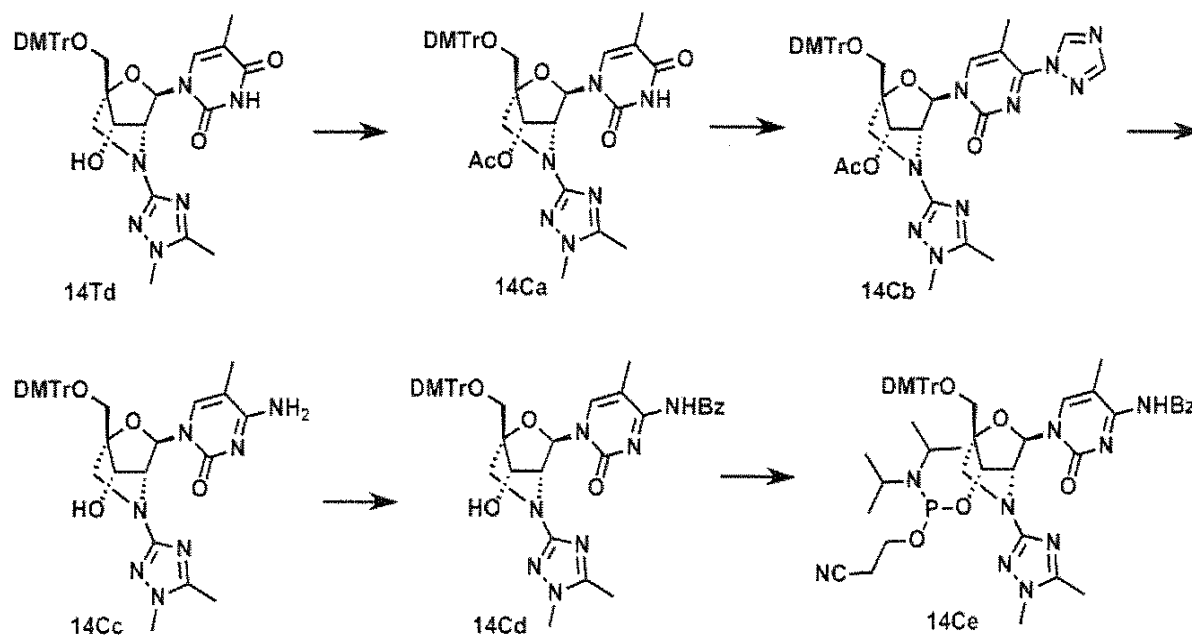
To a mixture of the compound 14Td (576 mg, 0.864 mmol),
dichromethane (8.6 mL) and DIPEA (451 μ L, 2.59 mmol) was
added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite
(385 μ L, 1.73 mmol), and the mixture was stirred at room
15 temperature under nitrogen atmosphere for 2 days. Saturated
aqueous sodium hydrogen carbonate solution was added to the
reaction solutions, and the mixture was extracted with
chloroform. The organic layer was passed through a phase
separator, and the solvents were evaporated. The residue
20 was purified by column chromatography (SiO_2 : hexane/ethyl
acetate = 50/50 to 0/100, diol: hexane/ethyl acetate = 50/50
to 0/100) to obtain a compound 14Te as the present compound
33 (517 mg, yield 69%).

HRMS(MALDI): calculated value as $C_{45}H_{55}N_8O_8P$ $[M+Na]^+$: 889.3773,
25 measured value: 889.3778

^{31}P -NMR (CDCl_3) δ : 148.40, 148.45

[0281]

Synthesis of ALNA[Trz]-mC



5 [0282]

Example 101 Synthesis of Compound 14Ca

[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]acetate

To a mixture of the compound 14Td (918 mg, 1.38 mmol), DMAP (16.9 mg, 0.138 mmol) and pyridine (6.9 mL) was added acetic anhydride (196 μL , 2.07 mmol), and the mixture was stirred at room temperature overnight. Ethyl acetate was added to the reaction solution, and the mixture was washed

with saturated aqueous sodium hydrogen carbonate solution, water, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and insoluble materials were filtered off, and the filtrates were then concentrated under reduced pressure to obtain a compound 14Ca (955 mg, yield 98%).

HRMS(MALDI): calculated value as $C_{38}H_{40}N_6O_8$ $[M+Na]^+$: 731.2800, measured value: 731.2791

[0283]

10 Example 102 Synthesis of Compound 14Cb

[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-3-(5-methyl-2-oxo-4-(1,2,4-triazol-1-yl)pyrimidin-1-yl]-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]acetate

15 To a mixture of the compound 14Ca (915 mg, 1.29 mmol), 1,2,4-triazole (802 mg, 11.6 mmol), DIPEA (2.25 mL, 12.9 mmol) and acetonitrile (12.9 mL) was added phosphoryl chloride (204 μ L, 2.19 mmol) under ice-cooling, and the mixture was stirred at room temperature under nitrogen atmosphere overnight. Ethyl acetate was added to the reaction solution, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and insoluble materials were filtered off, and the filtrates were concentrated under reduced pressure to obtain

20

25

a compound 14Cb (1.08 g) as a crude product.

HRMS(MALDI): calculated value as $C_{40}H_{41}N_9O_7$ $[M+Na]^+$: 782.3021,
measured value: 782.3019

[0284]

5 Example 103 Synthesis of Compound 14Cc

4-amino-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy]methyl]-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-7-
hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-
methylpyrimidin-2-one

10 To a mixture of the compound 14Cb (1.07 g, 1.29 mmol),
crude product) and acetonitrile (12.9 mL) was added 28%
ammonia water (8.6 mL), and the mixture was stirred at room
temperature overnight. Water was added to the reaction
solution, and the mixture was extracted with chloroform. The
15 organic layer was passed through a phase separator, and the
filtrates were concentrated to obtain a compound 14Cc (748
mg, yields over two steps 87%).

HRMS(MALDI): calculated value as $C_{36}H_{39}N_7O_6$ $[M+Na]^+$: 688.2854,
measured value: 688.2857

20 [0285]

Example 104 Synthesis of Compound 14Cd

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy]methyl]-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-7-
hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-
25 oxo-pyrimidin-4-yl]benzamide

To a mixture of the compound 14Cc (733 mg, 1.10 mmol) and pyridine (5.5 mL) was added benzoic anhydride (374 mg, 1.64 mmol), and the mixture was stirred at room temperature overnight. A 2N aqueous sodium hydroxide solution (5.5 mL) was added to the reaction solution, and the mixture was stirred at room temperature for 1 hour. Ethyl acetate was added to the reaction solution, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and insoluble materials were filtered off, and the filtrates were concentrated under reduced pressure. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 90/10) to obtain a compound 14Cd (744 mg, yield 88%).

HRMS (MALDI): calculated value as C₄₃H₄₃N₇O₇ [M+Na]⁺: 792.3116, measured value: 792.3111

[0286]

Example 105 Synthesis of Compound 14Ce

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 34)

To a mixture of the compound 14Cd (730 mg, 0.948 mmol), dichloromethane (9.5 mL) and DIPEA (0.495 mL, 2.84 mmol) was

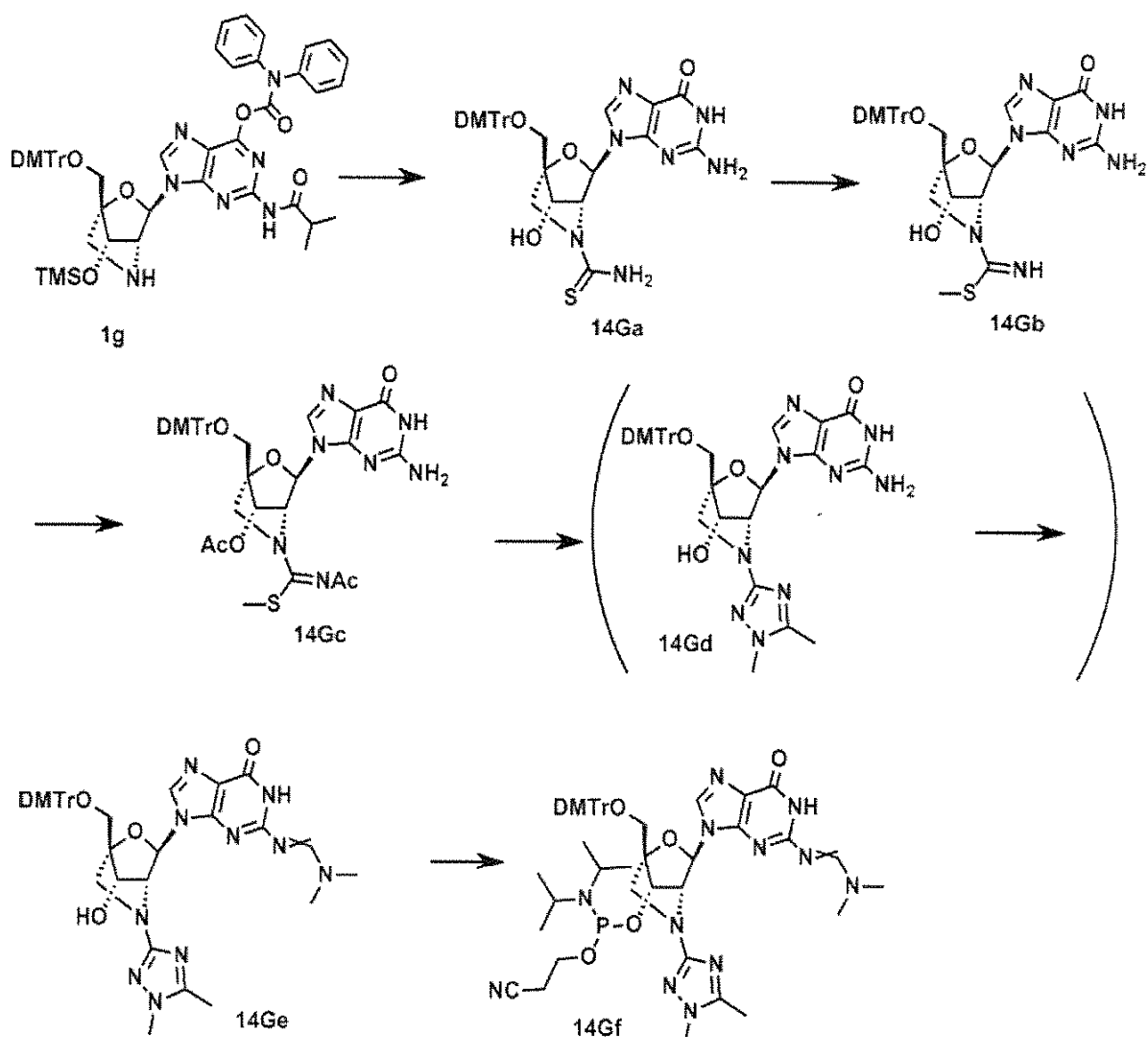
added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite
(0.423 mL, 1.90 mmol) under ice-cooling, and the mixture was
stirred at room temperature under nitrogen atmosphere
overnight. Saturated aqueous sodium hydrogen carbonate
5 solution was added to the reaction solution, and the mixture
was extracted with chloroform. The organic layer was passed
through a phase separator, and the solvents were evaporated.
The residue was purified by column chromatography (SiO₂:
hexane/ethyl acetate = 50/50 to 0/100, NH: hexane/ethyl
10 acetate = 50/50 to 0/100) to obtain a compound 14Ce as the
present compound 34 (672 mg, yield 73%).

HRMS(MALDI): calculated value as C₅₂H₆₀N₃O₈P [M+Na]⁺: 992.4195,
measured value: 992.4186

³¹P-NMR (CDCl₃) δ: 148.50, 148.58

15 [0287]

Synthesis of ALNA[Trz]-G



[0288]

Example 106 Synthesis of Compound 14Ga

(1R,3R,4R,7S)-3-(2-amino-6-oxo-1H-purin-9-yl)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptane-5-carbothioamide

To a solution of the compound 1g which was synthesized according to the method described in WO 2017/047816 A1 (55.0 g, 58.9 mmol) in THF (270 mL) was injected 1,1-thiocarbonyl

diimidazole (21.0 g, 118 mmol), and the mixture was stirred at room temperature for 2 hours. 28% Ammonia water (270 mL) was added to the reaction solution, and the mixture was stirred at room temperature for 30 minutes, followed by stirring at 50 °C for 46 hours. Chloroform (700 mL) was added to the reaction solution, and after stirring for a while, the organic layer was separated with a separatory funnel, and dried over anhydrous sodium sulfate. Insoluble materials were filtered off, and the solvents were then evaporated under reduced pressure, and the resulting residue was roughly purified by silica gel chromatography (SiO₂: chloroform/methanol = 100/0 to 90/10). The resulting residue was solubilized in ethyl acetate (250 mL), and thereto was added toluene (250 mL). The solvents were evaporated under reduced pressure to about 300 mL (solids were precipitated out), and to the resulting suspensions was added diisopropyl ether (IPE) (50 mL), and the mixture was stirred at room temperature for 30 minutes. The precipitated materials were collected by filtration, and the filtered materials were washed with IPE (50 mL), and then dried under reduced pressure to obtain a compound 14Ga (27.9 g, yields 72%).

MS (ESI): m/z = 656 [M+H]⁺

[0289]

Example 107 Synthesis of compound 14Gb

methyl (1R,3R,4R,7S)-3-(2-amino-6-oxo-1H-purin-9-yl)-1-

[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxyimidothioate

To a solution of the compound 14Ga (27.9 g, 42.5 mmol) in THF (400 mL) was added dropwise methyl iodide (4.24 mL, 68.1 mmol) under ice-cooling, and the mixture was stirred at room temperature for 24 hours. The reaction solution was added to a mixed solution of sodium bicarbonate water (which was prepared from saturated sodium bicarbonate water (400 mL) and water (400 mL)) and chloroform (1400 mL) while stirring. After stirring for a while, the organic layer was separated with a separatory funnel, and dried over anhydrous sodium sulfate. The mixture was filtered through Celite, and the filtrates were azeotroped with toluene to obtain a compound 14Gb (26.9 g, yield 82%).

MS (ESI): m/z = 670 $[M+H]^+$
[0290]

Example 108 Synthesis of Compound 14Gc

[(1R,3R,4R,7S)-5-(N-acetyl-C-methylsulfanylcarboxyimido-3-(2-amino-6-oxo-1H-purin-9-yl)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]acetate

To a mixture of the compound 14Gb (26.9 g, 34.7 mmol) in pyridine (100 mL) were added acetic anhydride (9.83 mL, 104 mmol) and DMAP (424 mg, 3.47 mmol), and the mixture was stirred at room temperature for 4 hours. To the reaction

solution were added ethyl acetate (400 mL) and saturated sodium bicarbonate water (200 mL), and the mixture was stirred vigorously for 20 minutes, and the organic layer was separated with a separatory funnel. The organic layer was washed with water (2 × 200 mL) and saturated brine (200 mL), and then dried over anhydrous sodium sulfate. After insoluble materials were filtered off, the solvents were evaporated under reduced pressure, and the resulting residue was azeotroped with toluene to obtain a compound 14Gc (30.5 g). The product was used as itself without purification in a next step.

MS (ESI): m/z = 754 $[M+H]^+$

[0291]

Example 109 Synthesis of Compound 14Gd

N'-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-N,N-dimethylformamidine

To a solution of the compound 14Gc (30.5 g, 35.9 mmol) in 1,4-dioxane (300 mL) was added methyl hydrazine (18.9 mL, 359 mmol), and the mixture was stirred at room temperature for 2 hours, followed by stirring at 50 °C for 3 hours. Water (100 mL) was added to the reaction solutions, the mixture was evaporated under reduced pressure. Chloroform (250 mL) was added thereto, and after stirring for a while,

the mixture was passed through a phase separator, and the organic layer was separated with a separatory funnel. After the solvents were evaporated, the solvents were azeotroped with toluene. The resulting residue was suspended with IPE
5 (100 mL), the mixture was placed in ultrasonic bath for 20 minutes, and the precipitated materials were then collected by filtration, and the filtered materials were air dried followed by drying under reduced pressure.

To the solution of the resulting residue in DMF (75 mL,
10 970 mmol) was added N,N-dimethyl formamide dimethylacetal (50 mL, 375 mmol), and the mixture was stirred at room temperature for 2 hours. The reaction solvents were evaporated under reduced pressure, and the resulting concentrated residue was purified roughly by silica gel
15 chromatography (SiO₂: chloroform/methanol = 100/0 to 90/10, SiO₂: ethyl acetate/methanol = 90/10 to 72/28). The crude products were washed by suspending in ethyl acetate (200 mL), and collected by filtration, and the filtered materials were dried under reduced pressure to a compound 14Gd (11.2 g,
20 yield 40%).

MS (ESI): m/Z = 748 [M+H]⁺

[0292]

Example 110 Synthesis of Compound 14Ge

N'-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
25 methoxy]methyl]-7-[2-cyanoethoxy-

(diisopropylamino)phosphanyl]oxy-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-N,N-dimethylformamidine (Present compound 35)

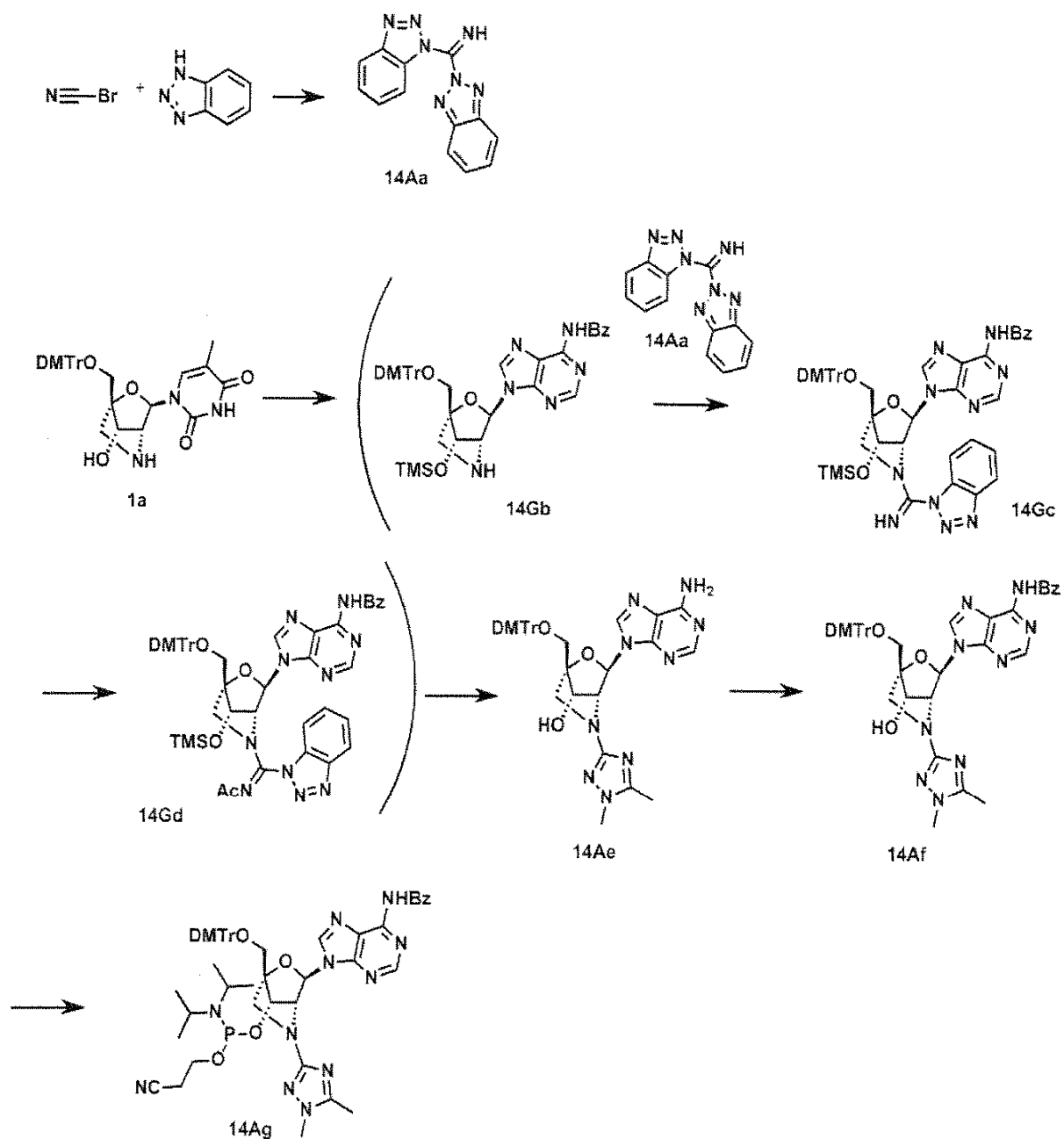
5 To a solution of the compound 14Gd (6.00 g, 7.60 mmol) in dichloromethane (60 mL) was added DIPEA (5.30 mL, 30.6 mmol) under ice-cooling, followed by adding dropwise 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (5.12 mL, 23.0 mmol), and the mixture was stirred at room temperature
10 for 6 hours. The reaction solution was cooled under ice-cooling, and thereto were added saturated sodium bicarbonate water (120 mL) and chloroform (160 mL), and the mixture was stirred vigorously for 15 minutes, and the organic layer was then separated with a separatory funnel, and dried over
15 anhydrous sodium sulfate. After insoluble materials were filtered off, the solvents were evaporated under reduced pressure, and the resulting residue was purified by silica gel chromatography (SiO₂: ethyl acetate/methanol = 100/0 to 84/16) to obtain a compound 14Ge as the present compound 35
20 (5.59 g, yield 74%).

MS (ESI): m/Z = 948 [M+H]⁺

³¹P-NMR (CDCl₃) δ: 148.52, 148.59

[0293]

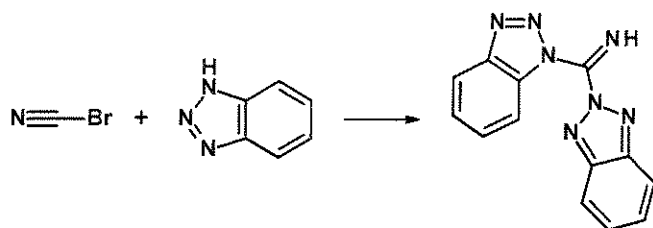
Synthesis of ALNA[Trz]-A



[0294]

Example 111 Synthesis of Compound 14Aa

Benzotriazol-1-yl (Benzotriazol-2-yl)methaneimine



To a solution of benzotriazole (7.70 g, 64.6 mmol) in EtOH (135 mL) was added dropwise a solution of cyanogen bromide (3.41 g, 32.2 mmol) in acetone (15 mL) slowly under ice-cooling, followed by adding dropwise a solution of 2N sodium hydroxide (16.2 mL, 32.4 mmol) slowly. After the mixture was stirred for 30 minutes under ice-cooling, the precipitated materials were collected by filtration, and the filtered materials were washed with cold ethanol (80 mL). The mixture was air dried followed by drying under reduced pressure gave a compound 14Aa (5.14 g, yield 61%).

¹H NMR (400 MHz, CHLOROFORM-*d*) δ ppm 6.92 - 6.98 (m, 1 H), 7.46 - 7.55 (m, 2 H), 7.58 - 7.65 (m, 1 H), 7.77 (ddd, J=8.35, 7.06, 1.03 Hz, 1 H), 8.17 - 8.22 (m, 1 H), 8.22 - 8.27 (m, 1 H), 8.31 - 8.37 (m, 1 H), 9.67 (s, 1 H)

MS (ESI): Non-detected

[0295]

Example 112 Synthesis of Compound 14Ae

(1R,3R,4R,7S)-3-(6-aminopurin-9-yl)-1-[[bis(4-

methoxyphenyl)-phenyl-methoxy]methyl]-5-(1,5-dimethyl-
1,2,4-triazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-7-ol

To a suspension of N-(9H-purin-6-yl)benzamide (5.15 g,
21.5 mmol) in toluene (200 mL) was injected the compound 1a
5 (10.3 g, 17.9 mmol) portionwise, followed by adding BSA (38
mL, 120 mmol), and the mixture was stirred vigorously at 60
°C for 30 minutes. Thereto was added TMSOTf (0.324 mL, 179
mmol) at the same temperature, and the mixture was stirred
at 60 °C for 15 minutes. The reaction solutions were cooled
10 under ice-cooling, and the reaction solutions were added
vigorously while stirring to a mixed solution of IPE (100
mL), water (100 mL) and saturated sodium bicarbonate water
(100 mL) that was cooled under ice-cooling. After the
mixture was stirred vigorously for 20 minutes, insoluble
15 materials were filtered off by Celite. The organic layer of
the filtrates were separated with a separatory funnel, washed
with saturated brine (200 mL) and then dried over anhydrous
sodium sulfate, and the solvents were evaporated to obtain
the residue of the crude compound 14Gb.

20 To the solution of the resulting crude residue in THF
(200 mL) was added the compound 14Aa (4.72 g, 17.9 mmol),
and the mixture was stirred at room temperature for 92 hours,
and further stirred at 50 °C for 9 hours. To the reaction
solution were added ethyl acetate (400 mL) and saturated
25 sodium bicarbonate water (200 mL), and after stirring for 20

minutes, the organic layer was separated with a separatory funnel. The organic layer was washed with water (300 mL) and saturated brine (100 mL), and then dried over anhydrous sodium sulfate, and insoluble materials were filtered off,
5 and the solvents were evaporated to obtain the residue of the crude compound 14Gc.

To a solution of the resulting residue in pyridine (65 mL) were added acetic anhydride (5.08 mL, 53.8 mmol) and DMAP (219 mg, 1.79 mmol), and the mixture was stirred at
10 room temperature for 3 hours. To the reaction solutions were added ethyl acetate (400 mL) and saturated sodium bicarbonate water (200 mL), and after stirring for 20 minutes, the organic layer was separated with a separatory funnel. The organic layer was washed with water (300 mL) and
15 saturated brine (100 mL) and then dried over anhydrous sodium sulfate. After insoluble materials were filtered off, the solvents were evaporated under reduced pressure, and the residue was azeotroped with toluene to obtain the residue of crude compound 14Gd.

20 To theae solution of the resulting residue in 1,4-dioxane (100 mL) was added dropwise methyl hydrazine (9.44 mL, 179 mmol) under ice-cooling using cold water, and the mixture was stirred at room temperature for 3 hours. The mixture was warmed to 50 °C, and stirred for 10 hours. Water
25 (100 mL) was added to the reaction solutions, and the

solvents were then evaporated under reduced pressure from the mixture, and the residue was extracted with chloroform (2 × 150 mL) using a separatory funnel. The organic layers were combined, the mixture was washed with saturated brine (100 mL), and then dried over anhydrous sodium sulfate. After insoluble materials were filtered off, the solvents were evaporated under reduced pressure, and the resulting residue was purified by silica gel chromatography (NH silica: chloroform/methanol = 100/0 to 99/1) to obtain a compound 14Ae (4.91 g, yield 41%).

MS (ESI): m/z = 675 $[M-H]^-$

[0296]

Example 113 Synthesis of Compound 14Af

N-[9-(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]purin-6-yl]benzamide

To a mixture of the compound 14Ae (4.80 g, 7.10 mmol) and pyridine (10 mL) was added benzoyl chloride (2.48 mL, 21.3 mmol) under ice-cooling, and the mixture was stirred under ice-cooling for 3 hours. To the reaction solutions was added a solution of 2N sodium hydroxide (25 mL, 50 mmol), and the mixture was stirred at room temperature for 2 hours. The reaction solutions were washed with chloroform (50 mL) four times, and the organic layer was passed through a phase

separator, and dried over anhydrous sodium sulfate. The solutions were filtered through Celite and concentrated. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 93/7) to obtain a compound
5 14Af (4.60 g, yield 83%).

MS (ESI): m/z = 781 [M+H]⁺

[0297]

Example 114 Synthesis of Compound 14Ag

N-[9-(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
10 methoxy]methyl]-7-[2-cyanoethoxy-
(diisopropylamino)phosphanyl]oxy-5-(1,5-dimethyl-1,2,4-
triazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]purin-6-
yl]benzamide (Present compound 36)

To a mixture of the compound 14Af (4.26 g, 5.46 mmol)
15 and dichloromethane (50 mL) were added DIPEA (3.78 mL, 21.8
mmol) and 2-cyanoethyl N,N-diisopropylchlorophosphoramidite
(3.66 mL, 16.4 mmol) under ice-cooling, and the mixture was
stirred at room temperature for 2 hours. The reaction
solution was cooled under ice-cooling, and saturated aqueous
20 sodium hydrogen carbonate solution (100 mL) was added to the
reaction solution, and the mixture was extracted with
chloroform (100 mL). The organic layer was dried over
anhydrous sodium sulfate, and insoluble materials were then
flited off, and the solvents were evaporated. The residue
25 was purified by column chromatography (SiO₂: ethyl

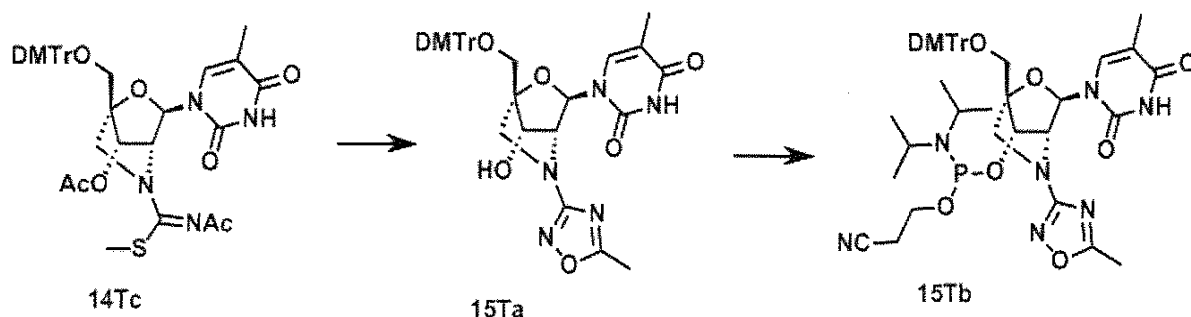
acetate/methanol = 100/0 to 94/6, NH silica: ethyl acetate/methanol = 100/0 to 93/7) to obtain a compound 14Ag as the present compound 36 (3.88 g, yield 67%).

MS (ESI): m/z = 980 $[M+H]^+$

5 ^{31}P -NMR (CDCl_3) δ : 148.84, 149.04

[0298]

Synthesis of ALNA[Oxz]-T



[0299]

10 Example 115 Synthesis of Compound 15Ta

1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-(5-methyl-1,2,4-oxadiazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2,4-dione

15 To a mixture of the compound 14Tc (2.10 g, 2.88 mmol) and EtOH (14.4 mL) was added hydroxyl amine (50% aqueous solution, 1.90 mL, 28.8 mmol), and the mixture was stirred at room temperature for 2 days. The reaction solution was concentrated, and the residue was purified by column chromatography (SiO_2 : chloroform/methanol = 100/0 to 90/10)

20

to obtain a compound 15Ta (1.34 g, yield 71%).

HRMS(MALDI): calculated value as $C_{35}H_{35}N_5O_8$ $[M+Na]^+$: 676.2378,
measured value: 676.2363

[0300]

5 Example 116 Synthesis of Compound 15Tb

3-[[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-(5-
methyl-1,2,4-oxadiazol-3-yl)-2-oxa-5-
azabicyclo[2.2.1]heptan-7-yl]oxy-

10 (diisopropylamino)phosphanyl]oxypropanenitrile (Present
compound 37)

To a mixture of the compound 15Ta (300 mg, 0.459 mmol),
dichloromethane (4.6 mL) and DIPEA (240 μ L, 1.38 mmol) was
added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite
15 (205 μ L, 0.918 mmol), and the mixture was stirred at room
temperature under nitrogen atmosphere for 5 days. Saturated
aqueous sodium hydrogen carbonate solution was added to the
reaction solution, and the mixture was extracted with
chloroform. The organic layer was passed through a phase
20 separator, and the solvents were evaporated. The residue
was purified by column chromatography (SiO_2 : hexane/ethyl
acetate= 60/40 to 10/90, Diol: hexane/ethyl acetate = 70/30
to 20/80, NH: chloroform/methanol = 100/0 to 95/5) to obtain
a compound 15Tb as the present compound 37 (214 mg, yield
25 55%).

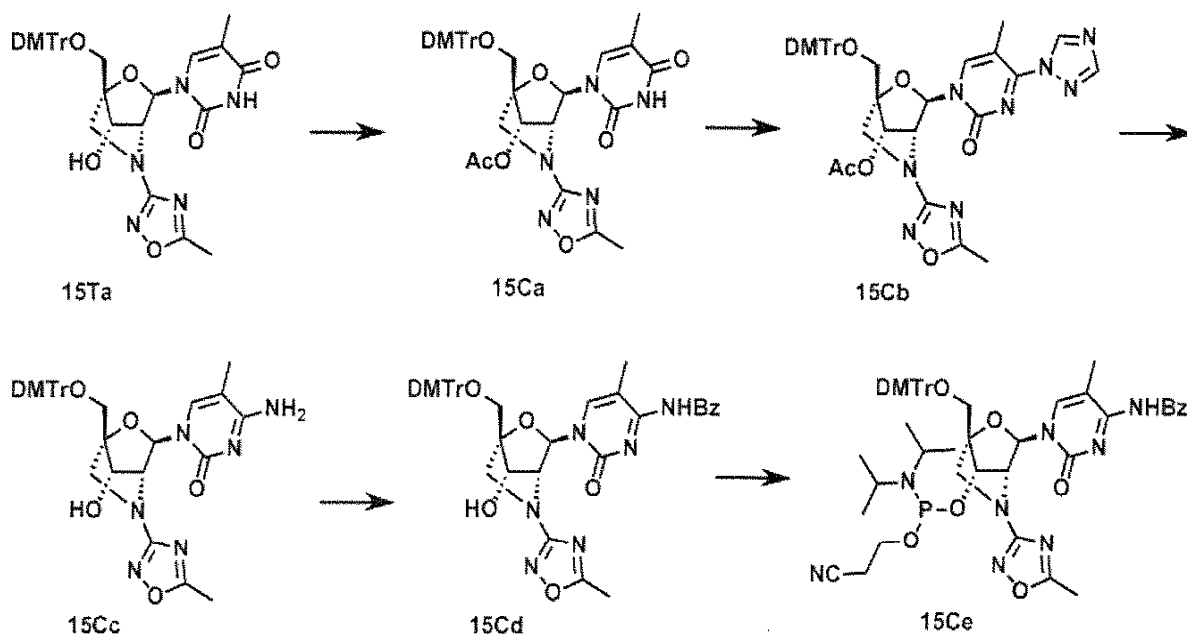
HRMS(MALDI): calculated value as $C_{44}H_{52}N_7O_9P$ $[M+Na]^+$: 876.3456,

measured value: 876.3441

^{31}P -NMR ($CDCl_3$) δ : 148.83, 149.09

[0301]

5 Synthesis of ALNA[Oxz]-mC



[0302]

Example 117 Synthesis of Compound 15Ca

[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-

10 methoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-(5-methyl-1,2,4-oxadiazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]acetate

To a mixture of the compound 15Ta (999 mg, 1.53 mmol),
 DMAP (18.7 mg, 0.153 mmol) and pyridine (7.7 mL) was added
 15 acetic anhydride (217 μ L, 2.30 mmol), and the mixture was

stirred at room temperature for 2 days. Ethyl acetate was added to the reaction solution, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, water, and saturated brine. The organic layer was dried
5 over anhydrous magnesium sulfate, and insoluble materials were filtered off, and concentrated under reduced pressure to obtain a compound 15Ca (1.10 g, the reaction has been proceeded quantitatively).

HRMS(MALDI): calculated value as $C_{37}H_{37}N_5O_9$ $[M+Na]^+$: 718.2483,
10 measured value: 718.2470

[0303]

Example 118 Synthesis of Compound 15Cb

[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-5-(5-methyl-1,2,4-oxadiazol-3-yl)-3-[5-
15 methyl-2-oxo-4-(1,2,4-triazol-1-yl)pyrimidin-1-yl]-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]acetate

To a mixture of the compound 15Ca (1.05 g, 1.51 mmol), 1,2,4-triazole (939 mg, 13.6 mmol), DIPEA (2.63 mL, 15.1 mmol) and acetonitrile (15 mL) was added phosphoryl chloride
20 (239 μ L, 2.57 mmol) under ice-cooling, and the mixture was stirred at room temperature under nitrogen atmosphere overnight. Ethyl acetate was added to the reaction solution, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, and saturated brine. The
25 organic layer was dried over anhydrous magnesium sulfate,

and insoluble materials were filtered off, and the filtrates were concentrated under reduced pressure to obtain a compound 15Cb (1.32 g) as a crude product.

HRMS(MALDI): calculated value as $C_{39}H_{38}N_8O_8$ $[M+Na]^+$: 769.2705,

5 measured value: 769.2707

[0304]

Example 119 Synthesis of Compound 15Cc

4-amino-1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-(5-methyl-1,2,4-oxadiazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-pyrimidin-2-one

10

To a mixture of the compound 15Cb (1.30 g, 1.51 mmol, crude product) and acetonitrile (15.1 mL) was added 28% ammonia water (10.1 mL), and the mixture was stirred at room temperature overnight. Water was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and concentrated to obtain a compound 15Cc (0.96 g, yields over 2 steps 97%).

15

HRMS(MALDI): calculated value as $C_{35}H_{36}N_6O_7$ $[M+Na]^+$: 675.2538, measured value: 675.2539

20

[0305]

Example 120 Synthesis of Compound 15Cd

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-(5-methyl-1,2,4-oxadiazol-3-

25

yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide

To a mixture of the compound 15Cc (0.93 g, 1.42 mmol) and pyridine (7.1 mL) was added benzoic anhydride (482 mg, 2.13 mmol), and the mixture was stirred at room temperature overnight. To the reaction solution was added 2N aqueous sodium hydroxide solution (7.1 mL), and the mixture was stirred at room temperature for 1 hour. Ethyl acetate was added to the reaction solution, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and insoluble materials were filtered off, and the filtrates were concentrated under reduced pressure. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 90/10) to obtain a compound 15Cd (958 mg, yield 89%).

HRMS(MALDI): calculated value as C₄₂H₄₀N₆O₈ [M+Na]⁺: 779.2800, measured value: 779.2794

[0306]

20 Example 121 Synthesis of Compound 15Ce

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(5-methyl-1,2,4-oxadiazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 38)

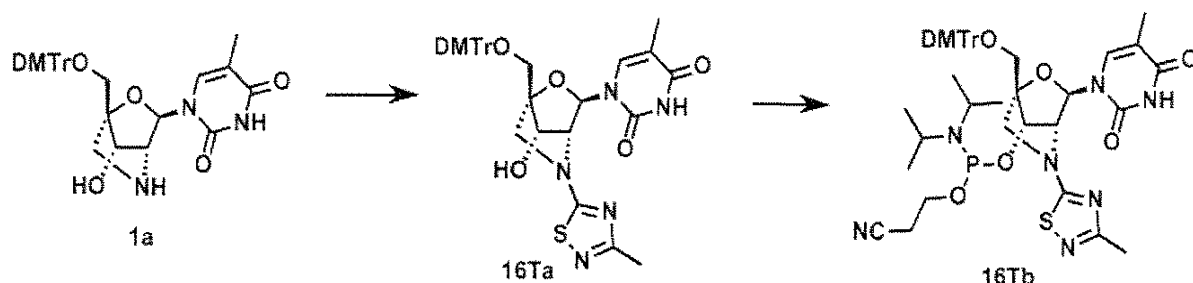
To a mixture of the compound 15Cd (949 mg, 1.25 mmol), dichloromethane (12.5 mL) and DIPEA (0.653 mL, 3.75 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (0.559 mL, 2.50 mmol) under ice-cooling, and the mixture was stirred at room temperature under nitrogen atmosphere overnight. Saturated aqueous sodium hydrogen carbonate solution was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and the solvents were evaporated. The residue was purified by column chromatography (SiO₂: hexane/ethyl acetate = 80/20 to 30/70, NH: hexane/ethyl acetate = 50/50 to 0/100) to obtain a compound 15Ce as the present compound 38 (834 mg, yield 70%).

HRMS(MALDI): calculated value as C₅₁H₅₇N₈O₉P [M+Na]⁺: 979.3878, measured value: 979.3876

³¹P-NMR (CDCl₃) δ: 148.98, 149.27

[0307]

Synthesis of ALNA[Tdz]-T



[0308]

Example 122 Synthesis of Compound 16Ta

1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-(3-methyl-1,2,4-thiadiazol-5-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-pyrimidin-2,4-dione

A mixture of the compound 1a (3.00 g, 5.25 mmol), 5-chloro-3-methyl-1,2,4-thiadiazole (848 mg, 6.30 mmol), DIPEA (2.72 mL, 15.7 mmol) and DMSO (26 mL) was stirred at 120 °C for 2 hours. Ethyl acetate was added to the reaction solution, and the mixture was washed with water and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and insoluble materials were filtered off, and the filtrates were concentrated under reduced pressure. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 95/5) to obtain a compound 16Ta (3.30 g, yield 94%).

MS(ESI): m/z = 670 (M+H)⁺

[0309]

Example 123 Synthesis of Compound 16Tb

3-[[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-(3-methyl-1,2,4-thiadiazol-5-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile (Present compound 39)

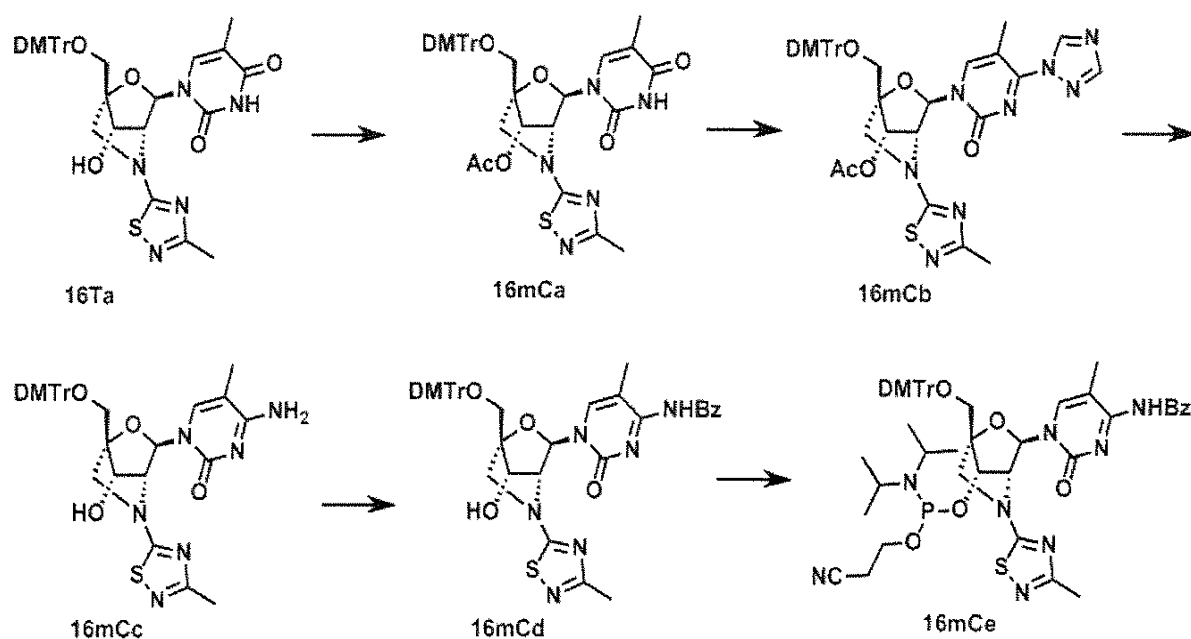
To a mixture of the compound 16Ta (2.26 g, 3.37 mmol), dichloromethane (17 mL), DIPEA (1.75 mL, 10.1 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (1.51 mL, 6.77 mmol), and the mixture was stirred at room temperature under nitrogen atmosphere overnight. Saturated aqueous sodium hydrogen carbonate solution was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and the solvents were evaporated. The residue was purified by column chromatography (SiO₂: hexane/ethyl acetate = 70/30 to 20/80, Diol: hexane/ethyl acetate = 70/30 to 20/80) to obtain a compound 16Tb as the present compound 39 (2.45 g, yield 84%).

MS(ESI): m/z = 870 (M+H)⁺

³¹P-NMR (CDCl₃) δ: 149.23, 149.51

[0310]

Synthesis of ALNA[Tdz]-mC



[0311]

Example 124 Synthesis of Compound 16mCa

[[(1R, 3R, 4R, 7S) -1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-(3-methyl-1,2,4-thiadiazol-5-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]acetate

To a mixture of the compound 16Ta (5.92 g, 8.84 mmol), DMAP (120 mg, 0.982 mmol) and pyridine (18 mL) was added acetic anhydride (1.25 mL, 13.2 mmol), and the mixture was stirred at room temperature overnight. The reaction solution was concentrated, and ethyl acetate was added to the residue, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate,

and insoluble materials were filtered off, and the filtrates were concentrated to obtain a compound 16mCa (5.69 g, yield 90%).

MS(ESI): m/z = 712 (M+H)⁺

5 [0312]

Example 125 Synthesis of Compound 16mCb

[[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-3-[5-methyl-2-oxo-4-(1,2,4-thiadiazole-1-yl)pyrimidin-1-yl]-5-(3-methyl-1,2,4-thiadiazol-5-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]acetate

10

To a mixture of the compound 16mCa (5.52 g, 7.75 mmol), 1,2,4-triazole (4.82 g, 69.8 mmol), DIPEA (13.4 mL, 77.5 mmol) and acetonitrile (39 mL) was added phosphoryl chloride (1.23 mL, 13.2 mmol) under ice-cooling, and the mixture was stirred at room temperature under nitrogen atmosphere for 2 hours. Ethyl acetate was added to the reaction solution, and the mixture was extracted with saturated aqueous sodium hydrogen carbonate solution, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and insoluble materials were filtered off, and the filtrates were concentrated to obtain a compound 16mCb as a crude product (6.18 g, containing impurities).

15

20

[0313]

Example 126 Synthesis of Compound 16mCc

25 4-amino-1-[[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-

methoxy]methyl]-7-hydroxy-5-(3-methyl-1,2,4-thiadiazol-5-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-pyrimidin-2-one

To a mixture of the compound 16mCb (6.18 g, 7.75 mmol, containing impurities) and acetonitrile (39 mL) was added 28% ammonia water (39 mL), and the mixture was stirred at room temperature for 4 days. Water was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was dried over anhydrous magnesium sulfate, and insoluble materials were filtered off, and the filtrates were concentrated. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 90/10) to obtain a compound 16mCc (4.50 g, yields over 2 steps 87%). MS(ESI): m/z = 667 (M-H)⁻

[0314]

Example 127 Synthesis of Compound 16mCd

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-(3-methyl-1,2,4-thiadiazol-5-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide

A mixture of the compound 16mCc (4.37 g, 6.53 mmol), benzoic anhydride (2.22 g, 9.81 mmol) and pyridine (13 mL) was stirred at room temperature overnight. To the reaction solution was added 2N aqueous sodium hydroxide solution (13 mL, 26 mmol), and the mixture was stirred at room temperature

for 1.5 hours. Ethyl acetate was added to the reaction solution, and the mixture was washed with saturated aqueous sodium hydrogen carbonate solution, and saturated brine. The organic layer was dried over anhydrous magnesium sulfate, and insoluble materials were filtered off, and the filtrates were concentrated. The residue was purified by column chromatography (SiO₂: chloroform/methanol = 100/0 to 90/10) to obtain a compound 16mCd (4.76 g, yield 94%).

MS(ESI): m/z = 773 (M+H)⁺

[0315]

Example 128 Synthesis of Compound 16mCe

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(3-methyl-1,2,4-thiadiazol-5-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide (Present compound 40)

To a mixture of the compound 16mCd (4.62 g, 5.98 mmol), dichloromethane (12 mL) and DIPEA (3.10 mL, 17.9 mmol) was added 2-cyanoethyl N,N-diisopropylchlorophosphoramidite (2.67 mL, 12.0 mmol), and the mixture was stirred at room temperature under nitrogen atmosphere overnight. Saturated aqueous sodium hydrogen carbonate solution was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and the solvents were evaporated. The residue

was purified by column chromatography (SiO₂: hexane/ethyl acetate = 80/20 to 30/70, NH: hexane/ethyl acetate = 70/30 to 20/80) to obtain a compound 16mCe as the present compound 40 (3.57 g, yield 61%).

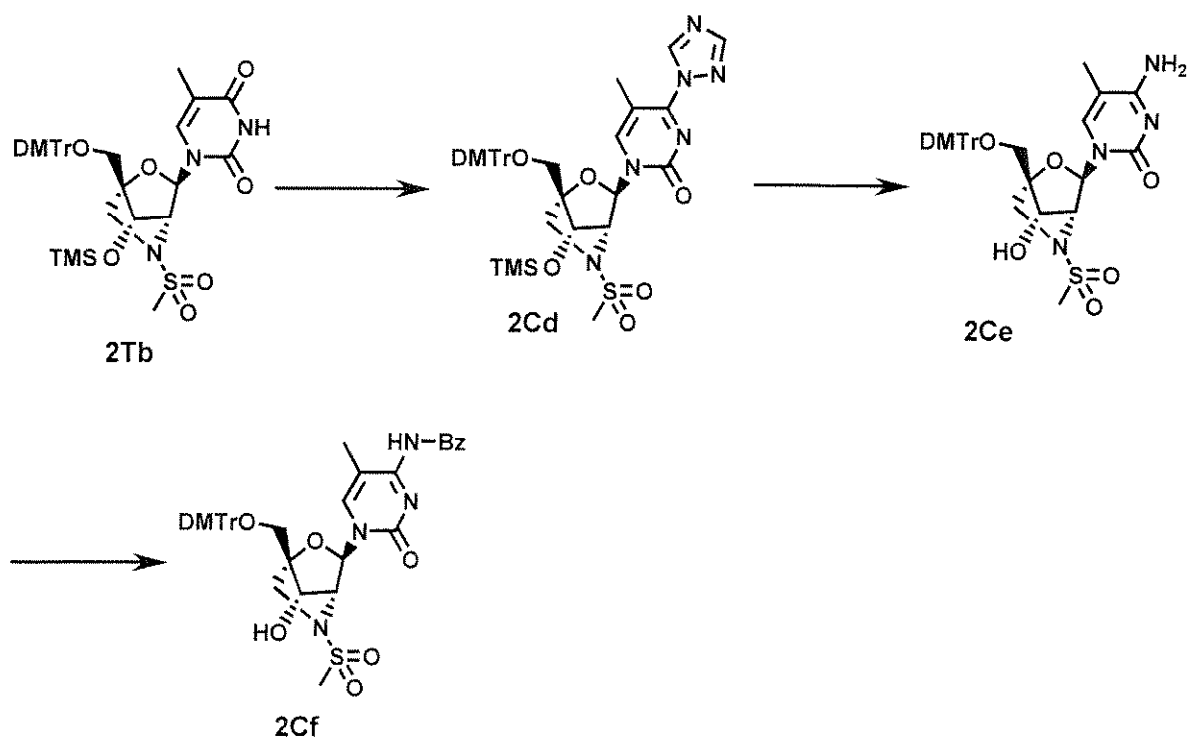
5 MS(ESI): m/z = 974 ($M+H$)⁺

³¹P-NMR (CDCl₃) δ : 149.45, 149.78

[0316]

ALNA[Ms]-mC can be also synthesized by transforming a nucleic acid base moiety of ALNA[Ms]-T nucleoside as follows.

10 Synthesis of Compound 2Cf



[0317]

Example 129 Synthesis of Compound 2Cd

3-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl]-

methoxy)methyl]-5-methylsulfonyl-7-trimethylsilyloxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-6-(1,2,4-triazol-1-yl)-1,6-dihydropyrimidin-2-one

Compound 2Tb (14.1 g, 0.019 mol) was solubilized in acetonitrile (310 mL), and thereto was added triethylamine (97.29 mL, 0.961 mol), and the mixture was stirred at 0 °C for 15 minutes. To the reaction solution was added 1,2,4-triazole (30.36 g, 0.439 mol), and the mixture was stirred at 0 °C for 15 minutes, and thereto was then added phosphoryl chloride (4.44 mL, 0.028 mol), and the mixture was stirred at 0 °C for 15 minutes, and warmed to room temperature for 2 hours. The reaction solution was concentrated under reduced pressure, and thereto were added ethyl acetate (200 mL) and water (100 mL), and the mixture was separated with a separatory funnel. The organic layer was washed with sodium carbonate (100 mL) twice, dried over sodium sulfate, and concentrated under reduced pressure to obtain a compound 2Cd as crude product (11.8 g).

[0318]

Example 130 Synthesis of Compound 2Ce

6-amino-3-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-pyrimidin-2-one

The crude compound 2Cd (11.8 g) was solubilized in 1,4-dioxane (118 mL), and 25% ammonia water (118 mL) was added

thereto, and the mixture was stirred at room temperature overnight. The reaction solution was concentrated under reduced pressure, and the resulting residue was purified by silica gel chromatography (dichloromethane/methanol, 95/5 to 94/6). The resulting mixture was stirred in a mixed solution of dichloromethane/hexane (10/90), and the precipitated materials were collected by filtration to obtain a compound 2Ce (8.3 g, yields over two steps 64%).

MS(ESI): $m/z = 649.3$ (M+H)⁺

[0319]

Example 131 Synthesis of Compound 2Cf

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]dihydropyrimidin-2-one

To a solution of the compound 2Ce (8.3 g, 0.012 mol) in DMF (63 mL) was added benzoic anhydride (2.57 g, 0.011 mol), and the mixture was stirred at room temperature for 24 hours. The reaction solution was diluted with ethyl acetate (200 mL), and the mixture was washed with water (100 mL) twice. The organic layer was washed with water (100 mL) and saturated brine (100 mL) successively, and dried over sodium sulfate. The resulting mixture was concentrated under reduced pressure, and the resulting residue was purified by silica gel chromatography (dichloromethane/methanol,

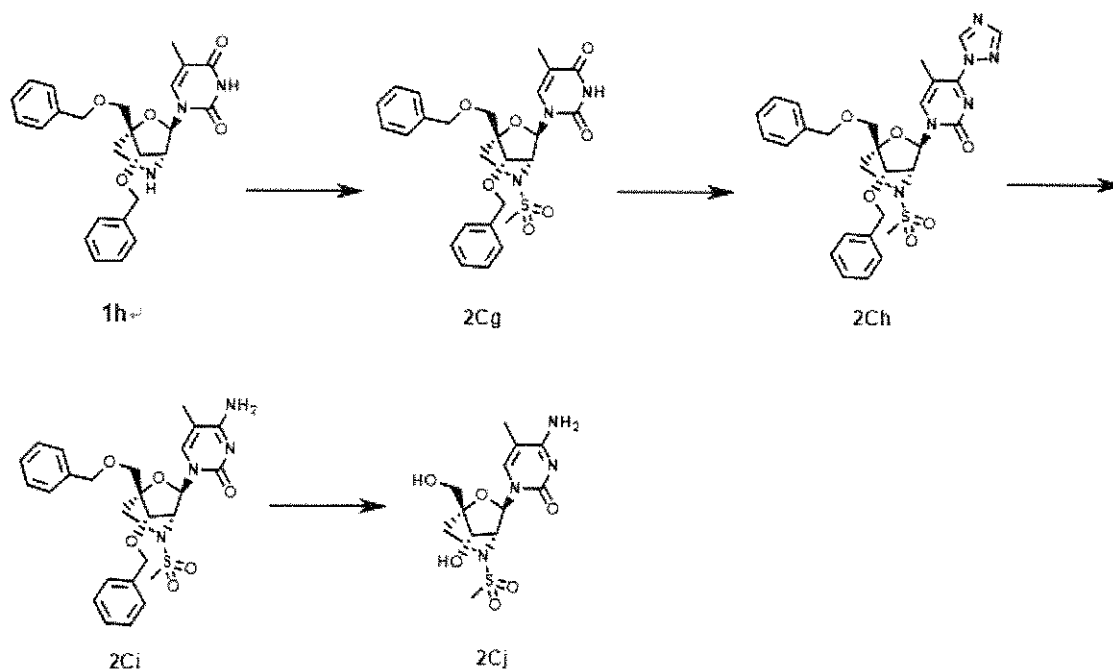
99.5/0.5 to 99/1). The resulting mixture was stirred in a mixed solution of dichloromethane/hexane (10/90), and the precipitated materials were collected by filtration to obtain the compound 2Cf (4.0 g, yield 47%).

5 MS(ESI): $m/z = 753.1$ ($M+H$)⁺

[0320]

Also ALNA[Ms]-mC nucleoside can be also synthesized as follows.

Synthesis of Compound 2Cj



10

[0321]

Example 132 Synthesis of Compound 2Cg

1-[(1R,3R,4R,7S)-7-benzyloxy-1-(benzyloxymethyl)-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2,4-dione

15

To a mixed solution of 1-[(1R,3R,4R,7S)-7-benzyloxy-1-(benzyloxymethyl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-2,4-dione (1h) (20.00 g, 21.61 mmol), which was synthesized according to the method described in WO 2017/047816 A1, in dichloromethane (200 mL) was added triethylamine (15.5 mL, 111 mmol), and thereto was added mesyl chloride (4.15 mL, 53.4 mmol) in ice bath, and the mixture was stirred in ice bath for 4.5 hours. 5% Sodium bicarbonate water (60 mL) was added thereto under ice-cooling, and the mixture was stirred and separated with a separatory funnel. The aqueous layer was extracted with dichloromethane (40 mL) and the extracts were combined and mixed with the dichloromethane organic layer. The organic layer was washed with 15% brine (60 mL), and the solvents were then evaporated. The concentrated residue was washed with methanol (90 mL) as a suspension. The resulting precipitated materials were collected by filtration, dried under vacuum to obtain a compound 2Cg (21.94 g, yield 94%).

MS(ESI): m/z = 528 (M+H)⁺

[0322]

Example 133 Synthesis of Compound 2Ch

1-[(1R,3R,4R,7S)-7-benzyloxy-1-(benzyloxymethyl)-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-4-(1,2,4-triazol-1-yl)pyrimidin-2-one

Compound 2Ch (3.4 g) as a crude product was obtained

from the compound 2Cg (3.0 g, 5.6 mmol) similarly to Example 129.

[0323]

Example 134 Synthesis of Compound 2Ci

5 4-amino-1-[(1R,3R,4R,7S)-7-benzyloxy-1-(benzyloxymethyl)-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-pyrimidin-2-one

Compound 2Ci (2.6 g, yield 87%) was obtained from the compound 2Ch (3.4 g, 5.8 mmol) similarly to Example 130.

10 MS(ESI): m/z = 527.2 (M+H)⁺

[0324]

Example 135 Synthesis of Compound 2Cj

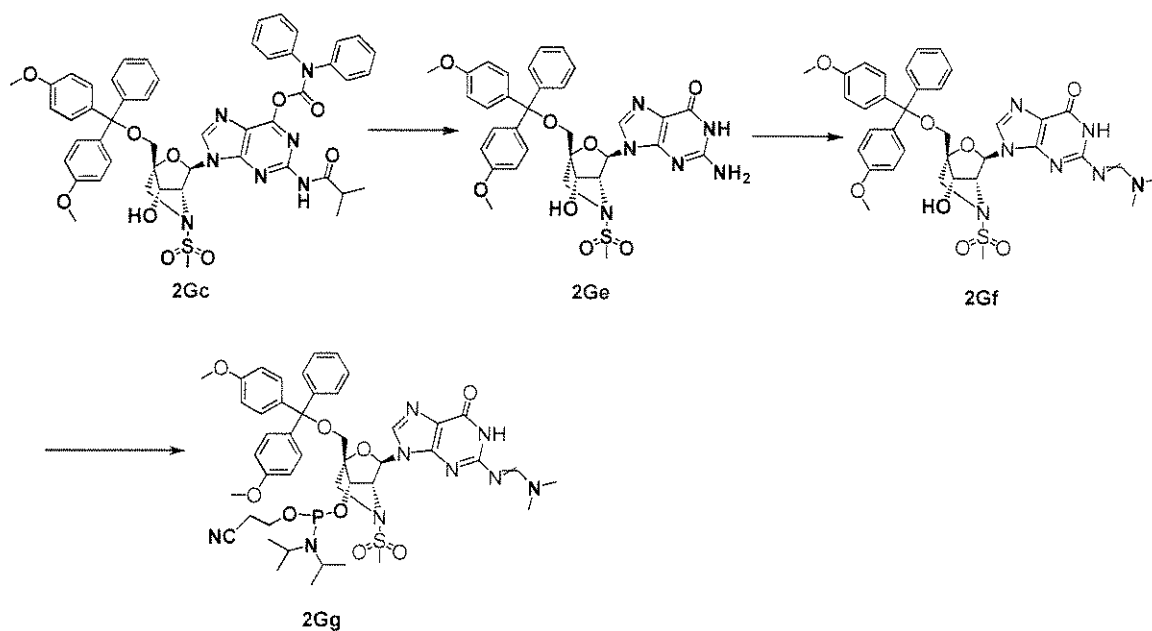
15 4-amino-1-[(1R,3R,4R,7S)-7-hydroxy-1-(hydroxymethyl)-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-pyrimidin-2-one

The compound 2Ci (2.6 g, 4.9 mmol) was solubilized in acetic acid (28.6 mL), and thereto was added 20% palladium/carbon (0.35 g), and the mixture was stirred at room temperature under hydrogen atmosphere for 16 hours.
20 Insoluble materials were filtered off, and filtrates were concentrated under reduced pressure. To the residue were added 1-butanol and IPE, and the mixture was stirred, and the precipitated materials were collected by filtration. The resulting solids were recrystallized with methanol to obtain
25 a compound 2Cj (1.0 g, yield 58%).

MS(ESI): $m/z = 346.9$ (M+H)⁺

[0325]

ALNA[Ms]-G wherein a guanine moiety is protected with a dimethylamino methylene group can be synthesized as follows.
 5 Here in the case where a monomer containing such a protection style is used in an oligomer synthesis, it is possible to control a production of by-products that may occur when oligomer synthesis is performed using Compound 2Gd wherein the 6-position of guanine is protected with DPC.



[0326]

Example 136 Synthesis of Compound 2Ge

2-amino-9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-1H-purin-6-one

To a solution of the compound 2Gc (1000 mg, 1.1 mmol)

in tetrahydrofuran (10 mL) was added 28% ammonia water (5 mL) at room temperature, the mixture was stirred at room temperature overnight. Thereto was further added 28% ammonia water (15 mL), and the mixture was stirred at 70 °C for 8 hours. Thereto were added water and dichloromethane at room temperature, and after stirring the mixture, the organic layer was recovered, and dried over sodium sulfate. The residue was filtered, and the solvents were evaporated, and the resulting mixture was purified by silica gel chromatography (chloroform/methanol, 97/3 to 92/8) to obtain a compound 2Ge (620 mg, yield 86%).

MS(ESI): $m/z = 675$ ($M+H$)⁺

[0327]

Example 137 Synthesis of Compound 2Gf

N'-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-N,N-dimethylformamidine

The compound 2Ge (150 mg, 0.22 mmol) was solubilized in tetrahydrofuran (2.22 mL), and thereto was added N,N-dimethyl formamide dimethylacetal (132 mg, 1.11 mmol) at room temperature, and the mixture was stirred. Ethyl acetate was added thereto at room temperature, and the mixture was extracted (2 mL) once. The resulting organic layer was washed with brine, and dried over sodium sulfate. The

residue was filtered, and the solvents were evaporated, and the residue was purified by silica gel chromatography (chloroform/methanol = 99/1, chloroform/methanol = 93/7) to obtain a compound 2Gf (145 mg, yield 89%).

5 MS(ESI): m/z = 728 (M-H)⁻

[0328]

Example 138 Synthesis of Compound 2Gg

N'-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-

10 cyanoethoxy(diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-N,N-dimethylformamidine

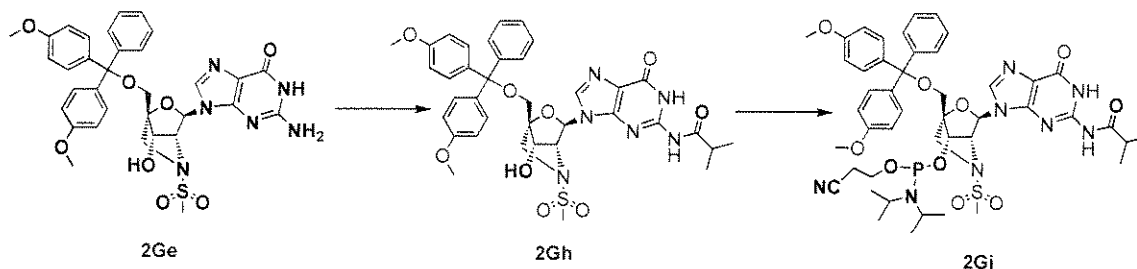
The compound 2Gf (900 mg, 1.2 mmol) was solubilized in dichloromethane (15 mL), and thereto were added N,N-
 15 diisopropyl ethylamine (747 mg, 5.8 mmol) and 2-cyanoethyl N,N-diisopropylchloro phosphoroamizide (1061 mg, 4.5 mmol) at room temperature, and the mixture was stirred overnight. Thereto were added saturated bicarbonate water and dichloromethane, and the organic layer was separated. The
 20 resulting organic layer was dried over sodium sulfate, and the residue was filtered. After the solvents were evaporated, the resulting mixture was purified by amino silica gel chromatography (chloroform/methanol, 100/0 to 97/3), followed by diol silica gel chromatography
 25 (chloroform/methanol, 100/0 to 97/3) to obtain a compound

2Gg (690 mg, yield 60%).

MS(ESI): $m/z = 930$ (M+H)⁺

[0329]

ALNA[Ms]-G wherein a guanine moiety is protected with
 5 only an isobutyloyl group can be synthesized as follows.
 Here in the case where a monomer containing such a protection
 style is used in an oligomer synthesis, it is possible to
 control a production of by-products that may occur when
 oligomer synthesis is performed using Compound 2Gd wherein
 10 the 6-position of guanine is protected with DPC.



[0329]

Example 139 Synthesis of Compound 2Gh

N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
 15 methoxy]methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-
 azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-2-
 methylpropane amide

The compound 2Ge (100 mg, 0.15 mmol) was solubilized in
 pyridine (1.4 mL), and thereto were added N,N-dimethyl-4-
 20 aminopyridine (29 mg, 0.24 mmol), and 2-methyl propanoyl-2-
 methyl propanoate (100 μ L, 0.6 mmol) at room temperature,
 and the mixture was stirred at 100 °C for 4 hours. Thereto

was added further 2-methyl propanoyl-2-methyl propanoate (100 μ L, 0.6 mmol), and the mixture was stirred at 100 $^{\circ}$ C for 2 hours. Thereto were added saturated bicarbonate water and sodium acetate at room temperature, and the organic layer was separated. The resulting organic layer was dried over sodium sulfate. The residue was filtered, and the solvents were evaporated, and purified by silica gel chromatography (hexane/ethyl acetate, 50/50 to 20/80). Further the obtained compound was solubilized in tetrahydrofuran (1 mL), and thereto was added 0.1 N aqueous sodium hydroxide solution (1 mL), and the mixture was stirred at 0 $^{\circ}$ C for 40 minutes. To the reaction solution were added water and ethyl acetate, and the organic layer was separated, and then washed with aqueous ammonium chloride solution, and the organic was separated again. After the resulting mixture was dried over sodium sulfate, the residue was filtered, and the solvents were evaporated to obtain a compound 2Gh (56 mg, yield 50%). MS(ESI): m/z = 745 (M+H) $^{+}$

[0331]

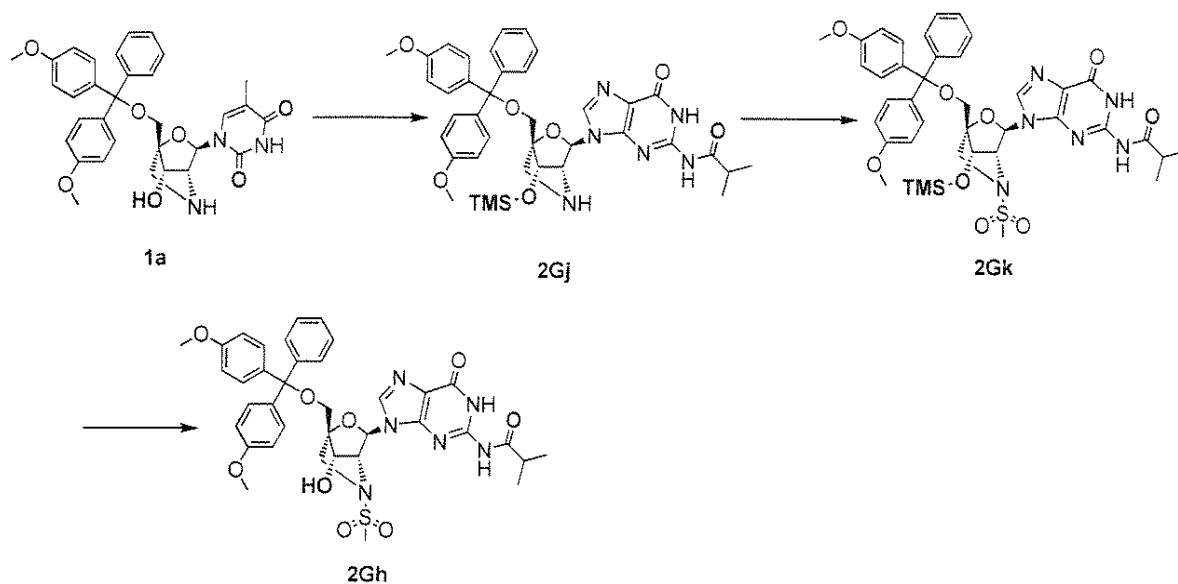
Example 140 Synthesis of Compound 2Gi

N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy(diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-2-methylpropane amide

Molecular sieve 3A was added to dichloromethane (3.6 mL) and the mixture was stirred for 1 hour, and thereto were added the compound 2Gh (500 mg, 0.67 mmol), 1-methyl imidazole (10.9 mg, 0.1 mol) and tetrazole (84 mg, 1.2 mmol),
5 and 2-Cyanoethyl N,N,N',N'-Tetraisopropylphosphordiamidite (949 mg, 3.1 mmol) at room temperature, and the mixture was stirred for 6 hours. Thereto was added saturated bicarbonate water at room temperature, and the mixture was extracted with dichloromethane twice. The resulting organic layer was
10 dried over sodium sulfate, and the residue was filtered. After the solvents were evaporated, the resulting residue was purified by silica gel chromatography (hexane/ethyl acetate, 50/50 to 25/75) to obtain a compound 2Gi (500 mg, yield 79%).

15 MS(ESI): $m/z = 946$ (M+H)⁺
[0332]

Also ALNA[Ms]-G monomer wherein a guanine moiety is protected with only an isobutyloyl group can be also synthesized directly by a transglycosylation from the
20 compound 1a as follows.



[0333]

Example 141 Synthesis of Compound 2Gj

N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
 5 methoxy)methyl]-7-trimethylsilyloxy-2-oxa-5-
 azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-2-
 methylpropane amide

To a mixed solution of the compound 1a (5.0 g, 8.75
 mmol) and 2-methyl N-(6-oxo-1,9-dihydropurin-2-yl)propane
 10 amide (3.87 g, 17.5 mmol) in 1,2-dichloroethane (30 mL) was
 added BSA (21.4 mL, 87.5 mmol), and the mixture was stirred
 at 60 °C for 30 minutes. Successively, thereto was added
 TMSOTf (0.254 mL, 1.31 mmol), and the mixture was stirred
 for 1 hour and a half. After allowed to stand to cool, to
 15 the reaction solutions were added chloroform (50 mL) and
 saturated aqueous sodium hydrogen carbonate solution (50 mL),

and the mixture was stirred. After suction filtering an insoluble materials off, the organic layer was washed with water and saturated brine, dried over magnesium sulfate, and concentrated. The resulting residue was purified by silica gel chromatography (chloroform/methanol, 100/0 to 90/10) to obtain a compound 2Gj (3.83 g, yield 59%).

MS(ESI): $m/z = 739.5$ (M+H)⁺

[0334]

Example 142 Synthesis of Compound 2Gk

N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-5-methylsulfonyl-7-trimethylsilyloxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-2-methylpropane amide

To a solution of the compound 2Gj (1.0 g, 1.35 mmol), pyridine (0.327 mL, 4.06 mmol) in dichloromethane (7 mL) was added methane sulfonic anhydride (284 mg, 1.63 mmol) under ice-cooling, and the mixture was stirred under ice-cooling for 45 minutes. Saturated aqueous sodium hydrogen carbonate solution was added to the reaction solution, and the mixture was extracted with chloroform. The organic layer was passed through a phase separator, and concentrated. The resulting residue was purified by silica gel chromatography (chloroform/methanol, 100/0 to 90/10) to obtain a compound 2Gk (657 mg, yield 59%).

MS(ESI): $m/z = 817.4$ (M+H)⁺

[0335]

Example 143 Synthesis of Compound 2Gh

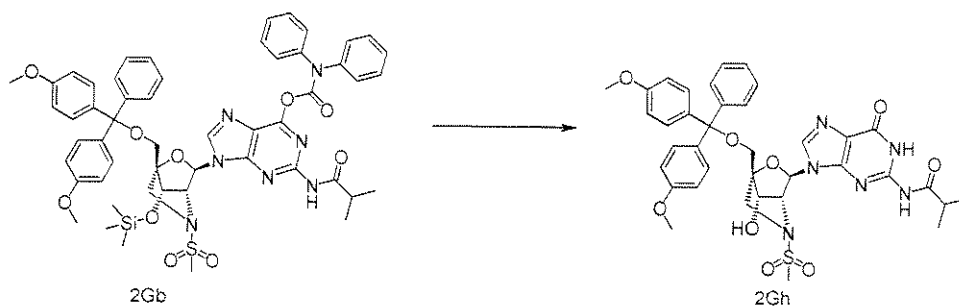
N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-2-methylpropane amide

To a solution of the compound 2Gk (3.16 g, 3.87 mmol) in THF (8 mL) was added TBAF (1 mol/L THF solution, 4.6 mL, 4.6 mmol) and the mixture was stirred at room temperature for 1 hour. The reaction solution was concentrated, and thereto was added ethyl acetate, and the organic layer was washed with water and saturated brine. The organic layer was dried over magnesium sulfate, and concentrated. The resulting residue was purified by silica gel chromatography (chloroform/methanol, 100/0 to 90/10) to obtain a compound 2Gh (2.57 g, yield 89%).

MS(ESI): $m/z = 745.5$ ($M+H$)⁺

[0336]

Also the compound 2Gh can be also synthesized from the compound 2Gb as follows.



[0337]

Example 144 Synthesis of Compound 2Gh

N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-
 5 azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-2-methylpropane amide

The compound 2Gb (1010 mg, 1.0 mmol) was solubilized in tetrahydrofuran (10 mL), and thereto were added tetrabutyl ammonium fluoride (1 mol/L tetrahydrofuran solution, 3 mL,
 10 3.0 mmol) at room temperature, and the mixture was stirred at 70 °C for 24 hours. Further, thereto was added tetrabutyl ammonium fluoride (1 mol/L tetrahydrofuran solution, 9 mL, 9.0 mmol) at room temperature, and the mixture was stirred at 80 °C for 8 hours. After the solvents were evaporated,
 15 the resulting mixture was purified by silica gel chromatography (chloroform/methanol, 98/2 to 93/7). The resulting mixture was further purified by silica gel chromatography (chloroform/methanol, 100/0 to 94/6) to obtain a compound 2Gh (570 mg, yield 77%).
 20 MS(ESI): m/z = 745 (M+H)⁺

[0338]

Example 145 Synthesis and Purification of Oligonucleotide analogues (in vitro)

Using each kind of amidite obtained in the above
 25 examples (present compounds 1 to 44), oligonucleotide

analogue compounds indicated in Tables 1 to 4 below were synthesized by DNA/RNA oligonucleotide automatic synthesizer nS-8II (manufactured by GeneDesign, Inc.) using 0.2 or 1.0 μ mol scale of CPG or polystyrene support. All of the
5 amidites were adjusted to 0.1 M acetonitrile solution, and the coupling time of non-natural type of nucleoside was 10 minutes, and any steps other than the step were performed under a standard condition of ns-811. Activator 42 (Sigma-Aldrich) as an activator, Sulfurizing ReagentII (Gren
10 Research Corporation) for thionylation were used. To the synthesized oligonucleotide was added 28% ammonia water, and the mixture was reacted at 60 to 65 °C for 8 hours to perform a cleavage from a support and a deprotection of a base moiety. Ammonia was concentrated and evaporated, and the resulting
15 mixture was subjected to a purification by reverse phase HPLC chromatography.

[0339]

Example 146 Synthesis and Purification of Oligonucleotide analogues (in vivo)

20 Using each kind of amidite obtained in the above examples (present compounds 1 to 44), oligonucleotide analogue compounds indicated in Table 5 below were synthesized by DNA/RNA oligonucleotide automatic synthesizer AKTA oligopilot plus 10 (manufactured by GE Healthcare Inc.)
25 using 20 μ mol scale of polystyrene support. DNA amidite was

adjusted to 0.1 M, and non-natural amidite was adjusted to 0.05 M acetonitrile solution, and the coupling recycle time of non-natural type of nucleoside was 20 minutes, and when the first base was introduced in a universal support, a coupling, a thiolation, a capping step was carried out twice in a row respectively. Any steps other than the step were performed under a standard condition of AKTA oligopilot plus 10. Activator 42 (Sigma-Aldrich) as an activator, Sulfurizing ReagentII (Gren Research Corporation) for thionylation were used. To the synthesized oligonucleotide was added 28% ammonia water, and the mixture was reacted at 60 to 65 °C for 8 hours to perform a cleavage from a support and a deprotection of a base moiety. Ammonia was concentrated and evaporated, and the resulting mixture was purified by an anion exchange column. Excess salts contained after anion exchange were removed by a desalting column.

[0340]

The purification and confirmation of purity of the synthesized oligonucleotide analogues were carried out by reverse phase HPLC chromatography under the following conditions.

Reverse HPLC

Mobile phase

A solution: 20 mM Cyclohexylammonium acetate aqueous

solution

B solution: acetonitrile

Gradient: A:B = 90:10 → 50:50 (40 min.) or A:B = 90:10
→ 40:60 (45 min.)

5 Used column:

Analysis: Waters XBridge[®] Oligonucleotide BEH C18
Column, 130 Å, 2.5 µm, 4.6 × 50 mm

Preparation: Waters XBridge[®] Oligonucleotide BEH C18
OBDM Prep Column, 130 Å, 2.5 µm, 10 × 50 mm

10 [0341]

Flow rate:

Analysis: 1 mL/min.

Preparation: 4 mL/min.

Column temperature: 60 °C

15 Detection: UV (260 nm)

[0342]

Anion exchange purification

Mobile phase

A solution: 20 mM Tris-HCl (pH 8.0) aqueous solution

20 B solution: 20 mM Tris-HCl (pH 8.0), 1 M NaBr aqueous
solution

Used column:

GE HiTrap CaptQ ImpRes (Anion exchange) 5 mL

Flow rate: 5 mL/min.

25 Column temperature: room temperature

Detection: UV (260 nm)

[0343]

Desalting column

Mobile phase

5 A solution: H₂O

 B solution: H₂O

Used column:

 GE HiPrep 26/10 Desalting

Flow rate: 10 mL/min.

10 Column temperature: room temperature

[0344]

 The molecular weight of the synthesized oligonucleotide analogues was determined by a Waters ZQ under the below-mentioned conditions.

15

Mobile phase

 A solution: 400 mM hexafluoroisopropanol, 15 mM triethylamine aqueous solution

 B solution: methanol

20 Gradient: A:B = 80:20 → 70:30 (2.5 min.) or A:B = 90:10
 → 40:60 (45 min.)

Used column:

 Analysis: Waters ACQUITY UPLC® Oligonucleotide BEH C18

Column, 130 Å, 1.7 µm, 2.1 × 100 mm

25 Flow rate: 0.2 mL/min.

Column temperature: 60 °C

Detection: UV (260 nm)

[0345]

Table 1 Mass spectrum measurement results of each Tm
5 sequence

gcggttT(X)tttgct (SEQ ID No. 1)

(wherein, small character represents each nucleic acid base,
T(X) represents each artificial nucleic acid species
containing thymine, and all phosphate bonds represent
10 phosphodiester.)

Oligonucleotide compound NO.	Artificial nucleic acid species contained in antisense strand	m/z
Tm-1 (SEQ ID No. 1-1)	ALNA[Me]	1223.8 [M-3H] ³⁻
Tm-1 (SEQ ID No. 1-2)	ALNA[Ms]	1245.3 [M-3H] ³⁻
Tm-3 (SEQ ID No. 1-3)	ALNA[Bs]	1266.4 [M-3H] ³⁻
Tm-4 (SEQ ID No. 1-4)	ALNA[mU]	1238.5 [M-3H] ³⁻
Tm-5 (SEQ ID No. 1-5)	ALNA[ipU]	1247.7 [M-3H] ³⁻
Tm-6 (SEQ ID No. 1-6)	ALNA[dmU]	1243.1 [M-3H] ³⁻
Tm-7 (SEQ ID No. 1-7)	ALNA[2Pym]	3736.52 [M-H] ¹⁻
Tm-8 (SEQ ID No. 1-8)	ALNA[4Pym]	3736.49 [M-H] ¹⁻
Tm-9 (SEQ ID No. 1-9)	ALNA[4CF3-2Pym]	3804.51 [M-H] ¹⁻
Tm-10 (SEQ ID No. 1-10)	ALNA[5Cl-2Pym]	3773.37 [M-H] ¹⁻
Tm-11 (SEQ ID No. 1-11)	ALNA[6NMe2-4Pym]	3777.99 [M-H] ¹⁻
Tm-12 (SEQ ID No. 1-12)	ALNA[2NMe2-4Pym]	3779.19 [M-H] ¹⁻
Tm-13 (SEQ ID No. 1-13)	ALNA[Prz]	3737.76 [M-H] ¹⁻
Tm-14 (SEQ ID No. 1-14)	ALNA[Trz]	3752.88 [M-H] ¹⁻
Tm-15 (SEQ ID No. 1-15)	ALNA[Oxz]	3743.19 [M-H] ¹⁻

[0346]

Table 2 Mass spectrum measurement results of each SRB1 (2-
10-2) sequence

T(X)C(X)agtcagtT(X)C(X) (SEQ ID No. 2)

15 (wherein, small character represents each nucleic acid base

of DNA, T(X) or C(X) each represents artificial nucleic acid species containing thymine, artificial nucleic acid species containing 5-methylcytosine, respectively, and all phosphate bonds represent phosphorothioate.)

5 [Table 2]

Oligonucleotide compound NO.	Artificial nucleic acid species contained in antisense strand	m/z
SRB1-1 (SEQ ID No. 2-1)	ALNA[Me]	1152.9 [M-4H] ⁴⁻
SRB1-2 (SEQ ID No. 2-2)	ALNA[Ms]	1217.0 [M-4H] ⁴⁻
SRB1-3 (SEQ ID No. 2-3)	ALNA[ipU]	1224.1 [M-4H] ⁴⁻
SRB1-4 (SEQ ID No. 2-4)	ALNA[2Pym]	4869.28 [M-H] ¹⁻
SRB1-5 (SEQ ID No. 2-5)	ALNA[Prz]	4871.85 [M-H] ¹⁻
SRB1-6 (SEQ ID No. 2-6)	ALNA[Trz]	4937.77 [M-H] ¹⁻
SRB1-7 (SEQ ID No. 2-7)	ALNA[Oxz]	4885.15 [M-H] ¹⁻

[0347]

Table 3 Mass spectrum measurement results of each SRB1 (3-10-3) sequence

T(X)T(X)C(X)agtcatgactT(X)C(X)C(X) (SEQ ID No. 3)

10 (wherein, small character represents each nucleic acid base of DNA, T(X) or C(X) each represents artificial nucleic acid species containing thymine, artificial nucleic acid species containing 5-methylcytosine, respectively, and all phosphate bonds represent phosphorothioate.)

15 [Table 3]

Oligonucleotide compound NO.	Artificial nucleic acid species contained in antisense strand	m/z
SRB1-8 (SEQ ID No. 3-1)	ALNA[Me]	1066.4 [M-5H] ⁵⁻
SRB1-9 (SEQ ID No. 3-2)	ALNA[Ms]	1143.3 [M-5H] ⁵⁻
SRB1-10 (SEQ ID No. 3-3)	ALNA[ipU]	1151.8 [M-5H] ⁵⁻

SRB1-11 (SEQ ID No. 3-4)	ALNA[2Pym]	5718.12 [M-H] ²⁻
SRB1-12 (SEQ ID No. 3-5)	ALNA[Prz]	5719.86 [M-H] ²⁻
SRB1-13 (SEQ ID No. 3-6)	ALNA[Trz]	5824.11 [M-H] ²⁻
SRB1-14 (SEQ ID No. 3-7)	ALNA[Oxz]	5741.47 [M-H] ²⁻

[0348]

Table 4 Mass spectrum measurement results of each anti-miR-21 sequence

A(m)C(X)atC(X)agtC(X)tgaT(X)aagC(X)tA(m) (SEQ ID No. 4)

5 (wherein, A(m) represents 2'-MOE adenosine, small character represents each nucleic acid base of DNA, T(X) or C(X) each represents artificial nucleic acid species containing thymine, artificial nucleic acid species containing 5-methylcytosine, respectively, and all phosphate bonds
10 represent phosphorothioate.)

[Table 4]

Oligonucleotide compound No.	Artificial nucleic acid species contained in antisense strand	m/z
anti-miR21-1 (SEQ ID No. 4-1)	ALNA[Me]	1081.4 [M-6H] ⁵⁻
anti-miR21-2 (SEQ ID No. 4-2)	ALNA[mU]	1117.6 [M-6H] ⁵⁻
anti-miR21-3 (SEQ ID No. 4-3)	ALNA[ipU]	1140.8 [M-6H] ⁵⁻
anti-miR21-4 (SEQ ID No. 4-4)	ALNA[dmU]	1129.1 [M-6H] ⁵⁻
anti-miR21-5 (SEQ ID No. 4-5)	ALNA[Trz]	6897.87 [M-H] ²⁻
anti-miR21-6 (SEQ ID No. 4-6)	ALNA[Oxz]	6832.17 [M-H] ²⁻

[0349]

Table 5 Mass spectrum measurement results of each Malat1 sequence

G(X)T(X)T(X)cactgaatG(X)C(X) (SEQ ID No. 5)

(wherein, small character represents each nucleic acid base of DNA, T(X),C(X) or G(X) each represents artificial nucleic acid species containing thymine, artificial nucleic acid species containing 5-methylcytosine, or artificial nucleic acid species containing guanine, respectively, and all phosphate bonds represent phosphorothioate.)

[Table 5]

Oligonucleotide compound No.	Artificial nucleic acid species contained in antisense strand	m/z
Malat1-1 (SEQ ID No. 5-1)	ALNA[Me]	1089.8 [M-4H] ⁴⁻
Malat1-2 (SEQ ID No. 5-2)	ALNA[Ms]	1169.3 [M-4H] ⁴⁻
Malat1-3 (SEQ ID No. 5-3)	ALNA[2Pym]	4680.86 [M-H] ¹⁻

[0350]

10 Example 147 Evaluation of binding ability to target strand and mismatch selectivity

Each of an antisense strand containing various artificial nucleic acid species (wherein sequence represents gcgttT(X)tttgct (the above-mentioned SEQ ID No. 1), small character represents each nucleic acid base of DNA, T(X) represents each artificial nucleic acid specie containing thymine); a sense strand, that is, DNA strand having a sequence complementary to the antisense strand (wherein sequence represents agcaaaaaacgc (SEQ ID No. 6), and small

character represents each nucleic acid base of DNA); a RNA strand having a complementary sequence (wherein sequence represents AGCAAAAACGC (SEQ ID No. 7), and large character represents each nucleic acid base.); and a RNA strand having
5 3 complementary sequences containing 1 base mismatch (wherein sequence represents AGCAAANAACGC (SEQ ID No. 8), and large character represents each nucleic acid base, and each N represents uracil (U), guanine(G), cytosine(C).) was annealed to form a double strand, and then T_m value that is
10 a temperature at which 50% of the double strand is deviated, was measured, and the double strand-forming capacity of oligonucleotide was examined.

Specifically, a sample solution (150 μ L) which contains aqueous sodium chloride solution 100 mmol/L, sodium
15 phosphate buffer (pH 7.4) 10 mmol/L, antisense strand 4 μ mol/L, and antisense strand 4 μ mol/L was warmed to 95 °C and then cooled to room temperature over 4 hours. Nitrogen stream was passed into a cell chamber of spectrophotometer (JASCO, V-730, PAC-743R) so as to prevent a condensation
20 phenomenon, and the sample solution was cooled to 10 °C gradually, and further kept at 10 °C for 1 minute, and the measurement was then started. The temperature was warmed to 80 °C at a rate of 0.5 °C per minute, and the ultraviolet absorption at 260 nm was measured at every timing of 0.5 °C
25 rise. Here in order to prevent a concentration change due

to a temperature rise, the cell with a lid was used. The measurement results are shown in Table 6.

[0351]

Table 6 Measurement results of each T_m value

5 [Table 6]

Structure of T(X) in antisense strand 5'- gcggttT(X)tttgct- 3'	T_m (°C)				
	v.s. DNA	v.s. RNA 5'-AGC AAA NAA CGC-3'			
		N=A T-A match	N=U T-U mismatch	N=G T-G mismatch	N=C T-C mismatch
DNA	51	48	32	42	30
ALNA[Me]	53	51	37	45	38
ALNA[Ms]	51	51	37	45	35
ALNA[Bs]	49	49	35	42	34
ALNA[mU]	51	51	38	44	36
ALNA[ipU]	52	51	38	42	35
ALNA[dmU]	50	50	39	42	35
ALNA[2Pym]	48	48	34	42	33
ALNA[4Pym]	52	52	39	43	36
ALNA[4CF3-2Pym]	47	49	35	42	32
ALNA[5Cl-2Pym]	47	48	34	42	32
ALNA[6NMe2-4Pym]	54	52	38	39	34
ALNA[2MNe2-4Pym]	52	50	38	40	35

ALNA[Prz]	50	51	38	43	34
ALNA[Trz]	48	50	36	42	34
ALNA[Oxz]	49	50	36	44	35

[0352]

As clearly shown in Table 6, the antisense strand containing various artificial nucleic acids has the same or higher sense strand binding ability and mismatch selectivity as compared with antisense strand containing natural DNA and ALNA[Me] as known ALNA.

[0353]

Example 147 In vitro SRB1 knockdown activity test

Hepa 1clc7 cells were seeded at 2.5×10^3 cells per well and cultured for 24 hours, and then various modified antisense oligonucleic acids having a complementary sequence to SRB1 were added at a final concentration of 100, 20, 4 nmol/L, and after 48 hours, a quantitative PCR was performed. The same conditions were carried out in 3 wells each. Regarding antisense activity, the inhibition rate of SRB1 expression when the antisense compound was added was set as 100% based on the amount ratio of SRB1 to GAPDH in a negative control. The results are shown in Table 7.

[0354]

Table 7 In vitro SRB1 knockdown activity test

[Table 7]

Artificial nucleic acid species	Knockdown activity (%)	
	3-10-3	2-10-2
ALNA[Me]	36	27
ALNA[Ms]	48	38
ALNA[ipU]	43	29
ALNA[2Pym]	57	19
ALNA[Prz]	40	44
ALNA[Trz]	60	39
ALNA[Oxz]	67	50

The target gene (SRB1) knockdown activity of the antisense oligonucleotide containing each artificial nucleic acid in the two sequence patterns of 3-10-3, and 2-10-2 at
5 Gymnosis 100 nM was shown.

The two array patterns are as follows.

3-10-3: T(X)T(X)C(X)agtcatgactT(X)C(X)C(X) (as the same as
SEQ ID No. 3 in the above Table 3)

2-10-2: T(X)C(X)agtcatgactT(X)C(X) (as the same as SEQ ID
10 No. 2 in the above Table 2)

Here small character represents each nucleic acid base, each of T(X) or C(X) represents artificial nucleic acid specie containing thymine, or artificial nucleic acid specie containing 5-methylcytosine, and all phosphate bonds
15 represent phosphorothioate.

[0355]

As clearly shown in Table 7, ALNA[Ms], ALNA[ipU], ALNA[2Pym], ALNA[Prz], ALNA[Trz], and ALNA[Oxz] had a strong in vitro target gene knockdown activity as compared to ALNA[Me] as a known ALNA.

5 [0356]

Example 148 In vitro miRNA-21 inhibitory activity test

3.0 × 10⁶ cells/ml of HEK293 cells were seeded in a 10 cm dish, and the cells were cultured overnight in a CO₂ incubator. Next day, using FuGENE HD Transfection Reagent, and cells were transfected with 10 µg of reporter plasmid in which the complementary sequence of miR-21 was cloned into the multicloning site of psiCHECK-2 vector (Promega), and were cultured in a CO₂ incubator for about 24 hours. These cells were recovered, and 2.0 × 10⁴ cells/well of cells were reseeded in a 96-well plate, and a miR-21 inhibitor into which each artificial nucleic acid was introduced was added. After cells were cultured in a CO₂ incubator for 96 hours. Using the Dual-Glo Luciferase Assay System, luminescence values of intracellular firefly luciferase and Renilla luciferase were detected with a plate reader. To correct the effects of transfection efficiency or cell number from the luminescence value due to Renilla luciferase activity, the ratio to the luminescence value due to firefly luciferase activity was calculated. Further, from the calculated ratio, the ratio in the cells into which the psiCHECK-2 vector was

10
15
20
25

introduced was set as 100% inhibition rate, and the ratio of cells into which the psiCHECK-2 vector into which the miR-21 complementary sequence was cloned was set as 0%, and each inhibition rate was calculated and the inhibitory activity was determined. The results are shown in Table 8.

[0357]

Table 8 In vitro miRNA-21 inhibitory activity

[Table 8]

Artificial nucleic acid species	miR-21 inhibition rate (%)	
	Gymnosis, 1μM	Gymnosis, 3μM
ALNA[Me]	6	19
ALNA[MU]	69	85
ALNA[ipU]	45	61
ALNA[dmU]	42	63
ALNA[Trz]	61	105
ALNA[Oxz]	31	63

Sequence of used antisense oligo is shown below.

A(m)C(X)atC(X)agtC(X)tgaT(X)aagC(X)tA(m) (the same as SEQ ID No. 4 in the above Table 4)

(wherein, A(m) represents 2'-MOE adenosine, small character represents each nucleic acid base of DNA, T(X) or C(X) each represents artificial nucleic acid species containing thymine, artificial nucleic acid species containing 5-methylcytosine, respectively, and all phosphate bonds

represent phosphorothioate.)

[0358]

As clearly shown in Table 8, ALNA[mU], ALNA[ipU], ALNA[dmU], ALNA[Trz], and ALNA[Oxz] had a strong in vitro
5 micro RNA inhibitory activity as compared to ALNA[Me] as a known ALNA.

[0359]

Example 149 In vivo MALAT1 knockdown test

Antisense oligonucleotides (10 mg/kg and 50 mg/kg)
10 targeting MALAT1 were prepared with PBS to make a concentration of 10 mg/mL and then diluted with PBS to make a 5 mL/kg, which was administered intravenously to 6 week-old C57BL/6J mice (male, Charles River Japan). After 72 hours, blood was collected from the abdominal vena cava of
15 the mouse under anesthesia of isoflurane (Japan Pharmacopoeia, manufactured by Pfizer, Inc.) and euthanized by exsanguination. Blood was coagulated and then centrifuged to prepare serum, which was then stored frozen (set at -80°C) until the time of measurement. After euthanizing by
20 exsanguination, liver, kidney, fat, skeletal muscle (quadriceps femoris, gastrocnemius muscle), lung, heart, stomach, jejunum, testis, skin, spleen, and brain were collected and frozen in liquid nitrogen. A homogenization buffer of Maxwell RSC simply RNA Tissue Kit was added to
25 frozen tissue, and the tissue was disrupted using a multi-

bead shocker, RNA was purified according to the protocol described in the kit. RT reaction and the subsequent reactions were performed duplicately, and the amount of RNA per reaction was used in 50 to 500 ng. Regarding knockdown activity, the inhibition rate of Malat1 expression was set as 100% based on the amount ratio of Malat1 to GAPDH in vehicle group. The results are shown in Tables 9 and 10.

[0360]

Table 9 Malat1 knockdown rate (%) in each organ when 50 mg/kg was administered

[Table 9]

Artificial nucleic acid	liver	kidney	quad- riceps femoris	gastro- cnemius muscle	brain	lung	heart	stomach	jejunum	testis	spleen	fat
ALNA[Me]	88	56	55	49	4	64	9	37	9	30	51	43
ALNA[Ms]	97	80	67	63	0	80	48	54	56	69	46	82
ALNA[2Pym]	83	46	28	20	0	51	34	0	49	28	26	47
ALNA[ipU]	89	62	0	24	0	82	39	63	60	49	84	57

Table 10 Malat1 knockdown rate (%) in each organ when 10 mg/kg was administered

[Table 10]

Artificial nucleic acid	liver	kidney	quad- riceps femoris	gastro- cnemius muscle	brain	lung	heart	stomach	jejunum	testis	spleen	fat
ALNA[Me]	47	41	0	21	10	50	12	1	0	0	43	3
ALNA[Ms]	90	66	22	47	0	49	17	32	10	39	19	70
ALNA[2Pym]	53	4	0	0	0	1	37	0	27	15	0	18
ALNA[ipU]	56	66	0	11	24	75	29	50	33	22	73	47

Sequence of used antisense oligonucleotide is shown below.

G(X)T(X)T(X)cactgaatG(X)C(X) (the same as SEQ ID No. 5 in the above Table 5)

(wherein, small character represents each nucleic acid base of DNA, T(X), C(X) or G(X) each represents artificial nucleic acid species containing thymine, artificial nucleic acid species containing 5-methylcytosine, or artificial nucleic acid species containing guanosine, respectively, and all phosphate bonds represent phosphorothioate.)

[0361]

10 As clearly shown in Tables 9 and 10, ALNA[Ms] had a strong target gene knockdown in many organs (liver, kidney, skeletal muscle, lung, heart, stomach, jejunum, testis, fat) as compared to ALNA[Me] as a known ALNA. Also ALNA[2Pym] had a strong target gene knockdown in particular
15 organs (heart, jejunum, testis, fat) as compared to ALNA[Me] as a known ALNA. Also ALNA[ipU] had a similar degree of target gene knockdown effect in liver as compared to ALNA[Me], whilst, had a remarkably stronger target gene knockdown effect in lung and spleen than ALNA[Me]. Since
20 various immune cells are accumulated in spleen, antisense oligos including ALNA[ipU] can be expected to act on immune cells, which are considered to have low migration with ordinary nucleic acid drugs. In addition, ALNA[ipU] can be expected to have a strong drug effect on lung or immune
25 system while reducing toxicity in liver.

[0362]

Example 150 MALAT1 knockdown activity test in each central tissue, liver, and kidney by intraventricular administration

5 7 week-old C57BL/6J (male, Japan Charles River) was fixed to a brain stereotaxic device under anesthesia by Isoflurane (manufactured by Pfizer Inc.). An incision was made in the scalp while lying on a heat insulating mat to expose the skull. After confirming the position of the
10 bregma, a hole was drilled in the skull 1 to 2 mm around the upper left ventricle (1.0 mm to the left and 0.4 mm behind the bregma) with a dental electric drill. A 27 gauge needle was inserted into the left ventricle (1.0 mm to the left of Bregma, 0.4 mm behind, 2.3 mm deep), and a single dose of
15 antisense oligo (5 μ g/10 μ L/head) targeting MALAT1 that was prepared to 5 μ g/10 μ L in saline using a needle-mounted microinfusion pump was administered at a flow rate of 2 μ L/min for 5 minutes. Thereafter, the hole in skull was closed with the medical cement Belfast (manufactured by
20 Mutsukagaku Kogyo), the incision was sutured, and gentamicin ointment 0.1% (manufactured by Takata Seiyaku) was applied to prevent infectious diseases. Seventy-two (72) hours after administration, blood was collected from the abdominal vena cava of mice under isoflurane anesthesia and euthanized by
25 exsanguination. Blood was coagulated and then centrifuged

to prepare serum, which was then stored frozen (set at -30°C) until the time of measurement. After euthanizing by exsanguination, brain, spinal cord, kidney, and liver were collected. For the brain, after euthanasia, skull and spine were separated, and then brain was taken out, and using Brain Matrices (RBM-2000S, manufactured by Bio Research Center Co., Ltd.), which is a slicer for making sagittal sections, and a brain is sliced from the median to the left brain side to a thickness of about 2 mm, and then an olfactory bulb, cerebral cortex, hippocampus, striatum, and cerebellum were taken out from the section. The rest of the section from which the 5 sites were removed was also collected as a complex site such as the brain stem. The spinal cord was separated at the boundary of T13, and two sites were collected as a part in the brain side from T13 (hereinafter, proximal spinal cord), and another part from T13 to L5 (hereinafter, distal spinal cord). The collected organs were immersed in RNA later (Invitrogen) and stored at -30 °C. The organs and QIAzol Lysis Reagent (QIAGEN) were added to the tube, and the tissue was disrupted using a multi-bead shocker, and chloroform (manufactured by jyunsei chemical) and mixed. The aqueous layer was recovered from this mixture by centrifuge separation. RNA was purified according to the protocol described in RNeasy 96 Universal Tissue Kit(QIAGEN). RT reaction and the subsequent reactions were performed by

using the amount of RNA per reaction in 2 to 7 ng. Regarding knockdown activity, the inhibition rate of Malat1 expression was set as 100% based on the amount ratio of Malat1 to GAPDH in vehicle group. Standard error was calculated based on individual values.

Table 11 MALAT1 knockdown rate (%) in each central tissue, liver, and kidney when 5 µg/head was administered

[Table 11]

Modified nucleic acid	olfactory bulb	cerebral cortex	hippo-campus	stria-tum	cerebe-llum	complex site including brain stem	proximal spinal cord	distal spinal cord	kidney	liver
ALNA[Me]	-5.0	26.2	11.2	37.1	-27.6	31.8	25.0	2.9	14.0	36.3
ALNA[Ms]	4.7	37.4	35.2	49.5	12.9	46.5	25.0	18.2	34.3	21.1
ALNA[mU]	8.5	29.8	13.3	24.0	-42.6	40.6	23.4	1.1	4.7	17.8
ALNA[Oxz]	8.0	28.2	18.7	33.1	-5.3	44.3	44.9	18.5	11.1	31.5

Sequence of used antisense oligonucleotide is shown below.

G(X)T(X)T(X)cactgaatG(X)C(X) (the same as SEQ ID No. 5 in the above Table 5)

(wherein, small character represents each nucleic acid base of DNA, T(X), C(X) or G(X) each represents artificial nucleic acid species containing thymine, artificial nucleic acid species containing 5-methylcytosine, or artificial nucleic acid species containing guanosine, respectively, and all phosphate bonds represent phosphorothioate.)

[0363]

As clearly shown in Table 11, ALNA[Ms], ALNA[mU], and

ALNA[Oxz] had stronger target gene knockdown activity in each central tissue as compared to ALNA[Me] as a known ALNA.

Industrial Applicability

5 [0364]

An amino bridged artificial nucleic acid (ALNA) of the present invention can be used in a preparation of novel artificial nucleic acid oligomer. Also an oligomer containing the amino bridged artificial nucleic acid of the
10 present invention has a strong target gene knockdown activity in particular organs and can be used as nucleic acid drugs.

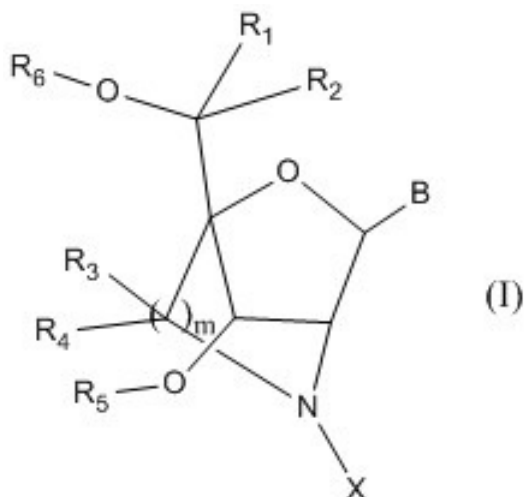
[Sequence Listing Free Text]

[0365]

15 SEQ ID Nos. 1 to 6 show DNA oligonucleotides.
SEQ ID Nos. 7 to 8 show RNA oligonucleotides.

CLAIMS

1. A compound represented by general formula I:



5 [wherein

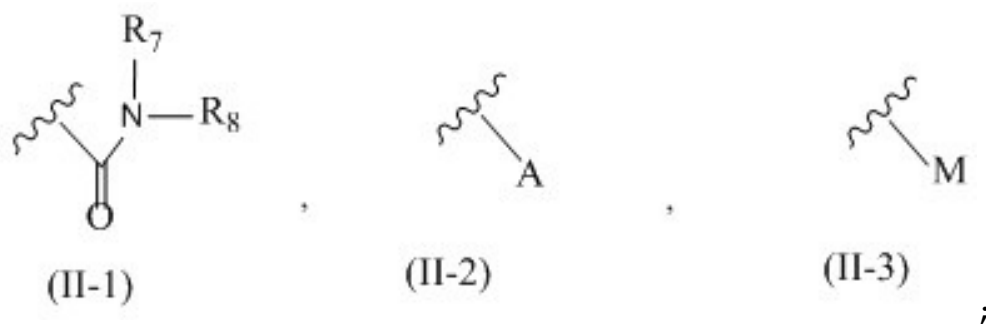
B represents a base moiety of nucleic acid wherein the base moiety may be optionally substituted with one or more substituents;

10 R_1 , R_2 , R_3 and R_4 represent independently of each other a hydrogen atom, or a C_{1-6} alkyl group which may be optionally substituted with one or more substituents;

15 R_5 and R_6 represent independently of each other a hydrogen atom, a protecting group for hydroxy group, or a phosphate group which may be optionally substituted with substituents;

m is an integer of 1 or 2;

X represents a group represented by the following formula (II-1), (II-2) or (II-3):



the symbols:



which is described in the formula (II-1), (II-2) or (II-3)

5 represents a binding point to 2'-amino group;

R_7 and R_8 represent independently of each other a hydrogen atom, a C_{1-6} alkyl group which may be optionally substituted with one or more substituents, a C_{2-6} alkenyl group which may be optionally substituted with one or more substituents, a C_{2-6} alkynyl group which may be optionally substituted with one or more substituents, or an aromatic group which may be optionally substituted with one or more substituents;

A represents an aromatic group;

15 M represents a sulfonyl group which is substituted with one substituent selected from the group consisting of a C_{1-6} alkyl group which may be optionally substituted with one or more substituents and an aromatic group which may be optionally substituted with one or more substituents.]

20 or salts thereof.

2. The compound according to claim 1 or salt thereof

wherein B represents an adeninyl group which may be optionally substituted with one or more substituents, a guaninyl group which may be optionally substituted with one or more substituents, a cytosinyl group which may be optionally substituted with one or more substituents, a 5-methylcytosinyl group which may be optionally substituted with one or more substituents, or an uracilyl group which may be optionally substituted with one or more substituents.

3. The compound according to claim 1 or 2 or salts thereof wherein R_1 , R_2 , R_3 and R_4 represent independently of each other a hydrogen atom; and

m is an integer of 1.

4. The compound according to any one of claims 1 to 3 or salts thereof wherein

15 X represents a group represented by formula (II-1); and

R_7 and R_8 represent independently of each other a hydrogen atom, a C_{1-6} alkyl group which may be optionally substituted with one or more substituents, or an aromatic group which may be optionally substituted with one or more substituents.

5. The compound according to claim 4 or salts thereof wherein R_7 and R_8 represent independently of each other a hydrogen atom, or a C_{1-3} alkyl group which may be optionally substituted with one or more substituents.

6. The compound according to claim 5 or salts thereof wherein one of R₇ and R₈ represents a hydrogen atom, and the other thereof represents a methyl group.

7. The compound according to claim 5 or salts thereof
5 wherein one of R₇ and R₈ represents a hydrogen atom, and the other thereof represents an isopropyl group.

8. The compound according to any one of claims 1 to 3 or salts thereof wherein

X represents a group represented by formula (II-2);

10 A represents a five or six membered heteroaryl group containing one or more heteroatoms selected from the group consisting of nitrogen atom, oxygen atom and sulfur atom, which may be optionally substituted with one or more substituents.

15 9. The compound according to claim 8 or salts thereof wherein A represents a five or six membered heteroaryl group containing two or three heteroatoms selected from the group consisting of nitrogen atom, oxygen atom and sulfur atom, provided that at least two nitrogen atoms are
20 included, which may be optionally substituted with one or more substituents, and the substituent is selected independently of each other from a group consisting of a C₁₋₃ alkyl group which may be optionally substituted with one or more halogen atoms, a halogen atom, and an amino
25 group which may be optionally substituted with one or more

C₁₋₃ alkyl group.

10. The compound according to claim 9 or salts thereof wherein the five or six membered heteroaryl group is selected from a group consisting of a triazolyl group, an
5 oxadiazolyl group, a thiadiazolyl group, a pyrimidinyl group, and a pyrazinyl group, each may be optionally substituted with one or more substituents.

11. The compound according to claim 10 or salts thereof wherein the triazolyl group which may be optionally
10 substituted with one or more substituents represents 1,5-dimethyl-1,2,4-triazol-3-yl group.

12. The compound according to claim 10 or salts thereof wherein the oxadiazolyl group which may be optionally substituted with one or more substituents represents 5-
15 methyl-1,2,4-oxadiazol-3-yl group.

13. The compound according to claim 10 or salts thereof wherein the thiadiazol group which may be optionally substituted with one or more substituents represents 3-methyl-1,2,4-thiadiazol-5-yl group.

20 14. The compound according to any one of claims 1 to 3 or salts thereof wherein

X represents a group represented by formula (II-3);
and

M represents a sulfonyl group which is substituted
25 with one or more substituents selected from the group

consisting of a C₁₋₆ alkyl group which may be optionally substituted with one or more substituents, and an aryl group which may be optionally substituted with one or more substituents.

5 15. The compound according to claim 14 or salts thereof wherein M represents a sulfonyl group which is substituted with one or more substituents selected from the group consisting of a methyl group and a phenyl group.

16. The compound according to according to any one of
10 claims 1 to 15 or salts thereof wherein R₆ represents a hydrogen atom or DMTr group, and R₅ represents a hydrogen atom or -P(O(CH₂)₂CN)(N(ipr)₂).

17. A compound selected from the group consisting of

3-[[(1R, 4R, 6R, 7S)-4-[[bis(4-methoxyphenyl)-phenyl-
15 methoxy]methyl]-6-(5-methyl-2,4-dioxo-pyrimidin-1-yl)-2-methylsulfonyl-5-oxa-2-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile;

N-[1-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-
20 (diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide;

[9-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-
25 (diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-

azabicyclo[2.2.1]heptan-3-yl]-2-(2-methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate;

N-[9-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-

5 (diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-purin-6-yl]benzamide;

3-[[(1R, 3R, 4R, 7S)-5-(benzenesulfonyl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-

10 yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile;

N-[1-[(1R, 3R, 4R, 7S)-5-(benzenesulfonyl)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-

15 yl]benzamide;

(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-methyl-3-(5-methyl-2,4-dioxo-pyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-5-

20 carboxamide;

(1R, 3R, 4R, 7S)-3-(4-benzamide-5-methyl-2-oxo-pyrimidin-1-yl)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-methyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide;

25 [9-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-

phenylmethoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(methylcarbamoyl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-2-(2-methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate;

5 (1R, 3R, 4R, 7S)-3-(6-benzamidepurin-9-yl)-1-[[bis(4-methoxyphenyl)phenylmethoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-methyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide;

(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)phenylmethoxy)methyl]-7-[2-cyanoethoxy-

10 (diisopropylamino)phosphanyl]oxy-N-isopropyl-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide;

(1R, 3R, 4R, 7S)-3-(4-benzamide-5-methyl-2-oxypyrimidin-

15 1-yl)-1-[[bis(4-methoxyphenyl)phenyl-methoxy)methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N-isopropyl-2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide;

[9-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-

20 (diisopropylamino)phosphanyl]oxy-5-(isopropylcarbamoyl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-2-(2-methylpropanoylamino)purin-6-yl]N,N-diphenylcarbamate;

(1R, 3R, 4R, 7S)-3-(6-benzamidepurin-9-yl)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy)methyl]-7-[2-cyanoethoxy-

25 (diisopropylamino)phosphanyl]oxy-N-isopropyl-2-oxa-5-

azabicyclo[2.2.1]heptane-5-carboxamide;

(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-
phenylmethoxy]methyl]-7-[2-cyanoethoxy-
(diisopropylamino)phosphanyl]oxy-N,N-dimethyl-3-(5-methyl-
5 2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptane-
5-carboxamide;

(1R, 3R, 4R, 7S)-3-(4-benzamide-5-methyl-2-oxo-pyrimidin-
1-yl)-1-[[bis(4-methoxyphenyl)-phenylmethoxy]methyl]-7-[2-
cyanoethoxy-(diisopropylamino)phosphanyl]oxy-N,N-dimethyl-
10 2-oxa-5-azabicyclo[2.2.1]heptane-5-carboxamide;

3-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy]methyl]-3-(5-methyl-2,4-dioxo-pyrimidin-1-yl)-5-
pyrimidin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-
(diisopropylamino)phosphanyl]oxypropanenitrile;

15 1-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy]methyl]-7-[2-cyanoethoxy-
(diisopropylamino)phosphanyl]oxy-5-pyrimidin-2-yl-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-
yl]benzamide;

20 N'-[9-(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy]methyl]-7-[2-cyanoethoxy-
(diisopropylamino)phosphanyl]oxy-5-pyrimidin-2-yl-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-N,N-
dimethyl-6-formamidine;

25 N'-[9-(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-

methoxy)methyl]-7-[2-cyanoethoxy-
(diisopropylamino)phosphanyl]oxy-5-pyrimidin-2-yl-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]purin-6-yl]benzamide;

3-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-
5 methoxy)methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-
pyrimidin-4-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-
(diisopropylamino)phosphanyl]oxypropanenitrile;

N-[1-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy)methyl]-7-[2-cyanoethoxy-
10 (diisopropylamino)phosphanyl]oxy-5-pyrimidin-4-yl-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-
yl]benzamide;

3-[[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy)methyl]-3-(5-methyl-2,4-dioxo-pyrimidin-1-yl)-5-[4-
15 (trifluoromethyl)pyrimidin-2-yl]-2-oxa-5-
azabicyclo[2.2.1]heptan-7-yl]oxy-
(diisopropylamino)phosphanyl]oxypropanenitrile;

N-[1-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy)methyl]-7-[2-cyanoethoxy-
20 (diisopropylamino)phosphanyl]oxy-5-[4-
(trifluoromethyl)pyrimidin-2-yl]-2-oxa-5-
azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-
yl]benzamide;

3-[[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-
25 methoxy)methyl]-5-(5-chloropyrimidin-2-yl)-3-(5-methyl-2,4-

dioxo-pyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile;

N-[1-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-5-(5-chloropyrimidin-2-yl)-7-[2-
 5 cyanoethoxy-(diisopropylamino)phosphanyl]oxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methylpyrimidin-4-yl]benzamide;

3-[[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-5-[6-(dimethylamino)pyrimidin-4-yl]-3-(5-
 10 methyl-2,4-dioxo-pyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamine)phosphanyl]oxypropanenitrile;

N-[1-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-
 15 cyanoethoxy(diisopropylamino)phosphanyl]oxy-5-[6-(dimethylamino)pyrimidin-4-yl]-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide;

3-[[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-5-[2-(dimethylamino)pyrimidin-4-yl]-3-(5-
 20 methyl-2,4-dioxo-pyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)oxypropanenitrile;

N-[1-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-

(diisopropylamino)phosphanyl]oxy-5-[2-(dimethylamino)pyrimidin-4-yl]-7-hydroxy-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide;

5 3-[1-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-pyrazin-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile;

 N-[1-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-10 methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-pyrazine-2-yl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide;

 3-[[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-15 methoxy]methyl]-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile;

 N-[1-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-20 methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide;

 N'-[9-[(1R, 3R, 4R, 7S)-1-[[bis(4-methoxyphenyl)-phenyl-25 methoxy]methyl]-7-[2-cyanoethoxy-

(diisopropylamino)phosphanyl]oxy-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-N,N-dimethylformamidine

N-[9-(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(1,5-dimethyl-1,2,4-triazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]purin-6-yl]benzamide;

3-[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-(5-methyl-1,2,4-oxadiazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile;

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-5-(5-methyl-1,2,4-oxadiazol-3-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide;

3-[[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-3-(5-methyl-2,4-dioxypyrimidin-1-yl)-5-(3-methyl-1,2,4-thiadiazol-5-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-7-yl]oxy-(diisopropylamino)phosphanyl]oxypropanenitrile;

N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-methoxy]methyl]-7-[2-cyanoethoxy-

(diisopropylamino)phosphanyl]oxy-5-(3-methyl-1,2,4-thiadiazol-5-yl)-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]benzamide;

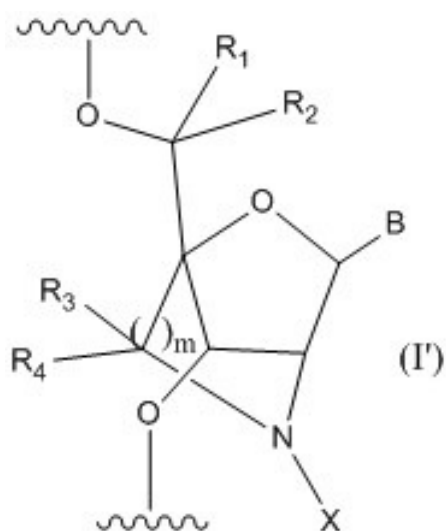
N-[1-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
5 methoxy]methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-5-methyl-2-oxo-pyrimidin-4-yl]dihydropyrimidin-2-one;

N'-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy]methyl]-7-[2-
10 cyanoethoxy(diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-N,N-dimethylformamidine;

N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy]methyl]-7-[2-cyanoethoxy-
15 (diisopropylamino)phosphanyl]oxy-5-methylsulfonyl-2-oxa-5-azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-2-methylpropane amide; and

N-[9-[(1R,3R,4R,7S)-1-[[bis(4-methoxyphenyl)-phenyl-
methoxy]methyl]-7-hydroxy-5-methylsulfonyl-2-oxa-5-
20 azabicyclo[2.2.1]heptan-3-yl]-6-oxo-1H-purin-2-yl]-2-methylpropane amide.

18. An oligonucleotide compound comprising one or more nucleosides represented by general formula (I'):



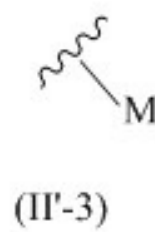
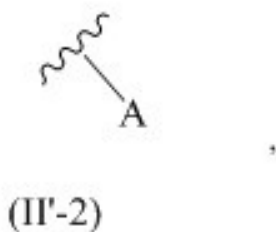
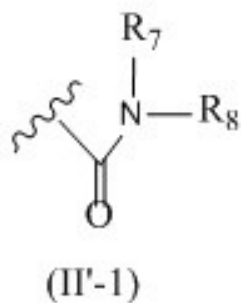
[wherein,

B represents a base moiety of nucleic acid wherein the base moiety may be optionally substituted with one or more substituents;

R₁, R₂, R₃ and R₄ represent independently of each other a hydrogen atom, or a C1-6 alkyl group which may be optionally substituted with one or more substituents;

m is an integer of 1 or 2;

X represents a group represented by the following formula (II'-1), (II'-2), or (II'-3):



;

the symbols:



which is described in the formula (II'-1), (II'-2) or (II'-3) represents a binding point to 2'-amino group;

A represents an aromatic group containing one or more
 5 heteroatoms selected from the group consisting of nitrogen atom, oxygen atom and sulfur atom, which may be optionally substituted with one or more substituents;

M represents a sulfonyl group which is substituted with one or more substituents selected from the group
 10 consisting of a C₁₋₆ alkyl group which may be optionally substituted with one or more substituents, and an aromatic group which may be optionally substituted with one or more substituents.]

or salts thereof.

15 19. The compound according to claim 18 or salts thereof wherein B represents an adeninyl group which may be optionally substituted with one or more substituents, a guaninyl group which may be optionally substituted with one or more substituents, a cytosinyl group which may be
 20 optionally substituted with one or more substituents, a 5-methylcytosinyl group which may be optionally substituted with one or more substituents or an uracil group which may be optionally substituted with one or more substituents.

20. The oligonucleotide compound according to claim 18 or
 25 19 or salts thereof wherein

R₁, R₂, R₃ and R₄ represent independently of each other a hydrogen atom; and

m is an integer of 1.

21. The oligonucleotide compound according to any one of
5 claims 18 to 20 or salts thereof, wherein

X represents a group represented by formula (II'-1);
and

R₇ and R₈ represent independently of each other a
hydrogen atom, a C1-6 alkyl group which may be optionally
10 substituted with one or more substituents, or an aromatic
group which may be optionally substituted with one or more
substituents.

22. The oligonucleotide compound according to claim 21 or
salts thereof wherein R₇ and R₈ represent independently of
15 each other a hydrogen atom, a C1-3 alkyl group which may be
optionally substituted with one or more substituents.

23. The oligonucleotide compound according to claim 22 or
salts thereof wherein one of R₇ and R₈ represents a
hydrogen atom, and the other thereof represents a methyl
20 group.

24. The oligonucleotide compound according to claim 23 or
salts thereof wherein one of R₇ and R₈ represents a
hydrogen atom, and the other thereof represents an
isopropyl group.

25. The oligonucleotide compound according to any one of

claims 18 to 20 or salts thereof wherein

X represents a group represented by formula (II'-2);
and

A represents a five or six membered heteroaryl group
5 containing one or more heteroatoms selected from the group
consisting of nitrogen atom, oxygen atom and sulfur atom,
which may be optionally substituted with one or more
substituents.

26. The oligonucleotide compound according to claim 25 or
10 salts thereof wherein

A represents a five or six membered heteroaryl group
containing two or three heteroatoms selected from nitrogen
atom, oxygen atom, and sulfur atom, provided that at least
two nitrogen atoms are included, which may be optionally
15 substituted with one or more substituents, and the
substituent is selected independently of each other from a
group consisting of a C1-3 alkyl group which may be
optionally substituted with one or more halogen atoms, a
halogen atom, and an amino group which may be optionally
20 substituted with one or more C1-3 alkyl groups.

27. The oligonucleotide compound according to claim 26 or
salts thereof wherein the five or six membered heteroaryl
group is selected from the group consisting of a triazolyl
group, an oxadiazolyl group, a thiadiazolyl group, a
25 pyrimidinyl group, and a pyrazinyl group, each may be

optionally substituted with one or more substituents.

28. The oligonucleotide compound according to claim 27 or salts thereof wherein the triazolyl group which may be optionally substituted with one or more substituents
5 represents 1,5-dimethyl-1,2,4-triazol-3-yl group.

29. The oligonucleotide compound according to claim 27 or salts thereof wherein the oxadiazolyl group which may be optionally substituted with one or more substituents represents 5-methyl-1,2,4-oxadiazol-3-yl group.

10 30. The oligonucleotide compound according to claim 27 or salts thereof wherein the thiadiazolyl group which may be optionally substituted with one or more substituents represents 3-methyl-1,2,4-thiadiazol-5-yl group.

31. The oligonucleotide compound according to any one of
15 claims 18 to 20 or salts thereof, wherein

X represents a group represented by formula (II'-3);
and

M represents a sulfonyl group which is substituted with one or more substituents selected from the group
20 consisting of a C1-6 alkyl group which may be optionally substituted with one or more substituents, and an aryl group which may be optionally substituted with one or more substituents.

32. The compound according to any one of claims 18 to 31
25 or salts thereof wherein M represents a sulfonyl group

which is substituted with one or more substituents selected from the group consisting of a methyl group and a phenyl group.

33. The oligonucleotide compound according to any one of
5 claims 1 to 32 or salts thereof wherein one or more phosphate bonds between nucleotides represent a phosphorothioate bond.