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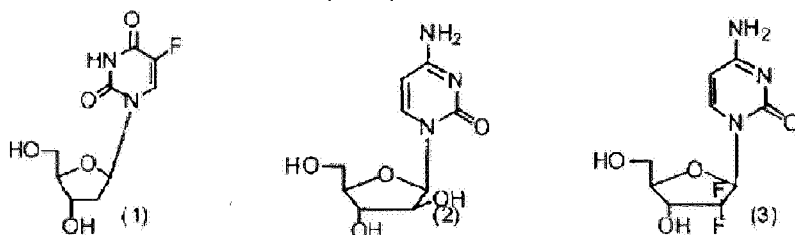
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ABRAHAM T W ET AL: "Synthesis and biological activity of aromatic amino acid phosphoramidates of 5-fluoro-2'-deoxyuridine and 1-beta-arabinofuranosylcytosine: evidence of phosphoramidase activity", JOURNAL OF MEDICINAL CHEMISTRY, AMERICAN CHEMICAL SOCIETY, US, vol. 39, 1 January 1996 (1996-01-01), pages 4569-4575, XP002147029, ISSN: 0022-2623, DOI: 10.1021/JM9603680

DESCRIPTION

Chemical Compounds

[0001] The present invention relates to nucleotide derivatives and their use in the treatment of cancer.

[0002] Nucleoside analogues such as fluorodeoxyuridine (1), cytarabine (2) and gemcitabine (3) are well established as anticancer agents. They function as inhibitors, of DNA synthesis after activation to their 5'-phosphate form.

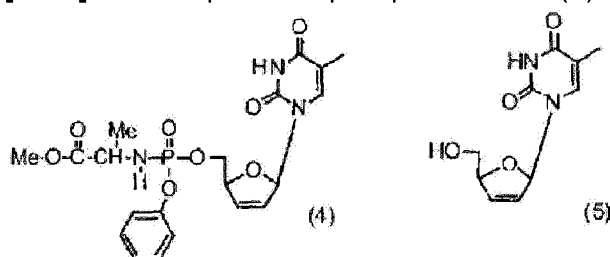


[0003] The free bioactive phosphate forms do not in general represent useful drugs due to their poor membrane permeation. In an effort to circumvent this a number of phosphate pro-drug approaches have been reported [Rosowsky et al, J. Med. Chem., 1982, 25, 171-8; Hong et al, J. Med. Chem., 1985, 28, 171-8; Kodama et al, Jpn. J. Cancer Res., 1989, 80, 679-85; Hong et al, 1979, 22, 1428-32; Ji et al, J. Med. Chem., 1990, 33, 2264-70; Jones et al, Nucleic Acids Res., 1989, 17, 7195-7201; Hunston et al, J. Med. Chem., 1984, 27, 440-4; Lorey et al, Nucleosides Nucleotides, 1997, 16, 1307-10; Farquhar et al, J. Med. Chem., 1983, 26, 1153-8; Shuto et al, Nucleosides Nucleotides, 1992, 11, 437-46; Le Bec et al, Tet. Letts., 1991, 32, 6553-6; Phelps et al, J. Med. Chem., 1980, 23, 1229-32].

[0004] In general the phosphate prodrugs have biological properties and therapeutic activities that are similar to, or somewhat lower than, the parent nucleoside analogue.

[0005] We have carried out extensive work in this area from an antiviral perspective, largely on dideoxy nucleosides, and have reported a phosphoramidate approach which has been widely adopted for the delivery of bio-active phosphates of antiviral nucleosides.

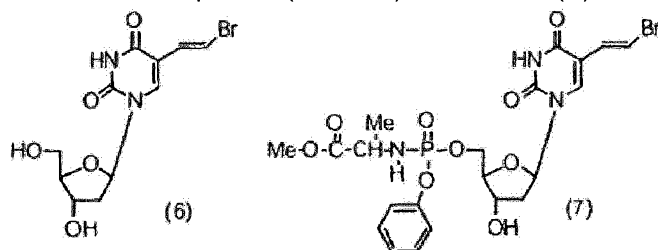
[0006] An example is the phosphoramidate (4) derived from anti-HIV d4T (5).



[0007] We observed the effect of variations in the ester [McGuigan et al, AVCC, 1998, 9, 473-9], amino acid [McGuigan et al, Antiviral Res., 1997, 35, 195-204; AVCC, 2000, 11, 111-6], and aryl [Siddiqui et al, J. Med. Chem., 1999, 42, 393-9] regions of the phosphoramidate, as well as the effect of amino acid stereochemistry [McGuigan et al, AVCC, 1996, 7, 184-8], phosphate stereochemistry [Allender et al, Analytica Chim. Acta, 2001, 435, 107-13] and nucleoside [Balzarini et al, BBRC, 1996, 225, 363-9; McGuigan et al, BioOrg. Med. Chem. Lett., 1996, 6, 2369-62; McGuigan et al, Bioorg. Med. Chem. Lett., 2000, 10, 645-7].

[0008] This work has lead to the optimal description of phenyl methoxyalaninyl phosphoramidate as the prototype pro-moiety for the intracellular delivery of bioactive nucleotides [Balzarini et al, PNAS, 1996, 93, 7295-9; McGuigan et al, J. Med. Chem., 1996, 39, 1748-53].

[0009] Lackey et al [Biochem Pharmacol., 2001, 61, 179-89] have reported the application of our phosphoramidate pro-drug method for antiviral nucleosides to the anti-herpetic agent bromovinyl-2'-deoxyuridine (BVDU) (6). In particular, they have found that the phenyl methoxyalaninyl phosphoramidate (7) has significant anti-cancer activity. This is in marked contrast to the parent (antiviral) nucleoside (6).



[0010] Limited SAR has been presented by this group, although in their patent applications [WO0239952, EP1200455, CA2317505, US6339151, EP116797, AU2451601] they claim a series of general variations in the base, and phosphate regions. However, based on our prior art, the phenyl methoxyalaninyl phosphoramidate (7) would be anticipated to be amongst the most optimal of structure.

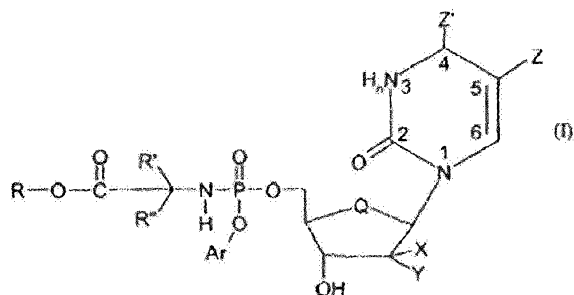
[0011] Surprisingly, it has now been found that other derivatives of oxyamino acid-phosphoramidate nucleoside analogues are significantly more potent in the treatment of cancer than the phenyl methoxyalaninyl phosphoramidate (7).

[0012] US 2003/109697 relates to phosphoramidate derivatives of nucleotides and their use in treating cells or tissue involved in a pathology characterized by hyper-proliferative cells such as cancer, infectious disease, autoimmune disorders and inflammatory conditions/diseases.

[0013] ABRAHAM T. W. et. al.: "Synthesis and Biological Activity of Aromatic Amino Acid Phosphoramidates of 5-fluoro-2'-deoxyuridine and 1-β-Arabinofuranosylcytosine: Evidence of Phosphoramidase Activity", JOURNAL OF MEDICINAL CHEMISTRY, AMERICAN CHEMICAL

SOCIETY, US, vol. 39, 1996, pages 4569-4575, XP002147029, ISSN: 0022-2623, DOI: 10.1021/JM9603680. This document relates to phosphoramidates of arabinofuranosylcytoside and FUDR for antitumor activity against L1210 mouse lymphocytic leukemia cells and CCRF-CEM human T-cell lymphoblastic leukaemia cells.

[0014] According to the first aspect of the present invention, there is provided a compound of formula I:



wherein:

R is selected from a C₁-C₁₆ branched or unbranched acyclic alkyl group, a C₃-C₁₂ cyclic alkyl group, a phenyl group or a naphthyl group and a benzyl group; and wherein R is unsubstituted;

R' and R'' are independently selected from H, methyl (-CH₃), benzyl (-CH₂C₆H₅) and secondary butyl (-CH₂CH(CH₃)₂), or, R' and R'' together with the C atom to which they are attached, provide a C₅₋₆ ring;

Q is -O-;

X and Y are both F;

Ar is naphthyl or phenyl; and one, two, three or four substituents, which may be the same or different, may be present on Ar and are selected from the group comprising halogen, -NO₂;-NH₂; -C₁₋₃alkyl; -C₁₋₃alkoxy; -SC₁₋₃alkyl; -CN; -COC₁₋₃alkyl; and -CO₂C₁₋₃alkyl; Z is H;

n is 0, Z' is -NH₂ and a double bond exists between position 3 and position 4; or a pharmaceutically acceptable salt, ester or salt of such ester of a compound on formula I.

[0015] Reference in the present specification to an alkyl group means a branched or unbranched, cyclic or acyclic, saturated or unsaturated (e.g. alkenyl or alkynyl) hydrocarbyl radical. Where cyclic, the alkylene group is preferably C₃ to C₁₂, more preferably C₅ to C₁₀, more preferably C₅ to C₇. Where acyclic, the alkyl group is preferably C₁ to C₁₆, more preferably C₁ to C₆.

[0016] Reference in the present specification to an aryl group means an aromatic group containing 5 to 14 ring atoms, for example phenyl or naphthyl. The aromatic group may be a heteroaromatic group containing one, two, three or four, preferably one, heteroatoms selected,

independently, from the group consisting of O, N and S. Examples of such heteroaromatic groups include pyridyl, pyrrolyl, furanyl and thiophenyl. Preferably, the aryl group comprises phenyl or substituted phenyl.

[0017] The alkyl and aryl groups may be substituted or unsubstituted. Where substituted, there will generally be one to three substituents present, preferably one substituent. Substituents may include halogen atoms, by which is meant F, Cl, Br and I atoms, and halomethyl groups such as CF₃ and CCl₃; oxygen containing groups such as oxo, hydroxy, carboxy, carboxyC₁₋₆alkyl, alkoxy, alkoyl, alkoyloxy, aryloxy, aryloyl and aryloyloxy; nitrogen containing groups such as amino, C₁₋₆alkylamino, diC₁₋₆alkylamino, cyano, azide and nitro; sulphur containing groups such as thiol, C₁₋₆alkylthiol, sulphonyl and sulphoxide; heterocyclic groups which may themselves be substituted; alkyl groups as defined above, which may themselves be substituted; and aryl groups as defined above, which may themselves be substituted, such as phenyl and substituted phenyl. Substituents on said heterocyclic, alkyl and aryl groups are as defined immediately above.

[0018] Reference in the present specification to alkoxy and aryloxy groups means, respectively, alkyl-O- (for example where alkyl is C₁ to C₁₆, preferably C₁ to C₆) and aryl-O- (for example where aryl is a 5 to 14 membered aromatic mono- or bifused ring moiety, optionally containing 1, 2, 3 or 4 heteroatoms selected, independently, from O, S and N, preferably aryl is phenyl).

[0019] Reference in the present specification to alkoyl and aryloyl groups means, respectively, alkyl-CO- (for example where alkyl is C₁ to C₁₆, preferably C₁ to C₆) and aryl-CO- (for example where aryl is a 5 to 14 membered aromatic mono or bifused ring moiety, optionally containing 1, 2, 3 or 4 heteroatoms selected, independently, from O, S and N, preferably aryl is phenyl).

[0020] Reference in the present specification to alkoyloxy and aryloyloxy means, respectively, alkyl-CO-O (for example where alkyl is C₁ to C₁₆, preferably C₁ to C₆) and aryl-CO-O (for example where aryl is a 5 to 14 membered mono- or bifused aromatic ring system, optionally containing 1, 2, 3 or 4 heteroatoms selected, independently, from O, S and N, preferably aryl is phenyl).

[0021] Reference in the present specification to heterocyclic groups means groups containing one or more, pyrrolyl, imidazolyl, pyrazolyl, thiazolyl, isothiazolyl, oxazolyl, pyrrolidinyl, pyrrolinyl, imidazolidinyl, imidazolynyl, pyrazolidinyl, tetrahydrofuranyl, pyranyl, pyronyl, pyridyl, pyrazinyl, pyridazinyl, piperidyl, piperazinyl, morpholinyl, thionaphthyl, benzofuranyl, isobenzofuryl, indolyl, oxyindolyl, isoindolyl, indazolyl, indolynyl, 7-azaindolyl, isoindazolyl, benzopyranyl, coumarinyl, isocoumarinyl, quinolyl, isoquinolyl, naphthridinyl, cinnolynyl, quinazolinyl, pyridopyridyl, benzoxazinyl, quinoxadinyl, chromenyl, chromanyl, isochromanyl and carbolinyl.

[0022] The group Ar comprises a substituted or unsubstituted aryl group, wherein the term

"aryl group" and the possible substitution of said group is as defined herein. Preferably, Ar is a substituted or unsubstituted phenyl group. Particularly preferred substituents are electron withdrawing groups such as halogen (preferably chlorine or fluorine), trihalomethyl (preferably trifluoromethyl), cyano and nitro groups. For example, Ar can be phenyl, 3,5-dichloro-phenyl, *p*-trifluoromethyl-phenyl, *p*-cyano-phenyl, or *p*-nitro-phenyl.

[0023] Suitably, R is a C₁₋₁₆ primary or secondary alkyl group, a C₅₋₇ carbocyclic aryl group or a C₁₋₆alkylC₅₋₁₁aryl group. More suitably, R is a C₁₋₁₀ alkyl group, a phenyl group or C₁₋₃ alkylC₅₋₇ aryl group. Preferably R is unsubstituted.

[0024] Preferably, R is methyl (-CH₃), ethyl (-C₂H₅), *n*- or *i*- propyl (-C₃H₇), *n*- or *i*- butyl (-C₄H₉) or benzyl (-CH₂C₆H₅). Most preferably, R is benzyl. Particularly, R is preferably benzyl when one of R' and R" is H and one of R' and R" is methyl (-CH₃).

[0025] Alternatively, preferably, R' and R" are, independently, H, methyl (-CH₃), secondary butyl (-CH₂-CH-(CH₃)₂), benzyl (-CH₂C₆H₅), or, together with the C atom to which they are attached, provide a C₅₋₆ ring.

[0026] Preferred compounds include those where R' and R" are both methyl, one of R' and R" is H and one of R' and R" is methyl, and R' and R", together with the C atom to which they are attached, provide a pentyl ring.

[0027] When R' and R" are different, the C atom to which they are attached is chiral. The present compounds can be L or D or a mixture of stereoisomers. Preferably they are L.

[0028] It will be appreciated that the moiety -O-C(O)-CR'R"-NH- corresponds to a carboxy-protected α-amino acid, R' and R" can thus correspond to the side chains of a naturally occurring amino acid.

[0029] For example, when one of R' and R" is H and one of R' and R" is Me or PhCH₂, the moiety corresponds to alanine or phenylalanine, respectively.

[0030] Preferably, the stereochemistry at the asymmetric centre -CR'R" corresponds to an L-amino acid. The stereochemistry at the asymmetric centre -CR'R" can, however, correspond to a D-amino acid. Alternatively, mixtures of compounds can be employed having asymmetric centres corresponding to L and D amino acids.

[0031] In the present specification by "naturally occurring amino acid" we mean Alanine, Arginine, Asparagine, Aspartic Acid, Cysteine, Cystine, Glycine, Glutamic Acid, Glutamine, Histidine, Hydroxylysine, Hydroxyproline, Isoleucine, Leucine, Lysine, Methionine, Phenylalanine, Proline, Serine, Threonine, Tryptophan, Tyrosine and Valine.

[0032] The present invention is not, however, limited to compounds having a moiety

corresponding to a naturally occurring amino acid. The present invention specifically includes compounds having a moiety which corresponds to a non-naturally occurring amino acid, such as, for example, those where $R'=R''=\text{alkyl}$, or, where together with the C atom to which they are attached, R' and R'' provide a cyclic moiety. Preferably with respect to the compound of formula I, the moiety ROCOCR'R''NH- corresponds to or is derived from a non-naturally occurring amino acid.

[0033] Z is H, Q is O, n is 0, X and Y are each F, and thus the base moiety of the compound of formula I corresponds to that of gemcitabine i.e. compound (3) above.

[0034] Compounds of formula 1 are those wherein n is 0 and X and Y are F. Compounds of formula I are those wherein n is 0, X and Y are F, Q is O and Z is H, thus corresponding to phosphoramidated gemcitabine.

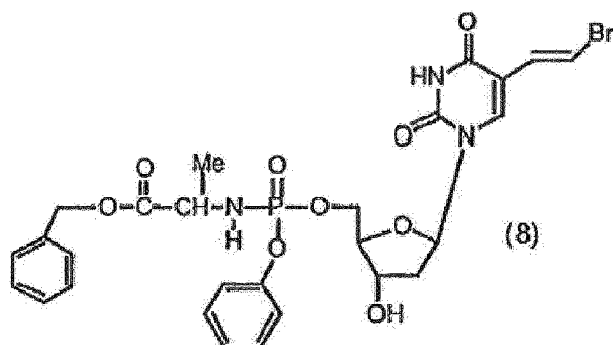
[0035] Preferably, Ar is an optionally substituted phenyl.

[0036] One, two, three or four substituents, which may be the same or different, may be present on Ar and are selected from the group comprising halogen, which may -F, -Cl, -Br or -I; $-\text{NO}_2$; $-\text{NH}_2$; optionally substituted $-\text{C}_{1-3}\text{alkyl}$; optionally substituted $-\text{C}_{1-3}\text{alkoxy}$, preferably methoxy ($-\text{OCH}_3$); optionally substituted $-\text{SC}_{1-3}\text{alkyl}$; $-\text{CN}$; optionally substituted $-\text{COC}_{1-3}\text{alkyl}$; and optionally substituted $-\text{CO}_2\text{C}_{1-3}\text{alkyl}$. The optional substituents are one or more up to six, preferably three, members selected from the group comprising halogen which may be F, Cl, Br and I and NO_2 . Preferred substituents on Ar include F, Cl, CF_3 , and NO_2 .

[0037] The substituents may be at any position on the ring moiety, Where the ring moiety is C_6 ie phenyl, a single substituent at the 2 (*ortho*) or 4 (*para*) position is preferred. Where Ar is phenyl, a single substituent at the 4 position is more preferred.

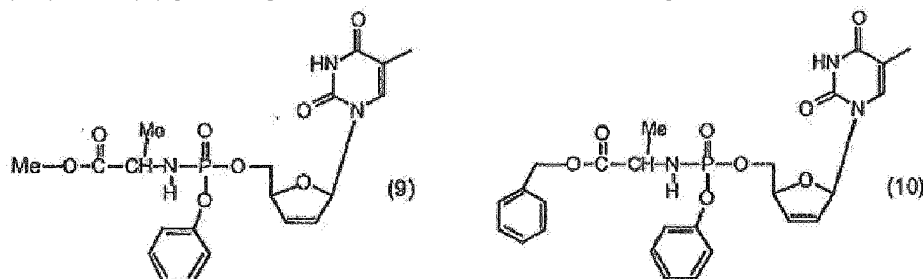
[0038] Preferably, Ar is an optionally substituted phenyl moiety. More preferably, Ar is selected from the group comprising: Ph- , $p\text{CF}_3\text{C}_6\text{H}_4\text{-}$, $p\text{FC}_6\text{H}_4\text{-}$, $p\text{NO}_2\text{C}_6\text{H}_4\text{-}$, $p\text{ClC}_6\text{H}_4\text{-}$ and $o\text{ClC}_6\text{H}_4\text{-}$.

[0039] Surprisingly, modifying the ester moiety in compound (7) has been found to show a marked increase in potency with respect to cancer cell lines. A compound of the present disclosure is the benzyl ester (8). It has surprisingly been found that the benzyl ester (8) is very significantly more potent against several cancer cell lines than the methyl ester (7):



[0040] Compound (8) inhibits the growth of colon cancer cell line HT115 by 50% at 1.4 μM , whilst (7) requires a concentration of 244 μM ; (8) is thus 174 times more potent. Compound (8) is also 8 times more potent than (7) versus prostate cancer cell line PC-3 (19 μM vs. 155 μM).

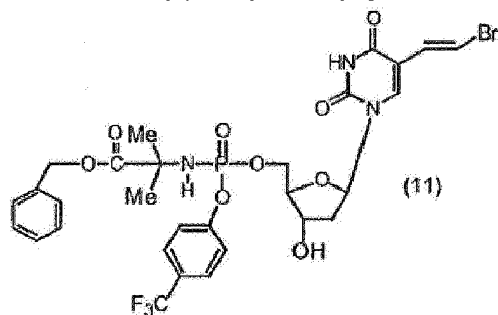
[0041] The degree of potency enhancement for (8) vs. (7) is surprising based on the prior art. Thus, comparing the equivalent phosphoramidates of d4T reveals a ca 4-fold potency boost of (10) over (9) [McGuigan et al, AVCC, 1998, 9, 473-9].



[0042] This would imply that the benzyl phosphoramidate motif in (10) is ca 4-fold more efficient at the intracellular delivery of the bio-active free phosphate forms of d4T than is the methyl ester (9). A person skilled in the art would anticipate a similar degree of enhancement for the benzyl phosphoramidate of BVDU (8) over the methyl ester (7) whilst we observed an almost 200-fold enhancement for colon cancer as noted above.

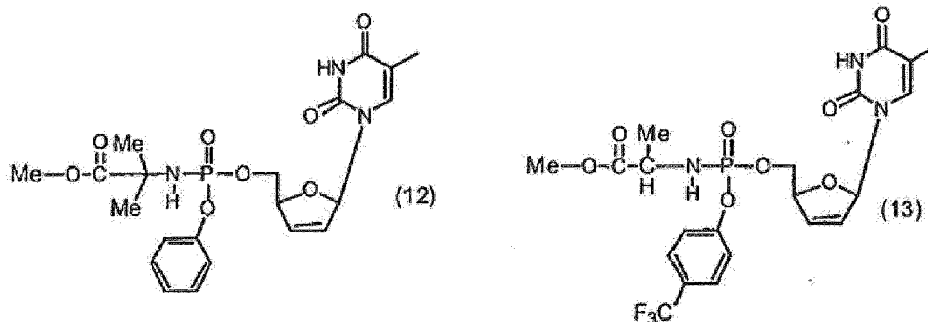
[0043] Surprising efficacy of modifications in the amino acid and aryl moieties of the BVDU phosphoramidate has also been found in compounds of the present disclosure.

[0044] Thus, compound (11) has simultaneous modification in these two regions, being the p-trifluoromethylphenyl benzyl [α,α -dimethylglycyl]phosphoramidate.



[0045] Compound 11 shows high potency against a range of cancer cell types and is significantly and surprisingly more potent than (7). Thus, for breast cancer (11) is 60-fold more active (1.3 μM vs 79 μM), and for prostate cancer (11) is 254-fold more potent (0.61 μM vs. 155 μM). Against colon cancer, (11) is 35-fold more potent (7 μM vs 244 μM). Again, the degree of enhancement of the analogue (11) vs. (7) is surprising based on prior art. Thus, comparing (12) [dimethyl glycine modification] and (13) [p- CF_3 phenyl modification] to (9) shows no

significant difference in potency.



[0046] Thus 50% effective doses vs HIV-1 for (9), (12) and (13) are: 0.075, 0.29, and 0.01 μ M respectively; within experimental error, (12) and (13) are identical in potency to (9). Thus a person skilled in the art would have predicted that (11) would show little enhancement over (7) as opposed to the 35 to 254-fold enhancements noted above.

[0047] Thus, compounds of the present disclosure and having variations in one or more of the ester (R), amino acid (R', R'') and aryl (Ar) region of the phosphoramidate structure compared to phenyl methoxyalaninyl phosphoramidate can give surprising and substantial potency boosts of pro-tides derived from BVDU against a range of cancer cell types.

[0048] According to a further aspect of the present invention there is provided a compound having formula I according to the present invention for use in a method of treatment, preferably in the prophylaxis or treatment of cancer.

[0049] According to a further aspect of the present disclosure there is provided a method of prophylaxis or treatment of cancer comprising administration to a patient in need of such treatment an effective dose of a compound having formula I according to the present invention.

[0050] According to a further aspect of the present disclosure there is provided use of a compound having formula I of the present invention in the manufacture of a medicament for use in the treatment or prophylaxis of cancer.

[0051] According to a further aspect of the present invention there is provided a pharmaceutical composition comprising a compound having formula I of the present invention in combination with a pharmaceutically acceptable excipient, carrier or diluent.

[0052] According to a further aspect of the present invention there is provided a method of preparing a pharmaceutical composition comprising the step of combining a compound having formula I of the present invention with a pharmaceutically acceptable excipient, carrier or diluent.

[0053] The present invention is particularly applicable for the treatment of a patient having breast cancer, colon cancer or prostate cancer. Examples of such cancers include breast MDA

MB231, colon HT115 and prostate PC-3.

[0054] The compound having formula I or pharmaceutical composition according to the present invention can be administered to a patient, which may be human or animal by any suitable means.

[0055] The medicaments employed in the present invention can be administered by oral or parenteral routes, including intravenous, intramuscular, intraperitoneal, subcutaneous, transdermal, airway (aerosol), rectal, vaginal and topical (including buccal and sublingual) administration.

[0056] For oral administration, the compounds of the invention will generally be provided in the form of tablets or capsules, as a powder or granules, or as an aqueous solution or suspension.

[0057] Tablets for oral use may include the active ingredient mixed with pharmaceutically acceptable excipients such as inert diluents, disintegrating agents, binding agents, lubricating agents, sweetening agents, flavouring agents, colouring agents and preservatives. Suitable inert diluents include sodium and calcium carbonate, sodium and calcium phosphate, and lactose, while cornstarch and alginic acid are suitable disintegrating agents. Binding agents may include starch and gelatin, while the lubricating agent, if present, will generally be magnesium stearate, stearic acid or talc. If desired, the tablets may be coated with a material such as glyceryl monostearate or glyceryl distearate, to delay absorption in the gastrointestinal tract.

[0058] Capsules for oral use include hard gelatin capsules in which the active ingredient is mixed with a solid diluent, and soft gelatin capsules wherein the active ingredient is mixed with water or an oil such as peanut oil, liquid paraffin or olive oil.

[0059] Formulations for rectal administration may be presented as a suppository with a suitable base comprising for example cocoa butter or a salicylate.

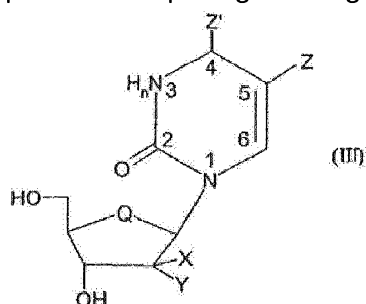
[0060] Formulations suitable for vaginal administration may be presented as pessaries, tampons, creams, gels, pastes, foams or spray formulations containing in addition to the active ingredient such carriers as are known in the art to be appropriate.

[0061] For intramuscular, intraperitoneal, subcutaneous and intravenous use, the compounds of the invention will generally be provided in sterile aqueous solutions or suspensions, buffered to an appropriate pH and isotonicity. Suitable aqueous vehicles include Ringer's solution and isotonic sodium chloride. Aqueous suspensions according to the invention may include suspending agents such as cellulose derivatives, sodium alginate, polyvinylpyrrolidone and gum tragacanth, and a wetting agent such as lecithin. Suitable preservatives for aqueous suspensions include ethyl and n-propyl p-hydroxybenzoate.

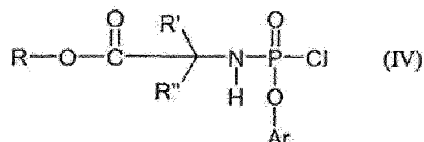
[0062] The compounds of the invention may also be presented as liposome formulations.

[0063] In general a suitable dose will be in the range of 0.1 to 300 mg per kilogram body weight of the recipient per day. A preferred lower dose is 0.5 mg per kilogram body weight of recipient per day, a more preferred lower dose is 6 mg per kilogram body weight of recipient per day, an even more preferred lower dose is 10 mg per kilogram body weight per recipient per day. A suitable dose is preferably in the range of 6 to 150 mg per kilogram body weight per day, and most preferably in the range of 15 to 100 mg per kilogram body weight per day. The desired dose is preferably presented as two, three, four, five or six or more sub-doses administered at appropriate intervals throughout the day. These sub-doses may be administered in unit dosage forms, for example, containing 10 to 1500 mg, preferably 20 to 1000 mg, and most preferably 50 to 700 mg of active ingredient per unit dosage form.

[0064] According to a further aspect of the present invention there is provided a process for the preparation of a compound having formula I according to the present invention, the process comprising reacting of a compound of formula (III):



with a compound of formula (IV):



wherein Ar, n, Q, R, R', R'', X, Y, Z' and Z have the meanings described above with respect to formula (I).

[0065] Embodiments of the present invention will now be described, by way of example only, with reference to the following examples, experimental procedures and experimental data.

[0066] Data are presented for a range of structures against tumour cell types representing a range of common cancers in man with un-met clinical need: breast MDA MB231, colon HT115, prostate PC-3. Data from these assays are presented as Table I.

Experimental Procedure

[0067] For the absence of doubt, compounds below labelled as "Illustrative Examples" do not form part of the invention.

General methods

[0068] The following anhydrous solvents and reagents were bought from Aldrich with sure stopper: dichloromethane (DCM), diethyl ether (Et₂O), tetrahydrofuran (THF), N-methylimidazole (NMI), methanol (MeOH), dimethylformamide (DMF), 1,4-dioxane. triethylamine was dried on molecular sieves of 4 Angstrom.

Thin Layer Chromatography

[0069] Thin layer chromatography (TLC) was performed on commercially available Merck Kieselgel 60 F₂₅₄ plates and separated components were visualized using ultraviolet light (254 nm and 366 nm).

Column Chromatography

[0070] Columns were performed using (Kieselgel 60, 35-70µm, Fluka) as the stationary phase, Samples were applied as a concentrated solution in the same eluent, or pre-adsorbed onto silica gel.

NMR Spectroscopy

[0071] ¹H, ¹³C and ³¹P-NMR were recorded on a Bruker Avance DPX300 spectrometer with operating frequencies of 300MHz, 75MHz and 121MHz respectively. ³¹P-MMR spectra are reported in units of δ relative to 85% phosphoric acid as external standard, positive shifts are downfield. The following abbreviations are used in the assignment of NMR signals: s (singlet), d (doublet), t (triplet), q (quartet), m (multiplet), bs (broad signal), dd (doublet of doublet), dt (doublet of triplet). Starred signal signal are splitted due to stereoisomeric mixtures.

Standard procedures

[0072] For practical purposes, standard procedures are given where applicable.

Standard procedure 1: Synthesis of Amino ester hydrochloride salts.

[0073] To a stirring solution of anhydrous alcohol (10 mol eq.) was added thionyl chloride (2 mol eq.) at 0° C, and the resulting solution stirred for 1 hr. After warming to room temperature, the appropriate amino acid (1 mol eq) was added and the reaction heated at reflux for 6-16

hrs. Removal of solvent and recrystallisation from methanol/ether gave the amino ester hydrochloride salts.

Standard procedure 2: Synthesis of Amino benzyl ester hydrochloride salts.

[0074] The appropriate amino acid (1.0 mol eq.), *p*-toluene sulfonic acid (1.0 mol eq.) and anhydrous benzyl alcohol (4.1 mol eq.) were heated at reflux in toluene (10 mol eq.) with Dean-Stark trap for 24 hrs. On cooling to room temperature, Et₂O was added and the mixture was left in ice bath for 1hr then filtrated and washed with Et₂O. The solid was dissolved in DCM and washed with 10% K₂CO₃ and water, The organic layer was dried over MgSO₄, filtered and the solvent removed under reduced pressure to give an oil. This was solubilized in acetone and neutralized with 1 M HCl. Et₂O was added and the solid was filtered and washed with Et₂O to give a white solid.

Standard procedure 3: Synthesis of Phosphorodichloridate species.

[0075] Phosphorus oxychloride (1.0 mol eq.) and the appropriate substituted phenol (1.0 mol) were stirred with anhydrous diethylether (31 mol eq.). To this was added anhydrous triethylamine (1.0 mol eq) at -80 °C and left to rise to room temperature over 16 hrs. the triethylamine hydrochloride salt was filtered off, and the filtrate reduced to dryness to give the crude product as a clear liquid.

Standard procedure 4: Synthesis of Phosphochloridate species.

[0076] Phosphodichloridate (1.0 mol eq.) and the appropriate amino ester hydrochloric salt (1.0 mol eq.) were suspended in anhydrous DCM. Anhydrous triethylamine was added dropwise at -80 °C and after 1hr the reaction was left to rise to room temperature. The formation of phosphochloridate was monitored by ³¹P-NMR. After 2-5 hrs the solvent was removed under reduced pressure and the solid obtained washed with anhydrous ether (2x20 ml), filtered, and the filtrate reduced to dryness to give the products as crude oil. These oils were usually used without further purification.

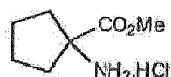
Standard procedure 5: Synthesis of Phosphoroamidate derivatives.

[0077] To a stirring solution of (E)-5-(2-bromovinyl)-2'-deoxyuridine (1.0 mol eq.) and the appropriate phosphochloridate (2.0- 3.0 mol eq) in anhydrous THF at -80°C was added dropwise over 1 min NMI (5.0 mol eq.). After 15 mins the reaction was left to rise to room temperature and stirred at room temperature for 2-19 hrs. The solvent was removed under

reduced pressure and the yellow oil obtained was dissolved in DCM, washed with 0.5 M HCl, and water. The organic layer is dried over MgSO_4 , filtered, reduced to dryness and purified by flash chromatography (Chloroform/Methanol 97/3, Dichloromethane/Methanol 97/3).

Synthesis of Methyl-1-amino-1-cyclopentanoate hydrochloride salt.

[0078] $\text{C}_6\text{H}_{14}\text{ClNO}_3$, MW=179.68.



[0079] This was synthesised according to **Standard Procedure 1**, using 1-amino-1-cyclopentanecarboxylic acid (3.876 g, 30 mmol) with thionyl chloride (4.44 mL, 45 mmol,) and anhydrous methanol (15.5 mL). The product was isolated as a white solid (4.81 g, yield 89%).

$^1\text{H-NMR}$ (CDCl_3 ; 300 MHz): δ 9.1 (3H, bs, NH_3^+Cl^-), 3.85 (3H, s, OCH_3), 2.3-2.2 (4H, m, 4H cyclopentane), 2.15 (2H, 2H cyclopentane), 1.95 (2H, m, 2H cyclopentane),

$^{13}\text{C-NMR}$ (CDCl_3 ; 75 MHz): δ 26.6 (2 CH_2 cyclopent), 38.1 (2 CH_2 cyclopent), 54.8 (CH_3O), 66.6 (C_α cyclopentane), 174.1 (COOMe).

Synthesis of Ethyl-1-amino-1-cyclopentanoate hydrochloride salt.

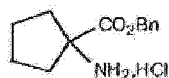
[0080] $\text{C}_8\text{H}_{16}\text{ClNO}_2$, MW=193.71.



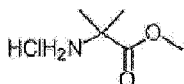
[0081] This was synthesised according to **Standard Procedure 1**, using 1-amino-1-cyclopentanecarboxylic acid (5.0 g, 38.6 mmol) with thionyl chloride (5.72 mL, 58 mmol) and anhydrous ethanol (29 mL). The product was isolated as a white solid (6.98 g, yield 93%).

$^1\text{H-NMR}$ (CDCl_3 ; 300 MHz): δ 9.0 (3H, bs, NH_3^+Cl^-), 4.3 (2H, q, $^3J=8$, OCH_2CH_3), 2.3-2.2 (4H, m, 4H cyclopentane), 2.15 (2H, 2H cyclopentane), 1.95 (2H, m, 2H cyclopentane), 1.4 (3H, t, $^3J=8$, OCH_2CH_3).

$^{13}\text{C-NMR}$ (CDCl_3 ; 75 MHz): δ 14.5 (CH_3CH_2), 25.8 (2 CH_2 cyclopent), 37.4 (2 CH_2 cyclopent), 63.0 (CH_3CH_2), 66.2 (C_α cyclopentane), 172.1 (COOEt).

Synthesis of Benzyl-1-amino-1-cyclopentanoate hydrochloride salt.**[0082]** C₁₄H₁₈ClNO₂, MW-255.78.

[0083] This was synthesised according to **Standard Procedure 2**, using 1-amino-1-cyclopentanecarboxylic acid (3.682 g, 28.5 mmol) with *p*-toluene sulfonic acid monohydrate (5.625 g, 29.55 mmol) and anhydrous benzylic alcohol (12 mL, 116 mmol), in Toluene (20 mL). The product was isolated as a white solid (6.441 g, yield 88.5%) **Hydrochloride salt**. ¹H-NMR (CDCl₃; 300 MHz): δ 9.05 (3H, bs, NH₃⁺Cl⁻), 7.4-7.25 (5H, m, Ph), 5.15 (2H, s, CH₂Ph), 2.3 (4H, m, 4H cyclopentane), 2.15 (2H, 2H cyclopentane), 1.95 (2H, m, 2H cyclopentane). ¹³C-NMR (CDCl₃; 75 MHz): δ 25.9 (2CH₂ cyclopent), 37.3 (2CH₂ cyclopent), 66.3 (Cq cyclopentane), 68.3 (CH₂Ph), 129.2, 129.0, 128.8 ('o', 'm', CH₂Ph), 135.5 ('p', CH₂Ph). 172.1 (COOBn).

Synthesis of methyl-2-amino-2-methylpropanoate hydrochloride salt**[0084]** C₅H₁₂ClNO₃, MW 153.61.

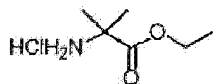
[0085] This was synthesised according to **Standard Procedure 1**, using 2-amino-isobutyric acid (5.102 g, 48.49 mmol) with thionyl chloride (11.538 g, 96.98 mmol, 7.04 mL) and anhydrous methanol (19.6 mL). The product was isolated as a white solid (6.036 g, yield 89.2%).

¹H-NMR (CDCl₃; 300 MHz): δ 8.81 (3H, bs, NH₃Cl), 3.83 (3H, s, OCH₃), 1.74 (6H, s, [CH₃]₂C).

¹³C-NMR (CDCl₃; 75 MHz): δ 24.1, 24.3 ([CH₃]₂C), 57.9 (C[CH₃]₂), 172.4 (COOCH₃).

Synthesis of ethyl-2-amino-2-methylpropanoate hydrochloride salt.

[0086] C₆H₁₄ClNO₂, MW 167.63.



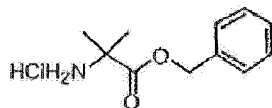
[0087] This was synthesized according to **Standard Procedure 1**, using 2-amino-isobutyric acid (5.102 g, 48.49 mmol) with thionyl chloride (11.772 g, 98.95 mmol, 7.2 mL) and anhydrous ethanol (29 mL). The product was isolated as a white solid (7.159 g, yield 86.3%).

¹H-NMR (CDCl₃; 300 MHz): δ 8.93 (3H, bs, NH₃⁺Cl⁻), 4.3 (2H, q, ³J=7.1 Hz, OCH₂CH₃), 1.75 (6H, s, [CH₃]₂C), 1.33 (3H, t, ³J=7.1 Hz, OCH₂CH₃).

¹³C-NMR (CDCl₃; 75 MHz): δ 14.4 (CH₃CH₂O), 24.3 ([CH₃]₂C), 57.9 (C[CH₃]₂), 63.1 (OCH₂CH₃), 171.6 (COOCH₂CH₃).

Synthesis of benzyl-2-amino-2-methylpropanoate hydrochloride salt.

[0088] C₁₁H₁₆ClNO₂, MW 229.70.



[0089] This was synthesised according to **Standard Procedure 2**, using 2-amino-isobutyric acid (1.960 g, 19.00 mmol) with *p*-toluene sulfonic acid monohydrate (3.750g, 19.7 mmol) and benzylic alcohol (8.360 g, 77.30 mmol, 8 mL), in toluene (20 mL). The product was isolated as a white solid (2.556 g, yield 87.4%)

***p*-toluenesulfonate salt:** ¹H-NMR (CDCl₃, 300 MHz): δ 8.40 (3H, bs, NH₃⁺Cl⁻), 7.79 (2H, d, ³J=8.0 Hz, 'm' *p*-TSA), 7.34 (5H, m, CH₂Ph), 7.14 (2H, d, ³J=8.0 Hz, 'o' *p*-TSA), 5.16 (2H, s, CH₂Ph), 2.38 (3H, s, CH₃ *p*-TSA), 1.57 (6H, s, [CH₃]₂C)

¹³C-NMR (CDCl₃; 75 MHz): δ 21.8 (CH₃, *p*-TSA), 23.9 ([CH₃]₂C), 57.8 (C[CH₃]₂), 68.3 (CH₂Ph), 126.55, 128.5, 128.8, 129.0, 129.3 (CH₂Ph+*p*-TSA), 135.4 ('ipso', CH₂Ph), 140.8 ('*p*', *p*-TSA), 141.9 ('ipso', *p*-TSA), 171.9 (COOCH₂Ph).

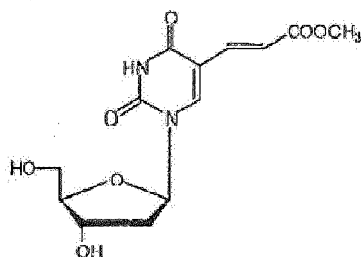
Hydrochloride salt: ¹H-NMR (CDCl₃; 300 MHz): δ 9.10 (3H, bs, NH₃⁺Cl⁻), 7.41-7.31 (5H, m, CH₂Ph), 5.27 (2H, s, CH₂Ph), 1.77 ([CH₃]₂C).

¹³C-NMR (CDCl₃; 75 MHz): δ 24.2 ([CH₃]₂C), 58.0 (C[CH₃]₂), 68.5 (CH₂Ph), 128.62, 129.0, 129.1 ('o', 'm', 'p', CH₂Ph), 135.2 ('ipso', CH₂Ph), 171.8 (COOCH₂Ph).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine

(E)-5-(2-Carbomethoxyvinyl)-2'-deoxyuridine

[0090]



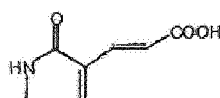
[0091] A mixture of $\text{Pd}(\text{OAc})_2$ (0.316 g, 1.41 mmol), PPh_3 (0.741 g, 2.82 mmol), and triethylamine (4.9 mL) in 1,4-dioxane (50 mL) was stirred at 70°C until an intense red colour had developed. To this 5-iodo-2'-deoxyuridine (10 g, 28.24 mmol) and methylacrylate (4.862 g, 56.48 mmol, 5.1 mL) in 1,4-dioxane (20 mL) were added and the mixture stirred at reflux for 30 mins. The reaction was filtered while still hot and the filtrate cooled over night at 4°C. The resulting pale yellow precipitate was filtered, washed with DCM and dried *in vacuo* to give the product as white solid (6.2 g, yield 70.7%).

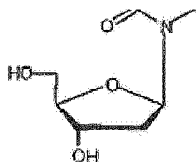
$^1\text{H-NMR}$ ($\text{DMSO-}d_6$; 300 MHz) δ 11.64 (1H, bs, NH-3), 8.42 (1H, s, H-6), 7.37 (1H, d, $^3J=15.8$ Hz, H vinylic), 6.86 (1H, d, $^3J=15.8$ Hz, H vinylic), 6.13 (1H, t, $^3J=6.5$ Hz, H-1'), 5.27-5.20 (2H, 2bs, OH-3', OH-5'), 4.27 (1H, m, H-3'), 3.81 (1H, m, H-4'), 3.68 (3H, s, CH_3), 3.60 (2H, m, H-5'), 2.18 (2H, m, H-2').

$^{13}\text{C-NMR}$ ($\text{DMSO-}d_6$; 75 MHz): δ 40.4 (C-2'), 51.6 (CH_3), 66.7 (C-5'), 70.0 (C-3'), 85.2 (C-4'), 88.0 (C-1'), 108.5 (C-5), 116.5 (C-5b), 138.5 (C-5a), 144.4 (C-6), 149.6, 162.1 (C-2, C-4), 167.6 (COO).

Synthesis of illustrative example (E)-5-(2-Carboxyvinyl)-2'-deoxyuridine

[0092]





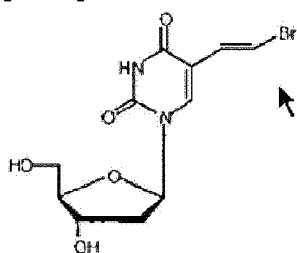
[0093] (E)-5-(2-carbomethoxyvinyl)-2'-deoxyuridine (6.0 g, 19.33 mmol) was dissolved in 300 mL of 1 M NaOH and the mixture stirred at room temperature for 3 hrs, filtered and the filtrate adjusted to pH 2 with 1M HCl. On cooling at 4°C a white precipitate formed. This was filtered off and washed with cold water (2x 20 ml) and acetone (2x20 mL) and dried to give a white solid (4.441g, yield 77.1%).

¹H-NMR (DMSO-*d*₆; 300 MHz): δ 12.18 (1H, bs, CO₂H), 11.64 (1H, s, NH-3), 8.40 (1H, s, H-6), 7.30 (1H, d, ³J=15.6 Hz, H vinyl), 6.78 (1H, d, ³J=15.8 Hz, H vinyl), 6.14 (1H, t, ³J=6.4 Hz, H-1'), 5.38-5.08 (2H, bs, OH-3', OH-5'), 4.26 (1H, m, H-3'), 3.80 (1H, m, H-4'), 3.64 (2H, m, H-5'), 2.18 (2H, m, H-2').

¹³C-NMR (DMSO-*d*₆; 75 MHz): δ 40.1 (C-2'), 61.2 (C-5'), 70.1 (C-3'), 85.1 (C-4'), 88.0 (C-1'), 108.7 (C-5), 118.0 (C-5b), 137.9 (C-5a), 143.9 (C-6), 149.6, 162.1 (C-2, C-4), 168.4 (COOH).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine

[0094]



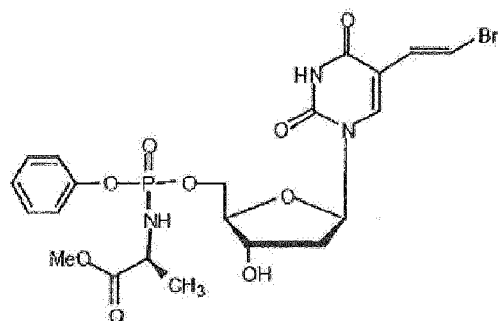
[0095] To a solution of (E)-5-(2-carboxyvinyl)-2'-deoxyuridine (5.777 g, 19.37 mmol) in dimethylformamide (29 mL) was added K₂CO₃ (5.890 g, 42.61 mmol) and the suspension stirred at room temperature for 15 mins. A solution of N-bromosuccinimide (3.655 g, 20.53 mmol) was added dropwise over 30 mins at 20°C. The resulting suspension was filtered and the solid washed with DMF. The combined filtrate and washings were evaporated to dryness *in vacuo* and the residue dissolved in MeOH. To this silica gel was added and the suspension evaporated to dryness and the solid applied to the top of chromatographic column. The column was eluted with chloroform/methanol 92/8 to give a white solid (5787g, 71.9%). Crystallisation from water gave a white powder.

^1H -NMR (DMSO- d_6 300 MHz) δ 11.59 (1H, bs, NH-3), 8.08 (1H, s, H-6), 7.25 (1H, d, $^3J=13.6$ Hz, H-5b), 6.85 (1H, d, $^3J=13.6$ Hz, H-5a), 6.13 (1H, t, $^3J=6.5$ Hz, H-1'), 5.29 (1H, bs, OH-3'), 5.13 (1H, bs, OH-5'), 4.24 (1H, m, H-3'), 3.79 (1H, m, H-4'), 3.66 (2H, m, H-5'), 2.51 (1H, m, H-2'), 2.14 (1H, m, H-2').

^{13}C -NMR (DMSO- d_6 ; 75 MHz): δ 40.2 (C-2'), 61.3 (C-5), 70.3 (C-4'), 84.8 (C-3'), 87.8 (C-1'), 108.9 (C-5b), 110.0 (C-5), 130.3 (C-5a), 149.6, 162.1 (C-2, C4).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[phenyl-(methoxy-L-alaninyl)]-phosphate (CPF 1).

[0096] $\text{C}_{21}\text{H}_{25}\text{BrN}_3\text{O}_9\text{P}$, MW 574.32.



[0097] This was synthesised according to *Standard procedure 5*, using BVdU (300 mg, 0.90 mmol), Phenyl-(methoxy-L-alaninyl)-phosphorochloridate (472 mg, 1.7 mmol), NMI (4.5 mmol, 378 μL) in THF (9 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (356 mg, yield 69%).

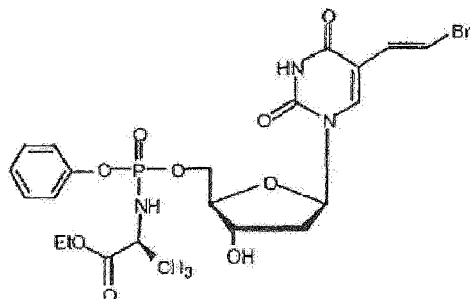
^{31}P -NMR (CDCl_3 , 121 MHz): δ 4.72, 4.40.

^1H -NMR (CDCl_3 , 300 MHz): δ 9.9 (1H, bs, H-3), 7.64 (1H, 2xs, H-6), 7.44-7.39 (1H, 2d, $^3J=14$ Hz, H-5b), 7.37-7.15 (5H, m, *OPh*), 6.75-6.67 (1H, 2d, $^3J=14$ Hz, H-5a), 6.30-6.21 (1H, 2t, $^3J=6$ Hz, H-1'), 4.57-4.29 (3H, m, H-5'+H-3'), 4.2-3.96 (3H, H-4', NH, CHala), 3.72 (3H, s, CH_3O), 2.49-2.40 (1H, m, one of H-2'), 2.12-2.01 (1H, m, one of H-2'), 1.38 (3H, d, $^3J=7$ Hz, CH_3_{ala}).

^{13}C -NMR (DMSO; 75 MHz): δ 22.4 (CH_3_{ala}), 41.9, 41.8 (C-2'), 51.9 ($\text{CH}[\text{CH}_3]$), 54.3 (CH_3O), 67.5 (C-5'), 72.3, 71.9 (C-3'), 87.3, 87.2, 86.9, 86.8 (C-1', C-4'), 110.6 (C-5b), 113.1 (C-5), 121.7 ('o', *OPh*), 127.0 ('p', *OPh*), 130.1 (C-5a), 131.5 ('m', *OPh*), 139.2 (C-6), 150.9 ('ipso', *OPh*), 151.9 (C-4), 163.2 (C-2), 175.7 (COOCH_3).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[phenyl-(ethoxy-L-alaninyl)]-phosphate (CPF 3).

[0098] $C_{22}H_{27}BrN_3O_9P$, MW=588.34.



[0099] This was synthesised according to *Standard procedure 5*, using BVdU (150 mg, 0.45 mmol), Phenyl(ethoxy-L-alaninyl)-phosphorochloridate (249 mg, 0.9 mmol), NMI (2.8 mmol, 190 μ L) in THF (4 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (145 mg, yield 55%).

^{31}P -NMR ($CDCl_3$, 121 MHz): δ 4.48, 4.86.

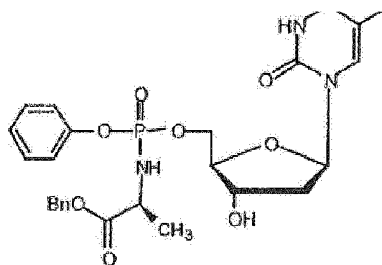
1H -NMR ($CDCl_3$, 300 MHz): δ 7.65 (1H, 2xs, H-6), 7.44-7.39 (1H, 2d, $^3J=13$ Hz, H-5b), 7.35-7.10 (5H, m, *OPh*), 6.78-6.65 (1H, 2d, $^3J=13$ Hz, H-5a), 6.35-6.25 (1H, 2t, $^3J=6$ Hz, H1'), 4.62-3.95 (8H, m, H-5', H-3', H-4', CHala, NH, CH_3CH_2O), 2.49-2.40 (1H, m, one of H-2'), 2.10-2.00 (1H, m, one of H-2'), 1.40 (3H, d, $^3J=7$ Hz, CH_3_{ala}), 1.25 (3H, 2t, $^3J=7$ Hz, CH_3CH_2O).

^{13}C -NMR ($CDCl_3$, 75 MHz): δ 14.5 (CH_3CH_2O) 21.2, 21.1 (CH_3ala), 40.9, 40.7 (C-2'), 50.8, 50.7 (CHala), 62.2, 62.1 (CH_3CH_2O), 66.5, 66.3 (C-5'), 70.9, 70.6 (C-3'), 86.0, 85.6 (C-1', C-4'), 110.1 (C-5b), 111.8 (C-5), 120.6 ('o', *OPh*), 125.0 ('p', *OPh*), 129.0 (C-5a), 130.2 ('m', *OPh*), 138.2 (C-6), 149.9 (C-4), 150.7 ('ipso', *OPh*), 162.3 (C-2), 174.2, 174.1 ($COOCH_2CH_3$).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[phenyl-(benzoxy-L-alaninyl)]-phosphate (CPF 2).

[0100] $C_{27}H_{29}BrN_3O_9P$, MW=649.08.





[0101] This was synthesized according to *Standard procedure 5*, using BVdU (150 mg, 0.45 mmol), Phenyl-(benzyloxy-L-alaninyl)-phosphorochloridate (249 mg, 0.9 mmol), NMI (2.8 mmol, 190 μ L) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (228 mg, yield 78%).

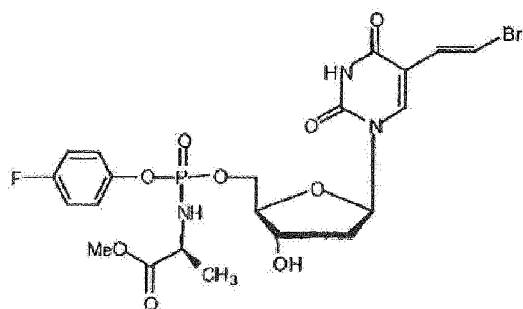
^{31}P -NMR (CDCl_3 , 121 MHz): δ 4,74, 4.44.

^1H -NMR (CDCl_3 , 300 MHz): δ 10.31 (1H, bs, H-3), 7.63 (1H, 2xs, H-6), 7.45-7.14 (11H, m, $\text{OPh} + \text{CH}_2\text{Ph}$, H-5b), 6.75-6.66 (1H, 2d, $^3J=14$ Hz, H-5a), 6.30-6.25 (1H, m, H-1'), 5.18-5.09 (1H, s, CH_2Ph), 4.70-4.04 (6H, m, H-3', H-5', H-4' NH , CHala), 2.42 (1H, m, one of H-2'), 2.02 (1H, m, one of H-2'), 1.40 (3H, d, $^3J=7$ Hz, CH_3ala).

^{13}C -NMR (CDCl_3 , 75 MHz): δ 20.7, 20.8 (CH_3ala), 40.4 (C-2'), 50.4 (CHala), 66.0 (C-5'), 67.4 (CH_2Ph), 70.6 (C-3'), 85.4, 85.5, 85.6, 85.8 (C-1', C-4'), 109.9 (C-5b), 111.5 (C-5b), 120.2 ('o', OPh), 125.4 ('p', OPh), 128.5, 128.6, 129.9 ('m' OPh , Bn, C-5a), 135.1 ('ipso', CH_2Ph), 137.8 (C-6), 149.8 (C-4), 150.2 ('ipso', OPh), 161.8 (C-2), 173.6 (COOBn).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-fluorophenyl-(methoxy-L-alaninyl)]-phosphate (CPF 5).

[0102] $\text{C}_{21}\text{H}_{24}\text{BrFN}_3\text{O}_9\text{P}$, MW=592.31.



[0103] This was synthesised according to *Standard procedure 5*, using BVdU (200mg, 0.60 mmol), para-fluorophenyl-(methoxy-L-alaninyl)-phosphorochloridate (442 mg, 1.5 mmol), NMI (4.98 mmol, 332 μ L) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (177 mg, yield 50%),

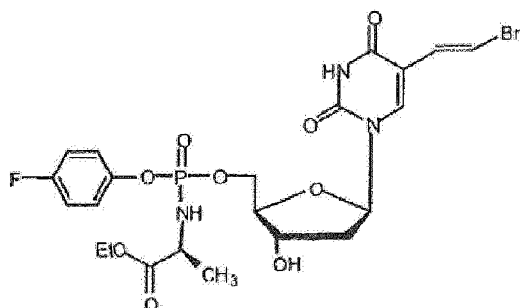
^{31}P -NMR (CDCl_3 , 121 MHz): δ 5.10, 4.81.

^1H -NMR (CDCl_3 ; 300 MHz): δ 10.1 (1H, bs, H-3), 7.60 (1H, 2xs, H-6), 7.39-7.32 (1H, 2d, $^3J=14$ Hz, H-5b), 7.20-6.95 (4H, m, *OPh*), 6.70-6.60 (1H, 2d, $^3J=14$ Hz, H-5a), 6.30-6.15 (1H, 2t, $^3J=6$ Hz, H1'), 4.55-4.29 (3H, m, H-5'+H-3'), 4.15 (1H, NH), 4.05-3.85 (2H, H-4', CHala), 3.72 (3H, 2s, CH_3O), 2.49-2.32 (1H, m, one of H-2'), 2.15-2.05 (1H, m, one of H-2'), 1.35 (3H, 2d, $^3J=6$ Hz, CH_3_{ala}).

^{13}C -NMR (DMSO; 75 MHz): δ 21.2 (CH_3_{ala}), 40.8 (C-2'), 50.8, 50.6 ($\text{CH}[\text{CH}_3]$), 53.2 (CH_3O), 66.7, 66.3 (C-5'), 71.9, 71.8 (C-3'), 86.1, 85.7, 85.8 (C-1', C-4'), 110.3 (C-5b), 111.9 (C-5), 117.0, 116.7 ('o', *OPh*), 122.0 ('m', *OPh*), 128.2 (C-5a), 138.2 (C-6), 149.0 ('ipso', *OPh*) 149.9 (C-4), 158.5 ('p', *OPh*), 163.2 (C-2), 175.1 (COOCH_3).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-fluorophenyl-(ethoxy-L-alaninyl)]-phosphate (CPF 6).

[0104] $\text{C}_{22}\text{H}_{26}\text{BrFN}_3\text{O}_9\text{P}$, MW-606.33.



[0105] This was synthesised according to *Standard procedure 5*, using BVdU (200 mg, 0.60 mmol), para-fluorophenyl-(ethoxy-L-alaninyl)-phosphorochloridate (464 mg, 1.5 mmol), NMI (4.98 mmol, 332 μ L) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (240 mg, yield 66%).

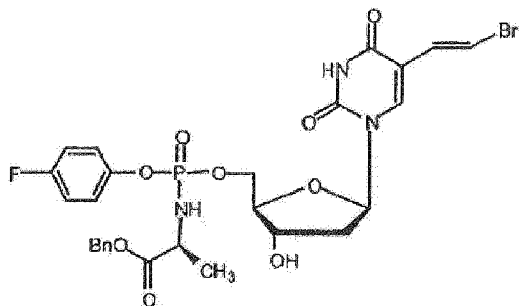
^{31}P -NMR (CDCl_3 121 MHz): δ 5.14, 4.88,

$^1\text{H-NMR}$ (CDCl_3 , 300 MHz): δ 10.25 (1H, bs, H-3), 7.85 (1H, 2xs, H-6), 7.44-7.39 (1H, 2d, $^3J=14$ Hz, H-5b), 7.3-7.0 (4H, m, *OPh*), 6.8-6.65 (1H, 2d, $^3J=14$ Hz, H-5a), 6.35-6.25 (1H, 2t, $^3J=6$ Hz, H1'), 4.6-4.1 (6H, m, H-5', H-3', CHala, NH, $\text{CH}_3\text{CH}_2\text{O}$), 4.02 (1H, m, H-4'), 2.55-2.45 (1H, m, one of H-2'), 2.20-2.10 (1H, m, one of H-2'), 1.40 (3H, d, $^3J=8$ Hz, CH_3ala), 1.25 (3H, 2t, $^3J=7$ Hz, $\text{CH}_3\text{CH}_2\text{O}$).

$^{13}\text{C-NMR}$ (CDCl_3 , 75 MHz): δ 14.5 ($\text{CH}_2\text{CH}_2\text{O}$) 21.3 (CH_3ala), 40.8, 40.7 (C-2'), 50.8, 50.7 (CHala), 62.3 ($\text{CH}_3\text{CH}_2\text{O}$), 66.7, 66.3 (C-5'), 71.1, 70.7 (C-3'), 86.1, 85.8, 85.6, 85.4 (C-1', C-4'), 110.4 (C-5b), 111.9 (C-5), 117.0 ('o', *OPh*), 122.2 'm', *OPh*, 128.9 (C-5a), 138.2 (C-6), 146.4 ('ipso', *OPh*), 149.9 (C-4), 158.5 ('p', *OPh*), 162.2, 161.8 (C-2), 174.2 ($\text{COOCH}_2\text{CH}_3$).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-fluorophenyl-(benzyloxy-L-alaninyl)]-phosphate (CPF 7).

[0106] $\text{C}_{27}\text{H}_{28}\text{BrFN}_3\text{O}_9\text{P}$, MW=668.40.



[0107] This was synthesised according to *Standard procedure 5*, using BVdU (200 mg, 0.60 mmol), para-fluorophenyl-(benzyloxy-L-alaninyl)-phosphorochloridate (556 mg, 1.5 mmol), NMI (4.98 mmol, 332 μL) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (256 mg, yield 64%).

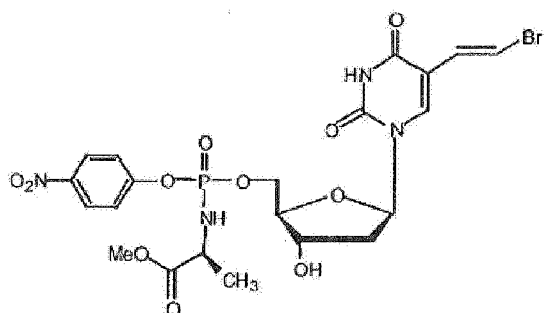
$^{31}\text{P-NMR}$ (CDCl_3 , 121 MHz): δ 4.74, 4.44.

$^1\text{H-NMR}$ (CDCl_3 , 300 MHz): δ 7.69 (1H, 2xs, H-6), 7.45-7.39 (1H, 2d, $^3J=14$ Hz, H-5b), 7.37-7.00 (9H, m, *OPh*+ CH_2Ph), 6.75-6.65 (1H, 2d, $^3J=14$ Hz, H-5a), 6.30-6.2 (1H, 2t, $^3J=6$ Hz, H-1'), 5.2 (1H, 2s, CH_2Ph), 4.85-4.00 (6H, m, H-3', H-5', H-4', NH, CHala), 2.47 (1H, m, one of H-2'), 2.0-2.15 (1H, m, one of H-2'), 1.38 (3H, d, $^3J=7$ Hz, CH_3ala).

^{13}C -NMR (CDCl_3 , 75 MHz): δ 21.2, 21.1 (CH_3ala), 40.7 (C-2'), 50.4 (CHala), 66.7, 66.4 (C-5'), 67.8 (CH_2Ph), 71.1, 70.7 (C-3'), 86.0, 85.7, 85.4, 85.3 (C-1', C-4'), 110.4 (C-5b), 111.9 (C-5), 117.0 ('o', OPh), 122.0 ('m', O-Ph), 128.1, 128.6 (Bn, C-5a), 135.4 ('ipso', CH_2Ph) 138.2 (C-6), 146.5 ('ipso', OPh), 149.9 (C-4), 158.5 'p' OPh), 162.2 (C-2), 173.9 (COOBn).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-nitrophenyl-(methoxy-L-alaninyl)]-phosphate (CPF 10).

[0108] $\text{C}_{21}\text{H}_{24}\text{BrN}_4\text{O}_{11}\text{P}$, MW=619.31.



[0109] This was synthesised according to **Standard procedure 5**, using BVdU (200 mg, 0.60 mmol), para-nitrophenyl-(methoxy-L-alaninyl)-phosphorochloridate (483 mg, 1.5 mmol), NMI (4.98 mmol, 332 μL) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (211 mg, yield 57%).

^{31}P -NMR (CDCl_3 , 121 MHz): δ 4.95.

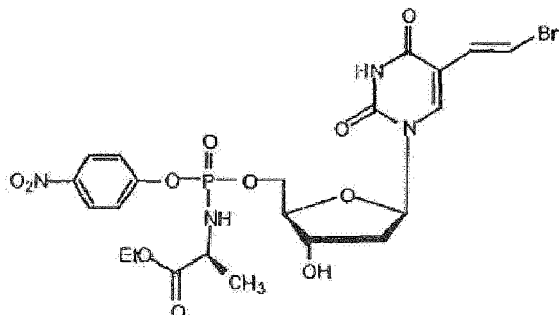
^1H -NMR (MeOD; 300 MHz): δ 8.3-8.2 (2H, m, OPh) 7.8-7.75 (1H, 2xs, H-6), 7.35-7.30, 7.55-7.4 (2H, m, OPh), 7.35-7.30 (1H, 2d, $^3J=14$ Hz, H-5b), 6.80-6.70 (1H, 2d, $^3J=14$ Hz, H-5a), 6.30-6.2 (1H, 2t, $^3J=6$ Hz, H1'), 4.5-4.3 (3H, m, H-5',H-3'), 4.2-4.0 (2H, m, H-4', CHala), 3.72 (3H, 2s, CH_3O), 2.35-2.15 (2H, m, 2 H-2'), 1.35 (3H, 2d, $^3J=7\text{Hz}$, CH_3ala).

^{13}C -NMR (DMSO; 75 MHz): δ 20.9 (CH_3ala), 41.6, 41.5 (C-2'), 52.0, 51.9 ($\text{CH}[\text{CH}_3]$), 53.4 (CH_3O), 68.5 (C-5'), 72.4, 72.3 (C-3'), 87.7, 87.4, 87.0, 86.9 (C-1', C-4'), 109.8 (C-5b), 112.8 (C-5), 122.6 ('o', OPh), 127.1 ('m', OPh), 130.8 (C-5a), 140.3 (C-6), 146.5 ('ipso', OPh), 151.4 (C-4), 157.2 ('p', OPh), 163.9 (C-2), 175.8, 175.5 (COOCH_3).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-

nitrophenyl-(ethoxy-L-alaninyl)]-phosphate (CPF 9).

[0110] $C_{22}H_{26}BrN_4O_{11}P$, MW=633.34.



[0111] This was synthesised according to. **Standard procedure 5**, using BVdU (200 mg, 0.60 mmol), para-nitrophenyl-(ethoxy-L-alaninyl)-phosphorochloridate (504 mg, 1.5 mmol), NMI (4.98 mmol, 332 μ L) in THF (5 mL) for 1 hr. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (232 mg, yield: 61%).

^{31}P -NMR ($CDCl_3$, 121 MHz): δ 4.28.

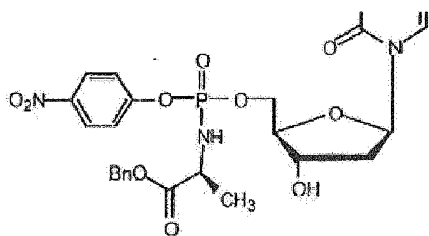
1H -NMR ($CDCl_3$, 300 MHz): δ 10.25 (1H, bs, H-3), 8.25-8.2 (2H, 2d, $^3J=9$ Hz, *OPh*), 7.7 (1H, 2xs, H-6), 7.5-7.45 (2H, 2d, $^3J=9$ Hz, *OPh*), 7.4-7.35 (1H, 2d, $^3J=14$ Hz, H-5b), 6.7-6.65 (1H, 2d, $^3J=14$ Hz, H-5a), 6.3-6.2 (1H, 2t, $^3J=6$ Hz, H1'), 4.8-4.1 (7H, m, H-5', H-4' H-3', CHala, NH, CH_3CH_2O), 2.45-2.4 (1H, m, one of H-2'), 2.20-2.10 (1H, m, one of H-2'), 1.40 (3H, d, $^3J=8$ Hz, CH_{3ala}), 1.3 (3H, 2t, $^3J=7$ Hz, CH_3CH_2O).

^{13}C -NMR ($CDCl_3$, 75 MHz): δ 14.5 (CH_3CH_2O) 21.1 (CH_{3ala}), 40.6 (C-2'), 50.8, 50.7 (CHala), 62.5 (CH_3CH_2O), 66.9, 66.8 (C-5'), 71.2, 70.9 (C-3'), 86.3, 85.9, 85.4, 85.3 (C-1', C-4'), 110.3 (C-5b), 111.8 (C-5), 121.3 ('o', *OPh*), 126.1 ('m', *OPh*), 128.8 (C-5a), 138.4 (C-6), 145.1 ('ipso', *OPh*), 149.9 (C-4), 155.5 ('p', *OPh*), 162.3 (C-2), 174.0, 173.9 ($COOCH_2CH_3$).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-nitrophenyl-(benzoxy-L-alaninyl)]-phosphate (CPF 8).

[0112] $C_{27}H_{28}BrN_4O_{11}P$, MW=695.41.





[0113] This was synthesized according to **Standard procedure 5**, using BVdU (200 mg, 0.60 mmol), para-nitrophenyl-(benzyloxy-L-alaninyl)-phosphorochloridate (597 mg, 1.5 mmol), NMI (4.98 mmol, 332 μ L) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (228 mg, yield 55%).

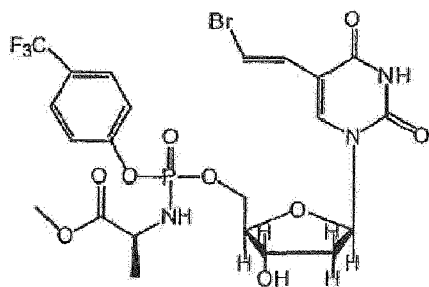
^{31}P -NMR (CDCl_3 , 121 MHz): δ 4,74, 4.44.

^1H -NMR (CDCl_3 , 300 MHz): δ 10.4-10.3 (1H, bs, H-3), 8.2-8.1 (2H, m, OPh), 7.69 (1H, 2xs, H-6), 7.4-7.2 (1H, 2d, $^3J=14$ Hz, H-5b), 7.37-7.00 (7H, m, $\text{OPh}+\text{CH}_2\text{Ph}$), 6.75-6.65 (1H, 2d, $^3J=14$ Hz, H-5a), 6.25-6.15 (1H, 2t, $^3J=6$ Hz, H-1'), 5.2 (1H, d, CH_2Ph), 4.87 (1H, m, H-3'), 4.6-4.2 (3H, m, H-5', CHala), 4.2-4.00 (2H, m, H-4', NH), 2.55-2.45 (1H, m, one of H-2'), 2.2-2.05 (1H, m, one of H-2'), 1.38 (3H, d, $^3J=7$ Hz, CH_3ala).

^{13}C -NMR (CDCl_3 , 75 MHz): δ 21.2, 21.1 (CH_3ala), 40.6 (C-2'), 50.9 (CHala), 67.1, 67.0 (C-5'), 68.0 (CH_2Ph), 71.3, 70.9 (C-3'), 86.3, 86.0, 85.3, 85.2 (C-1', C-4'), 110.4 (C-5b), 111.9, 111.8 (C-5), 121.3 ('o', OPh), 126.2-126.1 ('m', OPh), 129.1, 128.7, 128.6 (Bn, C-5a), 135.4 ('ipso', CH_2Ph), 138.3 (C-6), 145.1 ('ipso', OPh), 149.9 (C-4), 155.6 ('p' OPh), 162.2 (C-2), 173.8, 173.7 (COOBn).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine-5'-[para-(trifluoromethyl)-phenyl-(methoxy-L-alaninyl)]-phosphate (CPF 15).

[0114] $\text{C}_{22}\text{H}_{24}\text{BrF}_3\text{N}_3\text{O}_9$, MW=642.31.



[0115] This was synthesized according to **Standard procedure 5**, using BVdU (200 mg, 0.60 mmol), phenyl-(methoxy-L-alaninyl)-phosphorochloridate (518.8 mg, 1.5 mmol), NMI (246.3 mg, 3.0 mmol, 239 μ L) in THF (5 mL) for 4 hrs. The crude product was purified by column chromatography, eluting with chloroform/methanol 97:3 to give the pure product as a white foamy solid (211.1 mg, yield 34,7%).

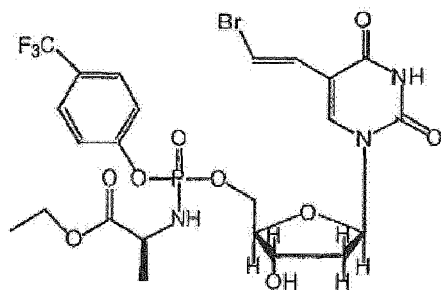
^{31}P -NMR (MeOD, 121 MHz): δ 5.23, 5.07.

^1H -NMR (MeOD, 300 MHz): δ 7.80 (1H, s, H-6), 7.70 (2H, d, $^3J=8.7$ Hz, *O*Ph), 7.47-7.42 (2H, m, *O*Ph), 7.37 (1H, d, $^3J=13.6$ Hz, H-5b), 6.82-6.78 (1H, d, $^3J=13.6$ Hz, H-5a), 6.30-6.23 (1H, m, H-1'), 4.52-4.29 (3H, m, H-3'+H-5'), 4.17-4.13 (1H, m, H-4'), 4.05-3.91 (1H, m, CHCH₃), 3.67 (3H, s, OCH₃), 2.35-2.32 (1H, m, one of H-2'), 2.23-2.16 (1H, m, one of H-2'), 1.37-1.34 (3H, d, $^3J=7.1$ Hz, CHCH₃).

^{13}C -NMR (MeOD, 75 MHz): δ 20.6, 20.7, 20.8, 20.9 (CHCH₃), 41.5, 41.7 (C-2'), 51.9, 52.0 (CHCH₃), 68.2, 68.3 (C-5'), 72.4, 72.5 (C-3'), 87.1, 87.2, 87.4, 87.6 (C-1', C-4'), 109.7 (C-5b), 112.6 (C-5), 122.5, 122.7 ('o', *O*Ph), 125.8 (CF₃, $J=269$ Hz), 128.7 ('m', *O*Ph), 128.8 ('p', $J=33$ Hz, *O*Ph), 130.9 (C-5a), 140.3 (C-6), 151.4, 151.5 ('ipso', *O*Ph), 155.1, 155.2 (C-4), 164.0 (C-2), 175.6, 175.9, (COOCH₃).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine-5'-[para-(trifluoromethyl)-phenyl-(ethoxy-L-alaninyl)]-phosphate (CPF 25).

[0116] $\text{C}_{23}\text{H}_{26}\text{BrF}_3\text{N}_3\text{O}_9\text{P}$, MW=656.34



[0117] This was synthesised according to **Standard procedure 5**, using BVdU (200 mg, 0.60 mmol), phenyl-(ethoxy-L-alaninyl)-phosphorochloridate (539.5 mg, 1.5 mmol), NMI (246.3 mg, 3.0 mmol, 239 μ L) in THF (5 mL) for 20 hrs. The crude product was purified by column chromatography, eluting with dichloromethane/methanol 95:5 to give the pure product as a white foamy solid (172.6 mg, yield 43.8%).

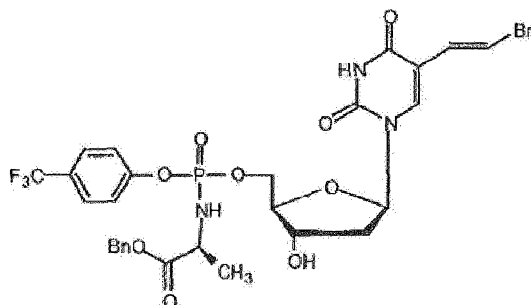
^{31}P -NMR (CDCl_3 , 121 MHz): δ 4.65, 4.35.

^1H -NMR (CDCl_3 , 300 MHz): δ 10.05 (1H, s, H-3), 7.69-7.64 (3H, m, H-6+O*Ph*), 7.46-7.39 (3H, m, O*Ph*+ H-5b), 6.70-6.68 (1H, 2d, $^3J=13.6$ Hz, H-5a), 6.34-6.25 (1H, m, H-1'), 4.57-4.35 (4H, m, H-3'+H-5'+NH), 4.27-4.13 (4H, m, H-4'+OCH₂CH₃+OH-3'), 4.12-3.98 (1H, m, CHCH₃), 2.53-2.47 (1H, m, one of H-2'), 2.21-2.12 (1H, m, one of H-2'), 1.43-1.40 (3H, d, $^3J=7.0$ Hz, CHCH₃), 1.28, 1.27 (3H, 2t, $^3J=7.0$ Hz, OCH₂CH₃)

^{13}C -NMR (CDCl_3 , 75 MHz): δ 14.5 (CH₃CH₂O), 21.2, 21.3 (CHCH₃), 40.7 (C-2'), 50.8, 50.9 (CHCH₃), 62.4 (CH₃CH₂O), 66.3, 66.7 (C-5'), 70.7, 71.1 (C-3'), 85.3, 85.4, 85.8, 86.1 (C-1', C-4'), 110.5 (C-5b), 112.0 (C-5), 122.0 ('o', O*Ph*), 124.2 (CF₃, $J=271$ Hz), 127.7, 127.8, 128.7 ('m', 'p', O*Ph*), 128.8 (C-5a), 138.0 (C6), 149.7 ('ipso', O*Ph*), 153.2 (C-4), 161.9 (C-2), 174.0, 174.1 (COOCH₂CH₃).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-trifluoromethyl-(benzyloxy-L-alaninyl)]-phosphate (CPF 4).

[0118] C₂₈H₂₈BrF₃N₃O₉P, MW=718.41.



[0119] This was synthesised according to **Standard procedure 5**, using BVdU (200 mg, 0.60 mmol), para-trifluorophenyl-(benzyloxy-L-alaninyl)-phosphorochloridate (632 mg, 1.5 mmol), NMI (4.98 mmol, 332 μL) in THF (6 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH₂Cl₂/Methanol 97:3 to give the pure product as a white foamy solid (308 mg, yield 71%).

^{31}P -NMR (CDCl_3 , 121 MHz): δ 5.31, 4.87.

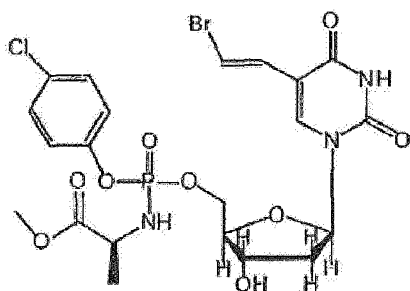
^1H -NMR (CDCl_3 , 300 MHz): δ 10.05 (1H, bs, H-3), 7.7, 7.25 (11H, m, H-5b, H-6 O*Ph*+CH₂*Ph*), 6.75-6.65 (1H, 2d, $^3J=14$ Hz, H-5a), 6.35-6.2 (1H, 2t, $^3J=6\text{Hz}$, H-1'), 5.15 (1H, 2s, CH₂Ph), 4.6-4.25 (4H, m, H-5', H-3', CHala) 4.2-4.00 (2H, m, H-4', NH), 2.55-2.4 (1H, m, one of H-2'), 2.2-

2.05 (1H, m, one of H-2'), 1.38 (3H, d, $^3J=7$ Hz, CH₃ala).

¹³C-NMR (CDCl₃, 75 MHz): δ 21.2, 21.1 (CH₃ala), 40.7 (C-2'), 50.9, 50.8 (CHala), 67.1, 67.0 (C-5'), 68.0 (CH₂Ph), 71.2, 10.9 (C-3'), 86.1, 85.8, 85.5, 85.4 (C-1', C-4'), 110.2 (C-5b), 111.9, 111.8 (C-5), 121.1 ('o', OPh), 125.1 (d, J=270Hz, CF₃), 127.6 ('m', OPh), 129.1, 128.7, 128.6 (Bn, C-5a), 130.1 ('p', q, J=32Hz, OPh) 135.4 ('ipso', CH₂Ph) 138.2 (C-6), 150.2, 150.1 (C-4), 153.6 ('ipso' OPh), 162.7 (C-2), 173.9, 173.6 (COOBn).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine-5'-[4-chlorophenyl-(methoxy-L-alaninyl)]-phosphate (CPF 13).

[0120] C₂₁H₂₄BrClN₃O₉P, MW=608.76.



[0121] This was synthesised according to **Standard procedure 5**, using BVdU (200 mg, 0.60 mmol), 4-chlorophenyl-(methoxy-L-alaninyl)-phosphorochloridate (374.5 mg, 1.2 mmol), NMI (246.3 mg, 3.0 mmol, 239 μ L) in THF (8 mL) for 5 hrs. The crude product was purified by column chromatography, eluting with Chloroform/Methanol 97:3 to give the pure product as a white foamy solid (139.0 mg, yield 38.0%).

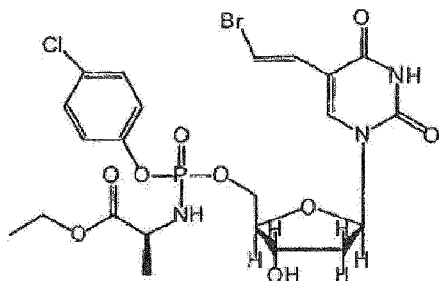
³¹P-NMR (CDCl₃, 121 MHz): δ 4.81, 4.54.

¹H-NMR (CDCl₃, 300 MHz): δ 10.11 (1H, bs, H-3), 7.68 (1H, s, H-6), 7.46-7.40 (1H, d, $^3J=13.6$ Hz, H-5b), 7.35-7.20 (4H, m, OPh), 6.76-6.67 (1H, 2d, $^3J=13.6$ Hz, H-5a), 6.34-6.24 (1H, m, H-1'), 4.58-4.40 (5H, m, H-3'+H-5'+NH), 4.36-4.19 (1H, m, H-4'), 4.07-3.99 (1H, m, CHCH₃), 3.75 (3H, s, OCH₃), 2.49-2.48 (1H, m, one of H-2'), 2.17-2.15 (1H, m, one of H-2'), 1.42-1.39 (3H, d, $^3J=7.0$ Hz, CHCH₃).

¹³C-NMR (CDCl₃, 75 MHz): δ 21.2 (CHCH₃), 40.7, 40.8 (C-2'), 50.6, 50.8 (CHCH₃), 53.2, 53.3 (OCH₃), 66.4, 66.7 (C-5'), 70.8, 71.2 (C-3'), 85.4, 85.5, 85.8, 86.2 (C-1', C-4'), 110.5 (C-5b), 111.9, 112.0 (C-5), 122.0 ('o', OPh), 128.9 (C-5a), 130.3 ('m', OPh), 131.1 ('p', OPh), 138.2 (C-6), 149.1, 149.2 ('ipso', OPh), 149.8 (C-4), 162.1, 162.2 (C-2), 174.5, 174.6 (COOCH₃).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine-5'-[4-chlorophenyl-(ethoxy-L-alaninyl)]-phosphate (CPF 11).

[0122] $C_{22}H_{26}BrN_3O_9P$, MW=622.79.



[0123] This was synthesised according to **Standard procedure 5**, using BVdU (300 mg, 0.90 mmol), 4-chlorophenyl-(ethoxy-L-alaninyl)-phosphorochloridate (557.7 mg, 1.71 mmol), NMI (221.7 mg, 2.7 mmol, 215 μ L) in THF (10 mL) for 16 hrs. The crude product was purified by column chromatography, eluting with dichloromethane/methanol 97:3 to give the pure product as a white foamy solid (168.4 mg, yield 30.0%).

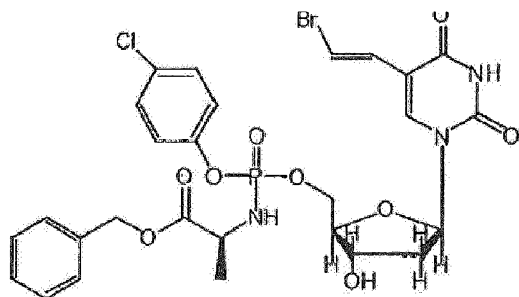
^{31}P -NMR ($CDCl_3$, 121 MHz): δ 4.88, 4.65.

1H -NMR ($CDCl_3$, 300 MHz): δ 9.51 (1H, bs, H-3), 7.69-7.68 (1H, 2s, H-6), 7.49-7.43 (1H, 2d, $^3J=13.6$ Hz, H-5h), 7.37-7.22 (4H, m, *OPh*), 6.79-6.71 (1H, 2d, $^3J=13.6$ Hz, H-5a), 6.33-6.24 (1H, m, H-1'), 4.62-4.34 (3H, m, H-3'+H-5'), 4.28-3.89 (5H, m, H-4'+ OCH_2CH_3 + $CHCH_3$ +NH), 2.59-2.45 (1H, m, one of H-2'), 2.22-2.14 (1H, m, one of H-2'), 1.43-1.41 (3H, d, $^3J=7.0$ Hz, $CHCH_3$), 1.33-1.28 (3H, 2t, $^3J=7.2$ Hz, OCH_2CH_3)

^{13}C -NMR ($CDCl_3$, 75 MHz): δ 14.5 (CH_3CH_2O), 21.2, 21.3 ($CHCH_3$), 40.7 (C-2'), 50.7, 50.8 ($CHCH_3$), 62.4(CH_3CH_2O), 66.7 (C-5'), 70.8, 71.2 (C-3'), 85.4, 85.8, 86.1 (C-1', C-4'), 110.4 (C-5b), 112.0 (C-5), 122.0, 122.1 ('o', *OPh*), 128.9 (C-5a), 130.3 ('m', *OPh*), 131.1 ('p', *OPh*), 138.2 (C-6), 149.2 ('ipso', *OPh*), 150.0 (C-4), 162.2 (C-2), 174.1, 174.2 ($COOCH_2CH_3$).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine-5'-[4-chlorophenyl-(benzoxy-L-alaninyl)] -phosphate (CPF 12).

[0124] $C_{22}H_{26}BrN_3O_9P$, MW=622.79.



[0125] This was synthesised according to **Standard procedure 5**, using BVdU (300 mg, 0.90 mmol), 4-chlorophenyl-(berizoxo-L-alaninyl)-phosphorochloridate (698.7 mg, 1.80 mmol), NMI (369,5 mg, 4.5 mmol, 358.7 μ L) in THF (10 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with dichloromethane/methanol 95:5 to give the pure product as a white foamy solid (310.0 mg, yield 50.3%).

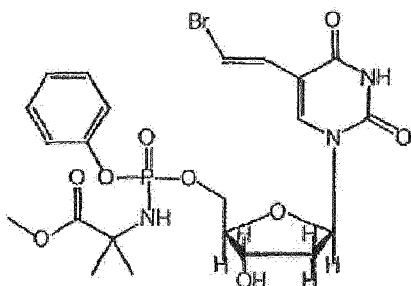
^{31}P -NMR (CDCl_3 , 121 MHz): δ 4.81, 4.53.

^1H -NMR (CDCl_3 , 300 MHz): δ 10.10 (1H, bs, H-3), 7.65-7.63 (1H, 2s, H-6), 7.69-7.68 (1H, 2s, H-6), 7.46, 7.41 (1H, 2d, $^3J=13.6$ Hz, H-5b), 7.40-7.17 (9H, m, *OPh*), 6.75-6.66 (1H, 2d, $^3J=13.6$ Hz, H-5a), 6.33-6.23 (1H, 2t, $^3J=6.0$ Hz, H-1'), 5.17 (2H, s, CH_2Ph), 4.60-4.23 (4H, m, H-3'+H-5'+NH), 4.20-3.97 (2H, m, H-4'+ CHCH_3), 2.48-2.44 (1H, m, one of H-2'), 2.15-2.05 (1H, m, one of H-2'), 1.43-1.40 (3H, d, $^3J=7.0$ Hz, CHCH_3)

^{13}C -NMR (CDCl_3 , 75 MHz): δ 21.2 (CHCH_3), 40.7 (C-2'), 50.8, 50.9 (CHCH_3), 66.6 (C-5'), 67.9 (CH_2Ph), 70.7, 71.1 (C-3'), 85.4, 85.5, 85.8, 86.1 (C-1', C-4'), 110.5 (C-5b), 111.9, 112.0 (C-5), 122.0, ('o', *OPh*), 128.7, 129.0, 129.1, 130.3 ('m', *OPh*+C-5a), 131.1 ('ipso', CH_2Ph), 135.4 ('p', *OPh*), 138.2 (C-6), 149.1 ('ipso', *OPh*), 150.0 (C-4), 162.1 (C-2), 173.9, 174.0 (COOCH_2Ph).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine-5'-[phenyl-(methoxy-a,a-dimethylglycyl)]-phosphate (CPF 26).

[0126] $\text{C}_{22}\text{H}_{27}\text{BrN}_3\text{O}_9\text{P}$, MW 588.34



[0127] This was synthesised according to **Standard procedure 5**, using BVdU (200 mg, 0.60 mmol), phenyl-(methyl-2-amino-2-methylpropanoate)-phosphorochloridate (437.5 mg, 1.5 mmol), NMI (246.3 mg, 3.0 mmol, 239.1 μ L) in THF (5 mL) for 4 hrs. The crude product was purified by column chromatography, eluting with chloroform/methanol 97:3 to give the pure product as a white foamy solid (117 mg, yield 33.1%).

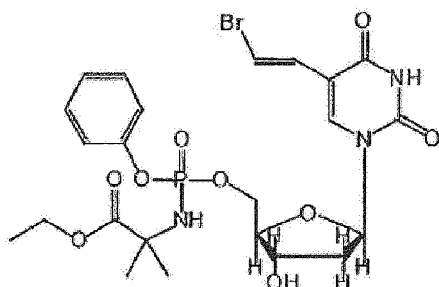
^{31}P -NMR (CDCl_3 , 121 MHz): δ 3.36, 3.14

^1H -NMR (CDCl_3 ; 300 MHz): δ 9.91 (1H, bs, H-3), 7.73, 7.65 (1H, 2s, H-6), 7.50-7.43 (1H, 2d, $^3J=13.6$ Hz, H-5b), 7.41-7.02 (5H, m, *OPh*), 6.81-6.71 (1H, 2d, $^3J=13.6$ Hz, H-5a), 6.34-6.28 (1H, m, H1'), 4.55-4.17 (6H, m, H-5'+H-4'+H-3', NH, OH-3'), 3.78 (3H, s, CH_3O), 2.53-2.39 (1H, m, one of H-2'), 2.25-1.99 (1H, m, one of H-2'), 1.60 (6H, s, $[\text{CH}_3]_2\text{C}$).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 27.5, 27.4, 27.2 ($[\text{CH}_3]_2\text{C}$), 40.7, 40.6 (C-2'), 53.5 (CH_3O), 57.6 ($[\text{CH}_3]_2\text{C}$), 66.5, 66.2 (C-5'), 70.7, 71.1 (C-3'), 85.4, 85.6, 85.5, 85.9 (C-1', C-4'), 110.4 (C-5b), 111.9 (C-5), 120.5, 120.6 ('o', *OPh*), 125.7 ('p', *OPh*), 128.9 (C-5a), 130.3 ('m', *OPh*), 138.0, 138.3 (C-6), 149.8 ('ipso', *OPh*) 150.9, 150.8 (C-4), 162.0, 162.1 (C-2), 176.4, 176.2 (COOCH_3).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine-5'-[phenyl-(ethoxy-a,a-dimethylglycyl)]-phosphate (CPF 27).

[0128] $\text{C}_{23}\text{H}_{29}\text{BrN}_3\text{O}_9\text{P}$, MW=602.37



[0129] This was synthesised according to **Standard procedure 5**, using BVdU (200 mg, 0.60 mmol), phenyl-(ethyl-2-amino-2-methylpropanoate)phosphorochloridate (458.0 mg, 1.5 mmol), NMI (246.3 mg, 3.0 mmol, 219.1 μ L) in THF (5 mL) for 5 hrs. The crude product was purified by column chromatography, eluting with chloroform/methanol 97:3 to give the pure product as a white foamy solid (106 mg, yield 29.3%).

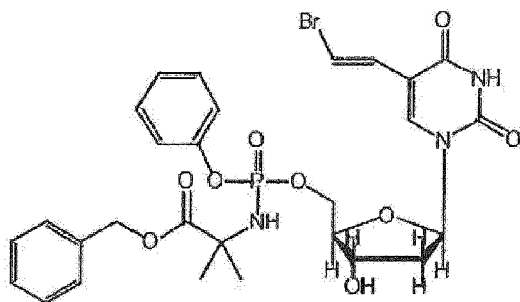
^{31}P -NMR (MeOD, 121 MHz): δ 3.91, 3.85

$^1\text{H-NMR}$ (MeOD, 300 MHz): δ 7.84, 7.81 (1H, 2s, H-6), 7.44-7.20 (6H, m, *OPh*+H-5b), 6.88-6.81 (1H, 2d, $^3J=13.6\text{Hz}$, H-5a), 6.34-6.28 (1H, m, H-1'), 4.50-4.34 (3H, m, H-5'+H-3'), 4.23-4.15 (3H, m, H-4'+ $\text{CH}_3\text{CH}_2\text{O}$), 2.38-2.28 (1H, m, one of H-2'), 2.22-2.09 (1H, m, one of H-2'), 1.51 (6H, s, $[\text{CH}_3]_2\text{C}$), 1.29 (3H, t, $^3J=7\text{ Hz}$, $\text{CH}_3\text{CH}_2\text{O}$)

$^{13}\text{C-NMR}$ (MeOD, 75 MHz): δ 14.9 ($\text{CH}_3\text{CH}_2\text{O}$) 27.9, 28.3 ($[\text{CH}_3]_2\text{C}$), 41.5 (C-2'), 58.51 ($\text{C}[\text{CH}_3]_2$), 63.1 ($\text{CH}_3\text{CH}_2\text{O}$), 68.2 (C-5'), 72.6 (C-3'), 87.1, 87.4 (C-1', C-4'), 109.6 (C-5b), 112.7 (C-5b), 122.0, 122.1, 122.2, ('o', *OPh*), 126.7 ('p', *OPh*), 131.0, 131.2 (C-5a, 'm' *OPh*), 140.4 (C-6), 151.4 ('ipso', *OPh*) 152.5 (C-4), 164.0 (C-2), 177.2 ($\text{COOCH}_2\text{CH}_3$).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine-5'-[phenyl-(benzoxo-a,a-dimethylglyciny)]-phosphate (CPF 14).

[0130] $\text{C}_{28}\text{H}_{31}\text{BrN}_3\text{O}_9\text{P}$, MW=664.44.



[0131] This was synthesized according to **Standard procedure 5**, using BVdU (242 mg, 0.73 mmol), phenyl-(benzyl-2-amino-2-methylpropanoate)-phosphorochloridate (533.0 mg, 2.0 mmol), NMI (298.0 mg, 3.63 mmol, 289 μL) in THF (5 mL) for 4 hrs. The crude product was purified by column chromatography, eluting with chloroform/methanol 97:3 to give the pure product as a white foamy solid (129.0 mg, yield 26.7%).

$^{31}\text{P-NMR}$ (CDCl_3 , 121 MHz): δ 3.30, 3.12.

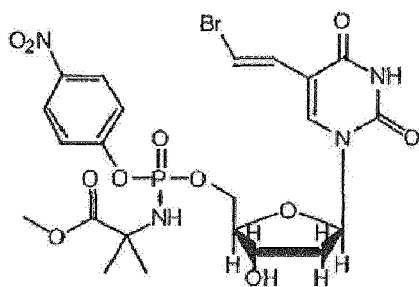
$^1\text{H-NMR}$ (CDCl_3 , 300 MHz): δ 9.92 (1H, bs, H-3), 7.67-7.60 (1H, 2s, H-6), 7.48-7.41 (1H, 2d, $^3J=13.6\text{ Hz}$, H-5b), 7.40-7.16 (10H, m, *OPh*+ CH_2Ph), 6.78-6.67 (1H, 2d, $^3J=13.6\text{ Hz}$, H-5a), 6.31-6.25 (1H, m, H-1'), 5.18 (1H, s, CH_2Ph), 4.50-4.09 (6H, m, H-3'+H-5'+H-4', *NH*, OH-3'), 2.48-2.25 (1H, m, one of H-2'), 2.16-1.82 (1H, m, one of H-2'), 1.60 (6H, s, $[\text{CH}_3]_2\text{C}$).

$^{13}\text{C-NMR}$ (CDCl_3 , 75 MHz): δ 27.3, 27.4, 28.5 ($[\text{CH}_3]_2\text{C}$), 40.6, 40.7 (C-2'), 57.6, 57.6 ($\text{C}[\text{CH}_3]_2$), 66.2, 66.5 (C-5'), 68.1 (CH_2Ph), 70.6, 71.1 (C-3'), 85.4, 85.5, 85.6, 85.8 (C-1', C-4'),

110.4 (C-5b), 112.0 (C-5), 120.4, 120.5, 120.6, 125.7, 128.4, 128.5, 128.8, 128.9, 130.3 (*O*Ph, C-5a), 135.7 ('*ipso*', CH₂Ph) 138.1, 138.3 (C-6), 149.8, 150.8, 150.9 ('*ipso*' *O*Ph, C-4), 162.1 (C-2), 177.5, 175.7 (COOCH₂Ph).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine-5'-[4-nitrophenyl-(methoxy- α,α -dimethylglycyl)]-phosphate (CPF 45).

[0132] C₂₂H₂₆BrN₄O₁₁P, MW=633.34.



[0133] This was synthesized according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), 4-nitrophenyl-(methyl-2-amino-2-methylpropanoate)-phosphorochloridate (378.8 mg, 1.13 mmol), NMI (184.7 mg, 2.25 mmol, 179.4 μ L) in THF (5 mL) for 3 hrs. The crude product was purified by column chromatography, eluting with dichloromethane/methanol 97:3 to give the pure product as a white foamy solid (145.7 mg, yield 50.9 %).

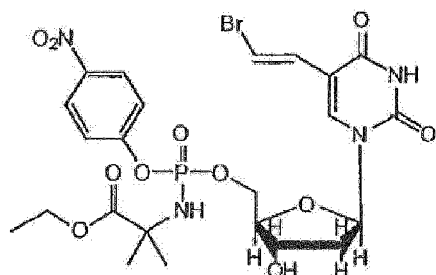
³¹P-NMR (MeOD, 121 MHz): δ 3.61, 3.56.

¹H-NMR (MeOD, 300 MHz): δ 8.30-8.25 (2H, d, ³J=9.0 Hz, *O*Ph), 7.79-7.78 (1H, 2s, H-6), 7.49-7.46 (2H, d, ³J=9.0 Hz, *O*Ph), 7.37-7.32 (1H, 2d, ³J=13.6 Hz, H-5b), 6.79-6.72 (1H, 2d, ³J=13.6 Hz, H-5a), 6.32-6.25 (1H, m, H-1'), 4.48-4.35 (3H, m, H.3'+H-5'), 4.15-4.14 (1H, m, H-4'), 3.71 (3H, s, CH₃O), 2.41-2.17 (2H, m, H-2'), 1.51 (6H, s, [CH₃]₂C).

³C-NMR (CDCl₃, 75 MHz): δ 28.0, 28.1, 28.2, 28.3 ([CH₃]₂C), 41.4, 41.5 (C-2'), 53.6 (CH₃O), 58.7 (C[CH₂]₂), 68.5 (C-5'), 72.3, 72.4 (C-3'), 86.9, 87.0, 87.4, 87.5 (C-1', C-4'), 109.7 (C-5b), 112.6 (C-5), 122.8, 122.9 ('*o*', *O*Ph), 127.0 ('*m*' *O*Ph), 130.9 (C-5a), 140.5 (C-6), 146.5 ('*p*', *O*Ph), 151.5 ('*ipso*', *O*Ph), 157.3 (C-4), 164.0 (C-2), 177.5 (COOCH₃).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine-5'-[4-nitrophenyl-(ethoxy- α,α -dimethylglycyl)]-phosphate (CPF 46).

[0134] $C_{23}H_{28}BrN_4O_{11}P$, MW=647.3.



[0135] This was synthesised according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), 4-nitrophenyl-(ethy)-2-amino-2-methylpropanoate)-phosphorochloridate (442.1 mg, 1.26 mmol), NMI (184.7 mg, 2.25 mmol, 179.4 μ L) in THF (5 mL) for 4 hrs. The crude product was purified by column chromatography, eluting with dichloromethane/methanol 97:3 to give the pure product as a white foamy solid (152.9 mg, yield 52,5 %).

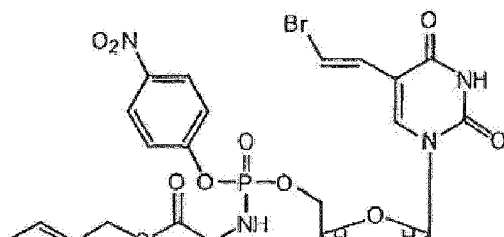
^{31}P -NMR ($CDCl_3$, 121 MHz): δ 3.00, 2.96.

1H -NMR ($CDCl_3$, 300 MHz): δ 10.28 (1H, bs, H-3), 8.25-8.12 (2H, d, $^3J=9.0$ Hz, *O*Ph), 7.68-7.67 (1H, 2s, H-6), 7.46-7.32 (3H, m, *O*Ph+H-5b), 6.69-6.67 (1H, 2d, $^3J=13.5$ Hz, H-5a), 6.32-6.26 (1H, m, H-1'), 4.75-4.36 (5H, m, H-3'+H-5'+OH-3'+NH), 4.25-4.17 (3H, m, *OCH*2CH₃, H-4'), 2.60-2.98 (1H, m, one of H-2'), 2.31-2.10 (1H, m, one of H-2'), 1.58 (6H, s, [*CH*3]₂C), 1.30-1.28 (3H, 2t, $^3J=7.1$ Hz, *OCH*2CH₃).

^{13}C -NMR ($CDCl_3$, 75 MHz): δ 14.5 (*CH*3CH₂O), 27.1, 27.2, 27.3, 27.4 ([*CH*3]₂C), 40.6 (C-2'), 57.7 (*CH*3CH₂O), 62.7 (CH₃*CH*2O), 67.0 (C-5'), 71.0, 71.2 (C-3'), 85.4, 85.9, 86.1 (C-1', C-4'), 110.3 (C-5b), 111.9 (C-5), 121.2, 121.3 ('o', *O*Ph), 126.2 ('m' *O*Ph), 128.8 (C-5a), 138.4 (C-6), 145.0 ('p', *O*Ph), 150.0 (C-4), 155.7-155.9 ('ipso', *O*Ph), 162.2 (C-2), 175.0-175.1 (*COOCH*2CH₃).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine-5'-[4-nitrophenyl-(benzoxy- α,α -dimethylglycyl)]-phosphate (CPF 47).

[0136] $C_{28}H_{30}BrN_4O_{11}P$, MW=709.44





[0137] This was synthesised according to **Standard procedure 5**, using BVdU (100 mg, 0.30 mmol), 4-nitrophenyl-(benzyl-2-amino-2-methylpropanoate)-phosphorochloridate (309.6 mg, 1.07 mmol), NMI (123.7 mg, 1.5 mmol, 120.1 μ L) in THF (5 mL) for 5 hrs. The crude product was purified by column chromatography, eluting with dichloromethane/methanol 97:3 to give the pure product as a white foamy solid (160.2 mg, yield 50.2 %).

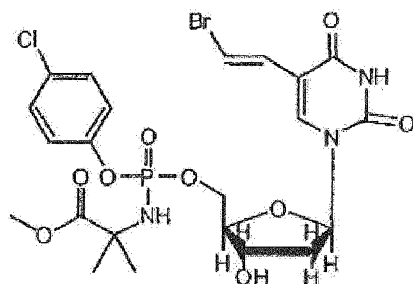
^{31}P -NMR (CDCl_3 , 121 MHz): δ 2.95, 2.89.

^1H -NMR (CDCl_3 300 MHz): δ 10.16 (1H, bs, H-3), 8.26-8.24 (2H, 2d, $^3J=9.1$ Hz, OPh), 7.71-7.69 (1H, 2s, H-6), 7.48-7.37 (8H, m, $\text{OPh}+\text{CH}_2\text{Ph}$, H-5b), 6.75-6.12 (1H, 2d, $^3J=13.5$ Hz, H-5a), 6.36-6.29 (1H, m, H-1'), 5.24 (2H, s, CH_2Ph), 4.81-4.40 (5H, m, H-3'+H-5'+OH-3', NH), 4.22-4.21 (1H, m, H-4'), 2.57-2.36 (1H, m, one of H-2'), 2.27-2.22 (1H, m, one of H-2'), 1.64 (6H, s, $[\text{CH}_3]_2\text{C}$).

^{13}C -NMR (CDCl_3 , 75 MHz): δ 27.4 ($[\text{CH}_3]_2\text{C}$), 40.6 (C-2'), 57.8 ($\text{C}[\text{CH}_3]_2$), 67.0 (C-5'), 68.2 (CH_2Ph), 71.1, 71.2 (C-3'), 85.3, 86.2 (C-1', C-4'), 110.5 (C-5b), 111.9 (C-5), 121.2, 126.2, 128.5, 128.8, 129.0, 129.1 ('o', 'm', 'p', $\text{CH}_2\text{Ph}+\text{OPh}+\text{C-5a}$), 135.5 ('ipso', CH_2Ph), (C-5a), 138.4 (C-6), 145.0 ('p', OPh), 150.0 (C-4), 155.7 ('ipso', OPh), 162.2 (C-2), 175.4-175.5 (COOCH_2Ph).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine-5'-[4-nitrophenyl-(methoxy- α,α -dimethylglycyl)]-phosphate (CPF 42).

[0138] $\text{C}_{22}\text{H}_{26}\text{BrClN}_3\text{O}_9\text{P}$, MW=622.79.



[0139] This was synthesised according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), 4-chlorophenyl-(methyl-2-amino-2-methylpropanoate)-phosphorochloridate (440.2 mg, 1.35 mmol), NMI (184.7 mg, 2.25 mmol, 179.4 μ L) in THF (5 mL) for 6 hrs. The crude product

was purified by column chromatography, eluting with dichloromethane/methanol 97:3 to give the pure product as a white foamy solid (146.7 mg, yield 56.5 %).

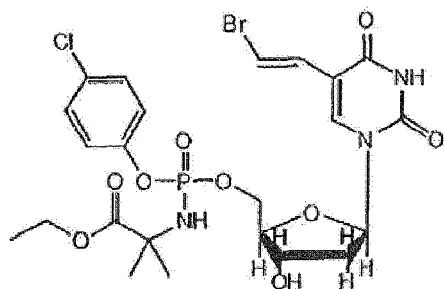
^3P -NMR (MeOD, 121 MHz): δ 3.98 (s).

^1H -NMR (MeOD, 300 MHz): δ , 7.71-7.69 (1H, 2s, H-6), 7.31-7.13 (5H, m, *O**Ph*+H-5b), 6.73-6.66 (1H, 2d, $^3J=13.6$ Hz, H-5a), 6.23-6.16 (1H, m, H-1'), 4.39-4.22 (3H, m, H-3'+H-5'), 4.05-4.03 (1H, m, H-4'), 3.61 (3H, s, *C**H**3*O), 2.29-2.19 (1H, m, one of H-2'), 2.15-2.05 (1H, m, one of H-2'), 1.38 (6H, s, [*C**H**3**2*]*C*).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 28.0, 28.2, 28.3, 28.4 ([*C**H**3**2*]*C*), 41.5, 41.6 (C-2'), 53.5, 53.6 (*C**H**3*O), 58.6 (*C**H**3*), 68.2 (C-5'), 72.4, 72.5 (C-3'), 87.1, 87.2, 87.3, 87.4 (C-1', C-4'), 109.7 (C-5b), 112.7 (C-5), 123.7, 123.8 ('o', *O**Ph*), 130.9, 131.1 ('m', *O**Ph*+C-5a), 131.9 ('p', *O**Ph*), 140.4 (C-6), 151.1, 151.2, 151.4 ('ipso', *O**Ph*+C-4), 164.0 (C-2), 177.6, 177.7 (*C**O**O**C**H**3*).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine-5'-[4-chlorophenyl-(ethoxy- α,α -dimethylglycyl)]-phosphate (CPF 43).

[0140] $\text{C}_{23}\text{H}_{28}\text{BrClN}_3\text{O}_9\text{P}$, MW=636.91.



[0141] This was synthesised according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), 4-chlorophenyl-(ethyl-2-amino-2-methylpropanoate)-phosphorochloridate (413.3 mg, 1.22 mmol), NMI (184.7 mg, 2.25 mmol, 179.3 μL) in THF (5 mL) for 16 hrs. The crude product was purified by column chromatography, eluting with dichloromethane/methanol 97:3 to give the pure product as a white foamy solid (74 mg, yield 25.8 %).

^{31}P -NMR (CDCl_3 , 121 MHz): δ 3.47, 3.33.

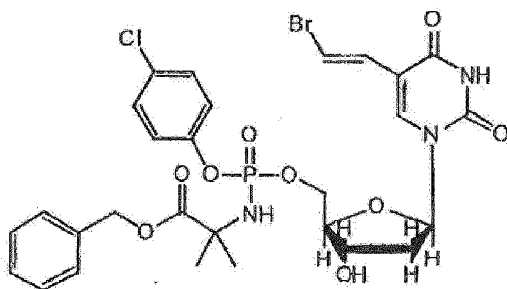
^1H -NMR (CDCl_3 , 300 MHz): δ 10.03-9.99 (1H, 2bs, H-3), 7.70-7.67 (1H, 2s, H-6), 7.47-7.43 (1H, 2d, $^3J=13.6$ Hz, H-5b), 7.35-7.20 (4H, m, *O**Ph*), 6.77-6.68 (1H, 2d, $^3J=13.6$ Hz, H-5a), 6.33-6.27 (1H, m, H-1'), 4.55-4.29 (5H, m, H-3'+H-5'+ OH-3'-N*H*), 4.22-4.17 (2H, q, $^3J=7.1$ Hz, O*C**H**2**C**H**3*+H-4'), 2.53-2.42 (1H, m, one of H-2'), 2.22-2.08 (1H, m; one of H-2'), 1.57-1.54 (6H,

2s, [$\underline{\text{CH}_3}_2\text{C}$), 1.31-1.30 (3H, 2t, $^3J=7.1$ Hz, $\text{OCH}_2\underline{\text{CH}_3}$).

^{13}C -NMR (CDCl_3 , 75 MHz): δ 14.5 ($\underline{\text{CH}_3\text{CH}_2\text{O}}$), 27.2, 27.3, 27.4 ($[\underline{\text{CH}_3}_2\text{C}]$), 40.7 (C-2'), 57.6 ($\underline{\text{C}}[\underline{\text{CH}_3}_2]$), 62.6 ($\text{CH}_3\underline{\text{CH}_2\text{O}}$), 66.5, 66.6 (C-5'), 70.8, 71.1 (C-3'), 85.5, 85.74, 86.0 (C-1', C-4'), 110.4 (C-5b), 112.0 (C-5), 121.9, 122.0, 122.1 ('o', $\text{O}\underline{\text{Ph}}$), 128.9, 130.2 ('m', $\text{O}\underline{\text{Ph}}$ +C-5a), 130.9 ('p', $\text{O}\underline{\text{Ph}}$), 138.3 (C-6), 149.4 ('ipso', $\text{O}\underline{\text{Ph}}$), 149.9 (C-4), 162.1, 162.2 (C-2), 175.7-175.9 ($\underline{\text{COOCH}_2\text{CH}_3}$).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine-5'-[4-chlorophenyl-(benzoxy- α,α -dimethylglycyl)]-phosphate (CPF 44).

[0142] $\text{C}_{28}\text{H}_{30}\text{BrClN}_3\text{O}_9\text{P}$, MW=698.88.



[0143] This was synthesised according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), 4-chlorophenyl-(benzyl-2-amino-2-methylpropanoate)-phosphorochloridate (505.0 mg, 1.25 mmol), NMI (184.7 mg, 2.25 mmol, 179.3 μL) in THF (5 mL) for 16 hrs. The crude product was purified by column chromatography, eluting with dichloromethane/methanol 97:3 to give the pure product as a white foamy solid (134.8 mg, yield 42.9%).

^{31}P -NMR (CDCl_3 , 121 MHz): δ 3.44, 3.26.

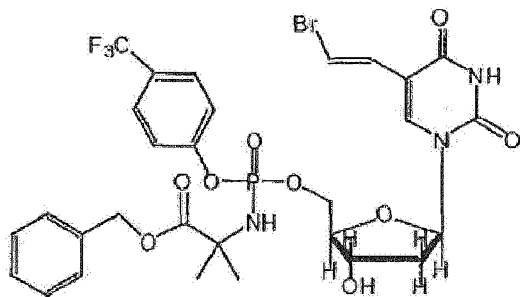
^1H -NMR (CDCl_3 , 300 MHz): δ 9.96-93 (1H, 2bs, H-3), 7.66-7.65 (1H, 2s, H-6), 7.47-7.41 (1H, 2d, $^3J=13.5$, H-5b), 7.39-7.18 (9H, m, $\text{O}\underline{\text{Ph}}$ + $\text{CH}_2\underline{\text{Ph}}$) 6.74-6.69 (1H, 2d, $^3J=13.5$ Hz, H-5a), 6.31-6.25 (1H, m, H-1'), 5.19 (2H, $\underline{\text{CH}_2\text{Ph}}$), 4.51-4.29 (4H, m, H-3'+H-5'+ NH), 4.15-4.12 (2H, m, H-4'+OH-3'), 2.48-2.40 (1H, m, one of H-2'), 2.18-2.05 (1H, m, one of H-2'), 1.60-1.59 (6H, 2s, [$\underline{\text{CH}_3}_2\text{C}$]).

^{13}C -NMR (CDCl_3 , 75 MHz): δ 27.1, 27.5 ($[\underline{\text{CH}_3}_2\text{C}]$), 40.7 (C-2'), 57.7 ($\underline{\text{C}}[\underline{\text{CH}_3}_2]$), 66.4, 66.6 (C-5'), 68.2 ($\underline{\text{CH}_2\text{Ph}}$), 70.7, 71.1 (C-3'), 85.4, 85.5, 85.7, 86.0 (C-1', C-4'), 110.5 (C-5b), 112.0 (C-5), 121.9, 122.0, 128.4, 128.5, 128.9, 129.1 ('o', 'm', 'p', $\text{CH}_2\underline{\text{Ph}}$ + $\text{O}\underline{\text{Ph}}$ +C-5a), 131.0 ('ipso', $\text{CH}_2\underline{\text{Ph}}$), 135.6 ('p', $\text{O}\underline{\text{Ph}}$), 138.1 (C-6), 149.3 ('ipso', $\text{O}\underline{\text{Ph}}$), 149.8 (C-4), 162.1 (C-2), 175.6

(COOCH₂Ph).

Synthesis of illustrative example (E)-5-(2-bromovinyl)-2'-deoxyuridine-5'-[para-(trifluoromethyl)-phenyl-(benzoxy- α,α -dimethylglycyl)]-phosphate (CPF 48).

[0144] C₂₉H₃₀BrF₃N₃O₉P, MW=732.44.



[0145] This was synthesised according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), 4-(trifluoromethyl)-phenyl-(benzyl-2-amino-2-methylpropanoate)-pbosporochloridate (529.4.5 mg, 1.22 mmol), NMI (184.7 mg, 2.25 mmol, 179.4 μ L) in THF (5 mL) for 4 hrs. The crude product was purified by column chromatography, eluting with dichloromethane/methanol 97:3 to give the pure product as a white foamy solid (142.1 mg, yield 43.1%).

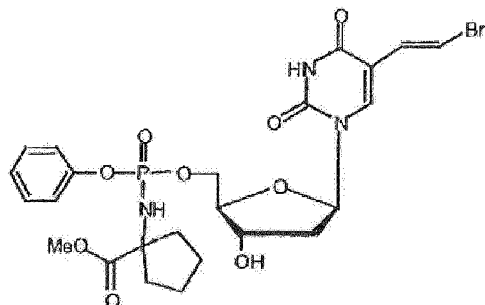
³¹P-NMR (CDCl₃, 121 MHz): δ 3.16, 3.01.

¹H-NMR (CDCl₃, 300 MHz): δ 10.06-10.02 (1H, 2bs, H-3), 7.67-7.66 (1H, s, H-6), 7.64-7.60 (2H, 2d, ³J=8.8 Hz, O_{Ph}), 7.46-7.32 (8H, m, O_{Ph}+CH₂Ph+H-5b), 6.77-6.68 (1H, 2d, ³J=13.6 Hz, H-5a), 6.31-6.26 (1H, m, H-1'), 5.18 (2H, s, CH₂Ph), 4.61-4.32 (4H, m, H-3'+H-5'+NH), 4.16-4.15 (2H, m, H-4'+OH-3'), 2.48-2.41 (1H, m, one of H-2'), 2.23-2.09 (1H, m, one of H-2'), 1.60-1.58 (6H, 2s, C[CH₃]₂)

¹³C-NMR (CDCl₃, 75 MHz): δ 27.0, 27.4, 27.5 (C[CH₃]₂), 40.6 (C-2'), 57.7, 57.8 (C[CH₃]₂), 66.8, 66.5 (C-5'), 68.2(CH₂Ph), 70.8, 71.1 (C-3'), 85.4, 85.7, 86.0 (C-1', C-4'), 110.4 (C-5b), 111.9 (C-5), 120.8, 120.9, 121.0, 127.6, 127.7, 128.0, 128.5, 128.8, 129.0 ('o', 'm', 'p', - O_{Ph}+CH₂Ph+C-5a), 124.2 (CF₃, J=267 Hz), 135.6 ('ipso', CH₂Ph), 138.2 (C-6), 149.9 (C-4), 153.3 ('ipso', O_{Ph}), 162.1 (C-2), 175.4 (COOCH₂Ph).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[phenyl-(methoxy- α,α -cycloleucynyl)]-phosphate (CPF 16).

[0146] $C_{24}H_{29}BrN_3O_9P$, MW-614.38.



[0147] This was synthesised according to **Standard procedure 5**, using BVdU (250 mg, 0.75 mmol), Phenyl-(methoxy- α,α -cycloleucynyl)-phosphorochloridate (589 mg, 1.87 mmol), NMI (6.2 mmol, 415 μ L) in THF (7 mL) for 3 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (234 mg, yield 51%).

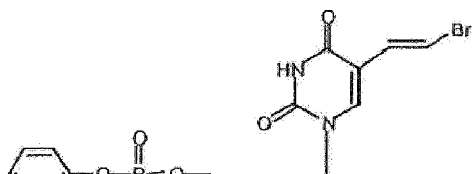
^{31}P -NMR ($CDCl_3$, 121 MHz): δ 3.87, 3.82.

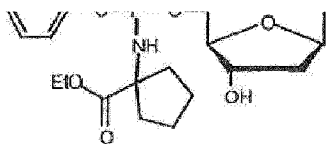
1H -NMR ($CDCl_3$; 300 MHz): δ 10.35-10.2 (1H, bs, H-3), 7.65 (1H, 2xs, H-6), 7.44-7.39 (1M, 2d, $^3J=13$ Hz, H-5b), 7.37-7.15 (5H, m, OPh), 6.8 (1H, 2d, $^3J=13$ Hz, H-5a), 6.30 (1H, 2t, $^3J=6$ Hz, H1'), 4.4-4.2 (4H, m, H-5', H-3', NH), 4.1 (1H, H-4'), 3.72 (3H, 2s, CH_3O), 2.49-2.40 (1H, m, one of H-2'), 2.35-2.01 (5H, m, one of H-2'+4H cyclopentane), 1.8-1.6 (4H, m, 4H cyclopentane).

^{13}C -NMR (DMSO; 75 MHz): δ 24.4, 24.3, 24.2 ($2CH_2$ cyclopent), 39.2, 38.6, 38.5 ($2CH_2$ cyclopent), 40.0 (C-2'), 53.2 (CH_3O), 66.4 ($C\alpha$ cyclopentane), 66.6 (C-5'), 70.9 (C-3'), 85.8, 85.6, 85.4, 85.3 (C-1', C-4'), 110.2 (C-5b), 111.9 (C-5), 120.7-120.6 ('o', OPh), 125.7 ('p', OPh), 129.0 (C-5a), 130.2 ('m', OPh), 138.5 (C-6), 149.9 (C-4), 150.9, 150.8 ('ipso', OPh), 162.3(C-2), 176.3, 176.2 ($COOCH_3$).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[phenyl-(ethoxy- α,α -cycloleucynyl)]-phosphate (CPF 17).

[0148] $C_{25}H_{31}BrN_3O_9P$, MW=628.41.





[0149] This was synthesised according to **Standard procedure 5**, using BVdU (250 mg, 0.75 mmol), Phenyl-(ethoxy- α,α -cycloleucynyl)-phosphorochloridate (642 mg, 1.87 mmol), NMI (6.2 mmol; 415 μ L) in THF (7 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (258 mg, yield 55%).

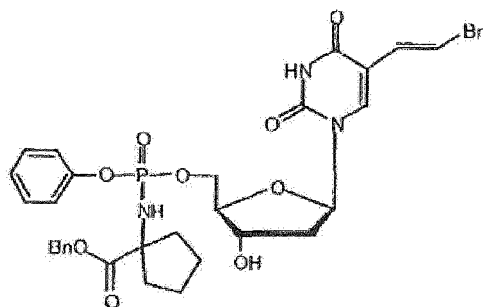
^{31}P -NMR (CDCl_3 , 121 MHz): δ 4.23, 4.1.

^1H -NMR (CDCl_3 , 300 MHz): δ 10.3-10.1 (1H, bs, H-3), 7.8-7.75 (1H, 2xs, H-6), 7.51 (1H, 2d, $^3J=14$ Hz, H-5b), 7.45-7.10 (5H, m, *OPh*), 6.8 (1H, 2d, $^3J=14$ Hz, H-5a), 6.22 (1H, 2t, $^3J=4$ Hz, H1'), 4.55-4.05 (7H, m, H-5', H-3', H-4', NH, $\text{CH}_3\text{CH}_2\text{O}$), 2.50-2.40 (1H, m, one of H-2'), 2.35-1.95 (5H, m, one of H-2'+4H cyclopentane), 1.95-1.75 (4H, m, 4H cyclopentane), 1.25 (3H, 2t, $^3J=7$ Hz, $\text{CH}_3\text{CH}_2\text{O}$).

^{13}C -NMR (CDCl_3 , 75 MHz): δ 14.5 ($\text{CH}_3\text{CH}_2\text{O}$), 24.5, 24.4 (2 CH_2 cyclopent), 39.2, 38.9 38.8, 38.4 (2 CH_2 cyclopent), 40.6 (C-2'), 62.2, 62.1 ($\text{CH}_3\text{CH}_2\text{O}$), 66.2 (*Cq* cyclopentane), 66.6 (C-5'), 70.8 (C-3'), 85.7, 85.5 (C-1', C-4'), 110.2 (C-5b), 111.5 (C-5), 120.7, 120.6 ('o', *OPh*), 125.6 ('p', *OPh*), 129.7 (C-5a), 130.2 ('m', *OPh*), 138.5, 138.3 (C-6), 149.7 (C-4), 150.9, 150.8 ('ipso', *OPh*), 162.3 (C-2), 176.3 ($\text{COOCH}_2\text{CH}_3$).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[phenyl-(benzoxycarbonyl- α,α -cycloleucynyl)]-phosphate (CPF 18).

[0150] $\text{C}_{30}\text{H}_{33}\text{BrN}_3\text{O}_9\text{P}$, MW=690.48.



[0151] This was synthesised according to **Standard procedure 5**, using BVdU (200 mg, 0.6

mmol), Phenyl-(benzyloxy- α,α -cycloleucynyl)-phosphorochloridate (589 mg, 1.5 mmol). NMI (4.98 mmol, 332 μ L) in THF (5 mL) for 10 hrs, The crude, product was purified by column chromatography, during with CH₂Cl₂/Methanol 97:3 to give the pure product as a white foamy solid (127 mg, yield 31%).

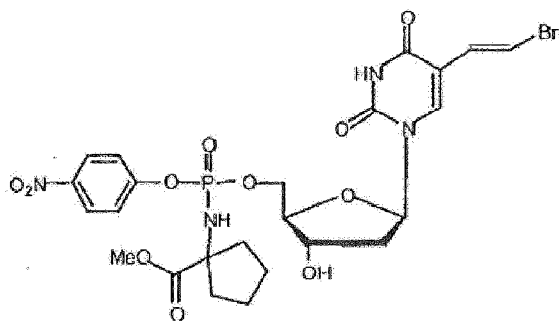
³¹P-NMR (CDCl₃, 121 MHz): δ 4,11, 4.01.

¹H-NMR (CDCl₃, 300 MHz): δ 10.2 (1H, bs, H-3), 7.8-7.6 (1H, 2xs, H-6), 7.45-7.4 (1H, 2d, ³J=14 Hz, H-5b), 7.40-7.10 (10H, m, OPh+CH₂Ph), 6.85 (1H, 2d, ³J=14 Hz, H-5a), 6.20 (1H, m, H-1'), 5.15 (1H, s, CH₂Ph), 4.4-4.2 (3H, m, H-3',H-4', NH), 4.1 (2H, m, H-5'), 2.45-2.35 (1H, m, one of H-2'), 2.35-1.95 (5H, m, one of H-2'+4H cyclopentane), 1.95-1.75 (4H, m, 4H cyclopentane).

¹³C-NMR (CDCl₃, 75 MHz): δ 24.4, 24.3, 24.2 (2CH₂ cyclopent), 39.9, 39.7 38.6, 38.5 (2CH₂ cyclopent), 40.5 (C-2'), 66.2 (C α cyclopentane), 66.5 (C-5'), 67.8 (CH₂Ph), 70.8, 70.7 (C-3'), 85.7, 85.6, 85.5, 85.4 (C-1', C-4'), 110.2 (C-5b), 111.8, 118.7 (C-5b), 120.7, 120.5 ('o', OPh), 125.7 ('p' OPh), 130.2, 129.0, 128.8, 128.7, 128.5 ('m' OPh, Bn, C-5a), 135.8('ipso', CH₂Ph) 138.4, 138.2 (C-6), 149.8 (C-4), 150.9, 150.8 ('ipso', OPh), 162.2 (C-2), 175.7, 175.5 (COOBn).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-nitrophenyl-(methoxy- α,α -cycloleucynyl)]-phosphate (CPF 19).

[0152] C₂₄H₂₈BrN₄O₁₁P, MW=659.38.



[0153] This was synthesized according to **Standard procedure 5**, using BVdU (200 mg, 0.60 mmol), para-nitrophenyl-(methoxy- α,α -cycloleucynyl)-phosphorochloridate (543 mg, 1.5 mmol), NMI (4.98 mmol, 332 μ L) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH₂Cl₂/Methanol 97:3 to give the pure product as a white foamy solid (239 mg, yield 60%),

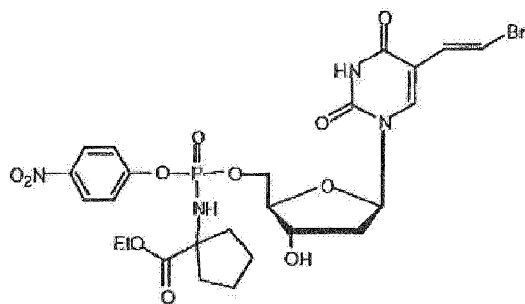
^{31}P -NMR (CDCl_3 , 121 MHz): δ 3.73.

^1H -NMR (CDCl_3 ; 300 MHz): δ 10.5-10.2 (1H, bs, H-3), 8.35-8.25 (2H, 2d, $^3J=6$ Hz *OPh*), 7.8-7.75 (1H, 2xs, H-6), 7.47 (2H, 2d, $^3J=6$ Hz, *OPh*), 7.45-7.35 (1H, 2d, $^3J=14$ Hz, H-5b), 6.75-6.67 (1H, 2d, $^3J=14$ Hz, H-5a), 6.30 (1H, 2t, $^3J=6$ Hz, H1'), 4.65-4.4 (3H, m, H-5', H-3'), 4.25-4.20 (1H, m, H-4'), 3.79 (3H, s, CH_3O), 2.6-2.4 (1H, m, one of H-2'), 2.3-1.98 (5H, m, one of H-2'+4H cyclopentane), 1.9-1.76 (4H, m, 4H cyclopentane).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 24.4, 24.3, 24.2 (2CH_2 cyclopent), 39.2, 39.1 (2CH_2 cyclopent), 40.5 (C-2'), 53.4, 53.3 (CH_3O), 66.8 (*Cq* cyclopentane), 67.1 (C-5'), 70.9 (C-3'), 86.1, 86.0, 85.5, 85.4 (C-1', C-4'), 110.2 (C-5b), 111.8 (C-5), 121.3, 121.2 ('o', *OPh*), 126.2 ('m' *OPh*), 128.9 (C-5a), 138.6 (C-6), 144.9 ('ipso', *OPh*) 149.9 (C-4), 155.9, 155.8 ('p' *OPh*), 162.3 (C-2), 176.3 (COOCH_3).

Synthesis of illustrative example (E)-S-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-nitrophenyl-(ethoxy- α,α -cycloleucynyl)]-phosphate (CPF 20).

[0154] $\text{C}_{25}\text{H}_{30}\text{BrN}_3\text{O}_{11}\text{P}$, MW=673.4.



[0155] This was synthesised according to **Standard procedure 5**, using BVdU (200 mg, 0.60 mmol), para-nitrophenyl-(ethoxy- α,α -cycloleucynyl)-phosphorochloridate (563 mg, 1.5 mmol), NMI (4.98 mmol, 332 μL) in THF (5 mL) for 1 hr. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (240 mg, yield: 59%),

^{31}P -NMR (CDCl_3 , 121 MHz): δ 3.83, 3.79,

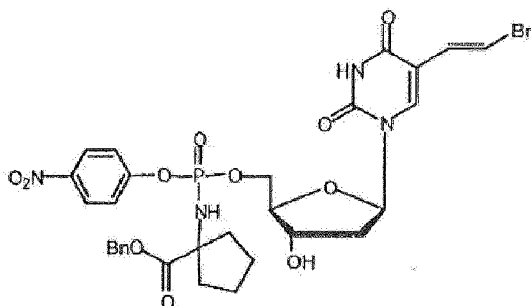
^1H -NMR (CDCl_3 , 300 MHz): δ 8.25-8.2 (2H, 2d, $^3J=9\text{Hz}$ *OPh*), 7.66 (1H, s, H-6), 7.4 (2H, 2d, $^3J=9\text{Hz}$, *OPh*), 7.3 (1H, 2d, $^3J=14$ Hz, H-5b), 6.85 (1H, 2d, $^3J=14$ Hz, H-5a), 6.3-6.2 (1H, m, H1'), 4.7-4.45 (4H, m, H-5', H-3', NH), 4.2-4.05 (3H, m, H-4', $\text{CH}_3\text{CH}_2\text{O}$), 2.55-2.4 (1H, m, one of H-2'), 2.2-1.95 (5H, m, one of H-2'+4H cyclopentane), 1.95-1.8 (4H, m, 4H cyclopentane),

1.2 (3H, 2t, $^3J=8$ Hz, $\text{CH}_3\text{CH}_2\text{O}$).

^{13}C -NMR (CDCl_3 , 75 MHz): δ 14.9 ($\text{CH}_3\text{CH}_2\text{O}$), 24.5, 24.4 (2CH_2 cyclopent), 39.1, 39.0, 38.8 (2CH_2 cyclopent), 40.7 (C-2'), 62.4 ($\text{CH}_3\text{CH}_2\text{O}$), 66.5 (Cq cyclopentane), 67.0 (C-5'), 70.9 (C-3'), 85.9, 85.4 (C-1', C-4'), 110.2 (C-5b), 111.8 (C-5), 121.3 ('o', OPh), 126.2 ('m', OPh), 128.8 (C-5a), 138.5 (C-6), 144.9 ('ipso', OPh), 149.9 (C-4), 155.5 ('p', OPh), 162.3 (C-2), 175.8, 175.7 ($\text{COOCH}_2\text{CH}_3$).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-nitrophenyl-(benzyloxy- α,α -cycloleucynyl)]-phosphate (CPF 21).

[0156] $\text{C}_{30}\text{H}_{32}\text{BrN}_4\text{O}_{11}\text{P}$, MW=735.47.



[0157] This was synthesised according to **Standard procedure 5**, using BVdU (200 mg, 0.60 mmol), para-nitrophenyl-(benzyloxy- α,α -cycloleucynyl)-phosphorochloridate (656 mg, 1.5 mmol), NMI (4.98 mmol, 332 μL) in THF (5 mL) for 3 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (269 mg, yield 61%).

^{31}P -NMR (CDCl_3 , 121 MHz): δ 3.72.

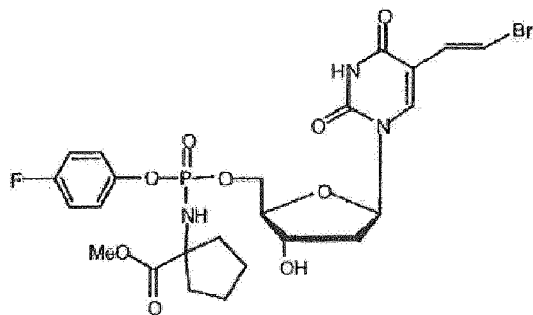
^1H -MMR (CDCl_3 , 300 MHz): δ 10.3 (1H, bs, H-3), 8.22-8.12 (2H, 2d, $^3J=7$ Hz, OPh), 7.65 (1H, 2xs, H-6), 7.45-7.30 (8H, m, H-5b+ OPh + CH_2Ph), 6.72-6.65 (1H, 2d, $^3J=14$ Hz, H-5a), 6.28 (1H, 2t, $^3J=6$ Hz, H-1'), 5.15 (1H, d, CH_2Ph), 4.6-4.35 (4H, m, H-3', H-5', H-4', NH), 2.55-2.4 (1H, m, one of H-2'), 2.3-1.92 (5H, m, one of H-2'+4H cyclopentane), 1.85-1.6 (4H, m, 4H cyclopentane).

^{13}C -NMR (CDCl_3 , 75 MHz): δ 24.4, 24.3, 24.2 (2CH_2 cyclopent), 39.1, 38.9, 38.7 (2CH_2 cyclopent), 40.5 (C-2), 66.9 (Cq cyclopentane), 67.1 (C-5'), 68.0 (CH_2Ph), 70.9 (C-3'), 85.3, 85.0 (C-1', C-4'), 110.3 (C-5b), 111.8 (C-5), 121.2 ('o', OPh), 126.1 ('m', OPh), 129.0, 128.8 (Bn, C-5a), 135.7 ('ipso', CH_2Ph), 138.5 (C-6), 144.9 ('ipso', OPh), 149.9 (C-4), 155.8 ('p'

OPh), 162.3 (C-2), 175.6 (COOBn).

Synthesis of illustrative example (E)-S-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-fluorophenyl-(methoxy- α,α -cycloleuciny)]-phosphate (CPF 22).

[0158] $\text{C}_{24}\text{H}_{28}\text{BrFN}_3\text{O}_9\text{P}$, MW=632.37.



[0159] This was synthesised according to **Standard procedure 5**, using BVdU (200 mg, 0.60 mmol), para-fluorophenyl-(methoxy- α,α -cycloleuciny)-phosphorochloridate (503 mg, 1.5 mmol), NMI (4.98 mmol, 332 μL) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (251 mg, yield 66%).

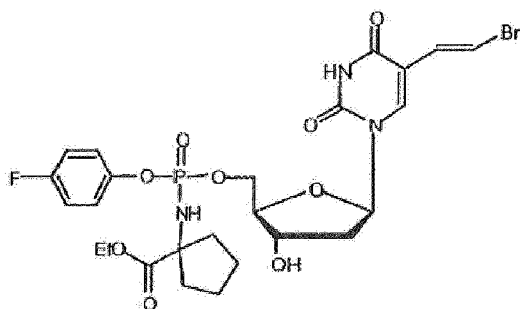
^{31}P -NMR (CDCl_3 , 121 MHz): δ 4.22.

^1H -NMR (CDCl_3 ; 300 MHz): δ 10.3 (1H, bs, H-3), 7.70 (1H, 2xs, H-6), 7.4 (1H, 2d, $^3J=14$ Hz, H-5b), 7.25-7.15 (2H, m, *OPh*), 7.1-6.95 (2H, m, *OPh*), 6.70 (1H, 2d, $^3J=14$ Hz, H-5a), 6.30-6.15 (1H, 2t, $^3J=5$ Hz, H1'), 4.55-4.05 (5H, m, H-5'+H-3', NH, H-4'), 3.72 (3H, 2s, CH_3O), 2.55-2.35 (1H, m, one of H-2'), 2.25-1.92 (5H, m, one of H-2'+4H cyclopentane), 1.85-1.6 (4H, m, 4H cyclopentane),

^{13}C -NMR (DMSO; 75 MHz): δ 24.4, 24.3, 24.2 (2CH_2 cyclopent), 39.3, 39.2, 38.9, 38.5 (2CH_2 cyclopent), 40.6 (C-2'), 53.3, 53.2 (CH_3O), 66.5 (*C α* cyclopentane), 66.7 (C-5'), 70.9 (C-3'), 85.8, 85.7, 85.4 (C-1', C-4'), 110.2 (C-5b), 111.9 (C-5), 116.9, 116.6 ('o', *OPh*), 122.2, 122.0 ('m', *OPh*), 128.5 (C-5a), 138.5 (C-6), 146.7 ('ipso', *OPh*), 149.9 (C-4), 158.5 ('p', *OPh*), 162.3 (C-2), 176.4, 176.3 (COOCH_3).

Synthesis of illustrative example (E)-S-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-fluorophenyl-(ethoxy- α,α -cycloleuciny)]-phosphate (CPF 23).

[0160] $C_{25}H_{30}BrFN_3O_9P$, MW=646.4.



[0161] This was synthesised according to **Standard procedure 5**, using BVdU (200 mg, 0.60 mmol), para-fluorophenyl-(ethoxy- α,α -cycloleucynyl)-phosphorochloridate (524 mg, 1.5 mmol), NMI (4.98 mmol, 332 μ L) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (274 mg, yield 71%).

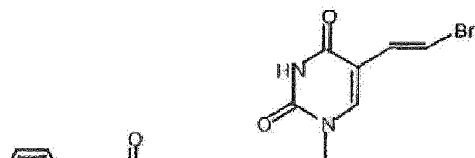
^{31}P -NMR ($CDCl_3$, 121 MHz): δ 5.30.

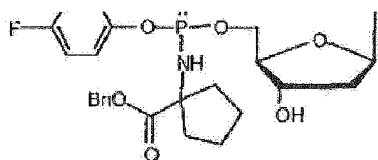
1H -NMR ($CDCl_3$, 300 MHz): δ 10.35 (1H, bs, H-3), 7.7 (1H, 2xs, H-6), 7.44 (1H, 2d, $^3J=14$ Hz, H-5b), 7.25-7.15 (2H, m, *OPh*), 7.1-6.95 (2H, m, *OPh*), 6.7 (1H, 2d, $^3J=14$ Hz, H-5a), 6.30 (1H, 2t, $^3J=6$ Hz, H1'), 4.55, 4.3 (3H, m, H-5', H-3'), 4.24, 4.1 (4H, m, NH, H-4', CH_3CH_2O), 2.55-2.4 (1H, m, one of H-2'), 2.22-1.90 (5H, m, one of H-2'+4H cyclopentane), 1.8-1.6 (4H, m, 4H cyclopentane), 1.3-1.2 (3H, 2t, $^3J=7$ Hz, CH_3CH_2O),

^{13}C -NMR ($CDCl_3$, 75 MHz): δ 14.5 (CH_3CH_2O), 24.6, 24.4, 24.3 (2 CH_2 cyclopent), 39.3, 39.2, 38.9, 38.6 (2 CH_2 cyclopent), 40.6 (C-2'), 62.2 (CH_3CH_2O), 66.5 (*Cq* cyclopentane), 66.7 (C-5'), 71.0 (C-3'), 85.8, 85.7, 85.5, 85.4 (C-1', C-4'), 110.2 (C-5b), 111.9 (C-5), 116.9, 116.5 ('o', *OPh*), 122.2, 122.1 ('m', *OPh*), 129.0 (C-5a), 138.5 (C-6), 146.8, 146.7 ('ipso', *OPh*), 149.9 (C-4), 158.5 ('p', *OPh*), 162.3 (C-2), 175.9, 175.8 (*COOCH_2CH_3*).

Synthesis of illustrative example (E)-S-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-fluorophenyl-(benzoyloxy- α,α -cycloleucynyl)]-phosphate (CPF 24).

[0162] $C_{30}H_{32}BrN_3O_9P$, MW=708.47.





[0163] This was synthesized according to **Standard procedure 5**, using BVdU (200 mg, 0.60 mmol), para-fluorophenyl-(benzyloxy- α,α -cycloleucynyl)-phosphorochloridate (616 mg, 1.5 mmol), NMI (4.98 mmol, 332 μ L) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, during with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (283 mg, yield 67%).

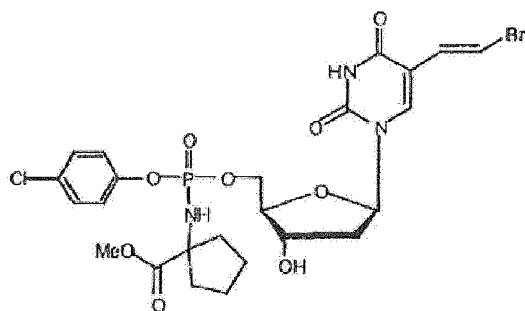
^{31}P -NMR (CDCl_3 , 121 MHz): δ 4.27.

^1H -NMR (CDCl_3 , 300 MHz): δ 10.3-9.85 (1H, bs, H-3), 7.65 (1H, 2xs, H-6), 7.45-7.35 (1H, 2d, $^3J=14$ Hz, H-5b), 7.40-7.30 (5H, m, CH_2Ph), 7.25-7.15 (2H, m, OPh), 7.05-6.95 (2H, m, OPh), 6.71 (1H, 2d, $^3J=14$ Hz, H-5a), 6.27 (1H, 2t, $^3J=6$ Hz, H-1'), 5.15 (1H, s, CH_2Ph), 4.45 (1H, m, H-3'), 4.40-4.30 (2H, m, H-5') 4.20-4.05 (2H, m, H-4', NH), 2.5-2.4 (1H, m, one of H-2'), 2.25-1.9 (5H, m, one of H-2'+4H cyclopentane), 1.8-1.6 (4H, m, 4H cyclopentane).

^{13}C -NMR (CDCl_3 , 75 MHz): δ 24.5, 24.3, 24.2 (2CH_2 cyclopent), 39.7, 39.6, 39.3, 39.2 (2CH_2 cyclopent), 40.5, 40.0 (C-2'), 66.6 ($\text{C}\alpha$ cyclopentane), 67.2, 66.7 (C-5'), 67.9 (CH_2Ph), 70.8, 70.7 (C-3'), 85.3, 85.7, 85.4, 85.3 (C-1', C-4'), 110.3 (C-5b), 111.8 (C-5), 116.9, 116.6 ('o', OPh), 122.2, 122.1 ('m', OPh), 129.0, 128.9, 128.6, 128.5 (Bn, C-5a), 135.8('ipso', CH_2Ph), 138.5 (C-6), 146.8, 146.7 ('ipso', OPh), 149.9 (C-4), 158.5 ('p' OPh), 162.2 (C-2), 175.7, 175.0 (COOBn),

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-chlorophenyl-(methoxy- α,α -cycloleucynyl)]-phosphate (CPF 32).

[0164] $\text{C}_{24}\text{H}_{28}\text{BrClN}_3\text{O}_9\text{P}$, MW=648,82.



[0165] This was synthesised according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), para-chlorophenyl-(methoxy- α,α -cycloleucynyl)-phosphorochloridate (475 mg, 1.35 mmol), NMI (4.5 mmol, 300 μ L) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH₂Cl₂/Methanol 97:3 to give the pure product as a white foamy solid (187 mg, yield 64%).

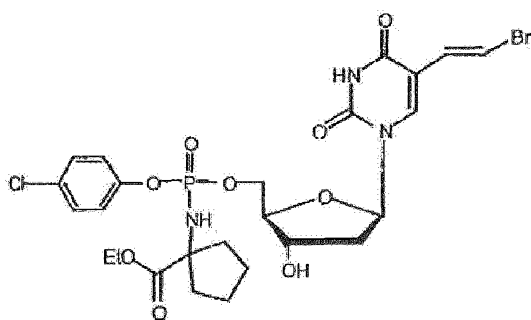
³¹P-NMR (MeOD, 121 MHz): δ 4.64.

¹H-NMR (MeOD; 300 MHz): δ 7.75 (1H, 2xs, H-6), 7.32 (1H, 2d, ³J=14 Hz, H-5b), 7.32-7.27(2H, m, *OPh*), 7.20-7.11 (2H,m, *OPh*), 6.72 (1H,2d, ³J=14 Hz, H-5a), 6.27-6.20 (1H, 2t, ³J=6 Hz, H1'), 4.35 (1H, m, H-3'), 4.30 (2H, m, H-5') 4.1 (2H, m, H-4'), 3.72 (3H, 2s, CH₃O), 2.32-2.20 (1H, m, one of H-2'), 2.20-1.92 (5H, m, one of H-2'+4H cyclopentane), 1.8-1.6 (4H, m, 4H cyclopentane).

¹³C-NMR (MeOD; 75 MHz): δ 25.7, 25.6 (2CH₂ cyclopent), 41.7, 41.6, 41.4, 41.3 (2CH₂ cyclopent), 42.7 (C-2'), 54.1, 53.9 (CH₃O), 67.8 (C α cyclopentane), 69.1, 69.0 (C-5'), 73.8 (C-3'), 88.4, 8.3, 88.2 (C-1', C-4'), 110.2 (C-5b), 111.8 (C-5), 122.1, 121.9 ('o', *OPh*), 128.9 (C-5a), 130.6 ('m', *OPh*), 130.8 ('p', *OPh*), 138.5 (C-6), 149.5, 149.4 ('ipso', *OPh*), 149.9 (C-4), 162.2(C-2), 175.6 (COOCH₃).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-chlorophenyl-(ethoxy- α,α -cycloleucynyl)]-phosphate (CPF 33).

[0166] C₂₅H₃₀BrClN₃O₉P, MW=662.85.



[0167] This was synthesized according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), para-chlorophenyl-(ethoxy- α,α -cycloleucynyl)-phosphorochloridate (495 mg, 1.35 mmol), NMI (4.5 mmol, 300 μ L) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH₂Cl₂/Methanol 97:3 to give the pure product as a white foamy solid (240 mg, yield 66%).

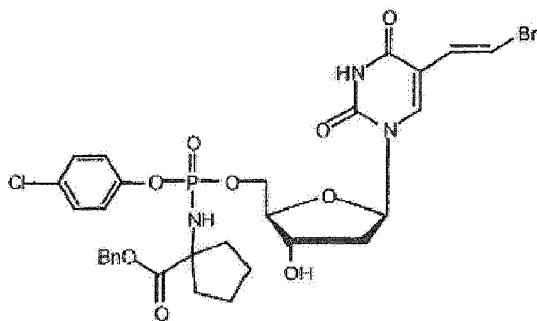
^{31}P -NMR (CDCl_3 , 121 MHz): δ 4.15.

^1H -NMR (CDCl_3 , 300 MHz): δ 10.25-10.1 (1H, bs, H-3), 7.65 (1H, 2xs, H-6), 7.4-7.3 (1H, 2d, $^3J=14$ Hz, H-5b), 7.25-7.20 (2H, m, *OPh*), 7.20-7.10 (2H, m, *OPh*), 6.75 (1H, 2d, $^3J=14$ Hz, H-5a), 6.20 (1H, m, H1'), 4.3-4.5 (3H, m, H-3', H-5'), 4.2-4.0 (4H, m, H-4', NH, $\text{CH}_3\text{CH}_2\text{O}$), 2.45-2.25 (1H, m, one of H-2'), 2.25-1.85 (5H, m, one of H-2'+4H cyclopentane), 1.75-1.55 (4H, m, 4H cyclopentane), 1.2 (3H, 2t, $^3J=7$ Hz, $\text{CH}_3\text{CH}_2\text{O}$).

^{13}C -NMR (CDCl_3 , 75 MHz): δ 14.5 ($\text{CH}_3\text{CH}_2\text{O}$), 24.5, 24.4 (2 CH_2 cyclopent), 39.3, 39.2, 38.8, 38.6 (2 CH_2 cyclopent), 40.5 (C-2'), 62.3 ($\text{CH}_3\text{CH}_2\text{O}$), 66.1 (*Cq* cyclopentane), 66.7 (C-5'), 70.8 (C-3'), 85.8, 85.4 (C-1', C-4'), 110.3 (C-5b), 111.9 (C-5), 122.1, 121.9 ('o', *OPh*), 129.0 (C-5a), 130.2 ('m', *OPh*), 130.8 ('p', *OPh*), 138.5 (C-6), 149.5, 149.4 ('ipso', *OPh*), 149.9 (C-4), 162.3 (C-2), 175.9 ($\text{COOCH}_2\text{CH}_3$).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-chlorophenyl-(benzyloxy- α,α -cycloleucynyl)]-phosphate (CPF 34).

[0168] $\text{C}_{30}\text{H}_{32}\text{BrClN}_3\text{O}_9\text{P}$, MW=724.92.



[0169] This was synthesised according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), para-chlorophenyl-(benzyloxy- α,α -cycloleucynyl)-phosphorochloridate (578 mg, 1.35 mmol), NMI (4.5 mmol, 300 μL) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (222 mg, yield 68%),

^{31}P -NMR (CDCl_3 , 121 MHz): δ 4.11, 4.05.

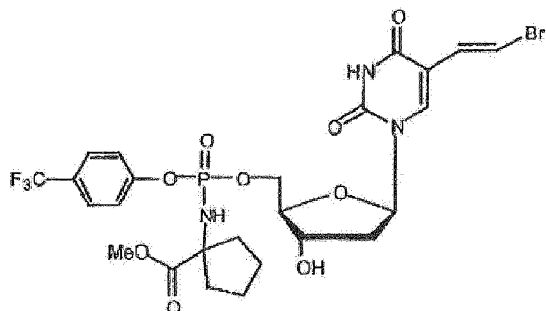
^1H -NMR (CDCl_3 , 300 MHz): δ 7.65 (1H, 2xs, H-6), 7.45-7.29 (10H, m, H-5b, 2H *OPh*+ CH_2Ph), 7.20-7.15 (2H, m, *OPh*), 6.75-6.67 (1H, 2d, $^3J=14$ Hz, H-5a), 6.28 (1H, 2t, $^3J=6\text{Hz}$, H-1'), 5.15 (1H, 2s, CH_2Ph), 4.5 (1H, m, H-3'), 4.35 (2H, m, H-5') 4.1 (H, m, H-4'), 4.00 (1H, m, NH), 2.48-

2.35 (1H, m, one of H-2'), 2.3-1.92 (5H, m, one of H-2'+4H cyclopentane), 1.8-1.6 (4H, m, 4H cyclopentane).

^{13}C -NMR (CDCl_3 , 75 MHz): δ 24.5, 24.4, 24.3, 24.2 (2CH_2 cyclopent), 39.3, 38.8, 38.6 (2CH_2 cyclopent), 40.5 (C-2'), 66.7 ($\underline{\text{Cq}}$ cyclopentane), 67.9 ($\underline{\text{CH}_2\text{Ph}}$), 68.4 (C-5'), 70.7 (C-3'), 85.7, 85.7, 85.4, 85.3 (C-1', C-4'), 110.3 (C-5b), 111.8 (C-5), 122.0, 121.9 ('o', $\underline{\text{OPh}}$), 129.1, 128.3, 128.2 (Bn, 'm', $\underline{\text{OPh}}$), 130.2 (C-5a), 135.8 ('ipso', $\text{CH}_2\underline{\text{Ph}}$), 136.3 ('p' OPh), 138.2 (C-6), 149.5, 149.3 ('ipso', $\underline{\text{OPh}}$), 149.9 (C-4), 162.2 (C-2), 175.7, 175.5 ($\underline{\text{COOBn}}$).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-trifluorophenyl-(methoxy- α,α -cycloleucynyl)]-phosphate (CPF 28).

[0170] $\text{C}_{25}\text{H}_{28}\text{BrF}_3\text{N}_3\text{O}_9\text{P}$, MW=682.38.



[0171] This was synthesized according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), para-trifluorophenyl-(methoxy- α,α -cycloleucynyl)-phosphorochloridate (521 mg, 1.35 mmol), NMI (4.5 mmol, 300 μL) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (199 mg, yield 65%).

^{31}P -NMR (CDCl_3 , 121 MHz): δ 3.80.

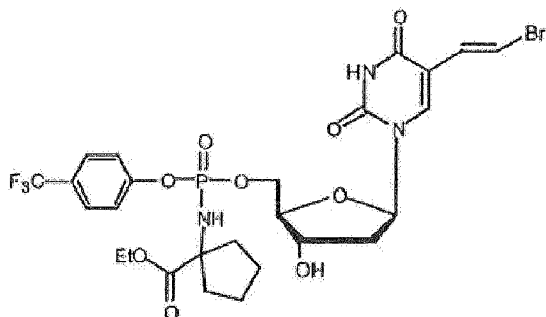
^1H -NMR (CDCl_3 ; 300 MHz): δ 7.70 (1H, 2s, H-6), 7.55 (1H, 2d, $^3J=14$ Hz, H-5b), 7.45-7.32 (4H, m, $\underline{\text{OPh}}$), 6.72 (1H, 2d, $^3J=14$ Hz, H-5a), 6.28 (1H, 2t, $^3J=6\text{Hz}$, H1'), 4.55 (1H, m, H-3'), 4.45 (2H, m, H-5'), 4.25 (1H, H-4'), 4.15 (1H, NH), 3.71 (3H, 2s, CH_3O), 2.6-2.4 (1H, m, one of H-2'), 2.3-1.9 (5H, m, one of H-2'+4H cyclopentane), 1.85-1.6 (4H, m, 4H cyclopentane).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 24.4, 24.3, 24.2 (2CH_2 cyclopent), 39.2, 39.1, 38.8, 38.6 (2CH_2 cyclopent), 40.5 (C-2'), 53.9 ($\underline{\text{CH}_3\text{O}}$), 66.3 ($\underline{\text{Cq}}$ cyclopentane), 66.8 (C-5'), 70.9 (C-3'), 85.8, 85.4 (C-1', C-4'), 110.3 (C-5b), 111.9 (C-5), 125.1 (d, $J=270\text{Hz}$, CF_3), 127.1, 127.0 ('o', $\underline{\text{OPh}}$), 127.8 ('m', $\underline{\text{OPh}}$), 128.9 (C-5a), 129.0 ('p', q, $J=32\text{Hz}$, $\underline{\text{OPh}}$), 138.5 (C-6), 149.9 (C-4), 153.5

('ipso', OPh), 162.2 (C-2), 176.3, 176.2 (COOCH₃).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-trifluorophenyl-(ethoxy- α,α -cycloleuciny)]-phosphate (CPF 29).

[0172] C₂₆H₃₀BrF₃N₃O₉P, MW=696.40.



[0173] This was synthesised according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), para-trifluorophenyl-(ethoxy- α,α -cycloleuciny)-phosphorochloridate (540 mg, 1.35 mmol), NMI (4.50 mmol, 300 μ L) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH₂Cl₂/Methanol 97:3 to give the pure product as a white foamy solid (185 mg, yield 59%).

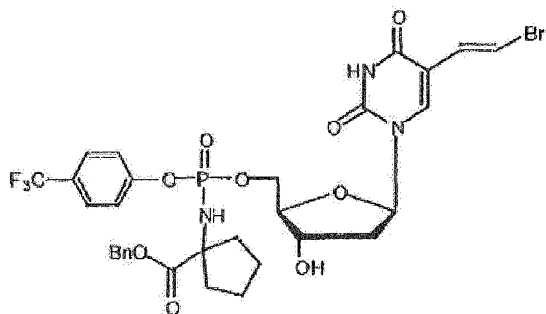
³¹P-NMR (CDCl₃, 121 MHz): δ 4.30.

¹H-NMR (CDCl₃, 300 MHz): δ 10.35 (1H, bs, H-3), 7.70 (1H, 2xs, H-6), 7.40 (1H, 2d, ³J=14 Hz, H-5b), 7.28-7.14 (2H, m, OPh), 7.05-6.95 (2H, m, OPh), 6.70 (1H, 2d, ³J=14 Hz, H-5a), 6.3 (1H, m, H1'), 4.55-4.3 (3H, m, H-5', H-3'), 4.2-4.1 (3H, m, H-4', CH₃CH₂O), 2.5-2.35 (1H, m, one of H-2'), 2.20-1.9 (5H, m, one of H-2'+4H cyclopentane), 1.85-1.6 (4H, m, 4H cyclopentane), 1.25 (3H, 2t, ³J=7 Hz, CH₃CH₂O).

¹³C-NMR (CDCl₃, 75 MHz): δ 14.5 (CH₃CH₂O), 24.5, 24.4 (2CH₂ cyclopent), 39.3, 39.2, 38.9, 38.5 (2CH₂ cyclopent), 40.6 (C-2'), 62.2 (CH₃CH₂O), 66.7 (Cq cyclopentane), 67.4, 67.3 (C-5'), 70.9 (C-3'), 85.8, 85.7 (C-1', C-4'), 110.2 (C-5b), 111.9 (C-5), 116.8, 116.5 ('o', OPh), 122.2, 122.1 ('m', OPh), 125.1 (d, J=270Hz, CF₃), 129.0 (C-5a), 131.1 ('p', q, J=32Hz, OPh), 138.5 (C-6), 146.8, 146.7 ('ipso', OPh), 149.9 (C-4), 162.3 (C-2), 175.9, 175.8 (COOCH₂CH₃).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-trifluorophenyl-(benzoxy- α,α -cycloleuciny)]-phosphate (CPF 30).

[0174] $C_{31}H_{32}BrF_3N_3O_9P$, MW=758.47.



[0175] This was synthesized according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), para-trifluorophenyl-(benzyloxy- α,α -cycloleucinyloxy)-phosphorochloridate (623 mg, 1.35 mmol), NMI (4.5 mmol, 300 μ L) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (218 mg, yield 64%).

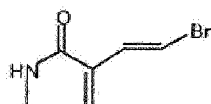
^{31}P -NMR ($CDCl_3$, 121 MHz): δ 4.30.

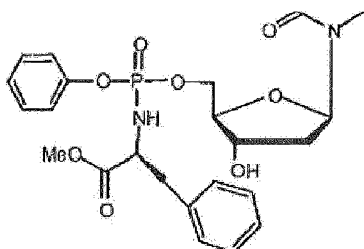
1H -NMR ($CDCl_3$, 300 MHz): δ 10.35 (1H, bs, H-3), 7.65 (1H, 2xs, H-6), 7.55 (2H, m, 2H *O*Ph), 7.45-7.25 (8H, m, 2H *O*Ph+ CH_2 Ph+H-5b), 6.7 (1H, 2d, $^3J=14$ Hz, H-5a), 6.30 (1H, 2t, $^3J=6$ Hz, H-1'), 5.15 (1H, 2.s, CH_2 Ph), 4.55-4.35 (3H, m, H-3'+ H-5'), 4.25 (1H, H-4'), 4.10 (1H, NH), 2.55-2.35 (1H, m, one of H-2'), 2.30-1.92 (5H, m, one of H-2'+4H cyclopentane), 1.8-1.6 (4H, m, 4H cyclopentane),

^{13}C -NMR ($CDCl_3$, 75 MHz): δ 25.5, 24.4, 24.3, 24.2 (2 CH_2 cyclopent), 39.2, 39.1, 38.7, 38.6 (2 CH_2 cyclopent), 40.5, 40.0 (C-2'), 66.4 (C α cyclopentane), 66.8 (C-5'), 68.0 (CH_2 Ph), 70.9 (C-3'), 86.0, 85.8, 85.4, 85.3 (C-1', C-4'), 110.3 (C-5b), 111.9 (C-5), 121.8, 120.8 ('o, *m*', *O*Ph), 125.2 (d, $J=270$ Hz, CF_3), 128.5, 127.7, 127.5 (Bn, C-5a), 129.2 ('*p*', *q*, $J=32$ Hz, *O*Ph), 135.4 ('*ipso*', CH_2 Ph), 138.5 (C-6), 149.9 (C-4), 153.5 ('*ipso*' *O*Ph), 162.2 (C-2), 175.6, 175.5 (COOBn).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[phenyl-(methoxy-L-phenylalaninyl)]-phosphate (CPF 36).

[0176] $C_{27}H_{29}BrN_3O_9P$, MW=650.41.





[0177] This was synthesised according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), Phenyl-(methoxy-L-phenylalaninyl)-phosphorochloridate (477 mg, 1.35 mmol), NMI (4.42 mmol, 190 μ L) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (169 mg, yield 58%),

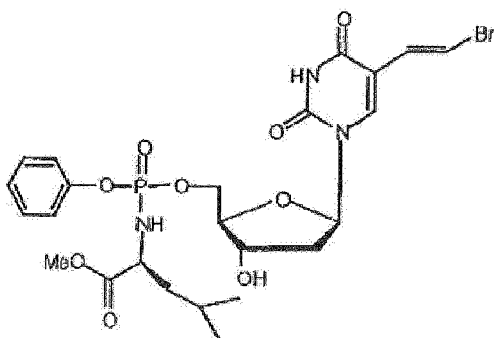
^{31}P -NMR (CDCl_3 , 121 MHz): δ 4.79, 4.71.

^1H -NMR (CDCl_3 ; 300 MHz): δ 9.95 (1H, bs, H-3), 7.60-7.55 (1H, 2xs, H-6), 7.48-7.4 (1H, 2d, $^3J=14$ Hz, H-5b), 7.3-7.1 (10H, m, $\text{CH}_2\text{Ph} + \text{OPh}$), 6.75-6.65 (1H, 2d, $^3J=14$ Hz, H-5a), 6.27-6.18 (1H, m, H1'), 4.57-4.29 (6H, m, H-5', H-3', H-4', NH, CHphenylala), 3.70 (3H, 2s, CH_3O), 3.01 (2H, m, CH_2Ph), 2.35-2.20 (1H, m, one of H-2'), 2.07-1.95 (1H, m, one of H-2').

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 36.3 ($\text{CH}_2\text{phenylalanine}$), 41.9, 41.8 (C-2'), 53.0 (CH_3O), 56.6, 56.1 (CHphenylala), 67.1 (C-5'), 71.3, 70.7 (C-3'), 85.7, 85.6, 85.5, 85.4 (C-1', C-4'), 110.4 (C-5b), 111.9 (C-5), 120.6, 120.5 ('o', OPh), 127.8 ('p', OPh), 130.1, 129.9, 129.8, 129.1 (CH_2Ph , C-5a, 'm' OPh), 138.0, 137.9 (C-6), 149.8 (C-4), 150.7, 150.6 ('ipso', OPh), 162.1, 162.0 (C-2), 173.5 (COOCH_3).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[phenyl-(methoxy-L-leucinyl)]-phosphate (CPF 35).

[0178] $\text{C}_{24}\text{H}_{31}\text{BrN}_3\text{O}_9\text{P}$, MW=616,40.



[0179] This was synthesised according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), Phenyl-(methoxy-L-leuciny)-phosphorochloridate (432 mg, 1.35 mmol), NMI (4.42 mmol, 190 μ L) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (167 mg, yield 60%).

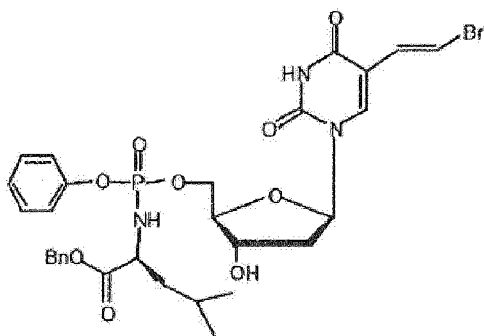
^{31}P -NMR (CDCl_3 , 121 MHz): δ 5.14, 4.60.

^1H -MMR (CDCl_3 ; 300 MHz): δ 10.1 (1H, bs, H-3), 7.75 (1H, 2xs, H-6), 7.45 (1H, 2d, $^3J=14$ Hz, H-5b), 7.4-7.2 (5H, m, *OPh*), 6.85 (1H, 2d, $^3J=14$ Hz, H-5a), 6.27-6.18 (1H, 2t, $^3J=6$ Hz, H1'), 4.5-4.2 (4H, m, H-5', H-3', NH), 4.1 (1H, m, H-4'), 3.95 (1H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 3.70 (3H, 2s, CH_3O), 2.40-2.20 (1H, m, one of H-2'), 2.05-1.95 (1H, m, one of H-2'), 1.8 (1H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 1.8-1.5 (2H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 1.0-0.9 (6H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 23.2, 23.1, 22.0, 21.9 (2C, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 24.9, 24.7 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 40.6 (C-2'), 43.7, 43.6 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 53.0 (CH_3O), 53.7, 53.6 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 66.6, 66.3 (C-5'), 71.1, 70.8 (C-3'), 86.0, 85.7, 85.6, 85.5 (C-1', C-4'), 110.4 (C-5b), 111.9 (C-5), 120.6, 120.5, 120.4 ('o', *OPh*), 125.8, 125.7 ('p', *OPh*), 128.9 (C-5a), 130.2 ('m' *OPh*), 138.1 (C-6), 149.9 (C-4), 150.8, 50.7 ('ipso', *OPh*), 162.2 (C-2), 175.1, 174.9 (COOH_3).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[phenyl-(benzoxy-L-leuciny)]-phosphate (CPF 37).

[0180] $\text{C}_{30}\text{H}_{35}\text{BrN}_3\text{O}_9\text{P}$, MW=692.49.



[0181] This was synthesised according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), Phenyl-(benzoxy-L-leuciny)-phosphorochloridate (534 mg, 1.35 mmol), NMI (4.42 mmol, 190 μ L) in THF (5 mL) for 2 hrs. The crude product was purified by column

chromatography, eluting with CH₂Cl₂/Methanol 97:3 to give the pure product as a white foamy solid (199 mg, yield 64%).

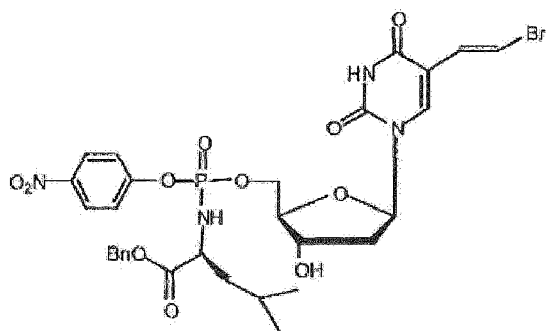
³¹P-NMR (CDCl₃, 121 MHz): δ 5.18, 4.54.

¹H-NMR (CDCl₃; 300 MHz): δ 9.95-9.85 (1H, bs, H-3), 7.55 (1H, 2xs, H-6), 7.38 (1H, 2d, ³J=14 Hz, H-5b), 7.3-7.1 (5H, m, CH₂Ph + *O**Ph*), 6.65 (1H, 2d, ³J=14 Hz, H-5a), 6.26-6.14 (1H, 2t, ³J=6 Hz, H1'), 5.1 (2H, 2s, CH₂Ph) 4.4-3.8 (6H, m, H-5', H-3', NH, H-4', CHCH₂CH(CH₃)₂), 2.35-2.25 (1H, m, one of H-2'), 1.95-1.85 (1H, m, one of H-2'), 1.6-1.4 (3H, m, CHCH₂CH(CH₃)₂), 0.8 (6H, m, CHCH₂CH(CH₃)₂).

¹³C-NMR (CDCl₃; 75 MHz): δ 23.2, 23.1, 22.0, 21.9 (2C, CHCH₂CH(CH₃)₂) 24.9, 24.7 (CHCH₂CH(CH₃)₂), 40.7 (C-2'), 43.9, 43.8 (CHCH₂CH(CH₃)₂), 53.9, 53.7 (CHCH₂CH(CH₃)₂), 66.4, 66.2 (C-5'), 67.8, 67.7 (CH₂Ph), 71.1, 70.7 (C-3'), 85.9, 85.6, 85.4, 85.3 (C-1', C-4'), 110.4 (C-5b), 111.9 (C-5), 120.6, 120.5 ('o', *O**Ph*), 125.8, 125.7 ('p', *O**Ph*), 130.2, 129.1, 128.9 (C-5a, CH₂Ph, 'm' *O**Ph*), 135.4 ('ipso', CH₂*Ph*), 138.1 (C-6), 149.8 (C-4), 150.2 ('ipso', *O**Ph*), 162.1 (C-2), 175.7, 174.6 (*COOBn*).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-nitrophenyl-(benzoxy-L-leucynyl)]-phosphate (CPF 38).

[0182] C₃₀H₃₄BrN₄O₁₁P, MW-737.49.



[0183] This was synthesised according to *Standard procedure 5*, using BVdU (150 mg, 0.45 mmol), para-nitrophenyl-(benzoxy-L-leucynyl)-phosphorochloridate (595 mg, 1.35 mmol), NM1 (4.42 mmol, 190 μL) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH₂Cl₂/Methanol 97:3 to give the pure product as a white foamy solid (176 mg, yield 53%).

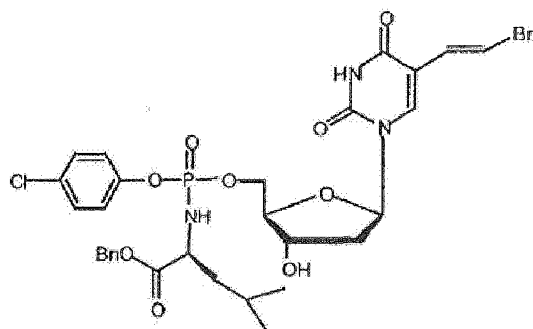
³¹P-NMR (CDCl₃, 121 MHz): δ 5.72, 4.35.

$^1\text{H-NMR}$ (CDCl_3 ; 300 MHz): δ 10.2 (1H, bs, H-3), 8.1(2H, m, 2H *OPh*), 7.65 (1H, 2xs, H-6), 7.45-7.2 (8H, m, H-5b, CH_2Ph + 2H *OPh*), 6.65 (1H, 2d, $^3J=14$ Hz, H-5a), 6.35-6.2 (1H, 2t, $^3J=6$ Hz, H1'), 5.15 (2H, 2s, CH_2Ph) 4.7-3.9 (6H, m, H-5', H-3', NH, H-4', $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 2.55-2.4 (1H, m, one of H-2'), 2.15-2.05 (1H, m, one of H-2'), 1.7-1.5 (3H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 0.95-0.8 (6H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$).

$^{13}\text{C-NMR}$ (CDCl_3 ; 75 MHz): δ 23.2, 23.1, 22.0, 21.9 (2C, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 24.9, 24.8 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 40.6 (C-2'), 43.7, 43.6 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 53.9, 53.7 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 66.9 (C-5'), 67.9 (CH_2Ph), 71.2, 70.8 (C-3'), 85.8, 85.3, 85.2 (C-1', C-4'), 110.6 (C-5b), 111.9 (C-5), 121.3 ('o', *OPh*), 129.2, 129.1, 128.8, 126.2 (C-5a, CH_2Ph , 'm' *OPh*), 135.4, 135.3 ('ipso', CH_2Ph), 138.2 (C-6), 145.2, 145.1 ('ipso', *OPh*), 149.9 (C-4), 155.5 ('p', *OPh*), 162.1 (C-2), 274.2 (COOBn).

Synthesis of illustrative example (E)-5-(2-Bromovinyl)-2'-deoxyuridine-5'-[para-chlorophenyl-(benzoxy-L-leuciny)]-phosphate (CPF 39).

[0184] $\text{C}_{30}\text{H}_{34}\text{BrClN}_3\text{O}_9\text{P}$, MW-726.94.



[0185] This was synthesized, according to **Standard procedure 5**, using BVdU (150 mg, 0.45 mmol), para-chlorophenyl-(benzoxy-L-leuciny)-phosphorochloridate (581 mg, 1.35 mmol.), NM1 (4.42 mmol, 190 μL) in THF (5 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 97:3 to give the pure product as a white foamy solid (221 mg, yield 68%).

$^{31}\text{P-NMR}$ (CDCl_3 , 121 MHz): δ 5.27, 4.76.

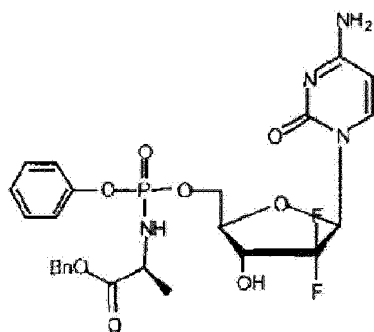
$^1\text{H-NMR}$ (CDCl_3 ; 300 MHz): δ 10.25-10.15 (1H, bs, H-3), 7.65 (1H, 2xs, H-6), 7.45 (1H, 2d, $^3J=14$ Hz, H-5b), 7.4-7.15 (9H, m, CH_2Ph + *OPh*), 6.7 (1H, 2d, $^3J=14$ Hz, H-5a), 6.35-6.2 (1H, 2t, $^3J=6$ Hz, H1'), 5.15 (2H, 2s, CH_2Ph) 4.55-3.9 (6H, m, H-5', H-3', NH, H-4',

$\text{CHCH}_2\text{CH}(\text{CH}_3)_2$, 2.5-2.4 (1H, m, one of H-2'), 2.15-2.0 (1H, m, one of H-2'), 1.7-1.45 (3H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 0.94-0.82 (6H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 23.1, 23.0, 22.2, 22.0 (2C, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 24.9, 24.7 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 40.7 (C-2'), 43.9, 43.8 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 53.9, 53.7 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 66.7, 66.3 (C-5'), 67.8 (CH_2Ph), 71.1, 70.7 (C-3'), 85.8, 85.7, 85.4 (C-1', C-4'), 110.5 (C-5b), 111.9 (C-5), 122.1, 122.0 ('o', OPh), 130.2, 129.1, 129.0 (C-5a, CH_2Ph , 'm' OPh), 131.1, 130.9 ('p', OPh), 135.5, 135.4 ('ipso', CH_2Ph), 138.2 (C-6), 149.2, 149.1 ('ipso', OPh), 149.2, 149.1 (C-4), 162.2 (C-2), 174.2, 174.2 (COOBn).

Synthesis of Gemcitabine-[phenyl-(benzoxy-L-alaninyl)]-phosphate.

[0186] $\text{C}_{25}\text{H}_{27}\text{F}_2\text{N}_4\text{O}_8\text{P}$, MW=580.47 (CPF 31).



[0187] This was synthesized according to **Standard procedure 5**, using gemcitabine (131 mg, 0.5 mmol), Phenyl-(benzoxy-L-alaninyl)-phosphorochloridate (529 mg, 1.5 mmol), NMI (4.42 mmol, 300 μL) in THF/pyridine (4/2 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 95:5 to give the pure product as a white foamy solid (46 mg, yield 16%).

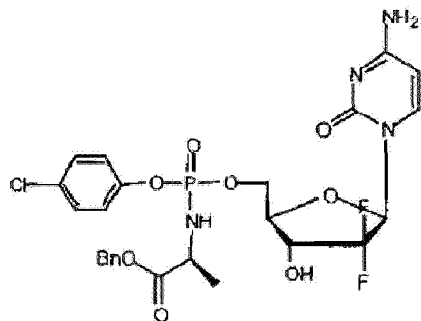
^{31}P -NMR (MeOD, 121 MHz): δ 5.05, 4.94.

^1H -NMR (MeOD, 300 MHz): δ 7.6-7.5 (1H, 2d, $^3J=7\text{Hz}$ H-6), 7.4-7.2 (10H, m, $\text{OPh}+\text{CH}_2\text{Ph}$), 6.25 (1H, m, H-1'), 5.95 (1H, 2d, $^3J=7\text{Hz}$, H-5), 5.19 (1H, 2s, CH_2Ph), 4.55-4.1 (3H, m, H-3', H-4', CHala), 4.05 (2H, m, H-5'), 1.20 (3H, 2t, $^3J=6\text{Hz}$, CH_3ala).

^{13}C -NMR (MeOD, 75 MHz): δ 20.8, 20.7 (CH_3ala), 52.2, 52.0 (CHala), 66.1 (C-5'), 68.4 (CH_2Ph), 71.9, 71.1 (C-3'), 80.6 (C-4'), 85.9 (C-1'), 97.1 (C-5), 121.8, 121.6 ('o', OPh), 123 (C-2'), 126.2 ('p', OPh), 131.8, 130.0, 129.7 ('m' OPh , Bn), 137.9 ('ipso', CH_2Ph), 142.7, 142.6 (C-6), 152.5, 152.4 ('ipso', OPh), 158.2 (C-2), 168.0 (C-4), 175.3, 174.9 (COOBn).

Synthesis of Gemcitabine-[para-chlorophenyl-(benzoxy-L-alaninyl)-phosphate,

[0188] $C_{25}H_{26}ClF_2N_4O_8P$, MW=614.92 (CPF 40).



[0189] This was synthesised according to **Standard procedure 5**, using gemcitabine (131 mg, 0.5 mmol), para-chlorophenyl-benzoxy-L-alaninyl)-phosphorochloridate (582 mg, 1.5 mmol), NMI (4.42 mmol, 300 μ L) in THF/pyridine (4/2 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 95:5 to give the pure product as a white foamy solid (76 mg, yield 25%),

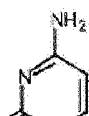
^{31}P -NMR (MeOD, 121 MHz): δ 5.08.

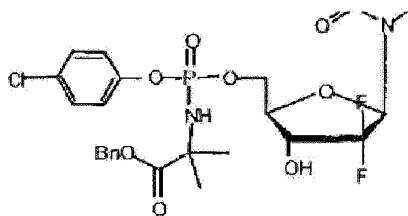
1H -NMR (MeOD, 300 MHz): δ 7.65 (1H, d, $^3J=7$ Hz H-6), 7.5-7.2 (9H, m, $O\text{Ph}+CH_2\text{Ph}$), 6.2 (1H, m, H-1'), 5.9 (1H, d, $^3J=7$ Hz, H-5), 5.12 (1H, 2s, $CH_2\text{Ph}$), 4.6-4.1 (3H, m, H-3', H-4', CHala), 4.05 (2H, m, H-5'), 1.45-1.35 (3H, 2t, $^3J=6$ Hz, $CH_3\text{ala}$).

^{13}C -NMR (MeOD, 75 MHz): δ 20.9, 20.7 ($CH_3\text{ala}$), 52.2, 52.0 ($\underline{CHala}$), 66.4, 66.2 (C-5'), 68.5 ($\underline{CH_2Ph}$), 71.5 (C-3'), 80.7 (C-4'), 86.4 (C-1'), 97.2 (C-5), 123.5 ('o', $O\text{Ph}$), 126.9 (C-2'), 131.2, 130.6, 130.3 ('m' OPh, Bn), 131.9 ('p', $O\text{Ph}$) 137.5 ('ipso', $CH_2\text{Ph}$), 142.8, 142.7 (C-6), 151.4, 151.0 ('ipso', $O\text{Ph}$), 158.2 (C-2), 166.9 (C-4), 175.1, 174.9 ($\underline{COOBn}$).

Synthesis of Gemcitabine-[para-chlorophenyl-(benzoxy- α,α -dimethylglycinyl)]-phosphate (CPF 41).

[0190] $C_{26}H_{28}ClF_2N_4O_8P$, MW=628.95.





[0191] This was synthesised according to **Standard procedure 5**, using gemcitabine (131 mg, 0.5 mmol), para-chlorophenyl-(benzoxy- α,α -dimethylglycyl)-phosphorochloridate (603 mg, 1.5 mmol), NM1 (4.42 mmol, 300 μ L) in THF/pyridine (4/3 mL) for 2 hrs. The crude product was purified by column chromatography, eluting with CH_2Cl_2 /Methanol 95:5 to give the pure product as a white foamy solid (163 mg, yield 52%).

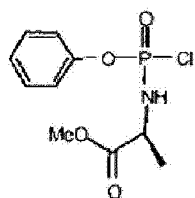
^{31}P -NMR (MeOD, 121 MHz): δ 3.56, 3.52.

^1H -NMR (MeOD, 300 MHz): δ 7.55 (1H, d, $^3J=7\text{Hz}$, H-6), 7.4-7.15 (9H, m, $\text{OPh}+\text{CH}_2\text{Ph}$), 6.25 (1H, m, H-1'), 5.85 (1H, d, $^3J=7\text{Hz}$, H-5), 5.15 (1R, 2s, CH_2Ph), 4.55-4.1 (3H, m, H-3', H-4'), 4.05 (2H, m, H-5'), 1.50 (6H, m, $^3J=6\text{ Hz}$, 2 CH_3 dimethygly).

^{13}C -NMR (MeOD, 75 MHz): δ 28.2, 28.0 (CH_3 dimethygly), 58.6 (Cq dimethygly), 66.2, 66.1 (C-5'), 66.7 (CH_2Ph), 71.5 (C-3'), 80.6 (C-4'), 86.4 (C-1'), 97.0 (C-5), 123.9, 123.6 ('o', OPh), 127.3 (C-2'), 130.0, 129.7 ('m' OPh , Bn), 131.8 ('p', OPh), 137.6 ('ipso', CH_2Ph), 142.8, 142.7 (C-6), 151.2, 151.1 ('ipso', OPh), 158.1 (C-2), 167.9 (C-4), 176.8, 176.7 (COOBn),

Synthesis of Phenyl-(methoxy-L-alaninyl)-phosphorochloridate.

[0192] $\text{C}_{10}\text{H}_{13}\text{ClNO}_4\text{P}$, MW-277.64.



[0193] This is synthesised according to **Standard procedure 4**, using L-alanine methyl ester hydrochloride (2 g, 14.3 mmol), phenyldichlorophosphate (3.02 g, 2.14 ml, 14.3 mmol), and TEA (2.9 g, 4.0 ml, 28.7 mmol) in DCM (60 mL), to yield 3.91 g (98%) of crude product used without further purification.

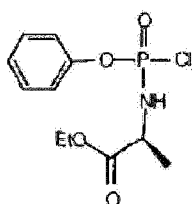
^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.28, 8.97.

$^1\text{H-NMR}$ (CDCl_3 ; 300 MHz): δ 7.39-7.34 (2H, m, 'o' OPh), 7.29-7.20 (2H, m, 'm+p' OPh), 4.98 (1H, bs, NH), 4.27-4.09 (1H, m, CHala), 3.78 (3H, s, OCH_3), 1.52-1.49 (3H, 2xd, $^3J=7\text{Hz}$, CH_3ala).

$^{13}\text{C-NMR}$ (CDCl_3 ; 75 MHz): δ 20.9 (CH_3ala), 51.0 (CHala), 53.6 (OCH_3), 120.9 ('o' OPh), 126.4 ('p', OPh), 130.2 ('m', OPh), 150.1 ('ipso', OPh), 173.6 (COOCH_3).

Synthesis of Phenyl-(ethoxy-L-alaninyl)-phosphorochloridate

[0194] $\text{C}_{11}\text{H}_{15}\text{ClNO}_4\text{P}$, MW-291.67.



[0195] This is synthesised according to **Standard procedure 4**, using L-alanine ethyl ester hydrochloride (770 mg, 5.01 mmol), phenyldichlorophosphate (1.12g, 5.01 mmol, 749 μL), and TEA (1.4 mL, 10.02 mmol) in DCM (40 mL). The crude was purified by flash chromatography (ethyl acetate/petroleum ether 7:3) affording 1.02 (69%) of oil.

$^{31}\text{P-NMR}$ (CDCl_3 , 121 MHz): δ 9.49, 9.07.

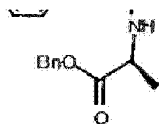
$^1\text{H-NMR}$ (CDCl_3 ; 300 MHz): δ 7.39-7.34 (2H, m, 'o' OPh), 7.29-7.20 (2H, m, 'm+p' OPh), 4.95 (1H, bs, NH), 4.3-4.1 (3H, m, OCH_2CH_3 , CHala), 1.50 (3H, 2xd, $^3J=7\text{Hz}$, CH_3ala), 1.30 (3H, t, $^3J=7.1\text{ Hz}$, $\text{OCH}_2\text{-CH}_3$).

$^{13}\text{C-NMR}$ (CDCl_3 ; 75 MHz): δ 14.5 (CH_3CH_2), 20.9 (CH_3ala), 51.0 (CHala), 62.6 (CH_3CH_2), 120.9 ('o' OPh), 126.5 ('p', OPh), 130.1 ('m', OPh), 150.1 ('ipso', OPh), 175.1 ($\text{COOCH}_2\text{CH}_3$).

Synthesis of Phenyl-(benzoxy-L-alaninyl)-phosphorochloridate.

[0196] $\text{C}_{16}\text{H}_{17}\text{ClNO}_4\text{P}$, MW= 353.74.





[0197] This is synthesised according to **Standard procedure 4**, using L-alanine benzyl ester hydrochloride (1.0 g, 4.64 mmol), phenyl-dichlorophosphate (980 mg, 0.69 ml, 4.64 mmol), and TEA (0.94 g, 1290 μ L, 9.27 mmol) in DCM (40 mL). The crude was purified by flash chromatography (ethyl acetate/petroleum ether 6:4) affording 1.61 (98%) of oil.

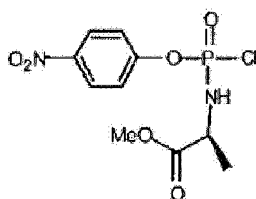
^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.41, 9.23.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.41-7.21 (10H, m, $\text{OPh}+\text{CH}_2\text{Ph}$), 5.24 (2H, s, CH_2Ph), 4.95-4.88 (1H, bs, NH), 4.36-4.15 (1H, m, CHala), 1.52-1.49 (3H, 2xd, $^3J=7\text{Hz}$, CH_3ala).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 20.8 (CH_3ala), 51.1 (CHala), 68.0 (CH_2Ph), 121.0 ('o' OPh), 126.4 ('p', OPh), 130.3, 129.0, 128.7 ('m'OPh, CH_2Ph), 135.5 ('ipso', CH_2Ph), 150.2 ('ipso' OPh), 172.9 (COOCH_2Ph).

Synthesis of p-nitrophenyl-(methoxy-L-alaninyl)-phosphorochloridate.

[0198] $\text{C}_{10}\text{H}_{12}\text{ClN}_2\text{O}_6\text{P}$, MW=322.64.



[0199] This is synthesised according to **Standard procedure 4**, using L-alanine methyl ester hydrochloride (0.70 g, 5.01 mmol), p-nitrophenyldichlorophosphate (1.362 g, 5.01 mmol), and TEA (1.4 ml, 10 mmol) in DCM (40 mL), to yield 1.60 g (99%) of crude product used without further purification.

^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.13, 9.03.

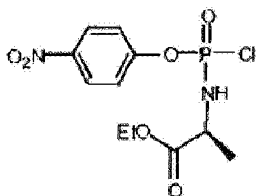
^1H -NMR (CDCl_3 ; 300 MHz): δ 8.1 (2H, 2d, $^3J=8\text{Hz}$, OPh), 7.3 (2H, 2d, $^3J=8\text{Hz}$, OPh), 5.0 (1H,bs, NH), 4.1 (1H, m, CHala), 3.75 (3H, s, OCH_3), 1.5-1.45 (3H, m, CH_3ala).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 20.8, 20.7 (CH_3ala), 51.1, 50.9 (CHala), 53.2, 53.2 (OCH_3),

121.8, 121.6 ('*o*' OPh), 126.5 ('*m*', OPh), 145.7 ('*ipso*', OPh), 154.7, 154.6 ('*p*', OPh), 173.4, 173.2 ($\underline{\text{C}}\text{OOCH}_3$).

Synthesis of p-nitrophenyl-(ethoxy-L-alaninyl)phosphorochloridate.

[0200] $\text{C}_{11}\text{H}_{14}\text{ClN}_2\text{O}_6\text{P}$, MW=336.67.



[0201] This is synthesised according to **Standard procedure 4**, using L-alanine ethyl ester hydrochloride (770 mg, 5.01 mmol), p-nitrophenyldichlorophosphate (1.362g, 5.01 mmol), and TEA (1.4 mL, 10.02 mmol) in DCM (40 mL), to yield 1.64 g (98%) of crude product used without further purification.

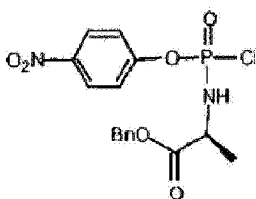
^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.06, 8.81.

^1H -NMR (CDCl_3 ; 300 MHz): δ 8.1 (2H, m, $\underline{\text{OPh}}$), 7.4 (2H, m, $\underline{\text{OPh}}$), 4.9-4.7 (1H, bs, $\underline{\text{NH}}$), 4.3-4.1 (3H, m, $\underline{\text{OCH}_2\text{CH}_3}$, $\underline{\text{CHala}}$), 1.55-1.45 (3H, 2xd, $^3J=7\text{Hz}$, $\underline{\text{CH}_3\text{ala}}$), 1.40 (3H, t, $^3J=7\text{Hz}$, $\underline{\text{OCH}_2\text{CH}_3}$).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 14.5 ($\underline{\text{CH}_3\text{CH}_2}$), 21.1, 20.9 ($\underline{\text{CH}_3\text{ala}}$), 51.2, 51.0 ($\underline{\text{CHala}}$), 62.6 ($\underline{\text{CH}_3\text{CH}_2}$), 121.7, 121.3 ('*o*' OPh), 126.2, 126.0 ('*m*', OPh), 145.7 ('*ipso*', OPh), 154.5 ('*p*', OPh), 173.4, 173.3 ($\underline{\text{C}}\text{OOCH}_2\text{CH}_3$).

Synthesis of p-nitrophenyl-(benzoxy-L-alaninyl)-phosphorochloridate.

[0202] $\text{C}_{16}\text{H}_{16}\text{ClN}_2\text{O}_6\text{P}$, MW= 398.04.



[0203] This is synthesised according to **Standard procedure 4**, using L-alanine benzyl ester hydrochloride (1.08 g, 5.01 mmol), para-nitrophenyl-dichloro phosphate (1.362 g, 5.01 mmol), and TEA (1.4 mL, 1.4 mmol) in DCM (40 mL), to yield 1.85 g (93%) of crude product used without further purification.

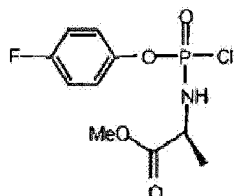
^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.15, 9.06.

^1H -NMR (CDCl_3 ; 300 MHz): δ 8.15 (2H, m, OPh), 7.45 (2H, m, OPh), 7.35-7.25 (5H, m, CH_2Ph), 5.2 (2H, 2s, CH_2Ph), 5.00 (1H, bs, NH), 4.2 (1H, m, CHala), 1.64 (3H, 2xd, $^3J=7\text{Hz}$, CH_3ala).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 20.8 (CH_3ala), 51.1 (CHala), 68.0 (CH_2Ph), 121.4 ('o' OPh), 126.1 ('m' OPh), 130.3, 129.0 (CH_2Ph), 145.7 ('ipso', CH_2Ph), 150.2 ('ipso', OPh), 154.6 ('p', OPh), 172.9 (COOCH_2Ph).

Synthesis of p-fluorophenyl-(methoxy-L-alaninyl)-phosphorochloridate.

[0204] $\text{C}_{10}\text{H}_{12}\text{ClFNO}_4\text{P}$, MW=295.63.

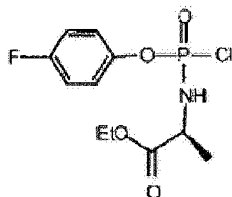


[0205] This is synthesised according to **Standard procedure 4**, using L-alanine methyl ester hydrochloride (0.70 g, 5.01 mmol), p-fluorophenyldichlorophosphate (1.210 g, 5.01 mmol), and TEA (1.4 mL, 10 mmol) in DCM (40 mL). The crude was purified by flash chromatography (ethyl acetate/petroleum ether 7:3) affording 1.11 g (75%) of oil.

^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.98, 9.96.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.1 (2H, m, OPh), 6.95 (2H, m, OPh), 5.0 (1H, bs, NH), 4.25-4.1 (1H, m, CHala), 3.78 (3H, 2s, OCH_3), 1.55 (3H, m, CH_3ala).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 20.8 (CH_3ala), 51.1, 50.9 (CHala), 53.3 (OCH_3), 117.1, 117.0 ('o' OPh), 122.6, 122.5 ('m', OPh), 146.0 ('ipso', OPh), 159.1, 159.0 ('p', OPh), 173.4, 173.2 (COOCH_3).

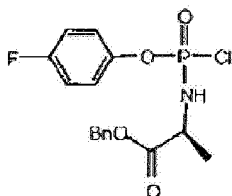
Synthesis of p-fluorophenyl-(ethoxy-L-alaninyl)-phosphorochloridate.**[0206]** $C_{11}H_{14}ClFNO_4P$, MW=309.66.

[0207] This is synthesised according to **Standard procedure 4**, using L-alanine ethyl ester hydrochloride (770 mg, 5.01 mmol), p-fluorophenyldichlorophosphate (1.210g, 5.01 mmol), and TEA (1.4 mL, 10.02 mmol) in DCM (40 ml). The crude was purified by flash chromatography (ethyl acetate/petroleum ether 7:3) affording 1.07 (69%) of oil.

^{31}P -NMR ($CDCl_3$, 121 MHz): δ 10.01, 9.95.

1H -NMR ($CDCl_3$; 300 MHz): δ 7.1 (2H, m, $O\text{Ph}$), 6.95 (2H, m, $O\text{Ph}$), 5.0 (1H, bs, NH), 4.25-4.1 (3H, m, OCH_2CH_3 , CH_{Hala}), 1.55 (3H, m, CH_3_{ala}), 1.40 (3H, t, $^3J=7\text{Hz}$, OCH_2CH_3).

^{13}C -NMR ($CDCl_3$; 75 MHz): δ 14.5 (CH_3CH_2), 21.1, 21.0 (CH_3_{ala}), 51.2, 51.1 (CH_{Hala}), 62.6 (CH_3CH_2), 117.3 ('o' $O\text{Ph}$), 122.2, 122.0 ('m', $O\text{Ph}$), 145.9, 145.8 ('ipso', $O\text{Ph}$), 159.0 ('p', $O\text{Ph}$), 173.6, 173.5 ($COOCH_2CH_3$).

Synthesis of p-fluorophenyl-(benzoxy-L-alaninyl)-phosphorochloridate.**[0208]** $C_{16}H_{16}ClFNO_4P$, MW=371.73.

[0209] This is synthesized according to **Standard procedure 4**, using L-alanine benzyl ester hydrochloride (1.08 g, 5.01 mmol), para-fluorophenyl-dichloro phosphate (1,210 mg, 5.01 mmol), and TEA (1.4mL, 1.4 mmol) in DCM (40 mL). The crude was purified by flash chromatography (ethyl acetate/petroleum ether 7:3) affording 1.599 (86%) of oil.

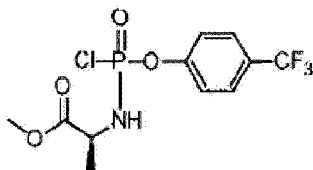
^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.15, 9.06.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.35-7.25 (5H, m, CH_2Ph), 7.1 (2H, m, OPh), 6.95 (2H, m, OPh), 5.2 (2H, 2s, CH_2Ph), 5.00 (1H, bs, NH), 4.25-4.1 (1H, m, CHala), 1.55 (3H, m, CH_3ala).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 20.8 (CH_3ala), 51.1, 51.0 (CHala), 68.1 (CH_2Ph), 117.0, 116.9 ('o' OPh), 122.6 ('m' OPh), 130.3, 129.0 (CH_2Ph), 135.7 ('ipso', CH_2Ph), 146.1, 146.0 ('ipso', OPh), 158.9 ('p', OPh), 173.1 (COOCH_2Ph).

Synthesis of 4-(trifluoromethyl)-phenyl-(methoxy-L-alaninyl)-phosphorochloridate.

[0210] $\text{C}_{11}\text{H}_{12}\text{ClF}_3\text{NO}_4\text{P}$, MW=345.64.



[0211] This is synthesised according to **Standard procedure 4**, using L-alanine methyl ester hydrochloride (1.0 g, 7.16 mmol), 4-(trifluoromethyl)-phenyl-phosphodichloridate (1.998 g, 7.16 mmol), and TEA (1.449 g, 14.32 mmol, 1916 μL) in DCM (30 mL), to yield 2.202 g (89.0%) of crude product used without further purification.

^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.36, 9.22.

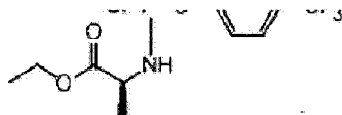
^1H -NMR (CDCl_3 ; 300 MHz): δ 7.66 (2H, d, $^3J=8.1$ Hz, OPh), 7.44-7.33 (2H, m, OPh), 5.10 (1H, bs, NH), 3.81-3.78 (3H, 2s, CH_3O), 3.77-3.68 (1H, m, CH_3CH), 1.56-1.52 (3H, m, CHCH_3).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 20.6, 20.7 (CH_3CH), 50.9, 51.1 (CHCH_3), 53.2 (CH_3O), 121.4 ('o', OPh), 124.1 (CF_3 , $J=270$ Hz), 128.0 ('m', OPh), 128.6 ('p', $J=34$ Hz), 152.4, 152.6 ('ipso', OPh), 173.4, 173.5 (COOCH_3).

Synthesis of 4-(trifluoromethyl)-phenyl-(ethoxy-L-alaninyl)phosphorochloridate.

[0212] $\text{C}_{12}\text{H}_{14}\text{ClF}_3\text{NO}_4$, MW-359.67.





[0213] This is synthesised according to **Standard procedure 4**, using L-alanine ethyl ester hydrochloride (1.0 g, 6.50 mmol), 4-(trifluoromethyl)-phenyl-phosphodichloridate (1.813 g, 6.50 mmol), and TEA (1.316 g, 13.00 mmol, 1740 μ L) in DCM (30 mL), to yield 2.150 g (92.2%) of crude product used without further purification.

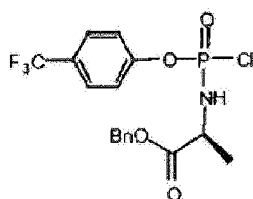
^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.33, 9.28.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.70 (2H, d, $^3J=8.2$ Hz, *OPh*), 7.46-7.39 (2H, m, *OPh*), 4.78 (1H, bs, *NH*), 4.33-4.17 (3H, m, $\text{CH}_3\text{CH}_2\text{O} + \text{CHCH}_3$), 1.59-1.55 (1H, m, CHCH_3), 1.56-1.52 (3H, m, CH_2CH_3).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 14.5 ($\text{CH}_3\text{CH}_2\text{O}$), 20.8, 20.9 (CH_3CH), 50.3, 50.9 (CHCH_3), 62.3, 62.5 ($\text{CH}_3\text{CH}_2\text{O}$), 121.4 ('*o*', *OPh*), 124.1 (*CF*₃, $J=270$ Hz), 127.7 ('*m*', *OPh*), 128.7 ('*p*', $J=33$ Hz), 152.4 ('*ipso*', *OPh*), 172.9 ($\text{COOCH}_2\text{CH}_3$).

Synthesis of p-trifluorophenyl-(benzoyl-L-alaninyl)-phosphorochloridate.

[0214] $\text{C}_{17}\text{H}_{16}\text{ClF}_3\text{NO}_4\text{P}$, MW= 421.73.



[0215] This is synthesized according to **Standard procedure 4**, using L-alanine benzyl ester hydrochloride (1.08 g, 5.01 mmol), para-trifluorophenyl-dichloro phosphate (1.490 mg, 5.01 mmol), and TEA (1.4 mL, 1.4 mmol) in DCM (40 mL). The crude was purified by flash chromatography (ethyl acetate/petroleum ether 6:4) affording 1.80 (85%) of oil.

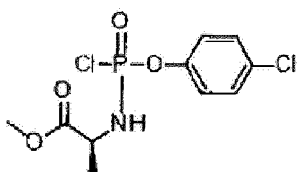
^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.11, 8.84.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.65 (2H, m, *OPh*), 7.4-7.2 (7H, m, $\text{CH}_2\text{Ph} + 2\text{H } \text{OPh}$), 5.25 (2H, 2s, CH_2Ph), 4.75-4.55 (1H, bs, *NH*), 4.25-4.1 (1H, m, CHala), 1.60-1.55 (3H, 2d, $^3J=7\text{Hz}$, CH_3ala).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 20.9 ($\underline{\text{CH}}_3\text{ala}$), 51.3, 51.0 ($\underline{\text{CH}}\text{ala}$), 68.2, 68.1 ($\underline{\text{CH}}_2\text{Ph}$), 121.4, 120.9 ('o', OPh), 125.2 (d, $J=270\text{Hz}$, CF_3), 126.6 ('m', OPh), 129.1, 128.8, 127.8 (Bn), 130.0 ('p', q, $J=32\text{Hz}$, OPh), 135.4 ('ipso', CH_2Ph), 153.0 ('ipso', OPh), 172.8 ($\underline{\text{COOCH}}_2\text{Ph}$).

Synthesis of 4-chlorophenyl-(methoxy-L-alaninyl)-phosphorochloridate.

[0216] $\text{C}_{10}\text{H}_{12}\text{Cl}_2\text{NO}_4\text{P}$, MW-312.09.



[0217] This is synthesised according to **Standard procedure 4**, using L-alanine methyl ester hydrochloride (1.0 g, 7.16 mmol), 4-chlorophenylphosphorodichloridate (1.757 g, 7.16 mmol), and TEA (1.449 g, 14.32 mmol, 1995 μL) in DCM (30 mL), to yield 1.621 g (72,5%) of crude product used without further purification.

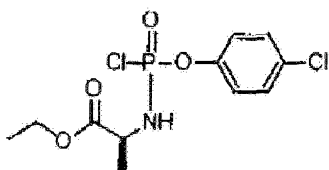
^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.36, 9.07.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.35-7.15 (4H, m, OPh), 4.48-4.36 (1H, bs, NH), 4.12-4.04 (1H, m, $\underline{\text{CHCH}}_3$), 3.76-3.74 (3H, 2s, OH_3O), 1.49-1.46 (3H, m, CHCH_3).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 21.0 ($\underline{\text{CH}}_3\text{CH}$), 50.8, 51.1 ($\underline{\text{CHCH}}_3$), 53.4 ($\underline{\text{CH}}_3\text{O}$), 121.9, 122.1, 122.1, 122.4 ('o', OPh), 130.6, 130.4, 130.2 ('m', OPh), 132.0 ('p', OPh), 148.6 ('ipso', OPh), 173.5 ($\underline{\text{COOCH}}_3$).

Synthesis of 4-chlorophenyl-(ethoxy-L-alaninyl)-phosphorochloridate.

[0218] $\text{C}_{11}\text{H}_{14}\text{Cl}_2\text{NO}_4\text{P}$, MW=326.11.



[0219] This is synthesized according to **Standard procedure 4**, using L-alanine ethyl ester hydrochloride (1.000 g, 6.50 mmol), 4-chlorophenylphosphorodichloride (1.595 g, 6.50 mmol), and TEA (1.315 g, 13.00 mmol, 1810 μ L) in DCM (20 mL), to yield 1.794 mg (yield 84.7%) of product.

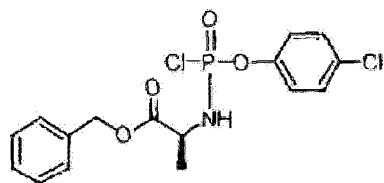
^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.54, 9.25.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.44-7.21 (4H, m, OPh), 4.59 (1H, bs, NH), 4.33-4.13 (3H, m, $\text{OCH}_2\text{CH}_3 + \text{CHCH}_3$), 1.57-1.56 (3H, m, CH_3CH), 1.43-1.21 (3H, m, OCH_2CH_3).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 14.5, 14.6 (OCH_2CH_3), 21.0, 21.5 (CH_3CH), 50.9, 51.2 (CHCH_3), 62.4, 62.5 (OCH_2CH_3), 122.04, 122.3, 122.4 ('o', OPh), 130.4 ('m', OPh), 131.9 ('p', OPh), 148.5, 148.6 ('ipso', OPh), 173.0, 173.1 ($\text{COOCH}_2\text{CH}_3$).

Synthesis of 4-nitrophenyl-(benzyl-2-amino-2-methylpropanoate)-phosphorochloridate.

[0220] $\text{C}_{16}\text{H}_{16}\text{Cl}_2\text{NO}_4\text{P}$, MW=388.18.



[0221] This is synthesized according to **Standard procedure 4**, using L-alanine benzyl ester hydrochloride (1.000 g, 4.63 mmol), 4-chlorophenylphosphodichloride (1.136 g, 4.63 mmol), and TEA (937.0 mg, 9.26 mmol, 1290 μ L) in DCM (40 mL), to yield 1534 mg (yield 86.5%) of crude product used without further purification.

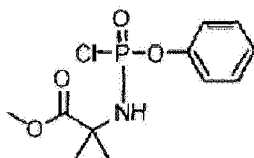
^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.43, 9.16.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.42-7.08 (9H, m, $\text{OPh} + \text{CH}_2\text{Ph}$), 5.19 (2H, s, CH_2Ph), 4.61-4.54 (1H, bs, NH), 4.26-4.10 (1H, m, CHCH_3), 1.42-1.38 (3H, m, CH_3CH).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 20.9, 21.0 (CH_3CH), 51.0, 51.2 (CHCH_3), 68.1, 68.2 (OCH_2Ph), 122.3, 122.4 ('o', OPh), 128.8, 129.1, 130.4 ('o', 'm', 'p', $\text{CH}_2\text{Ph} + \text{OPh}$), 131.9 ('ipso', CH_2Ph), 135.3 ('p', OPh), 148.5 ('ipso', OPh), 172.7, 172.8 (COOCH_2Ph).

Synthesis of phenyl-(methyl-2-amino-2-methylpropanoate)-phosphorochloridate.

[0222] $C_{11}H_{15}ClNO_4P$, MW=291.67.



[0223] This is synthesised according to **Standard procedure 4**, using 2-aminoisobutyrate methyl ester hydrochloride (583.5 mg, 3.75 mmol), phenyl dichlorophosphate (791.1 mg, 3.75, 560 μ L), and TEA (758.9 mg, 7.5 mmol, 1045 μ L) in DCM (20 mL), to yield 1.041 g (95.2%) of crude product used without further purification.

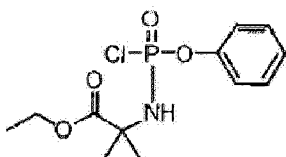
^{31}P -NMR ($CDCl_3$, 121 MHz): δ 6.99 (s).

1H -NMR ($CDCl_3$; 300 MHz): δ 7.41-7.17 (5H, m, $O\text{Ph}$), 4.98 (1H, bs, NH), 3.80 (3H, s, OCH_3), 1.71-1.69 (6H, 2s, $[CH_3]_2C$).

^{13}C -NMR ($CDCl_3$; 75 MHz): δ 27.3, 27.2, 27.0 ($[CH_3]_2C$), 53.6 (OCH_3), 58.8 ($C[CH_3]_2$), 120.0, 121.1 ('*o*' $O\text{Ph}$), 126.2 ('*p*', $O\text{Ph}$), 130.3 ('*m*', $O\text{Ph}$), 145.7 ('*p*', $O\text{Ph}$), 150.2, 150.3 ('*ipso*', $O\text{Ph}$), 175.6, 175.7 ($COOCH_3$).

Synthesis of phenyl-(ethyl-2-amino-2-methylpropanoate)-phosphorochloridate.

[0224] $C_{12}H_{17}ClNO_4P$, MW=305.69.



[0225] This is synthesised according to **Standard procedure 4**, using 2-aminoisobutyrate ethyl ester hydrochloride (628.6 mg, 3.75 mmol), phenyl dichlorophosphate (791.1 mg, 3.75, 560 μ L), and TEA (758.9 mg, 7.5 mmol, 1045 μ L) in DCM (20 mL), to yield 1.018 g (88.8%) of crude product used without further purification,

^{31}P -NMR ($CDCl_3$, 121 MHz): δ 7.02 (s)

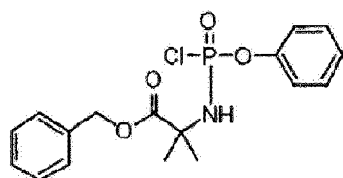
1H -NMR ($CDCl_3$; 300 MHz): δ 7.23-7.37 (5H, m, $O\text{Ph}$), 4.98 (1H, bs, NH), 4.24 (2H, q, $^3J=7.1$

Hz, OCH_2CH_3), 1.70, 1.68 (6H, 2s, $[\text{CH}_3]_2\text{C}$), 1.30 (3H, t, $^3J=7.1$ Hz, OCH_2CH_3).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 14.5 ($\text{CH}_3\text{CH}_2\text{O}$), 27.3, 26.9 ($[\text{CH}_3]_2\text{C}$), 58.7 ($\text{C}[\text{CH}_3]_2$), 62.7 (OCH_2CH_3), 121.1, 121.0 ('o', OPh), 127.6 ('p', OPh), 130.7 ('m', OPh), 150.4 ('ipso', OPh), 175.2, 175.1 ($\text{COOCH}_2\text{CH}_3$).

Synthesis of phenyl-(benzyl-2-amino-2-methylpropanoate)-phosphorochloridate.

[0226] $\text{C}_{17}\text{H}_{19}\text{ClINO}_4\text{P}$, MW=367.76.



[0227] This is synthesised according to **Standard procedure 4**, using 2-aminoisobutyrate benzyl ester hydrochloride (861.4 mg, 3.75 mmol), phenyl dichlorophosphate (791.1 mg, 3.75, 560 μL), and TEA (758.9 mg, 7.5 mmol, 1045 μL) in DCM (30 mL). The crude was purified by flash chromatography (ethyl acetate/petroleum ether 6:4) affording 580 mg (42.2%) of oil.

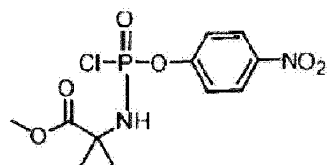
^{31}P -NMR (CDCl_3 , 121 MHz): δ 6.79 (s)

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.45-7.27 (10H, m, OPh+CH₂Ph), 5.28 (2H, s, CH₂Ph), 4.81, 4.78 (1H, 2bs, NH), 1.78, 1.75 (6H, 2s, $[\text{CH}_3]_2\text{C}$).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 27.3, 26.9 ($[\text{CH}_3]_2\text{C}$), 53.9 ($\text{C}[\text{CH}_3]_2$); 60.9 (CH₂Ph), 121.0, 126.3, 129.6, 129.0, 129.1, 130.3, 135.5 (OPh, CH₂Ph), 135.5 ('ipso', CH₂Ph), 150.3, 150.2 ('ipso', OPh), 175.0, 175.2 (COOCH_2Ph).

Synthesis of 4-nitrophenyl-(methyl-2-amino-2-methylprepanoate)-phosphorochloridate.

[0228] $\text{C}_{11}\text{H}_{14}\text{ClN}_2\text{O}_6\text{P}$, MW=336.67.



[0229] This is synthesised according to **Standard procedure 4**, using 2-aminoisobutyrate methyl ester hydrochloride (290.0mg, 1.89 mmol), 4-nitrophenylphosphodichloride (483.3 mg, 1.89 mmol), and TEA (382.5 mg, 3.78 mmol, 526.9 μ L) in DCM (15 mL), to yield 486 mg (yield 76.4%) of crude product used without further purification.

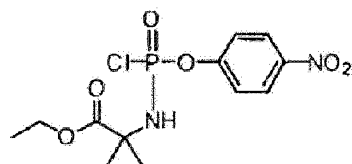
^{31}P -NMR (CDCl_3 , 121 MHz): δ 6.61 (s)

^1H -NMR (CDCl_3 ; 300 MHz): δ 8.25 (2H, d, $^3J=9.0$ Hz, $\text{O}Ph$), 7.43 (2H, d, $^3J=9.0$ Hz, $\text{O}Ph$), 4.91-4.87 (1H, 2bs, NH), 3.79 (3H, s, OCH_3), 1.69-1.66 (6H, 2s, $[\text{CH}_3]_2\text{C}$).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 27.0, 27.1, 27.3 ($[\text{CH}_3]_2\text{C}$), 53.8 (OCH_3), 59.2 ($\text{C}[\text{CH}_3]_2$), 121.7, 121.8 ('o' $\text{O}Ph$), 126.2 ('m', $\text{O}Ph$), 145.7 ('p', $\text{O}Ph$), 154.8, 154.7 ('ipso', $\text{O}Ph$), 175.4, 175.6 (COOCH_3).

Synthesis of 4-nitrophenyl-(ethyl-2-amino-2-methylpropanoate)-phosphorochloridate.

[0230] $\text{C}_{12}\text{H}_{16}\text{ClN}_2\text{O}_6\text{P}$, MW=350.69.



[0231] This is synthesised according to **Standard procedure 4**, using 2-aminoisobutyrate ethyl ester hydrochloride (270.0 mg, 1.61 mmol), 4-nitrophenylphosphodichloride (412.3 mg, 1.61 mmol), and TEA (325.8 mg, 3.22 mmol, 448.8 μ L) in DCM (15 mL), to yield 500 mg (yield 88.5%) of crude product used without further purification.

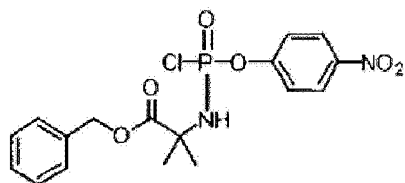
^{31}P -NMR (CDCl_3 , 121 MHz): δ 6.64 (s)

^1H -NMR (CDCl_3 ; 300 MHz): δ 8.35 (2H, d, $^3J=9.0$ Hz, $\text{O}Ph$), 7.53 (2H, d, $^3J=9.0$ Hz, $\text{O}Ph$), 4.99-4.96 (1H, 2bs, NH), 4.34 (2H, q, $^3J=7.1$ Hz, OCH_2CH_3), 1.79-1.76 (6H, 2s, $[\text{CH}_3]_2\text{C}$), 1.40 (3H, t, $^3J=7.1$ Hz, OCH_2CH_3).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 14.5 (OCH_2CH_3), 27.0, 27.3 ($[\text{CH}_3]_2\text{C}$), 59.1, 59.2 ($\text{C}[\text{CH}_3]_2$), 62.9, 63.0 (OCH_2CH_3), 121.7, 121.8 ('o' $\text{O}Ph$), 126.2 ('m', $\text{O}Ph$), 145.7 ('p', $\text{O}Ph$), 154.7, 154.8 ('ipso', $\text{O}Ph$), 175.4, 175.6 ($\text{COOCH}_2\text{CH}_3$).

Synthesis of 4-nitrophenyl-(benzyl-2-amino-2-methylpropanoate)-phosphorochloridate.

[0232] $C_{17}H_{18}ClN_2O_6P$, MW=412.76.



[0233] This is synthesised according to **Standard procedure 4**, using 2-aminoisobutyrate benzyl ester hydrochloride (578 mg, 2.52 mmol), 4-nitrophenylphosphodichloride (645 mg, 2.52 mmol), and TEA (510 mg, 5.04 mmol, 702.5 μ L) in DCM (20 mL), to yield 936 mg (yield 90.0%) of crude product used without further purification.

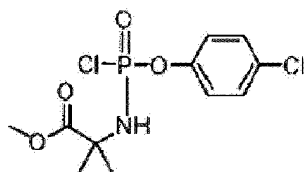
^{31}P -NMR ($CDCl_3$, 121 MHz): δ 6.56 (s)

1H -NMR ($CDCl_3$; 300 MHz): δ 8.29 (2H, d, $^3J=9.0$ Hz, $O\text{Ph}$), 7.47 (2H, d, $^3J=9.0$ Hz, $O\text{Ph}$), 7.40-7.37 (5H, m, $CH_2\text{Ph}$), 5.27 (2H, s, $CH_2\text{Ph}$), 5.04-5.01 (1H, 2bs, NH), 1.77-1.74 (6H, 2s, $[CH_3]_2C$).

^{13}C -NMR ($CDCl_3$; 75 MHz): δ 27.0, 27.3, ($[CH_3]_2C$), 59.2 ($C[CH_3]_2$), 68.5 (OCH_2Ph), 121.6, 121.7, 126.2, 128.6, 129.1, ('o', 'm', 'p', $CH_2Ph + OPh$), 135.7 ('ipso', CH_2Ph), 145.7 ('p', OPh), 154.7, 154.8 ('ipso', OPh), 175.8, 175.9 ($COOCH_2Ph$).

Synthesis of 4-chlorophenyl-(methyl-2-amino-2-methylpropanoate)-phosphorochloridate.

[0234] $C_{11}H_{14}Cl_2NO_4P$, MW=326.11



[0235] This is synthesized according to **Standard procedure 4**, using 2-aminoisobutyrate methyl ester hydrochloride (280.0 mg, 1.82 mmol), 4-chlorophenylphosphodichloride (447.4 mg, 1.82 mmol), and TEA (368.3 mg, 3.64 mmol, 507.3 μ L) in DCM (20 mL), to yield 554 mg

(yield 91.1%) of crude product used without further purification.

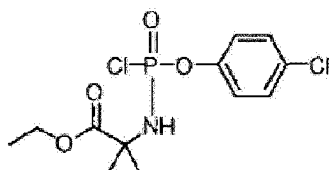
^{31}P -NMR (CDCl_3 , 121 MHz): δ 7.05 (s)

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.38 (2H, d, $^3J=9.0$ Hz, *OPh*), 7.28-7.24 (2H, 2d, $^3J=9.0$ Hz, *OPh*), 4.87-4.83 (1H, 2bs, *NH*), 3.84 (3H, s, *OCH₃*), 1.73.1-71 (6H, 2s, [*CH₃*]₂C).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 27.0, 27.3, ([*CH₃*]₂C), 53.7 (*OCH₃*), 58.9 (*C[CH₃]₂*), 122.5 ('*o*', *OPh*), 129.7 ('*m*', *OPh*), 131.8 ('*p*', *OPh*) 148.7, 148.9 ('*ipso*', *OPh*), 175.5, 175.7 (*COOCH₃*).

Synthesis of 4-chlorophenyl-(ethyl-2-amino-2-methylpropanoate)-phosphorochloridate.

[0236] $\text{C}_{12}\text{H}_{16}\text{Cl}_2\text{NO}_4\text{P}$, MW=340.14.



[0237] This is synthesised according to **Standard procedure 4**, using 2-aminoisobutyrate ethyl ester hydrochloride (293.4 mg, 1.75 mmol), 4-chlorophenylphosphodichloride (430.0 mg, 1.75 mmol), and TEA (354.2 mg, 3.50 mmol, 488.0 μL) in DCM (15 mL), to yield 571.7 mg (yield 96.1%) of crude product used without further purification.

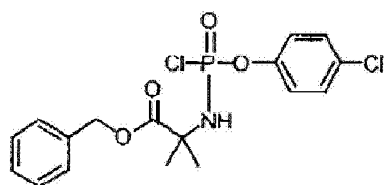
^{31}P -NMR (CDCl_3 , 121 MHz): δ 7.09 (s)

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.38 (2H, d, $^3J=9.1$ Hz, *OPh*), 7.26 (2H, d, $^3J=9.1$ Hz, *OPh*), 4.88-4.84 (1H, 2bs, *NH*), 4.29 (2H, q, $^3J=7.1$ Hz, *OCH₂CH₃*), 1.74-1.70 (6H, 2s, [*CH₃*]₂C), 1.35 (3H, t, $^3J=7.1$ Hz, *OCH₂CH₃*).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 14.5 (*OCH₂CH₃*), 27.0, 27.3 ([*CH₃*]₂C), 58.9 (*C[CH₃]₂*), 62.8 (*OCH₂CH₃*), 122.5 ('*o*', *OPh*), 130.4 ('*m*', *OPh*), 131.8 ('*p*', *OPh*), 148.7, 148.8 ('*ipso*', *OPh*), 175.1, 175.3 (*COOCH₂CH₃*).

Synthesis of 4-chlorophenyl-(benzyl-2-amino-2-methylpropanoate)-phosphorochloridate.

[0238] $C_{17}H_{18}Cl_2NO_4P$, MW=402.21.



[0239] This is synthesised according to **Standard procedure 4**, using 2-aminoisobutyrate benzyl ester hydrochloride (402.0 mg, 1.75 mmol), 4-chlorophenylphosphodichloride (430 mg, 1.75 mmol), and TEA (354.2 mg, 3.50 mmol, 488.0 μ L) in DCM (15 mL), to yield 657.9 mg (yield 93.5%) of crude product used without further purification.

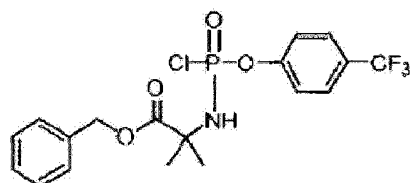
^{31}P -NMR ($CDCl_3$, 121 MHz): δ 7.00 (s)

1H -NMR ($CDCl_3$; 300 MHz): δ 7.39-7.12 (9H, m, $CH_2Ph + OPh$), 5.18 (2H, s, CH_2Ph), 4.75-4.72 (1H, 2bs, NH), 1.68-1.65 (6H, 2s, $[CH_3]_2C$).

^{13}C -NMR ($CDCl_3$; 75 MHz): δ 27.0, 27.3, ($[CH_3]_2C$), 59.0 ($C[CH_3]_2$), 68.4 (OCH_2Ph), 122.5, 128.6, 129.1, 130.7 ('o', 'm', 'p', $CH_2Ph + OPh$), 131.8 ('p', CH_2Ph), 135.4 ('p', OPh), 148.6, 148.7 ('ipso', OPh), 174.9, 175.1 ($COOCH_2Ph$).

Synthesis of 4-(trifluoromethyl)phenyl-(benzyl-2-amino-2-methylpropanoate)-phosphorochloridate.

[0240] $C_{18}H_{18}ClF_3NO_4P$, MW=435.76.



[0241] This is synthesised according to **Standard procedure 4**, using 2-aminoisobutyrate benzyl ester hydrochloride (341.0 mg, 1.49 mmol), 4-(trifluoromethyl)-phenylphosphodichloridate (414.3 mg, 1.49 mmol), and TEA (300.5 mg, 2.97 mmol, 413.9 μ L) in DCM (15 mL), to yield 623.9 mg (96.4%) of crude product used without further purification.

^{31}P -NMR ($CDCl_3$, 121 MHz): δ 6.74 (s)

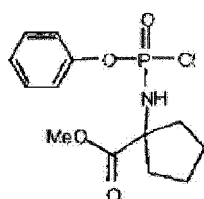
1H -NMR ($CDCl_3$; 300 MHz): δ 7.66 (2H, d, $^3J=8.8$ Hz, OPh), 7.42-7.30 (7H, m, $OPh + CH_2Ph$),

5.25 (2H, s, CH_2Ph), 4.95-4.91 (1H, 2bs, NH), 1.75-1.72 (6H, 2s, $[\text{CH}_3]_2\text{C}$).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 26.9, 27.0, 27.3 ($[\text{CH}_3]_2\text{C}$), 59.1 ($\text{C}[\text{CH}_3]_2$), 68.4 (CH_2Ph), 121.1, 121.4, 127.7, 128.4, 128.5, 128.6, 128.9 ('o', 'm', 'p', $\text{OPh}+\text{CH}_2\text{Ph}$), 124.2 (CF_3 , $J=265$ Hz), 135.4 ('ipso', CH_2Ph), 152.6, 152.7 ('ipso', OPh), 174.9, 175.0 (COOCH_2Ph).

Synthesis of Phenyl-(methoxy- α,α -cycloleucynyl)-phosphorochloridate.

[0242] $\text{C}_{13}\text{H}_{17}\text{ClINO}_4\text{P}$, MW=317.70.



[0243] This is synthesised according to **Standard procedure 4**, using methyl-1-amino-1-cyclopentanoate hydrochloride salt (0.885 g, 5.01 mmol), phenyldichlorophosphate (1.12 g, 0.749 ml, 5.01 mmol), and TEA (1.4 ml, 10 mmol) in DCM (40 mL), to yield 1.266 g (81%) of crude product used without further purification.

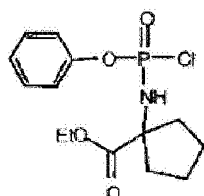
^{31}P -NMR CCl_4 , 121 MHz): δ 7.90.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.4-7.2 (5H, m, OPh), 4.3 (1H, bs, NH), 3.75 (3H, 2s, OCH_3), 2.15 (4H, m, 4H cyclopentane), 1.9-1.7 (4H, m, 4H cyclopentane)..

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 24.4 (2CH_2 cyclopent), 38.8, 38.7, 38.6 (2CH_2 cyclopent), 53.3, 53.2 (CH_3O), 66.6 (C_α cyclopentane), 121.1, 121.0 ('o' OPh), 126.3 ('p', OPh), 130.3, 130.2 ('m', OPh), 150.2 ('ipso', OPh), 174.8 (COOCH_3).

Synthesis of Phenyl-(ethoxy- α,α -cycloleucynyl)-phosphorochloridate.

[0244] $\text{C}_{14}\text{H}_{19}\text{ClINO}_4\text{P}$, MW=331.73.



[0245] This is synthesised according to **Standard procedure 4**, using ethyl-1-amino-1-cyclopentanoate hydrochloride salt (955 mg, 5.01 mmol), phenyldichlorophosphate (1.12g, 5.01 mmol, 749 μ L), and TEA (1.4 mL, 10.02 mmol) in DCM (40 mL). The crude was purified by flash chromatography (ethyl acetate/petroleum ether 7:3) affording 1.457 g (89%) of oil.

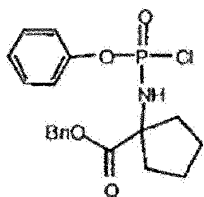
^{31}P -NMR (CDCl_3 , 121 MHz): δ 8.04, 7.97.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.4-7.1 (5H, m, OPh), 4.7 (1H, bs, NH), 4.2 (2H, 2q, $^3J=7.1$ Hz, OCH_2CH_3), 2.15 (4H, m, 4H cyclopentane), 1.9-1.7 (4H, m, 4H cyclopentane), 1.30 (3H, t, $^3J=7.1$ Hz, OCH_2CH_3).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 14.5 (CH_3CH_2), 24.5 (2 CH_2 cyclopent), 38.8, 38.7, 38.6, 38.5 (2 CH_2 cyclopent), 62.0 (CH_3CH_2), 68.3 (C_α cyclopentane), 120.9 ('o' OPh), 126.3 ('p', OPh), 130.3 ('m', OPh), 150.3-150.2 ('ipso', OPh), 174.9-174.8 ($\text{COOCH}_2\text{CH}_3$).

Synthesis of Phenyl-(benzoxy- α,α -cycloleucinyl)-phosphorochloridate.

[0246] $\text{C}_{19}\text{H}_{21}\text{ClINO}_4\text{P}$, MW=393.80.



[0247] This is synthesised according to **Standard procedure 4**, using benzyl-1-amino-1-cyclopentanoate hydrochloride salt (0.984 g, 3.84 mmol), phenyl-dichlorophosphate (0.577 ml, 3.84 mmol), and TEA (1.08 mL, 7.69 mmol) in DCM (30 mL), to yield 1.485 g (98%) of crude product used without further purification.

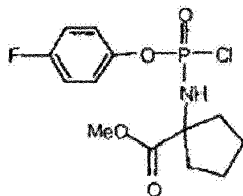
^{31}P -NMR (CDCl_3 , 121 MHz): δ 7.85.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.3-7.0 (10H, m, $\text{OPh}+\text{CH}_2\text{Ph}$), 5.2 (2H, s, CH_2Ph), 4.95-4.65 (1H, bs, NH), 2.25-2.1 (4H, m, 4H cyclopentane), 1.9-1.7 (4H, m, 4H cyclopentane).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 24.4, 24.3 (2 CH_2 cyclopent), 38.8, 38.7, 38.5 (2 CH_2 cyclopent), 67.3 (C_α cyclopentane), 68.0 (CH_2Ph), 121.0 ('o' OPh), 126.4 ('p', OPh), 130.1, 129.0, 128.8 ('m' OPh, CH_2Ph), 135.4 ('ipso', CH_2Ph), 150.1 ('ipso', OPh), 173.4 (COOCH_2Ph).

Synthesis of p-fluorophenyl-(methoxy- α,α -cycloleuciny)-phosphorochloridate.

[0248] $C_{13}H_{16}ClNO_4P$, MW=335.70.



[0249] This is synthesised according to **Standard procedure 4**, using methyl-1-amino-1-cyclopentanoate hydrochloride salt (0.885 g, 5.01 mmol), para-fluorophenyldichlorophosphate (1.21 g, 5.01 mmol), and TEA (1.4 ml, 10 mmol) in DCM (40 mL), to yield 1.65 g (99%) of crude product used without further purification.

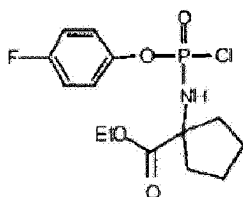
^{31}P -NMR ($CDCl_3$, 121 MHz): δ 8.61.

1H -NMR ($CDCl_3$; 300 MHz): δ 7.3-7.2 (2H, m, *O*Ph), 7.1-7.0 (2H, m, *O*Ph), 4.7 (1H, bs, *NH*), 3.78 (3H, 2s, *OCH*3), 2.25-2.15 (4H, m, 4H cyclopentane), 2.0-1.8 (4H, m, 4H cyclopentane)..

^{13}C -NMR ($CDCl_3$; 75 MHz): δ 24.4 (2CH₂ cyclopent), 38.7, 38.6, 38.5 (2CH₂ cyclopent), 53.3 (*C*H₃*O*), 66.3-66.2 (*C*q cyclopentane), 117.1-116.8 ('*o*' *O*Ph), 122.6-122.5 ('*m*', *O*Ph), 146.1-145.9 ('*ipso*', *O*Ph), 159.0 ('*p*', *O*Ph), 175.3-175.2 (*C*OOCH₃).

Synthesis of p-fluorophenyl-(ethoxy- α,α -cycloleuciny)-phosphorochloridate.

[0250] $C_{14}H_{18}ClFNO_4P$, MW=349.72.



[0251] This is synthesised according to **Standard procedure 4**, using ethyl-1-amino-1-cyclopentanoate hydrochloride salt (955 mg, 5.01 mmol), para-fluorophenyldichlorophosphate (1.21g, 5.01 mmol), and TEA (1.4 mL, 10.02 mmol) in DCM (40 mL), to yield 1.64 g (94%) of

crude product used without further purification.

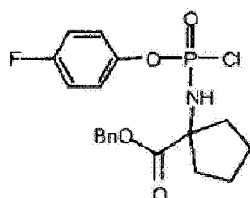
^{31}P -NMR (CDCl_3 , 121 MHz): δ 8.70.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.3-7.2 (2H, m, *OPh*), 7.1-7.0 (2H, m, *OPh*), 4.8 (1H, bs, *NH*), 4.2 (2H, 2q, $^3J=7.1$ Hz, *OCH₂CH₃*), 2.25-2.1 (4H, m, 4H cyclopentane), 2.0-1.8 (4H, m, 4H cyclopentane), 1.4 (3H, t, $^3J=7.1$ Hz, *OCH₂CH₃*).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 14.4 (*CH₃CH₂*), 24.4 (2CH₂ cyclopent), 38.8, 38.7, 38.6, 38.5 (2CH₂ cyclopent), 62.3 (*CH₃CH₂*), 68.3 (*C_q* cyclopentane), 117.4, 117.0 ('*o*' *OPh*), 122.7, 122.6 ('*m*', *OPh*), 146.1, 146.0 ('*ipso*', *OPh*), 159.0 ('*p*', *OPh*), 174.9 (*COOCH₂CH₃*).

Synthesis of p-fluorophenyl-(benzoxy- α,α -cycloleuciny)-phosphorochloridate.

[0252] $\text{C}_{19}\text{H}_{20}\text{ClFNO}_4\text{P}$, MW=4011.79.

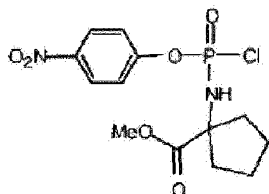


[0253] This is synthesized according to **Standard procedure 4**, using benzy]-1-amino-1-cyclopentanoate hydrochloride salt (1.281 g, 5.01 mmol), para-fluorophenyl-dichlorophosphate (1.21 g, 5.01 mmol), and TEA (1.4 mL, 10 mmol) in DCM (40 mL), to yield 1.85 g (90%) of crude product used without further purification.

^{31}P -NMR (CDCl_3 , 121 MHz): δ 7.85.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.65-7.4 (5H, m, *CH₂Ph*), 7.3-7.2 (2H, m, *OPh*), 7.1-7.0 (2H, m, *OPh*), 5.2 (2H, s, *CH₂Ph*), 4.6 (1H, bs, *NH*), 2.2-2.1 (4H, m, 4H cyclopentane), 2.0-1.8 (4H, m, 4H cyclopentane).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 24.5 (2CH₂ cyclopent), 38.9, 38.8, 38.6, 38.5 (2CH₂ cyclopent), 68.1 (*C_q* cyclopentane), 68.4 (*CH₂Ph*), 117.0, 116.8 ('*o*' *OPh*), 122.6, 122.5 ('*m*' *OPh*), 129.1, 129.0, 128.8, 128.7 (*CH₂Ph*), 135.7 ('*ipso*', *CH₂Ph*), 146.1, 145.9 ('*ipso*', *OPh*), 159.0 ('*p*', *OPh*), 174.6 (*COOCH₂Ph*).

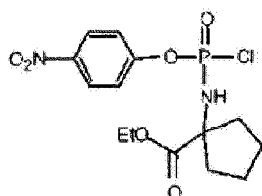
Synthesis of p-nitrophenyl-(methoxy- α,α -cycloleuciny)-phosphorochloridate.**[0254] $C_{13}H_{16}ClN_2O_6P$, MW-362.70.**

[0255] This is synthesised according to **Standard procedure 4**, using methyl-1-amino-1-cyclopentanoate hydrochloride salt (0.885 g, 5.01 mmol), para-nitrophenyldichlorophosphate (1.632 g, 5.01 mmol), and TEA (1.4 ml, 10 mmol) in DCM (40 mL), to yield 1.601 g (90%) of crude product used without further purification.

^{31}P -NMR ($CDCl_3$, 121 MHz): δ 8.02.

1H -NMR ($CDCl_3$; 300 MHz): δ 8.2 (2H, 2d, $^3J=8$ Hz, $O\text{Ph}$), 7.32 (2H, 2d, $^3J=8$ Hz, $O\text{Ph}$), 4.9 (1H, bs, NH), 3.71 (3H, s, OCH_3), 2.25-2.00 (4H, m, 4H cyclopentane), 1.95-1.7 (4H, m, 4H cyclopentane)..

^{13}C -NMR ($CDCl_3$; 75 MHz): δ 24.3 (2CH₂ cyclopent), 38.7, 38.6 (2CH₂ cyclopent), 53.3 ($\underline{C}H_3O$), 68.6 ($\underline{C}\alpha$ cyclopentane), 121.8, 121.7 ('o' OPh), 126.0 ('m', OPh), 145.6 ('ipso', OPh), 154.8, 154.7 ('p', OPh), 175.1-175.0 ($\underline{C}OOCH_3$).

Synthesis of p-nitrophenyl-(ethoxy- α,α -cycloleuciny)-phosphorochloridate,**[0256] $C_{14}H_{18}ClN_2O_6P$, MW=376.73.**

[0257] This is synthesised according to **Standard procedure 4**, using ethyl-1-amino-1-cyclopentanoate hydrochloride salt (**955** mg, 5.01 mmol), para-nitrophenyldichlorophosphate (1.362 g, 5.01 mmol), and TEA (1.4 mL, 10.02 mmol) in DCM (40 mL), to yield 1.669 g (90%) of crude product used without further purification.

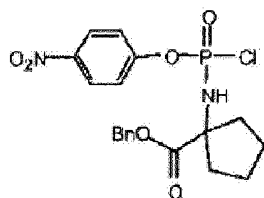
^{31}P -NMR (CDCl_3 , 121 MHz): δ 7.95.

^1H -NMR (CDCl_3 ; 300 MHz): δ 8.1 (2H, 2d, $^3J=8$ Hz, OPh), 7.28 (2H, 2d, $^3J=8$ Hz, OPh), 4.8 (1H, bs, NH), 4.2 (2H, 2q, $^3J=7.1$ Hz, OCH_2CH_3), 2.2-2.0 (4H, m, 4H cyclopentane), 1.95-1.7 (4H, m, 4H cyclopentane), 1.27 (3H, t, $^3J=7.1$ Hz, OCH_2CH_3).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 14.4 (CH_3CH_2), 24.4 (2 CH_2 cyclopent), 38.8, 38.7 (2 CH_2 cyclopent), 62.4 (CH_2CH_2), 68.5 (C_α cyclopentane), 121.8, 121.1 ('o' OPh), 126.1, 125.9 ('m', OPh), 145.6 ('ipso', OPh), 154.8 ('p', OPh), 174.9 ($\text{COOCH}_2\text{CH}_3$).

Synthesis of p-nitrophenyl-(benzoxy- α,α -cycloleucinyI)-phosphorochloridate.

[0258] $\text{C}_{19}\text{H}_{20}\text{ClN}_2\text{O}_6\text{P}$, MW=438.80.



[0259] This is synthesised according to **Standard procedure 4**, using benzyl-1-amino-1-cyclopentanoate hydrochloride salt (0.835 g, 3.25 mmol), para-nitrophenyl-dichlorophosphate (0.85 g, 3.25 mmol), and TEA (0.91 mL, 6.7 mmol) in DCM (30 mL), to yield 1.215 g (85%) of crude product used without further purification.

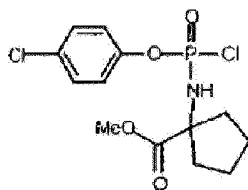
^{31}P -NMR (CDCl_3 , 121 MHz): δ 7.99, 7.90.

^1H -NMR (CDCl_3 ; 300 MHz): δ 8.1 (2H, 2d, $^3J=8$ Hz, OPh), 7.4-7.2 (7H, m, $\text{OPh} + \text{CH}_2\text{Ph}$), 5.18 (2H, s, CH_2Ph), 5.0 (1H, bs, NH), 2.2-2.0 (4H, m, 4H cyclopentane), 1.95-1.75 (4H, m, 4H cyclopentane).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 24.4 (2 CH_2 cyclopent), 38.8, 38.7, 38.6, 38.5 (2 CH_2 cyclopent), 68.0 (CH_2Ph), 68.6 (C_α cyclopentane), 121.8, 121.7 ('o' OPh), 126.1, 125.9 ('m' OPh), 129.1, 129.0, 128.8, 128.6 (CH_2Ph), 135.7 ('ipso', CH_2Ph), 145.6 ('ipso', OPh), 154.8, 154.7 ('p', OPh), 174.5, 174.4 (COOCH_2Ph).

Synthesis of p-chlorophenyl-(methoxy- α,α -cycloleucinyI)-phosphorochloridate.

[0260] $C_{13}H_{16}Cl_2NO_4P$, MW=352.15.



[0261] This is synthesised according to **Standard procedure 4**, using methyl-1-amino-1-cyclopentanoate hydrochloride salt (0.443 g, 2.5 mmol), para-chlorophenyldichlorophosphate (0.613 g, 2.5 mmol), and TEA (0.7 ml, 5 mmol) in DCM (20 mL), to yield 0.852 g (98%) of crude product used without further purification.

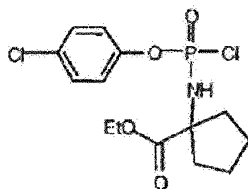
^{31}P -NMR ($CDCl_3$, 121 MHz): δ 9.55, 9.5.

1H -NMR ($CDCl_3$; 300 MHz): δ 7.35-7.15 (4H, m, OPh), 4.95 (1H, bs, NH), 3.78 (3H, s, OCH_3), 2.2-2.00 (4H, m, 4H cyclopentane), 1.95-1.7 (4H, m, 4H cyclopentane).

^{13}C -NMR ($CDCl_3$; 75 MHz): δ 24.3 ($2CH_2$ cyclopent), 38.7 ($2CH_2$ cyclopent), 53.3 (CH_3O), 68.6 ($C\alpha$ cyclopentane). 122.0 ('o' OPh), 130.1 ('m', OPh), 133.2 ('p', OPh), 149.9 ('ipso', OPh), 175.1-175.0 ($COOCH_3$).

Synthesis of p-chlorophenyl-(ethoxy- α,α -cyclopentyl)-phosphorochloridate.

[0262] $C_{14}H_{18}Cl_2NO_4P$, MW=366.18.



[0263] This is synthesised according to **Standard procedure 4**, using ethyl-1-amino-1-cyclopentanoate hydrochloride salt (0.477 g, 2.5 mmol), para-chlorophenyldichlorophosphate (0.613 g, 2.5 mmol), and TEA (0.7 mL, 5 mmol) in DCM (20 mL), to yield 0.880 g (97%) of crude product used without further purification.

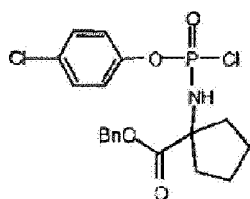
^{31}P -NMR ($CDCl_3$, 121 MHz): δ 9.85, 9.70.

$^1\text{H-NMR}$ (CDCl_3 ; 300 MHz): δ 7.35-7.15 (4H, m, OPh), 4.9 (1H, bs, NH), 4.22 (2H, 2q, $^3J=7.1$ Hz, OCH_2CH_3), 2.2-2.0 (4H, m, 4H cyclopentane), 1.95-1.7 (4H, m, 4H cyclopentane), 1.27 (3H, t, $^3J=7$ Hz, OCH_2CH_3).

$^{13}\text{C-NMR}$ (CDCl_3 ; 75 MHz): δ 14.4 (CH_3CH_2), 24.4 (2CH_2 cyclopent), 38.8, 38.7 (2CH_2 cyclopent), 62.5, 62.4 (CH_3CH_2), (68.) (C_α cyclopentane), 122.2, 122.1 ('o' OPh), 130.1 ('m', OPh), 133.2 ('p', OPh), 149.8 ('ipso', OPh), 174.8 ($\text{COOCH}_2\text{CH}_3$).

Synthesis of p-chlorophenyl-(benzoxy- α,α -cycloleucinyI)-phosphorochloridate.

[0264] $\text{C}_{19}\text{H}_{20}\text{Cl}_2\text{NO}_4\text{P}$, MW=428.25.



[0265] This is synthesised according to **Standard procedure 4**, using benzyl-1-amino-1-cyclopentanoate hydrochloride salt (0.640 g, 2.5 mmol), para-chlorophenyldichlorophosphate (0.613 g, 2.5 mmol), and TEA (0.7 mL, 5 mmol) in DCM (20 mL), to yield 1.041 g (97%) of crude product used without further purification.

$^{31}\text{P-NMR}$ (COCl_3 , 121 MHz): δ 9.39, 8.95.

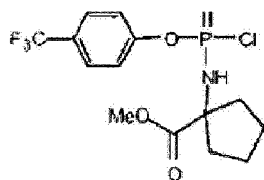
$^1\text{H-NMR}$ (CDCl_3 ; 300 MHz): δ 7.4-7.15 (9H, m, OPh + CH_2Ph), 5.20 (2H, s, CH_2Ph), 5.0 (1H, bs, NH), 2.2-2.0 (4H, m, 4H cyclopentane), 1.95 -1.75 (4H, m, 4H cyclopentane).

$^{13}\text{C-NMR}$ (CDCl_3 ; 75 MHz): δ 24.4 (2CH_2 cyclopent), 38.8, 38.7, 38.6 (2CH_2 cyclopent), 68.1, 68.0 (CH_2Ph), 68.2 (C_α cyclopentane), 121.9, 121.8 ('o' OPh), 130.5, 130.4, 129.3, 129.2 ('m' OPh, CH_2Ph), 133.2 ('p', OPh), 135.7 ('ipso', CH_2Ph), 149.9 ('ipso', OPh), 174.3, 174.2 (COOCH_2Ph).

Synthesis of p-trifluorophenyl-(methoxy- α,α -cycloleucinyI)-phosphorochloridate.

[0266] $\text{C}_{14}\text{H}_{16}\text{ClF}_3\text{NO}_4\text{P}$, MW=385.70.





[0267] This is synthesised according to **Standard procedure 4**, using methyl-1-amino-1-cyclopentanoate hydrochloride salt (0.443 g, 2.5 mmol), paratrifluorophenyldichlorophosphate (0.700 g, 2.5 mmol), and TEA (0.7 ml, 5 mmol) in DCM (20 mL), to yield 0.931 g (97%) of crude product used without further purification.

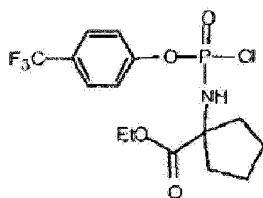
^{31}P -NMR (CDCl_3 , 121 MHz): δ 8.80, 8.62.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.65 (2H, 2d, $^3J=8$ Hz, O Ph), 7.35 (2H, 2d, $^3J=8$ Hz O Ph), 5.02 (1H, bs, NH), 3.78 (3H, s, OCH $_3$), 2.25-2.05 (4H, m, 4H cyclopentane), 1.95-1.7 (4H, m, 4H cyclopentane).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 22.8 (2CH $_2$ cyclopent), 37.5, 37.2 (2CH $_2$ cyclopent), 51.5 (CH $_3$ O), 68.4 (C α cyclopentane), 120.0 (' σ ', O Ph), 124.8 (d, $J=270\text{Hz}$, CF $_3$), 126.6 (' m ', O Ph), 129.5 (' p ', q, $J=32\text{Hz}$, O Ph), 152.8 (' $ipso$ ', OPh), 175.2 (COOCH $_3$).

Synthesis of p-trifluorophenyl-(ethoxy- α,α -cycloleuciny)-phosphorochloridate.

[0268] C $_{15}$ H $_{18}$ ClF $_3$ NO $_4$ P, MW=399.73.



[0269] This is synthesised according to **Standard procedure 4**, using ethyl-1-amino-1-cyclopentanoate hydrochloride salt (0.477 g, 2.5 mmol), para-trifluorophenyldichlorophosphate (0.700 g, 2.5 mmol), and TEA (0.7 mL, 5 mmol) in DCM (20 mL), to yield 0.950 g (89%) of crude product used without further purification.

^{31}P -NMR (CDCl_3 , 121 MHz): δ 8.49.

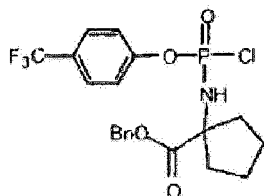
^1H -NMR (CDCl_3 ; 300 MHz): δ 7.45 (2H, m, O Ph), 7.2 (2H, m, O Ph), 5.12 (1H, bs, NH), 4.05 (2H, m, OCH $_2$ CH $_3$), 2.15-2.0 (4H, m, 4H cyclopentane), 1.9-1.65 (4H, m, 4H cyclopentane), 1.2

(3H, 2t, $^3J=7$ Hz, OCH_2CH_3).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 14.3 (CH_3CH_2), 24.2, 24.1 (2CH_2 cyclopent), 38.6, 38.5, 38.4 (2CH_2 cyclopent), 62.0 (CH_3CH_2), 68.4 (C_α cyclopentane), 121.5 ('o', OPh), 125.0 (d, $J=270\text{Hz}$, CF_3), 127.5 ('m', OPh), 129.9 ('p',q, $J=32\text{Hz}$, OPh), 152.8, 152.7 ('ipso', OPh), 174.9, 174.6 ($\text{COOCH}_2\text{CH}_3$).

Synthesis of p-trifluorophenyl-(benzoxy- α,α -cycloleucinyI)-phosphorochloridate.

[0270] $\text{C}_{20}\text{H}_{20}\text{ClF}_3\text{NO}_4\text{P}$, MW= 461.80.



[0271] This is synthesized according to **Standard procedure 4**, using benzyl-1-amino-1-cyclopentanoate hydrochloride salt (0.700 g, 2.73 mmol), para-trifluorophenyldichlorophosphate (0.75 g, 2.73 mmol), and TEA (0.75 mL, 5.47 mmol) in DCM (25 mL), to yield 1.089 g (86%) of crude product used without further purification.

^{14}P -NMR (CDCl_3 , 121 MHz): δ 9.39, 8.95.

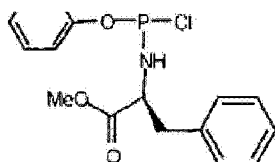
^1H -NMR (CDCl_3 ; 300 MHz): δ 7.50 (2H, m, OPh), 7.4-7.15 (7H, m, $\text{OPh}+\text{CH}_2\text{Ph}$), 5.20 (2H, s, CH_2Ph), 4.95 (1H, bs, NH), 2.2-2.0 (4H, m, 4H cyclopentane), 1.95-1.75 (4H, m, 4H cyclopentane),

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 24.3 (2CH_2 cyclopent), 38.8, 38.7, 38.6 (2CH_2 cyclopent), 68.1, 68.0 (CH_2Ph), 68.2 (C_α cyclopentane), 121.4, 121.3 ('o', OPh), 125.1 (d, $J=270\text{Hz}$, CF_3), 126.6 ('m' OPh), 129.2, 128.8, 127.8 (Bn), 129.8 ('p',q, $J=32\text{Hz}$, OPh), 135.7 ('ipso', CH_2Ph), 151.5 ('ipso', OPh), 174.5, 174.4 (COOCH_2Ph).

Synthesis of Phenyl-(methoxy-L-phenylalaninyI)-phosphorochloridate.

[0272] $\text{C}_{16}\text{H}_{17}\text{ClNO}_4\text{P}$, MW=353.14.





[0273] This is synthesised according to **Standard procedure 4**, using L-phenylalanine methyl ester hydrochloride (1.08 g, 5 mmol), phenyldichlorophosphate (1.12 g, 0.75 ml, 5 mmol), and TEA (1.4ml, 10 mmol) in DCM (40 mL), to yield 1.626 g (92%) of crude product used without further purification.

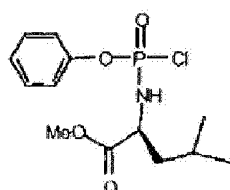
^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.1, 8.95.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.3-7.1 (10H, m, $\text{CH}_2\text{Ph} + \text{OPh}$), 5.00 (1H, bs, NH), 4.35 (1H, m, CHphenylala), 3.79 (3H, 2s, CH_3O), 3.00 (2H, m, CH_2Ph)

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 36.3 ($\text{CH}_2\text{phenylalanine}$), 53.0 (CH_3O), 56.6, 56.5 (CHphenylala), 121.0 ('o' OPh), 126.4 ('p' OPh), 130.2 ('m', OPh), 150.2 ('ipso', OPh), 174.1 (COOCH_3),

Synthesis of Phenyl-(methoxy-L-leucynyl)-phosphorochloridate

[0274] $\text{C}_{13}\text{H}_{19}\text{ClINO}_4\text{P}$, MW=319.72.



[0275] This is synthesised according to **Standard procedure 4**, using L-leucine methyl ester hydrochloride (0.91 g, 5 mmol), phenyldichlorophosphate (1.12 g, 0.75 ml, 5 mmol), and TEA (1.4 ml, 10 mmol) in DCM (40 mL), to yield 1.58 g (99%) of crude product used without further purification.

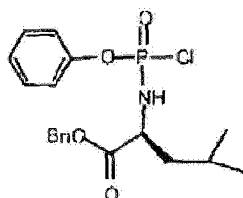
^{31}P -MMR (CDCl_3 , 121 MHz): δ 9.45. 9.35.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.4-7.2 (5H, m, OPh), 4.90 (1H, bs, NH), 3.95 (1H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 3.78 (3H, s, OCH_3), 1.8 (1H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 1.8-1.5 (2H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 1.0-0.9 (6H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 23.2, 23.1, 22.4, 22.3 (2C, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 24.9, 24.8 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 43.6 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 53.2 (CH_3O), 53.7, 53.6 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 120.9 ('o' OPh), 126.4 ('p', OPh), 130.2 ('m', OPh), 150.1 ('ipso', OPh), 173.6 (COOCH_3).

Synthesis of Phenyl-(benzoxy-L-leucinyl)-phosphorochloridate

[0276] $\text{C}_{19}\text{H}_{23}\text{ClINO}_4\text{P}$, MW=395.82.



[0277] This is synthesised according to **Standard procedure 4**, using L-leucine benzyl ester hydrochloride (1.29 g, 5.0 mmol), phenyl-dichlorophosphate (1.12 g, 0.75 ml, 5.0 mmol), and TEA (1.4 mL, 10.0 mmol) in DCM (40 mL), to yield 1.88 g (95%) of crude product used without further purification.

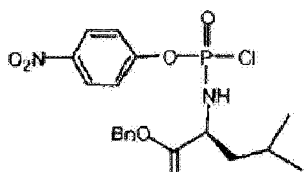
^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.93, 9.57.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.5-7.2 (10H, m, $\text{OPh} + \text{CH}_2\text{Ph}$), 5.2 (2H, 2s, CH_2Ph), 4.95 (1H, bs, NH), 4.2-4.1 (1H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 1.95-1.80 (1H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 1.7 (2H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 1.0-0.9 (6H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 23.2, 23.1, 22.4, 22.3 (2C, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 24.9 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 43.5 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 53.8, 53.3 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 67.8, 67.7 (CH_2Ph), 120.7 ('o' OPh), 126.4 ('p', OPh), 130.2, 129.1, 128.8, 128.7 ('m' OPh, CH_2Ph), 135.8 ('ipso', CH_2Ph), 150.2 ('ipso', OPh), 174.1 (COOCH_2Ph).

Synthesis of p-nitrophenyl-(benzoxy-L-leucinyl)-phosphorochloridate.

[0278] $\text{C}_{19}\text{H}_{22}\text{ClIN}_2\text{O}_6\text{P}$, MW=440.81.



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[0279] This is synthesized according to **Standard procedure 4**, using L-leucine benzyl ester hydrochloride (1.08 g, 5.01 mmol), para-nitrophenyl-dichloro phosphate (1.362 g, 5.01 mmol), and TEA (1.4 mL, 1.4 mmol) in DCM (40 mL), to yield 2.08g (95%) of crude product used without further purification.

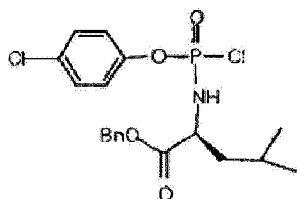
^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.87, 9.38.

^1H -NMR (CDCl_3 ; 300 MHz): δ 8.25-8.10 (2H, m, OPh), 7.35-7.25 (7H, m, OPh + CH_2Ph), 5.15 (2H, 2s, CH_2Ph), 4.95 (1H, bs, NH), 4.15 (1H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 1.95 (1H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 1.7 (2H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 1.0-0.9 (6H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 23.2, 23.1, 22.1, 22.0 (2C, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 24.8 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 43.4, 43.3 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 54.2, 53.9 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 68.0, 67.9 (CH_2Ph), 121.6 ('o' OPh), 126.2, 126.1 ('m' OPh), 129.2, 129.0 (CH_2Ph), 135.4, 135.3 ('ipso', CH_2Ph), 145.8, 145.7 ('ipso', OPh), 154.7, 154.5 ('p', OPh), 173.0, 172.8 (COOCH_2Ph).

Synthesis of p-chlorophenyl-(benzoxy-L-leuciny)-phosphorochloridate.

[0280] $\text{C}_{19}\text{H}_{22}\text{Cl}_2\text{NO}_4\text{P}$, MW=430.26.



[0281] This is synthesised according to **Standard procedure 4**, using L-leucine benzyl ester hydrochloride (0.644 g, 2.5 mmol), para-chlorophenyl-dichlorophosphate (0.613 g, 2.5 mmol), and TEA (0.7 mL, 5 mmol) in DCM (20 mL), to yield 0.968 g (90%) of crude product used without further purification.

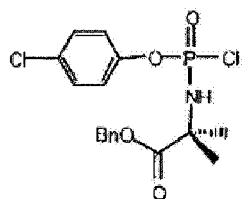
^{31}P -NMR (CDCl_3 , 121 MHz): δ 9.71, 9.55.

^1H -MMR (CDCl_3 ; 300 MHz): δ 7.4-7.0 (9H, m, OPh + CH_2Ph), 5.15 (2H, s, CH_2Ph), 4.5 (1H, d, $^3J=\text{Hz}$, NH), 4.0 (1H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 1.9-1.8 (1H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 1.7 (2H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 0.85 (6H, m, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 23.4, 23.3, 22.5, 22.4 (2C, $\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 25.0 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 45.8, 43.7 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 54.0, 53.8 ($\text{CHCH}_2\text{CH}(\text{CH}_3)_2$), 68.2 (CH_2Ph), 122.5 ('o' OPh), 130.5, 130.4, 129.3, 129.2 ('m'OPh, CH_2Ph), 133.2 ('p', OPh), 135.7 ('ipso', CH_2Ph), 149.9, 149.8 ('ipso', OPh), 173.4, 173.2 (COOCH_2Ph).

Synthesis of 4-chlorophenyl-(methyl-2-amine-2-methylpropanoate)-phosphorochloridate.

[0282] $\text{C}_{11}\text{H}_{14}\text{Cl}_2\text{NO}_4\text{P}$, MW=326.11.



[0283] This is synthesised according to **Standard procedure 4**, using 2-aminoisobutyrate methyl ester hydrochloride (280.0mg, 1.82 mmol), 4-chlorophenylphosphodichloride (447.4 mg, 1.82 mmol), and TEA (368.3 mg, 3.64 mmol, 507.3 μL) in DCM (20 mL), to yield 554 mg (yield 91.1%) of crude product used without further purification.

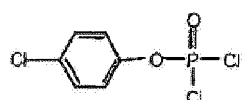
^{31}P -NMR (CDCl_3 , 121 MHz): δ 7.05 (s)

^1H -MMR (CDCl_3 ; 300 MHz): δ 7.38 (2H, d, $^3J=9.0$ Hz, OPh), 7.29-7.24 (2H, 2d, $^3J=9.0$ Hz, OPh), 4.87-4.83 (1H, 2bs, NH), 3.84 (3H, s, OCH₃), 1.73-1.71 (6H, 2s, [CH₃]₂C).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 27.0, 27.3, ([CH₃]₂C), 53.7 (OCH₃), 58.9 (C[CH₃]₂), 122.5 ('o', OPh), 129.7 ('m', OPh), 131.8 ('p', OPh) 148.7, 148.9 ('ipso', OPh), 175.5, 175.7 (COOCH_3).

Synthesis of 4-chlorophenyl-phosphodichloridate.

[0284] $\text{C}_6\text{H}_4\text{Cl}_3\text{O}_2\text{P}$, MW=245.43.



[0285] This was synthesised according to **Standard procedure 3**, using phosphorus-

oxychloride (1533 mg, 10.00 mmol, 932 μ L), 4-chlorophenol (1.285 g, 10.00 mmol) and TEA (1,011 g, 10.00 mmol, 1394 μ L) in ethylether (100 mL) to give an oil (1.897 g, 77.3 % yield).

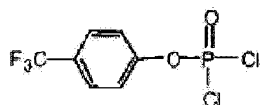
^{31}P -NMR (CDCl_3 , 121 MHz): δ 5.18.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.45 (2H, d, $^3J=9.0$ Hz, OPh), 7.30 (2H, d, $^3J=9.0$ Hz, OPh).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 122.5 ('o', OPh), 130.6 ('m', OPh), 133.2 ('p', OPh), 148.5 ('ipso', OPh).

Synthesis of 4-(trifluoromethyl)-phenyl-phosphodichloridate.

[0286] $\text{C}_7\text{H}_4\text{ClF}_3\text{O}_3\text{P}$, MW=278.99.



[0287] This was synthesised according to **Standard procedure 3**, using phosphorus-oxychloride (1.570 mg, 10.24 mmol, 954.5 μ L), 4-trifluoromethylphenol (1660 g, 10.24 mmol) and TEA (1.036 g, 10.24 mmol, 1427 μ L) in ethylether (100 mL) to give an oil (2.521 g, 88.2% yield).

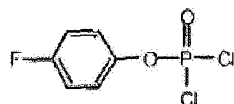
^{31}P -NMR (CDCl_3 , 121 MHz): δ 4.75.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.77 (2H, d, $^3J=8.4$ Hz, OPh), 7.49 (2H, d, $^3J=8.4$ Hz, OPh).

^{13}C -NMR (CDCl_3 ; 75 MHz): δ 121.6 ('o', OPh), 123.6 (CF_3 , $J=271$ Hz, OPh), 128.2 ('m', OPh), 129.7 ('p', $J=33$ Hz), 152.7 ('ipso', OPh).

Synthesis of 4-fluorophenyl-phosphodichloridate.

[0288] $\text{C}_6\text{H}_4\text{Cl}_2\text{FO}_2\text{P}$, MW=228.97.



[0289] This was synthesised according to **Standard procedure 3**, using phosphorus-oxychloride (1.395 mL, 15.00 mmol), 4-chlorophenol (1.68 g, 15.00 mmol) and TEA (2.1 mL, 15.00 mmol) in ethylether (140 mL) to give an oil (3.96 g, 96 % yield).

^{31}P -NMR (CDCl_3 , 121 MHz); δ 5.52.

^1H -NMR (CDCl_3 ; 300 MHz): δ 7.15 (2H, d, $^3J=8.0$ Hz, *OPh*), 7.05 (2H, d, $^3J=8.0$ Hz, *OPh*).

^{13}C -NMR. (CDCl_3 ; 75 MHz): δ 116.8 ('o', *OPh*), 122.1 ('m', *OPh*), 146.7 ('p', *OPh*), 158.7 ('ipso', *OPh*).

[0290] Experimental data are given in Table I illustrating the activity of compounds embodying the present invention, and of some comparative compounds, with respect to human breast cancer cell line MDA MB231, human colon cancer cell line HT115 and human prostate cancer cell line PC-3. The compounds include those whose preparations are described above and compounds made by preparative methods corresponding to the methods described above.

[0291] The experimental procedures used human colon cancer cell line (HT115), human prostate cancer cell line (PC-3), human breast cancer cell line (MDA MB 231) and normal human umbilical vein endothelial cell (HUVEC). Compounds were diluted over a range of concentrations and added to cells over 1 to 3 days. The cytotoxicity was determined using a MTT assay at the end of each experiment.

[0292] In the Table:

ArO refers to Ar as defined above with respect to formula I:

J refers to the moiety of the present compounds represented by, respectively, $\text{ROCOCR}'\text{R}''\text{NH}-$, as defined above with respect to formula I, or, with respect to Examples 51, 52 and 53, $\text{HOCO R}'\text{R}''\text{NH}-$, as defined above with respect to formula II; and

B refers to the base moiety of the compounds:

BVU stands for 2-bromovinyl uridine.

5-(C=CC[O]O)MeU stands for methyl propenoate-2'-deoxyuridine.

GemCyt stands for Gemcitabine.

Examples A, 1, 67 and G are further comparative examples.

Example A is 5-(2-Bromovinyl)-2'-deoxyuridine.

Example 1 is Example 1 above corresponding to compound (7) above.

Example 67 is propenoate-2'-deoxyuridine.

Example G is gemcitabine.

TABLE

Ex	ArO	J	B	EC50/ μ M	EC50/ μ M	EC50/ μ M
				Breast	Colon	Prostate
				MDA MB231	HT115	PC-3
A	-	-	BVU	125	78.7	120
1	PhO	MeAlaNH	BVU	79	244.5	155
2	PhO	BnAlaNH	BVU	34	1.4	19
3	PhO	EtAlaNH	BVU	56	52	36
4	p-CF ₃ PhO	BnAlaNH	BVU	31	7.4	9.3

5	p-FPhO	MeAlaNH	BVU	159	17	58
6	p-FPhO	EtAlaNH	BVU	46	11	42
7	p-FPhO	BnAlaNH	BVU	17	3.5	16
8	p-NO ₂ PhO	BnAlaNH	BVU	28	-	9
9	p-NO ₂ PhO	EtAlaNH	BVU	177	118.7	365
10	p-NO ₂ PhO	MeAlaNH	BVU	105	96.7	10.4
11	p-ClPhO	EtAlaNH	BVU	28.7	14.9	3.4
12	p-ClPhO	BnAlaNH	BVU	6.2	3.4	2.4
13	p-ClPhO	MeAlaNH	BVU	61	70.2	13
14	PhO	Bn(Me ₂ Gly)NH	BVU	19	14.5	5.1
15	p-CF ₃ PhO	MeAlaNH	BVU	47	79.2	15
16	PhO	Me(cPntGly)NH	BVU	79	77	16
17	PhO	Et(cPntGly)NH	BVU	44	81.3	41
18	PhO	Bn(cPntGly)NH	BVU	78	9.7	33
19	p-NO ₂ PhO	Me[cPntGly]NH	BVU	56	38.2	88
20	p-NO ₂ PhO	Et[cPntGly]NH	BVU	13	57.3	15
21	p-NO ₂ PhO	Bn[cPntGly]NH	BVU	8.4	17.2	2.2
22	PFPhO	Me[cPntGly]NH	BVU	57	59.7	51
23	PFPhO	Et[cPntGly]NH	BVU	9.9	18.1	2.7
24	PFPhO	Bn[cPntGly]NH	BVU	9.4	17	3.7
25	p-CF ₃ PhO	EtAlaNH	BVU	33.8		4.6
26	PhO	Me(Me ₂ Gly)NH	BVU	41.1	77.9	1.5
27	PhO	Et(Me ₂ Gly)NH	BVU	217.9	39.7	76.1
28	p-CF ₃ PhO	Me(cPntGly)NH	BVU	28.8	21.2	-
29	p-CF ₃ PhO	Et(cPntGly)NH	BVU	45.6	15.1	4.3
30	p-CF ₃ PhO	Bn(cPntGly)NH	BVU	6.9	6.4	-
32	p-ClPhO	Me[cPntGly]NH	BVU	2.6	99.3	52.2
33	p-ClPhO	Et[cPntGly]NH	BVU	12	97.9	83.2
34	p-ClPhO	Bn[cPntGly]NH	BVU	3.9	8.9	6.3
35	PhO	MeLeuNH	BVU	18.5	7.7	75.7
36	PhO	Me[Phc]NH	BVU	19.8	32.1	86.9
37	PhO	BnLeuNH	BVU	2.8	7	7.16
38	p-NO ₂ PhO	BnLeuNH	BVU	6.3	10.7	7.2

39	p-ClPhO	BnLeuNH	BVU	4.3	288.5	193.1
42	p-ClPhO	Me(Me2Gly)NH	BVU	8.7	183.4	441.6
43	p-ClPhO	Et(Me2Gly)NH	BVU	5.9	174.3	1.15
44	p-ClPhO	Bn(Me2Gly)NH	BVU	2.3	4.5	9.12
45	p-NO2PhO	Me(Me2Gly)NH	BVU	9.4	24.7	222.8

46	p-NO2PhO	Et(Me2Gly)NH	BVU	2	224	82.4
47	p-NO2PhO	Ba(Me2Gly)NH	BVU	4.5	16.7	27.2
48	p-CF3PhO	Bn(Me2Gly)NH	BVU	1.3	7	0.61
49	o-ClPhO	BnAlaNH	BVU	5.4	16.2	5.4
50	o-ClPhO	Bn(Me2Gly)NH	BVU	5.7	3.9	6.59
51	-	L-AlaNH	BVU		295.4	
52	-	LeuNH	BVU		438.1	
53	-	PhAlaNH	BVU		66	
54	PhO	Bn[PhAla]NH	BVU		5.1	
55	PhO	Me[D-Ala]NH	BVU		392.7	
56	PhO	Bn[D-Ala]NH	BVU		20.8	
57	p-NO2PhO	Bn[D-Ala]NH	BVU		20.2	
58	p-CF3	Me(Me2Gly)NH	BVU		83.6	
59	p-CF3	Et(Me2Gly)NH	BVU		24.7	
60	p-FPhO	Et(Me2Gly)NH	BVU		86.8	
61	p-CF3PhO	Bn[L-PhAla]NH	BVU		6.3	
62	p-CF3PhO	Bn[L-Leu]NH	BVU		1.9	
63	PhO	tBu[L-Ala]NH	BVU		31.5	
64	p-NO2PhO	Bn[L-PhAla]NH	BVU		16.6	
65	p-FPhO	Me(Me2Gly)NH	BVU			
66	p-NO2PhO	Me(Me2Gly)NH	5-(C=CC(O)O) Me)U		20.7	
67	-	-	5-(C=CC(O)O) Me)U		93.7	
69	PhO	MeMetNH	BVU	-	-	6.3
70	PhO	MeIrpNH	BVU	-	-	16
71	PhO	BnMetNH	BVU	-	-	6.3
72	PhO	BnIleNH	BVU	-	-	1.6
73	PhO	EtIleNH	BVU	-	-	30.6
74	PhO	MeGlyNH	BVU	-	-	31
75	PhO	BnGlyNH	BVU	-	-	29
77	p-Cl PhO	BnGlyNH	BVU	-	-	150
78	p-CF3 PhO	BnValNH	BVU	-	-	1.6
80	PhO	Me2AspNH	BVU	-	-	158
81	PhO	EtGluNH	BVU	-	-	31
82	m-ClPhO	BnAlaNH	BVU	-	-	21
83	m-ClPhO	BaMe2GlyNH	BVU	-	-	6.3
84	p-FphO	BnMe2GlyNH	BVU	-	-	4.5

85	PhO	BnValNH	BVU	-	-	31.2
86	p-ClPhO	BnValNH	BVU	-	-	0.9
87	p-FphO	BnValNH	BVU	-	-	1.6

88	PhO	BnPhGlyNH	BVU	-	-	0.75
89	p-ClPhO	BnPhGlyNH	BVU	-	-	6.5
91	p-Cl ₂ PhO	BnPhGlyNH	BVU	-	-	0.7
94	PhO	i-BuAlaNH	BVU	-	-	51
95	PhO	2-BuAlaNH	BVU	-	-	6.8
G	-	-	GemCyt	2.8	606.1	3.12
31	PhO	BnAlaNH	GemCyt	42.6	5.7	0.22
40	p-ClPhO	BnAlaNH	GemCyt	9.2	16.1	15.4
41	p-ClPhO	Bn[Me2Gly]NH	GemCyt	3.1	317.1	68.8

[0293] Gemcitabine (Example G in the Table) and compound CPF31 (Example 31 in the Table: gemcitabine-[phenyl-(benzoxy-L-alaninyl)]-phosphate) were compared in a mouse model with xenografts of human cancer (colon HT115 and prostate PC3).

[0294] Mice were dosed daily at a range of concentrations (0.01-10 μ M) and tumour volume assessed versus control.

[0295] Kaplan-Meier statistics were computed regarding incident-free survival.

[0296] In the attached drawings:

Figure 1 shows for the mouse xenograft the tumour volume for prostate data at day 13 using Gemzar™ (gemcitabine available ex. Lilly);

Figure 2 shows for the mouse xenograft the tumour volume for prostate data at day 13 using CPF31;

Figure 3 shows the incident free survival functions v. day for each of CPF31 and gemcitabine; and

Figure 4 shows for the mouse xenograft the tumour volume for colon data at day 24 using, respectively, Gemzar and compound CPF31.

[0297] Referring to the drawings, CPF31 can be seen to be significantly less toxic than gemcitabine.

[0298] CPF31 was significantly effective at reducing prostate and colon tumour volume relative to control at daily dosing of 5 and 10 μ M (3 and 6 μ g/ml). Gemcitabine was not effective at the highest non-toxic concentration.

[0299] Gemzar is seen from Figure 1 to be toxic above 1 μ M. In contrast, CPF31 is seen from Figure 2 to have substantially lower toxicity.

[0300] Figure 3 shows that CPF31 has significantly lower side effects on a comparable basis: 3 animals show serious toxicity (10% body mass loss) in GMZ and in CPF31 on day 10, collectively 4 in GMZ and 1 in CPF31 on day 11 and 5 in GMZ and 1 in CPF on day 13. Using Chi square analysis by combining 5 and 10µM groups, the significance is $p=0.193$, 0.078 and 0.0289 on day 10, 11 and 13. It is clear that by day 13, CPF31 displayed significantly less side effects, and the anti-cancer effects continue to exceed that of Gemzar,

[0301] Figure 3 shows the Kaplan-Meier survival curve, incidence free survival: based on the loss according to weight loss. A Cox proportion analysis shows that CPF31 is far less toxic than GMZ based on the weight-loss calculated loss ($p=0.043$).

[0302] CPF31 was found to be active at 5µM *in vitro*, whereas Gemzar was found to be active at 600µM, with respect to the same colon cell line. Figure 4 shows the results of testing both *in vivo* at 5µM. The greater activity of CPF31 in reducing tumour volume is shown in Figure 4.

REFERENCES CITED IN THE DESCRIPTION

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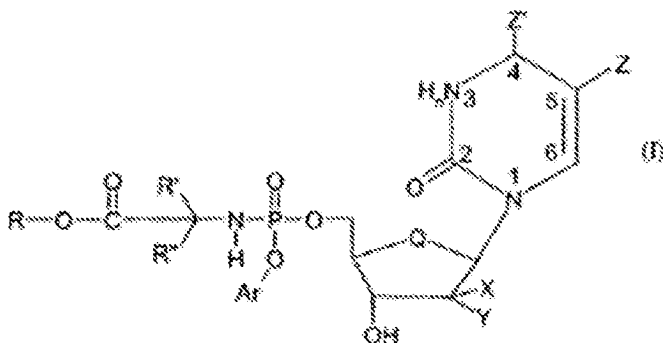
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PATENTKRAV

1. Kemisk forbindelse med formel I:



hvor:

- 5 R er valgt fra en forgrenet eller ikke-forgrenet, acyklisk C₁-C₁₆-alkylgruppe, en cyklisk C₃-C₁₂-alkylgruppe, en phenylgruppe eller en naphthylgruppe og en benzylgruppe; og hvor R er usubstitueret;
- R' og R'' er uafhængigt valgt fra H, methyl (-CH₃), benzyl (-CH₂C₆H₅) og sekundær butyl (-CH₂CH(CH₃)₂), eller R' og R'' sammen med det C-atom, hvortil de er bundet, danner en C₅₋₆-ring;
- 10 Q er -O-;
- X og Y begge er F;
- Ar er naphthyl eller phenyl; og én, to, tre eller fire substituenten, der kan være ens eller forskellige, kan være til stede på Ar og er valgt fra gruppen, der omfatter halogen, -NO₂; -NH₂;
- 15 -C₁₋₃-alkyl; -C₁-C₃-alkoxy; -SC₁₋₃-alkyl; -CN; -COC₁₋₃-alkyl og -CO₂C₁₋₃-alkyl;
- Z er H;
- n er 0, Z' er -NH₂ og en dobbeltbinding eksisterer mellem position 3 og position 4; eller et farmaceutisk acceptabelt salt, en ester eller salt af sådan ester af en forbindelse med formel I.
2. Forbindelse ifølge krav 1, hvor R er en forgrenet eller ikke-forgrenet, acyklisk C₁-C₆-alkylgruppe.
- 20 3. Forbindelse ifølge krav 1, hvor R er cyklisk C₅-C₇-alkylgruppe.
4. Forbindelse ifølge krav 1, hvor R er en benzylgruppe.
5. Forbindelse ifølge et hvilket som helst af krav 1 til 4, hvor én af R' og R'' er H og én af R' og R'' er methyl.
- 25 6. Forbindelse ifølge et hvilket som helst af kravene 1 til 5, hvor Ar er naphthyl.
7. Forbindelse ifølge et hvilket som helst af kravene 1 til 6, hvor Ar er phenyl.
8. Forbindelse ifølge et hvilket som helst af kravene 1 til 7, hvor Ar er usubstitueret.
9. Forbindelse ifølge et hvilket som helst af kravene 1 til 7, hvor substituenterne på Ar kan

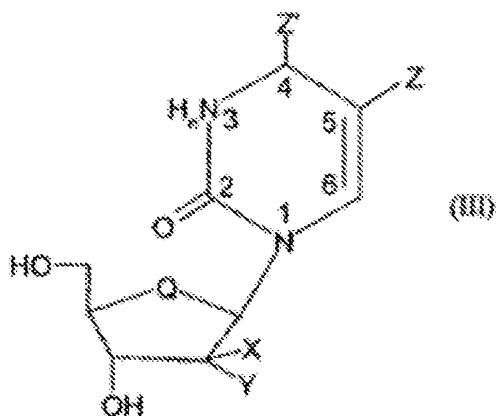
være valgt fra F, Cl, CF₃ og NO₂.

10. Forbindelse ifølge et hvilket som helst af krav 1 til 9 til anvendelse i en fremgangsmåde til profylakse eller behandling af cancer.

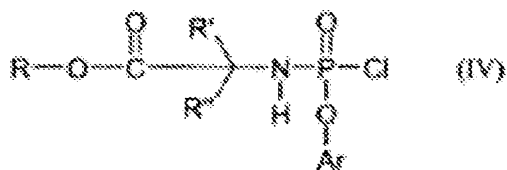
11. Farmaceutisk sammensætning, der omfatter en forbindelse ifølge et hvilket som helst af krav 1 til 9 i kombination med en farmaceutisk acceptabel bærer, fortynder eller excipients.

12. Fremgangsmåde til fremstilling af en farmaceutisk sammensætning, der omfatter trinnet med kombinerende af en forbindelse ifølge et hvilket som helst af krav 1 til 9 med en farmaceutisk acceptabel excipients, bærer eller fortynder.

13. Fremgangsmåde til fremstilling af en forbindelse med formel I ifølge krav 1, hvilken fremgangsmåde omfatter omsætning af en forbindelse med formel (III):



med en forbindelse med formel (IV)



15 hvor Ar, n, Q, R, R', R'', X, Y, Z og Z' har de betydninger, der er beskrevet i krav 1, og der eksisterer en dobbeltbinding mellem position 3 og position 4.

DRAWINGS

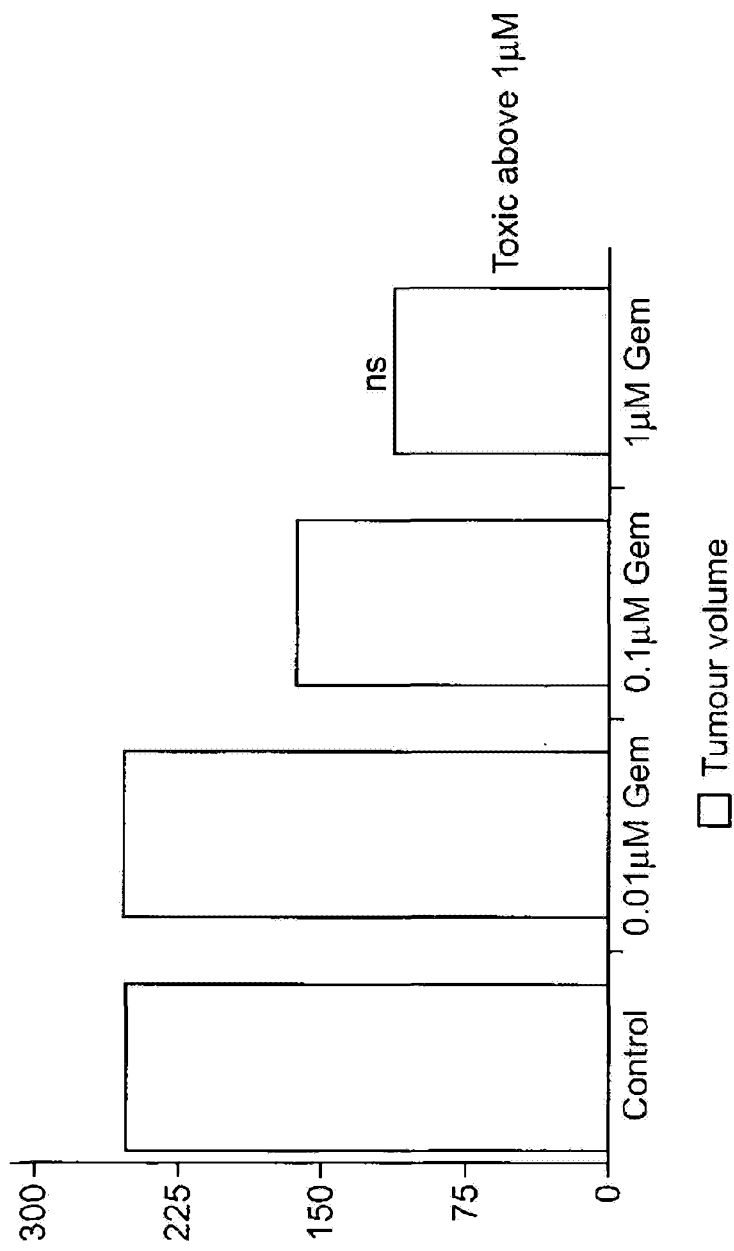


FIG. 1

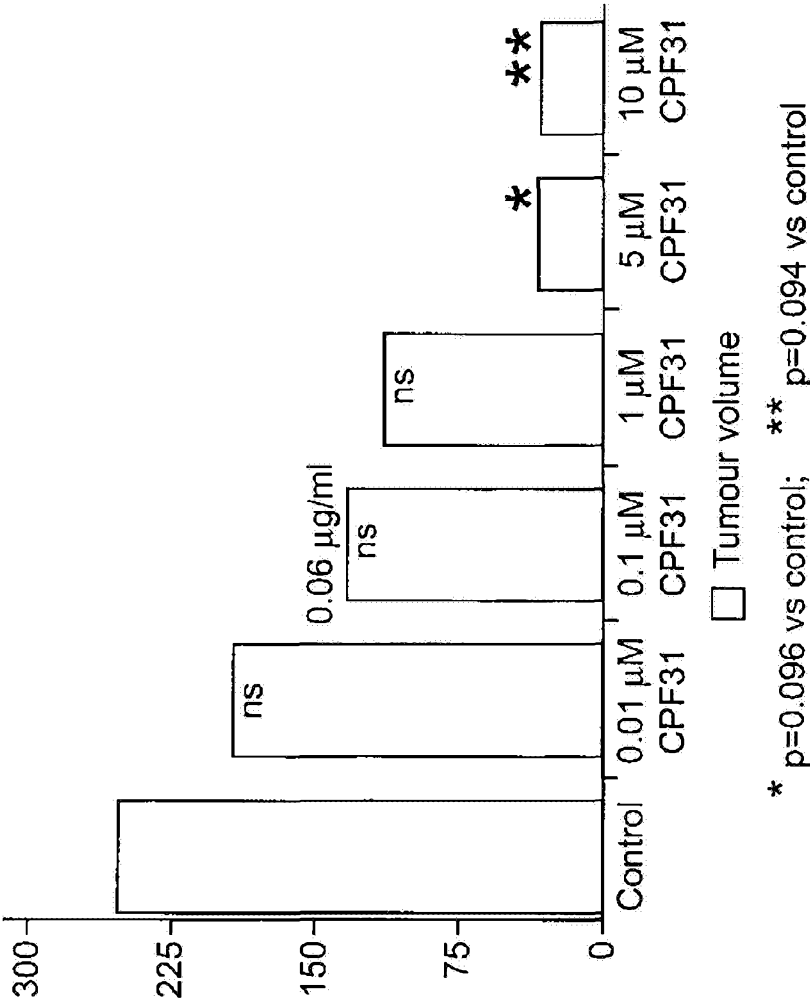


FIG. 2

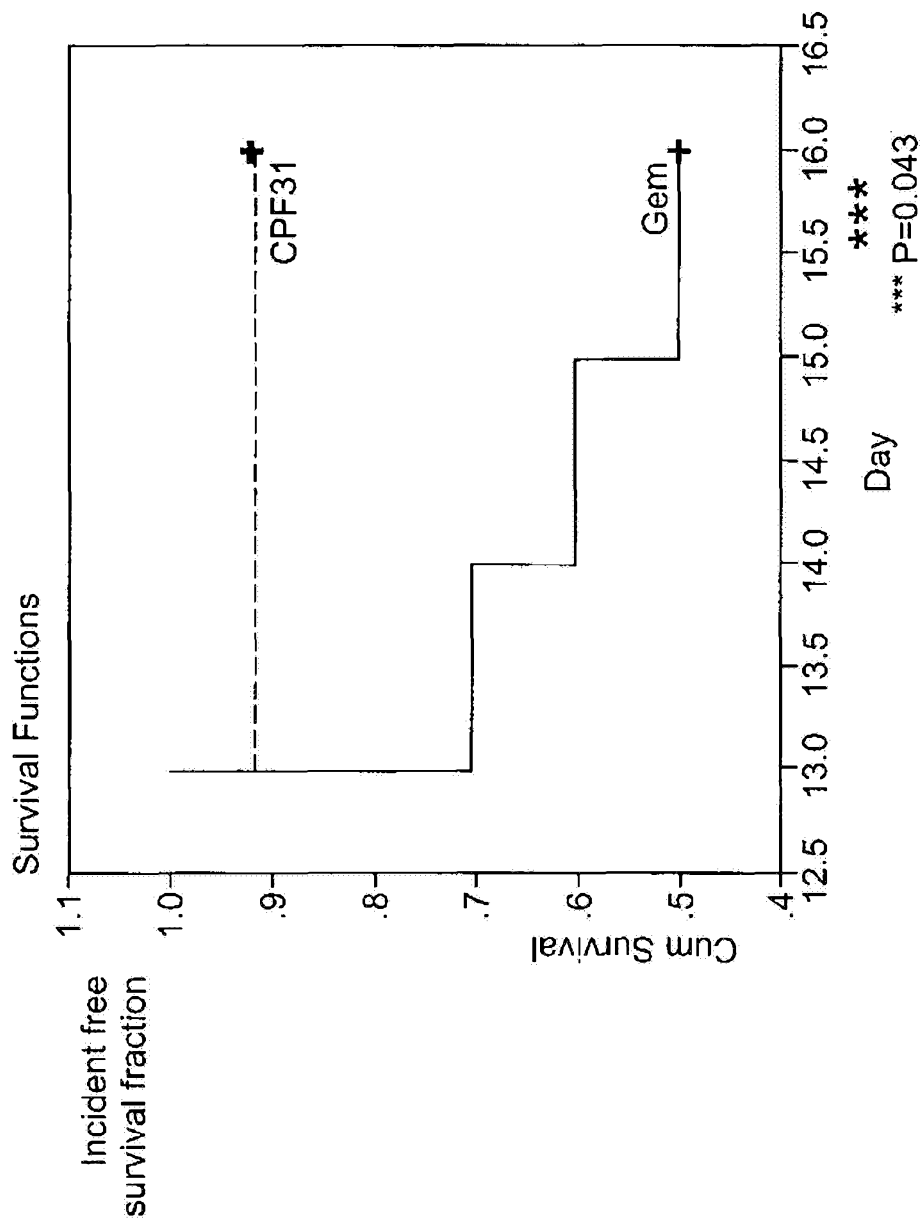
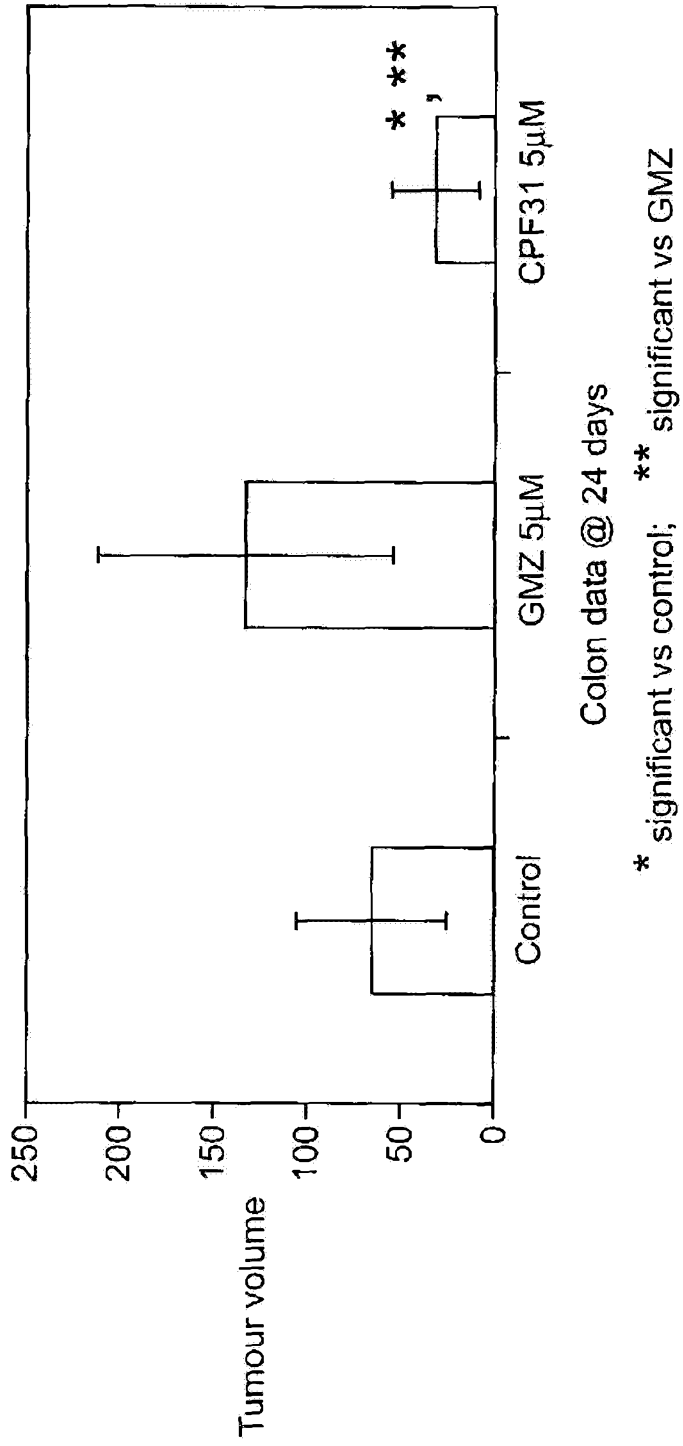


FIG. 3

**FIG. 4**