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OLIGONUKLEOTIDE ZUR MODULATION VON GENEXPRESSION UND DEREN ANWENDUNGEN
 OLIGONUCLÉOTIDES POUR MODULER L'EXPRESSION DE GÈNES ET LEURS UTILISATIONS

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Description

[0001] The present invention relates to oligonucleotides for modulating the expression of a gene, in particular for modulating a gene responsible for a pathology of genetic, tumoural or viral origin.

5 [0002] Furthermore, the present invention relates to the use of said oligonucleotides, possibly chemically modified, for the treatment and/or the diagnosis of said diseases. Oligonucleotides are short sequences of natural RNA or DNA nucleic acids or of synthetic nucleic acids, e.g. PNA (Peptide Nucleic Acid), LNA (Locked Nucleic Acid) and morpholinos.

10 [0003] Oligonucleotides have experimentally demonstrated to be very effective in modulating the expression of a gene at the level of both transcription and translation. Because of this ability, oligonucleotides represent a valid means for the treatment of numerous pathologies, in particular diseases of genetic, tumoural or viral origin.

[0004] Modulating the expression of a gene can imply inhibiting or activating it.

15 [0005] For example, oligonucleotides are known which are capable of inhibiting the transcription of a gene by forming a complementary bond with the antisense strand of the gene (Hélène C, *Bioch Bioph Acta* 1990, 1049(2):99-125) or modifying the state of the chromatin in the regulatory regions of the gene of interest (Rossi JJ, *Nat Chem Biol* 2007, 3(3):136-7). Other oligonucleotides are capable of inhibiting gene translation, for example, via a complementary bond with the target Messenger RNA (mRNA). In the case of single-stranded oligonucleotides, this bond causes enzymatic degradation of the mRNA by the RNase H complex. In the case in which the oligonucleotides are molecules of double-stranded "interfering" RNA, the complementary bond of the oligonucleotides with the target mRNA causes degradation of the messenger by the "slicer" enzyme of the RISC complex. In the latter case, the oligonucleotides can also be 20 oligonucleotides equivalent to an endogenous microRNA capable of associating, by virtue of imperfect complementarity, with the 3'UTR region (3' untranslated region) of the target mRNA, causing the translation of that mRNA to be blocked.

25 [0006] Oligonucleotides can also induce the activation of a gene or an increase in the transcription thereof, for example via a complementary bond with long antisense non-coding RNA (Morris KV, *Epigenetics*, 2009, 4(5): 296-301), or by inhibiting the complementary microRNA, with a consequent increase in the translation of the target mRNA of the micro-RNA.

30 [0007] Oligonucleotides can be chemically modified with the aim of increasing their effectiveness in therapeutic and/or diagnostic terms. For example, an oligonucleotide can be modified in order to improve its specificity and/or the force with which it pairs with the complementary sequence, or else an oligonucleotide can be modified in order to make it less sensitive to enzymatic degradation, to improve its pharmacokinetic/pharmacodynamic profile, or to facilitate its passage through cell membranes.

35 [0008] In addition to oligonucleotides of natural nucleic acids (i.e. short sequences of DNA and/or RNA), there exist oligonucleotides of synthetic nucleic acids, e.g. peptide nucleic acids (PNA) and locked nucleic acids (LNA), which have been greatly studied and characterized, above all with the aim of modulating the expression of a gene by means of a an antigene strategy (i.e. designed to strike the gene directly). PNA and LNA oligonucleotides, like all modified oligonucleotides, are in general much more chemically stable than DNA or RNA oligonucleotides. Their stability can be further improved by synthesizing chimera oligonucleotides. A chimera oligonucleotide is, for example, an oligonucleotide sequence in which both classic monomers (deoxyribonucleotides or ribonucleotides) and synthetic nucleobases (monomers), for example LNA monomers, are inserted.

40 [0009] LNAs are used with an antisense strategy to silence genes by inhibiting the transcript of the target gene (Braasch DA, *Nucl Acids Res*, 2002, 30(23):5160-7). Alternatively, LNA oligonucleotides can also be used in an antigene strategy, as was done by Smith and colleagues (Ge R, *Faseb J*, 2007, 1902-14), who designed the oligonucleotide sequence in such a way as to enable interaction on both strands of the nucleic acid by means of a "strand invasion" mechanism, so as to form a "Z-shaped" structure (defined as "Zorro" oligonucleotides). PNA oligonucleotides are enzymatically more stable when compared with other oligonucleoside structures. PNAs can bind to double-stranded DNA (DNAds) through "strand invasion", or can pair, in a complementary manner, with a molecule of single-stranded DNA (DNAss), or else they can bind to strands of RNA, giving rise to a hybrid double-helix PNA/DNA or PNA/RNA structure which is thermodynamically very stable compared to "homoduplex" structures (such as a DNA/DNA double strand).

45 [0010] PNAs represent a highly advantageous system for modulating the expression of a gene, above all using an antigene strategy. In fact, it has been demonstrated that PNAs exhibit high specificity for the target sequences and thus enable the expression of the protein to be inhibited in an efficient manner.

50 [0011] Therefore, PNAs represent a promising therapeutic approach for the treatment of genetic or viral diseases.

[0012] The only disadvantage of PNAs is the fact that they have a limited ability to pass through cell membranes. However, this limitation has been solved by conjugating oligonucleotides in general, and PNAs in particular, with molecules capable of rendering the passage through cell membranes more effective (carriers).

55 [0013] In fact, oligonucleotides, and PNAs specifically, can in general be administered using carriers (or "tags") conjugated to them, for example peptide sequences with a length varying from 1 to 30 amino acids.

[0014] One particular application of oligonucleotides regards modulating the expression of the genes which are activated or repressed in tumours.

[0015] It is well known that tumours are caused by the dysregulation of various genes. Usually, the damage affects proto-oncogenes (or also simply oncogenes) like the MYC genes (tra cui MYC, MYCN, MYC1), survivin (BIRC5), BCL2, PLK4, ALK and PKM2, which are activated or overexpressed in the tumour. Moreover, in a tumour, antitumour or oncosuppressor genes such as caspase-8 and RASSF1 are usually also inactivated.

5 [0016] In particular, the oncogenes of the MYC family are involved in the development of numerous human tumours and are among the genes most responsible for the onset and progression of neoplasms. Amplification and/or overexpression of these genes is almost always associated with tumours, both of the paediatric type (e.g. neuroblastoma, medulloblastoma and rhabdomyosarcoma) and adult type (e.g. small cell lung cancer, or glioblastoma)(Pession A, Cur Cancer Drug Target, 2005, 5(4):273-83). In fact, they act by means of mechanisms which are fundamental for tumour 10 growth, such as induction of cell proliferation, resistance to apoptosis, formation of metastasis and resistance to chemotherapy drugs.

[0017] Many oligonucleotides with an antitumour effect are known in the literature.

15 [0018] For example, there exist oligonucleotides which are directed against the MYC, MYCN, BCL2, BIRC5 genes in antisense strategies (EV Prochownik, Exp Rev Antic Ther 2004, 4(2):289-302; Felsher DW, Drug News Persp, 2003, 16(6):370-4; CF Bennet, Exp Opin Investig Drugs, 1999, 8(3):237-53) or oligonucleotides directed against MYCN and MYC with an antigene effect (LC Boffa, Oligonucleotides 2005, 15(2):85-93).

20 [0019] DNA-based phosphorothioate antisense oligonucleotides have also been generated with the aim of inhibiting the translation of the MYCN gene into neuroblastoma cells (Berkhout CA, JNCI, 2003, 95(18):1394-403) and antisense oligonucleotides have been generated through "small interfering RNA (siRNA)" to inhibit the translation of the MYCN 25 gene into neuroblastoma cells (Kang JH, Bioch Bioph Res Com, 2006, 351(1):192-197). WO 03/070917 WO 02/092617, WO 2009/009739, and WO 01/61030 disclose antisense and/or siRNA compounds targeted to the c-myc gene. WO 2004/096826 discloses inhibition of N-MYC expression by PNA anti-gene oligonucleotides. However, there is still a strong felt need to identify further oligonucleotides capable of modulating the expression of a gene in an increasingly specific and selective manner so as to be able to have a potent therapeutic and/or antitumour effect.

30 [0020] In order to obtain oligonucleotides utilizable as therapeutic and/or diagnostic means, it is necessary to identify the sequences of a target gene, or a target messenger RNA and/or their respective regulatory sequences which are capable of determining a significant and selective modulation effect on the expression of the gene itself in terms of both transcription and translation.

35 [0021] Thus there is also a felt need to define the general rules governing the process of identifying - within the sequence of a gene, or the sequence of a messenger RNA or the regulatory sequences thereof - the oligonucleotide sequences that are most promising for the purposes of modulating the transcription/translation of the gene itself.

40 [0022] The needs in this sector as just described are met by the present invention, which, refers to a single strand anti-gene oligonucleotide complementary to the anti-sense DNA strand of a target gene, comprising 6-30 nucleotides, preferably 12-24 nucleotides, said oligonucleotide being characterized by a sequence comprising at least three groups of at least two consecutive guanines. Preferably the groups of at least two consecutive guanines are continuous with each other or spaced apart by at least one nucleotide, said nucleotide not being a guanine, more preferably the groups of at least two consecutive guanines are at least four, five or six in number. According to a preferred embodiment said oligonucleotide is conjugated with a carrier sequence at the 3' and/or 5' end of said oligonucleotide, preferably a carrier selected from the group consisting of: SEQ ID NO: 47-56. According to a preferred embodiment said oligonucleotide is at least one molecule of natural nucleic acid, preferably DNA or RNA, possibly chemically modified, or synthetic nucleic acid, preferably PNA, LNA or morpholino, possibly chemically modified, or a combination of said natural nucleic acid and said synthetic nucleic acid. Preferably 1) said at least one DNA molecule comprises at least one LNA, methylphosphonate-LNA, BNA, RNG, DNG, GNA, UNA, ENA, ANA, F-ANA, PNA, G-PNA or morpholino nucleotide; or comprises a 2'-O-Methyl, 2'-O-Methoxyethyl or 2'-fluoro RNA nucleotide; or 2) said at least one RNA molecule comprises at least one RNA nucleotide selected from among: a 2'-O-Methyl nucleotide, a 2'-O-Methoxyethyl nucleotide, a 2'-fluoro nucleotide; or at least one nucleotide of a nucleic acid selected from among: LNA, Methylphosphonate LNA, BNA, RNG, DNG, GNA, UNA, ENA, ANA, F-ANA, PNA, G-PNA or morpholino. More preferably said at least one DNA molecule or said at least one RNA molecule comprises at least one nucleotide modified at the level of the phosphodiester bond as a phosphorothioate bond. According to a preferred embodiment said PNA has a modified backbone wherein the alpha carbon has the side chain of arginine or lysine as a substituent. According to a preferred embodiment oligonucleotide is directed against at least one gene responsible for a disease of genetic or viral origin or against at least one tumour gene, wherein said gene is preferably selected from the group consisting of: at least one gene belonging to the MYC family, preferably MYC, MYCN or MYCL1, BIRC5, BCL2, PLK4, ALK, PKM2, CASP8 and RASSF1. According to a preferred embodiment the oligonucleotide is selected from SEQ ID NO: 2, 4, 5, 6, 68 and 69 directed against the MYCN gene. In another preferred embodiment, the oligonucleotide is selected from SEQ ID NO:

55 70-74 directed against the gene MYC; SEQ ID NO: 75, 76 directed against the gene BIRC5; SEQ ID NO: 77-79 directed against the gene ALK; SEQ ID NO: 80-82 directed against the gene BCL2; or SEQ ID NO: 83, 84 directed

against the gene PLK4. A further aspect of the present invention refers to a composition comprising at least one oligonucleotide mentioned above and at least one pharmacologically acceptable excipient. Preferably the composition comprising at least one oligonucleotide mentioned above, in combination with a selected compound with pharmacological action, preferably in the group consisting of: NGF, somatostatin, retinoic acid, actinomycin D, asparaginase, bleomycin, busulphan capecitabine, carboplatin, cyclophosphamide, cyclosporine, cisplatin, cytarabine, clorambucil, dacarbazine, daunorubicin, docetaxel, doxorubicin hydrochloride, epirubicin hydrochloride, etoposide, fludarabine phosphate, fluorouracil, gemcitabine, idarubicin hydrochloride, hydroxyurea, ifosfamide, irinotecan hydrochloride, melphalan, mercaptopurine, methotrexate, mitomycin, mitoxantrone, oxaliplatin, paclitaxel, procarbazine, raltitrexed, streptozocin, tegafur-uracil, temozolamide, thioguanine, thiotepa, topotecan, vinblastine, vincristine sulphate, vindesine and vinorelbine; wherein said combination is preferably: SEQ ID: NO 5 and carboplatin, or etoposide or cisplatin or vincristine.

[0023] A further aspect of the present invention refers to the oligonucleotide or to the composition disclosed above for use as a medicament.

[0024] A further aspect of the present invention refers to the oligonucleotide or to the composition disclosed above for the treatment of a tumour wherein said tumour is an adult or a pediatric tumour, preferably selected from the group consisting in: neuroblastoma, retinoblastoma, medulloblastoma, ependymoma, pheochromocytoma, embryonal carcinoma, germ cell tumour, alveolar rhabdomyosarcoma, embryonal rhabdomyosarcoma, Wilms' tumour, clear cell sarcoma of the kidney, synovial sarcoma, hepatoblastoma, acute lymphoid leukaemia, chronic lymphoid leukaemia, acute lymphoblastic leukaemia, chronic lymphoblastic leukaemia, Burkitt's lymphoma, acute myeloid leukaemia, chronic myeloid leukaemia, acute megakaryoblastic leukaemia, B chronic lymphoid leukaemia, T-cell leukaemia, lymphomas, small cell lung cancer (microcytoma), lung adenocarcinoma, squamous cell lung carcinoma, typical and atypical primary lung cancer, large cell lung carcinoma, large-cell neuroendocrine lung carcinoma, glioblastoma, hepatocarcinoma, basal cell carcinoma, ovarian tumour, breast tumour and colon cancer.

[0025] In the case of natural nucleic acids (DNA and RNA), each monomer (nucleotide) consists of a nitrogenous base, a sugar and a triphosphate. The base is selected from among adenine, guanine, thymine, cytosine and uracil (only in RNA). The sugar is deoxyribose in the case of DNA and ribose in the case of RNA. The monomers are linked in the polymer by a phosphodiester bond.

[0026] In further embodiments the at least three groups of guanines preferably comprises at least four, five or six groups of at least two consecutive guanines.

[0027] Alternatively the at least three groups of guanines preferably comprises at least one group of at least two consecutive guanines and at least two groups of at least three consecutive guanines.

[0028] Alternatively the at least three groups of guanines preferably comprises at least one group of at least three consecutive guanines and at least two groups of at least two consecutive guanines.

[0029] Alternatively the at least three groups of guanines preferably comprises at least one group of at least two consecutive guanines, at least one group of at least three consecutive guanines and/or at least one group of six consecutive guanines.

[0030] In general, the oligonucleotides of the present invention are perfectly complementary to the target sequence and, preferably, the groups of consecutive guanines can be consecutive to one another so that, for example, three groups of 2 consecutive guanines is a group of 6 consecutive guanines. Alternatively, the groups of consecutive guanines can be spaced apart by at least one nucleotide.

[0031] In general, the at least three groups of at least two consecutive guanines according to the present invention can be located near the 5' end of the oligonucleotide, or near the 3' end of the oligonucleotide, or else they can be located at the centre of the oligonucleotide sequence.

[0032] In preferred embodiments of the present invention said oligonucleotide is conjugated, preferably at its 3' and/or 5' end, with a carrier sequence, which is preferably a short amino acid sequence.

[0033] Said short amino acid sequence (carrier) preferably consists of a number of amino acids ranging from 1 to 30, preferably 1 to 10, even more preferably 1 to 7. The amino acids can be in L or D form, preferably in D form.

[0034] The carriers that are preferred for the purposes of the present invention are selected from the group consisting of: SEQ ID NO: 47 (PKKKRKV); SEQ ID NO: 48 (VKKKKKP); SEQ ID NO: 49 (KKKKKK); SEQ ID NO: 50 (PKRKRKV); SEQ ID NO: 51 (KRKRKRK) ; SEQ ID NO: 52 (KKKRKV) ; SEQ ID NO: 53 (PKKKRK) ; SEQ ID NO: 54 (KKKRK); SEQ ID NO: 55 (RRRR) and SEQ ID NO: 56 (PKKKRKVHHHHH).

[0035] The carrier that is particularly preferred for the purposes of the present invention is the peptide having the SEQ ID NO: 47.

[0036] In the context of the present invention, "carrier" means a peptide capable of favourably modifying the pharmacokinetic and/or pharmacodynamic profile and/or cellular and/or nuclear penetration of an oligonucleotide.

[0037] In the context of the present invention, "modulating the expression of a gene" means inhibiting or activating (increasing) the expression of a gene. Said inhibition or activation of (increase in) gene expression can occur on a

transcription or translation level.

[0038] The inhibition or activation of gene expression can be achieved on a transcription level by means of oligonucleotides which act with an antigene mechanism (or antigene oligonucleotide, i.e. directed against the antisense strand of the gene, that is, antigene strategy). The parameters or rules or prerequisites that an oligonucleotide must satisfy in order to effectively modulate the expression of a gene, identified by the Applicant and stated above, are valid for any gene whatsoever and can thus be applied, for example, for the purpose of identifying sequences of oligonucleotides which are capable of modulating the expression of the gene(s) responsible for diseases of genetic and/or viral origin or the genes involved in the onset of tumour pathologies.

[0039] The oligonucleotides thus identified can be used, preferably as drugs, in therapeutic approaches for the treatment of specific genetic, viral or tumoural diseases.

[0040] Alternatively, the oligonucleotides can be utilized for diagnostic purposes.

[0041] In fact, the disclosure relates to the use of the oligonucleotides of the invention, possibly chemically modified, for therapeutic and/or diagnostic purposes.

[0042] The oligonucleotides of the present invention are short oligonucleotides of 6-30, preferably 12-24 residues (nucleotides or monomers). The oligonucleotides can consist of a natural nucleic acid base, for example DNA or RNA, or a synthetic nucleic acid base, for example PNA, LNA or morpholino. Alternatively, the oligonucleotides can comprise a combination of DNA, RNA and/or synthetic nucleic acids, preferably PNA or LNA (hybrid or chimeric oligonucleotides).

[0043] In some embodiments of the present invention, the oligonucleotides can be chemically modified, for example for the purpose of improving their therapeutic and/or diagnostic effectiveness.

[0044] In preferred embodiments of the present invention, the oligonucleotides can be PNA molecules with a backbone in which the carbon in the alpha position ($C_{\alpha(\omega)}$) is bound to substituents other than the typical hydrogen atom of glycine. For example, instead of the side chain of glycine the side chain of another amino acid of natural or synthetic origin can be used, preferably selected from the group consisting in: arginine, lysine, histidine, leucine, isoleucine, tyrosine, asparagine, serine, threonine, glutamine, valine, alanine, cysteine, methionine, phenylalanine, glutamate, aspartate, proline, tryptophan and ornithine. Said amino acid can be of the dextrorotatory configuration (D) or the levorotatory configuration (L).

[0045] In some instances, said "small interfering RNA" comprises RNA monomers (ribonucleotides) and at least one modified monomer in the ribose 2' position, preferably a 2'-O-methoxyethyl, 2'-O-methyl or 2'-fluoro monomer; or said "small interfering RNA" comprises RNA monomers (ribonucleotides) and at least one monomer of a synthetic nucleic acid preferably selected from among: LNA, Methylphosphonate LNA, BNA (bridged nucleic acid), UNA (unlocked nucleic acid), ENA (ethylene-bridged nucleic acid), ANA (arabinose nucleic acid) and F-ANA (fluoro-arabinoside nucleic acid).

[0046] In preferred instances, said "small interfering RNA" is designed in such a way that at the ends of the complementary double strand, only one of the two strands has at least one, preferably two, monomers of natural or synthetic nucleic acids which protrude, i.e. are not paired. In further embodiments, natural or synthetic protruding nucleic acids are preferably selected from among: a 2'-O-methoxyethyl, 2'-O-methyl and 2'-fluoro monomer or a monomer of: LNA, Methylphosphonate LNA, BNA, UNA (unlocked nucleic acid), ENA (ethylene-bridged nucleic acid), ANA (arabinose nucleic acid) and F-ANA (fluoro-arabinoside nucleic acid).

[0047] In further embodiments, the oligonucleotides are defined as hybrid or chimeric and are single-stranded comprising RNA monomers (ribonucleotides) and LNA monomers (this oligonucleotide is represented as RNA/LNA).

[0048] Alternatively, the hybrid oligonucleotides can comprise the RNA monomers (ribonucleotides) and at least one RNA monomer selected from among: a 2'-O-Methoxyethyl (MOE) monomer, a 2'-O-methyl monomer and a 2'-fluoro monomer; or the hybrid oligonucleotides can comprise RNA monomers (ribonucleotides) and at least one monomer of a synthetic nucleic acid preferably selected from among: LNA, methylphosphonate LNA, UNA (unlocked nucleic acid), BNA, ENA (ethylene nucleic acid), ANA (arabinose nucleic acid) and F-ANA (fluoro-arabinoside nucleic acid).

[0049] In a further embodiment, the oligonucleotides based on single-stranded RNA can comprise classic ribonucleotides (i.e. not chemically modified) and ribonucleotides or deoxyribonucleotides which have been modified at the level of the phosphodiester bond, for example by means of a phosphorothioate bond, or DNG (deoxyribonucleic guanidine), RNG (ribonucleic guanidine), GNA (glycerol nucleic acid), G-PNA (gamma-PNA) or PMO (Morpholino).

[0050] In preferred embodiments of the invention, the oligonucleotides are chimeric single-stranded sequences comprising DNA monomers (deoxyribonucleotides) and LNA monomers.

[0051] For an antigene strategy, which means modulating the expression of a gene on the transcription level, one can preferably employ:

- PNA-based oligonucleotides, optionally conjugated with a carrier (generally consisting of 1 to 30 residues), preferably at the 3' end and/or 5'; or
- PNA-based oligonucleotides, said PNA comprising at least one alpha carbon ($C_{\alpha(\omega)}$) of the backbone with a substituent other than the H atom of the canonical glycine; or
- single-stranded oligonucleotides comprising RNA monomers (the classic ribonucleotides) and optionally at least

one modified nucleotide (monomer) (for example a 2'-O-Methyl RNA monomer, a 2'-Fluoro RNA monomer), or at least one monomer of a nucleic acid selected from among: LNA, methylphosphonate LNA, BNA, UNA, GNA, ANA, FANA, ENA, DNG and RNG, or a ribonucleotide modified at the level of the phosphodiester bond; or

- 5 • mutually complementary double-stranded RNA-based oligonucleotides (siRNA); or
- partially complementary double-stranded chimeric oligonucleotides comprising RNA monomers (the classic ribonucleotides) and at least one LNA monomer; or
- double-stranded chimeric oligonucleotides comprising RNA monomers (the classic ribonucleotides) and at least one 2'-O-(2-Methoxyethyl) RNA monomer; or
- 10 • single-stranded chimeric oligonucleotides comprising DNA monomers (the classic deoxyribonucleotides) and at least one LNA monomer; or
- oligonucleotides comprising DNA monomers (the classic deoxyribonucleotides) and at least one 2'-Fluoro RNA monomer or at least one monomer of a nucleic acid selected from among: LNA, Methylphosphonate LNA, BNA, UNA, GNA, ENA, ANA, FANA, DNG and RNG.

15 [0052] Preferably, the oligonucleotides of the present invention are directed against a gene involved in the development of a disease of genetic and/or viral origin or a tumour. Said gene is preferably selected from the group consisting of: genes of the MYC family (preferably MYC, MYCN, MYCL1), survivin genes (BIRC5), BCL2, PLK4, ALK, PKM2, caspase-8 and RASSF1.

20 [0053] In particular, the oligonucleotides of the present invention are directed against the genes of the MYC family, preferably against MYCN. Also disclosed are oligonucleotides selected from the group consisting of: SEQ ID NO: 2-15, 66-84, SEQ ID NO: 24, 25, 31 and 32, the pair of complementary oligonucleotides having SEQ ID NO: 26 and 57, the pair of complementary oligonucleotides having SEQ ID NO: 27 and 58, the pair of complementary oligonucleotides having SEQ ID NO: 28 and 59, the pair of complementary oligonucleotides having SEQ ID NO: 29 and 60, the pair of complementary oligonucleotides having SEQ ID NO: 30 and 61, the pair of complementary oligonucleotides having SEQ 25 ID NO: 33 and 62, the pair of complementary oligonucleotides having SEQ ID NO: 34 and 63, the pair of complementary oligonucleotides having SEQ ID NO: 35 and 64 and the pair of complementary oligonucleotides having SEQ ID NO: 36 and 65.

30 [0054] In some embodiments, said oligonucleotides are PNA oligonucleotides, preferably said PNAs are directed against MYCN.

35 [0055] In some instances the PNAs are selected from the group consisting of: SEQ ID NO: 2-15.

[0056] In further instances the PNAs are selected from the group consisting of: SEQ ID NO: 66-84.

[0057] In further instances the PNAs are selected from the group consisting of: SEQ ID NO: 2-15, 66-84.

40 [0058] Preferably, the PNA oligonucleotides are selected from the group consisting of: SEQ ID NO: 2-13, more preferably SEQ ID NO: 2-8, even more preferably SEQ ID NO: 2-6. The PNA oligonucleotide that is particularly preferred for the purposes of the present invention is SEQ ID NO: 5. Preferably, SEQ ID NO: 5 is conjugated at the 5' or 3' end with SEQ ID NO: 47. More preferably, SEQ ID NO: 47 consists of amino acids in D form.

[0059] The PNAs SEQ ID NO: 2-15 are preferably directed against MYCN and, more preferably, they modulate the expression of MYCN with an antigene strategy.

45 [0060] The PNAs SEQ ID NO: 66-69 are also preferably directed against MYCN and, more preferably, they modulate the expression of MYCN with an antigene strategy.

[0061] The PNAs SEQ ID NO: 70-74 are preferably directed against MYC and, more preferably, they modulate the expression of MYC with an antigene strategy.

[0062] The PNAs SEQ ID NO: 75, 76 are preferably directed against BIRC5 and, more preferably, they modulate the expression of BIRC5 with an antigene strategy.

50 [0063] The PNAs SEQ ID NO: 77-79 are preferably directed against ALK and, more preferably, they modulate the expression of ALK with an antigene strategy.

[0064] The PNAs SEQ ID NO: 80-82 are preferably directed against BCL2 and, more preferably, they modulate the expression of BCL2 with an antigene strategy.

[0065] The PNAs SEQ ID NO: 83, 84 are preferably directed against PLK4 and, more preferably, they modulate the expression of PLK4 with an antigene strategy.

[0066] In further embodiments, said oligonucleotides are DNA-LNA chimeric oligonucleotides, preferably said oligonucleotides are directed against MYCN.

[0067] In other instances DNA-LNA chimeric oligonucleotides are selected from the group consisting of: SEQ ID NO: 24 and 25.

55 [0068] SEQ ID NO: 24 and 25 are directed against the MYCN gene.

[0069] SEQ ID NO: 24 and 25 modulate the gene's expression through an antigene strategy.

[0070] In further embodiments, said oligonucleotides are single-stranded chimeric oligonucleotides comprising DNA monomers and/or at least one phosphorothioate DNA monomer. Said oligonucleotides are preferably directed against

MYCN.

[0071] A further aspect of the disclosure relates to the use of the above-described oligonucleotides for therapeutic and/or diagnostic purposes.

5 [0072] In particular, the oligonucleotides can be used individually or combined together for the treatment of diseases of genetic and/or viral origin, in particular for the treatment of genetic diseases caused either by overexpression or inhibition of a gene, i.e. genetic diseases which require the modulation of the expression of a gene which is overexpressed or inhibited.

10 [0073] The oligonucleotides disclosed can be used for the therapeutic treatment of a genetic disease, preferably selected from the group consisting of: Gorlin syndrome, Down syndrome, Feingold syndrome, Hirschsprung's disease, Von Hippel Lindau syndrome, Ataxia Telangiectasia, Li-Fraumeni syndrome, Turcot syndrome, familial tumours and Parkinson's disease.

15 [0074] Furthermore, the oligonucleotides of the invention are used for the therapeutic treatment of tumoural pathologies in children or adults. In particular, the tumours to which reference is made are caused, preferably, by the overexpression of a gene or oncogene selected from the group consisting of: MYC, MYCN, MYCL1, survivin (BIRC5), BCL2, PLK4, ALK and PKM2. Alternatively, the tumours are caused, preferably, by the inhibition (inactivation) of an oncosuppressor or anti-tumour gene and preferably selected from the group consisting of: caspase-8 and RASSF1.

20 [0075] The tumours to which reference is made are preferably selected from the group consisting of: neuroblastoma, retinoblastoma, medulloblastoma, ependymoma, pheochromocytoma, embryonal carcinoma, germ cell tumour, alveolar rhabdomyosarcoma, embryonal rhabdomyosarcoma, Wilms' tumour, clear cell sarcoma of the kidney, synovial sarcoma, hepatoblastoma, acute lymphoid leukaemia, chronic lymphoid leukaemia, acute lymphoblastic leukaemia, chronic lymphoblastic leukaemia, Burkitt's lymphoma, acute myeloid leukaemia, chronic myeloid leukaemia, acute megakaryoblastic leukaemia, B chronic lymphoid leukaemia, T-cell leukaemia, lymphomas, small cell lung cancer (microcytoma), lung adenocarcinoma, squamous cell lung carcinoma, typical and atypical primary lung cancer, large cell lung carcinoma, large-cell neuroendocrine lung carcinoma, glioblastoma, hepatocarcinoma, basal cell carcinoma, ovarian tumour, breast tumour and colon cancer.

25 [0076] Particularly preferred for the purposes of the present invention are the tumours selected from the group consisting of: neuroblastoma, retinoblastoma, rhabdomyosarcoma, Wilms' tumour, medulloblastoma, small cell lung cancer and basal cell carcinoma.

30 [0077] The subject matter of the present invention further relates to a composition comprising at least one oligonucleotide according to the present invention and at least one pharmacologically accepted excipient. Preferably, said at least one oligonucleotide is a PNA, preferably selected from the group consisting of: SEQ ID NO: 2, 4, 5, 6, 68-84 more preferably SEQ ID NO: 2, 4, 5, 6. The oligonucleotide that is particularly preferred is SEQ ID NO: 5. Preferably, SEQ ID NO: 5 is conjugated at the 5' or 3' end with SEQ ID NO: 47. More preferably, SEQ ID NO: 47 consists of amino acids in the D form.

35 [0078] Said PNA is preferably conjugated at its 5' or 3' end with a carrier which is preferably selected from the group consisting of: SEQ ID NO: 47-56.

40 [0079] The disclosure further relates to a combination comprising at least one oligonucleotide including the oligonucleotide having SEQ ID NO: 1, at least one compound, preferably at least one compound with a pharmacological effect, more preferably a chemotherapeutic agent, and, optionally, at least one pharmacologically accepted excipient.

45 [0080] In some instances said at least one compound is at least one additional antigen and/or antisense oligonucleotide, or at least one pharmacological agent, or at least one compound of biological or biotechnological origin or deriving from chemical synthesis or combinations thereof. Said compound of biological or biotechnological origin or deriving from chemical synthesis is preferably selected from the group consisting of: a monoclonal antibody, a chemotherapeutic agent, an immunomodulating agent, a growth factor, a cytokine, a peptide, an angiogenesis inhibitor, a tumour growth inhibitor, a steroid hormone and/or a non-steroid hormone and vitamins.

50 [0081] Examples of compounds that are particularly preferred for the purposes of the present invention are selected from the group consisting of: nerve growth factor (NGF), somatostatin, retinoic acid, actinomycin D, asparaginase, bleomycin, busulphan, capecitabine, carboplatin, cyclophosphamide, cyclosporine, cisplatin, cytarabine, clorambucil, dacarbazine, daunorubicin, docetaxel, doxorubicin hydrochloride, epirubicin hydrochloride, etoposide, fludarabine phosphate, fluorouracil, gemcitabine, idarubicin hydrochloride, hydroxyurea, ifosfamide, irinotecan hydrochloride, melphalan, mercaptopurine, methotrexate, mitomycin, mitoxantrone, nutline, oxaliplatin, paclitaxel, procarbazine, raltitrexed, streptozocin, tegafur-uracil, temozolomide, thioguanine, thiotepa, topotecan, vinblastine, vincristine, vindesine and vinorelbine and combinations thereof.

55 [0082] More preferably, the compounds are selected from the group consisting of: carboplatin, cisplatin, etoposide, vincristine, cyclophosphamide and combinations thereof.

[0083] The Applicant has found that the administration of at least one oligonucleotide according to the invention in concomitance with at least one compound, preferably at least one chemotherapeutic agent, as described above, makes it possible to reduce the concentration of said compound to be administered, while at the same time guaranteeing an

increase in the therapeutic effectiveness and lower toxicity.

[0084] The Applicant has found that, under these conditions, the reduction in the concentration of said compound depends on the particular pathology; in particular, the concentration of the chemotherapeutic compound depends on the type of tumour. For some tumours, such as: neuroblastoma, retinoblastoma, medulloblastoma, small cell lung cancer, Wilms' tumour, alveolar rhabdomyosarcoma and embryonal rhabdomyosarcoma, the concentration of the at least one chemotherapeutic agent, administered in combination with at least one oligonucleotide according to the present invention, can be reduced by up to 10 times while guaranteeing the same therapeutic effect as a normal dose of a chemotherapeutic agent.

[0085] Particularly effective as a pharmaceutical combination (in terms of improved therapeutic effect) is the combination of at least one PNA according to the present disclosure, preferably at least one PNA selected from the group consisting of: SEQ ID NO: 1-15, 66-84, preferably SEQ ID NO: 1-13, more preferably SEQ ID NO: 1-8, more preferably SEQ ID NO: 1-6, even more preferably SEQ ID NO: 1 and/or 5, and at least one compound, preferably a chemotherapeutic agent, more preferably selected from the group consisting of: etoposide (VP16), carboplatin, cisplatin or vincristine, cyclophosphamide and combinations thereof.

[0086] Particularly disclosed is a combination selected from among: SEQ ID: NO 1 and carboplatin, or etoposide or cisplatin or vincristine; or SEQ ID: NO 5 and carboplatin or etoposide or cisplatin or vincristine.

[0087] Said PNA is preferably conjugated at its 3' and/or 5' end with a carrier, which is preferably selected from the group consisting in: SEQ ID NO: 47-56.

[0088] The improvement effect is preferably to be found both when the combination is administered simultaneously and when said at least one compound is administered at successive times, preferably at intervals, more preferably at regular intervals of 3 hours, 6 hours, 12 hours, 24 hours, 48 hours or 72 hours.

[0089] In preferred embodiments, at least one oligonucleotide of the invention, can be administered conjugated or complexed, preferably with at least one vehicle particle, at least one vehicle polymer or at least one self-assembled vehicle oligonucleotide (also known as aptamers).

[0090] In further preferred embodiments, at least one oligonucleotide of the invention, can be conjugated or complexed and administered with at least one liposomal micelle, at least one micro-particle or at least one nanoparticle such as to favour permeation of the target tissue.

[0091] Said particle, usually spherical in shape and used as a means of specific delivery, can be formulated with many different chemical compounds. For example, said particle can be a formulation or co-formulation of polymeric compounds such as: chitosan, hyaluronic acid, polyethylene glycol (PEG), polyethyleneimine (PEI), polylactic acid (PLA), poly(lactic-co-glycolic acid) (PLGA), hydroxyapatite (HAP), polyunsaturated fatty acids, saturated fatty acids, cationic lipids, HAP-PLA, HAP-PLA/PGA and derivatives thereof. In further preferred embodiments, at least one oligonucleotide of the invention can be administered conjugated or complexed, preferably with at least one particle of the previously described type, and at least one ligand or at least one portion of a ligand for a specific receptor of the target cells (such as, for example, GD2, folic acid, TRAIL, NGF), either of chemical or biotechnological origin, may be present in the polymeric membranes as an adjuvant, useful for favouring the internalization of said oligonucleotide of the invention in the target cells.

[0092] In further preferred embodiments, at least one oligonucleotide of the invention can be conjugated with at least one ligand or a portion of a ligand (such as, for example, GD2 (ganglioside GD2), folic acid, TRAIL (TNF-RELATED APOPTOSIS-INDUCING LIGAND), NGF (Nerve Growth Factor) specific for a receptor of the target cells.

[0093] In further preferred embodiments, at least one oligonucleotide of the invention, can be administered, on its own or in a combination, in association with at least one further medical application in order to enhance their effectiveness, preferably also by facilitating permeation of the target cells and/or tissues.

[0094] Said medical application is preferably selected from the group consisting of: oxygen therapy, magnetotherapy, thermotherapy, electrostimulation, ultrasound, radiotherapy, chemotherapy and phototherapy.

Example 1

[0095] Chemical synthesis of the oligonucleotides.

[0096] The chemical synthesis of the oligonucleotides is based on the use of DNA nucleoside phosphoroamidites modified with a protecting group, 4,4'-dimethoxytrityl (DMTr) on 5'-OH and β -cyanoethyl on the 3'-phosphate group; protecting groups are also used for the primary amines (nucleobase heterocycles), which are otherwise too reactive.

[0097] The chemical synthesis of DNA oligonucleotides takes place in a 3'-5' direction. Use is made of a CPG (acronym of controlled pore glass) resin or a polystyrene support, functionalized with the first nucleotide base. The synthesis begins with a step of deprotecting the 5'-dimethoxytrityl group using a solution of 3% trichloroacetic acid (TCA) in dichloromethane (DCM). This is followed by activation, using ethylthiotetrazole (ETT) or benzylthiotetrazole (BTT) 0,3M, of the phosphoroamidite corresponding to the second base to be inserted in sequence, which will then be coupled with the previously deprotected 5'OH, thereby forming a phosphodiester bond.

[0098] The next step is "capping", which serves to acetylate the 5'OH groups that have not reacted. Capping is carried out using 2 solutions, one containing tetrahydrofuran (THF)/lutidine/acetic anhydride (8:1:1) and the other containing a 10% solution of methylimidazole in THF. The unstable trivalent bond of the phosphite triesters is stabilized by iodine in a THF/pyridine solution which oxidizes them to pentavalent phosphodiesters.

5 [0099] After oxidization, the cycle is repeated, starting with detritylation of the second unit introduced and so forth. This cycle is repeated for the number of times necessary, depending on how many bases it is desired to insert in sequence. Finally, the final 5'-DMTr group is removed by means of a treatment with an acid at room temperature.

10 [0100] Depending on the protecting groups present on the bases (which in turn depends on the chemistry of the bases selected, PTO, 2'OMe, etc.) it can be left at 55 °C for 16 hours with ammonium hydroxide or at 55 °C for 35 minutes with an ammonium hydroxide/methylamine (AMA) solution in order to deprotect the phosphors by β -elimination of the cyanoethyl groups and remove the protecting groups on the nucleobase heterocycles.

15 [0101] Alternatively, the 5'-DMTr group can be maintained throughout the phase of analysis (HPLC, MS) and preparative chromatography in order to better purify the final product from the by-products and finally removed by means of a treatment with acetic acid

[0102] The chemical synthesis of RNA oligonucleotides differs from that of DNA oligonucleotides because of the 2'OH group present on the ribose and thus the presence of an additional protecting group for each phosphoroamidite.

20 [0103] Consequently, the synthesis of RNA oligonucleotides requires a longer coupling time and further steps to deprotect that group.

[0104] The same protocol as described above is used for the synthesis of oligonucleotides with chemically modified monomers, such as phosphorothioates (PTO), 2'0-Methyl (2'OMe), 2'Fluoro (2'-F), arabinoside nucleic acid (ANA), and locked nucleic acid (LNA).

25 [0105] The details of the specific techniques for each modified base are provided by the company the monomer molecules are purchased from (Link Technologies Ltd.).

[0106] The morpholinos were purchased from the manufacturer (Gene Tools, LLC).

25 [0107] The synthesis of PNA oligonucleotides was carried out on a 10 micromole scale, and included a purification and characterization step.

30 [0108] The synthesis of the molecule was carried out in the solid phase, using a Rink Amide-Chemmatrix® resin and a Syro automatic synthesizer (MultiSynTech). The first monomer of the synthesis is manually bound to the resin. Each automatic synthesis cycle is divided into three steps. The first step is deprotection, carried out using a solution of 20% piperidine in DMF (dimethylformamide).

35 [0109] The second step is the coupling reaction between the entering monomer and the growing chain. This reaction is carried out by adding 5 0.22M equivalents (eq) of monomer (FMOC-PNA-G(Bhoc)-OH, FMOC-PNA-A(Bhoc)-OH, FMOC-PNA-C(Bhoc)-OH, FMOC-PNA-T-OH) in NMP (N-methylpyrrolidone) and 4.5 0.32M eq of an activator in DMF (in this case HATU) in an alkaline environment with an 8% solution of 2,6-lutidine and DIPEA (N,N-diisopropylethylamine) in DMF. The coupling reaction is repeated, in duplicate, at the point of attachment of the first and second monomer on the preloaded resin, in the passage from the peptide chain to the PNA one, and on the last monomer.

40 [0110] The third step is the "capping" reaction, which serves to block, by acetylation, the sites that have not reacted during the coupling step. The reaction is achieved using a solution containing 6% 2,6-lutidine and 5% acetic anhydride in DMF. Upon completion of the synthesis, the molecules are removed from the solid support.

[0111] This reaction is obtained with a solution of TFA (trifluoroacetic acid) and meta-cresol in 4:1 ratios.

[0112] The molecule thus obtained is collected by precipitation in diethyl ether.

45 [0113] Once recovered in water, it is purified in HPLC. The column used for purification is a C18 300A 5u Jupiter (© Phenomenex, Inc.). Purification is carried out using a linear gradient from 100% A (water 95%; acetonitrile 5%; 0.1% TFA)- 0% B (water 60%; acetonitrile 40%; 0.1% TFA) to 60% A-40% B in 30 minutes. The complete gradient used is 0-5 minutes 0% B; 5-35 minutes 40% B; 35-37 100% B; 37-42 100% B; 42-44 0% B.

[0114] Finally, the purified product is analyzed by ESI mass spectroscopy (© Waters).

Antigene oligonucleotides for blocking gene transcription.

50 [0115] For the purpose demonstrating that the oligonucleotides of the invention, selected and designed according to the parameters described in the present invention, work to selectively block the transcription of a gene, PNA-based oligonucleotides directed against the MYCN gene were designed and synthesized; they are shown in Table 1.

TABLE 1

| SEQ ID NO | PNA SEQUENCE | % GC | mRNA Inhibition (%) | Cell prolif. (%) | Phoenix prolif. (%) | NIH-3T3 prolif. (%) |
|---------------|-----------------------------------|------|---------------------|------------------|---------------------|---------------------|
| SEQ ID NO: 1 | ATGCC <u>GGG</u> CATGATCT | 56.3 | 42 | 70 | 100 | 100 |
| SEQ ID NO: 2 | <u>GGG</u> TGGATGC <u>GGGGGG</u> | 81.3 | 75 | 32 | 100 | 100 |
| SEQ ID NO: 3 | GATGC <u>GGGGGG</u> CTCCT | 75 | 68 | 41 | 100 | 100 |
| SEQ ID NO: 4 | GTC <u>GGCGGGAGG</u> TAAG | 68.8 | 65 | 48 | 100 | 100 |
| SEQ ID NO: 5 | GCT <u>GGGTGG</u> ATGC <u>GGG</u> | 75 | 62 | 53 | 100 | 100 |
| SEQ ID NO: 6 | <u>TGG</u> ACGCG <u>CTGGGTGG</u> | 75 | 60 | 54 | 100 | 100 |
| SEQ ID NO: 7 | CGCG <u>CTGGGTGG</u> ATGC | 75 | 58 | 57 | 100 | 100 |
| SEQ ID NO: 8 | GT <u>CTGGACGCGC</u> T <u>GGG</u> | 75 | 54 | 59 | 100 | 100 |
| SEQ ID NO: 9 | CC <u>CTGCAGTCGGCGGG</u> | 81.3 | 51 | 64 | 100 | 100 |
| SEQ ID NO: 10 | <u>CGGCCGCGGGGCCGCCA</u> | 93.8 | 51 | 65 | 100 | 100 |
| SEQ ID NO: 11 | <u>GGGAACTGTGTTGGAG</u> | 56.3 | 48 | 68 | 100 | 100 |
| SEQ ID NO: 12 | TGT <u>CTGGACGCGC</u> T <u>GG</u> | 68.8 | 47 | 69 | 100 | 100 |
| SEQ ID NO: 13 | ACG <u>CTCAGGGACCACG</u> | 68.8 | 48 | 66 | 100 | 100 |
| SEQ ID NO: 14 | <u>CCCGGACGAAGATGAC</u> | 62.5 | 40 | 70 | 100 | 100 |

| | | | | | | | |
|----|---------------|--------------------------|------|----|-----|-----|-----|
| 5 | SEQ ID NO: 15 | ACTGTGTT <u>GGAGCCGA</u> | 56.3 | 37 | 77 | 100 | 100 |
| 10 | SEQ ID NO: 16 | CCTGTCGTAGACAGCT | 56.3 | 14 | 90 | 100 | 100 |
| 15 | SEQ ID NO: 17 | TGTGACAGTCATCTGT | 56.3 | 10 | 98 | 100 | 100 |
| 20 | SEQ ID NO: 18 | GTGACAGTCATCTGTC | 50 | 10 | 98 | 100 | 100 |
| 25 | SEQ ID NO: 19 | GACAGTCATCTGTCTG | 50 | 5 | 100 | 100 | 100 |
| | SEQ ID NO: 20 | CGTCGATTTCTTCCTC | 50 | 5 | 100 | 100 | 100 |
| | SEQ ID NO: 21 | CTCGAGTTTGA <u>CTCGC</u> | 56.3 | 1 | 100 | 100 | 100 |
| | SEQ ID NO: 22 | GCGCCTCCCC <u>TGATTT</u> | 62.5 | 2 | 100 | 100 | 100 |
| | SEQ ID NO: 23 | ATATCCCC <u>GAGCTTC</u> | 56.3 | 2 | 100 | 100 | 100 |

[0116] The PNA oligonucleotides were tested both individually and conjugated with a carrier at the 3' and/or 5' end with the aim of favouring cell membrane permeation. In particular, the oligonucleotides were conjugated to the carboxyl terminus at 3' with the amino acid sequence SEQ ID NO: 43, i.e. proline-lysine-lysine-lysine-arginine-lysine-valine.

[0117] The oligonucleotide having SEQ ID NO: 1 is the sequence that was the subject of patent EP 1618195 and represents the control sequence and sequence used for comparison.

[0118] The oligonucleotides having SEQ ID NO: 2-15 contain a group of two consecutive guanines (SEQ ID NO: 14, 15), or a group of three consecutive guanines (SEQ ID NO: 13), or two groups of two consecutive guanines (SEQ ID NO: 12), or a group of two guanines and a group of three consecutive guanines (SEQ ID NO: 11, 10, 9, 8 and 7), or two groups of two consecutive guanines and a group of three consecutive guanines (SEQ ID NO: 6 and 4), or a group of two consecutive guanines and two groups of three consecutive guanines (SEQ ID NO: 5), or a group of six consecutive guanines (SEQ ID NO: 3), or a group of six consecutive guanines, a group of three consecutive guanines and a group of two consecutive guanines (SEQ ID NO: 2).

[0119] The groups of consecutive guanines are shown underlined in Table 1.

[0120] The oligonucleotides having SEQ ID NO: 16-23 do not have groups of consecutive guanines, being negative controls. Moreover, for the purpose of selectively modulating the transcription of a gene, oligonucleotides having SEQ ID NO: 24-25 were also designed and synthesized; these are shown in Table 2.

Table 2

| | | |
|---------------|--------------|---|
| SEQ ID NO: 24 | DNA-LNA 25-1 | <u>ATGCCGGGCATGATC</u> T |
| SEQ ID NO: 25 | DNA-LNA 25-2 | <u>ATGCCGGGCATGATC</u> T |

[0121] In particular, single-stranded chimeric oligonucleotides, comprising DNA monomers and LNA monomers (SEQ ID NO: 24 and 25) were designed and synthesized.

[0122] The DNA bases in the oligonucleotide sequence are shown as bases in boldface type, whereas the LNA monomers are underlined. Each chimeric oligonucleotide molecule was designed and synthesized by inserting the LNA

monomers spaced apart by 1, 2 or 3 DNA bases in order to avoid rapid degradation by the endogenous nucleases, as has been reported in the literature (Koch T, Biochem J 2001, 354 (Pt 3):481-4; Koji Nagahama, Bioorg Med Chem Lett, 2009, 19(10):2707-9).

5 Antisense oligonucleotides for blocking gene translation.

10 [0123] For the purpose demonstrating that the oligonucleotides of the invention, selected and designed according to the parameters described in the present invention, work to selectively block the translation of the target gene, antisense oligonucleotides having SEQ ID NO: 26-36, directed against the MYCN gene, were designed, synthesized and experimentally analyzed in vitro. They are shown in Table 3.

Table 3

| SEQ ID NO | | SENSE SEQUENCE | | ANTISENSE SEQUENCE |
|---------------------|-------------------------|---|------------------|---|
| 15 SEQ ID NO: 26 | siMYCN (795) | UGAAGAUGAUGAAG <u>AGGAA</u> | SEQ ID NO: 57 | UUCCUCUUCAUCAUC UUCA |
| 20 SEQ ID NO: 27 | siMYCN (799) | GAUGAUGAAG <u>AGGAAGA</u> UG | SEQ ID NO: 58 | CAUCUCCUCUUCAU CAUC |
| 25 SEQ ID NO: 28 | siMYCN (801) | UGAUGAAG <u>AGGAAGA</u> UGAA | SEQ ID NO: 59 | UUCAUCUUCCUCUUC AUCA |
| 30 SEQ ID NO: 29 | siMYCN (808) | <u>GAGGAAGAUGAAGAGGAAG</u> | SEQ ID NO: 60 | CUUCCUCUUCAUCUU CCUC |
| 35 SEQ ID NO: 30 | siMYCN (810) | <u>GGAAGAUGAAGAGGAAGAA</u> | SEQ ID NO: 61 | UUCUCCUCUUCAUC UUCC |
| 40 SEQ ID NO: 31 | MYCN-PTO (1) as | | | CGT<u>GG</u>GAGCAGCTCGG CAT |
| 45 SEQ ID NO: 32 | MYCN-PTO (763) as | | | CAG<u>GG</u>GTGTCCCTCTCC GGA |
| 50 SEQ ID NO: 33 | siRNA-2'-OMe-RNA 808 | GAG<u>GAAGAUGAAGAGGAAG</u> TT | SEQ ID NO: 62 | CUUCCUCUUCAUCUU CCUC |
| 55 SEQ ID NO: 34 | siRNA-2'F-RNA 808 | CAG<u>GAAGAUGAAGAGGAAG</u> UU | SEQ ID NO: 63 | GUUCCUCUUCAUCUU CCUCUU |
| 60 SEQ ID NO: 35 | siRNA-LNA 808 | GAG<u>GAAGAUGAAGAGGAAG</u> TT | SEQ ID NO: 64 | CUUCCUCUUCAUCUU CCUC |
| 65 SEQ ID NO: 36 | siRNA-ANA 808 | CAG<u>GAAGAUGAAGAGGAAG</u> AA | SEQ ID NO: 65 | GUUCCUCUUCAUCUU CCUCAA |

[0124] Complementary double-stranded oligonucleotides based on RNA (small interfering RNA (siRNA) (SEQ ID NO: 26-30) were produced to selectively modulate the translation of the target gene, MYCN.

[0125] Moreover, oligonucleotides based on DNA SEQ ID NO: 31 and 32, in which the phosphodiester bond was modified into phosphorothioate, were produced.

[0126] Double-stranded chimeric oligonucleotides: based on RNA monomers and monomers 2'0-Methyl (SEQ ID NO: 33); based on RNA monomers and 2'-Fluoro monomers (SEQ ID NO: 34); based on RNA monomers and LNA monomers (SEQ ID NO: 35) and based on RNA monomers and arabinoside RNA monomers (SEQ ID NO: 36), were also produced. In the oligonucleotide sequences SEQ ID NO: 33-36, the 2'0-Methyl, 2'-Fluoro, LNA and arabinoside (ANA) monomers are shown in boldface, whereas each group of at least two consecutive guanines is underlined.

[0127] As the siRNA of the oligonucleotides is double-stranded, the chimera was designed and synthesized in such a manner as to leave two 2'-O-Me monomers, or two 2'-Fluoro monomers, or two LNA monomers, or two RNA arabinoside monomers, both in 3' and in 5', unpaired to the complementary strand and thereby avoid rapid degradation by the cell enzymes.

5

Treatments with Oligonucleotides - QT-PCR

[0128] For the purpose of assessing the ability of the oligonucleotides of the present invention to modulate the expression of the target genes, their ability to reduce the quantity of messenger RNA was analyzed using the Real-Time PCR technique.

10

[0129] For this purpose, use was made of 24-well plates, which were seeded with 5.0x10⁴ cells with 0.3 ml of OPTI-MEM (GIBCO BRL) medium, 4% FBS and 2mM L-glutamine (experiments in triplicate) per well.

[0130] The cells were incubated for 24 hours at 37 °C, in an atmosphere containing 5% CO₂ to permit adherence to the base of the wells.

15

[0131] Each oligonucleotide analyzed, except for the PNA oligonucleotides, was incubated beforehand with 2µl of Lipofectamine 2000 (Invitrogen), with 0.3 mL of serum-free OPTI-MEM (GIBCO BRL) medium.

[0132] For each well, the oligonucleotides were analyzed at the following final concentrations:

20

- the antisense oligonucleotides siRNA and siRNA gapmer (i.e. the oligonucleotides containing one or more monomers of chemically modified nucleic acids at the 3' or 5' end, while in the central portion they have monomers of nucleic acids that have not been modified or have been modified at the level of the phosphodiester bond as a phosphorothioate bond) were used at 200nM;
- the antisense oligonucleotides containing phosphorothioate DNA monomers, the RNA antigenic oligonucleotides (agRNA) and the oligonucleotides containing DNA monomers and LNA monomers were used at 10 µM;
- the morpholino oligonucleotides were used at a concentration of 1 µM, and
- the PNA oligonucleotides were administered at a concentration of 1 µM.

25

[0133] The cells were treated with the oligonucleotides, adding FBS up to 4% 6 hours after administration of the same. After 24 hours the total RNA was extracted from each well and purified using the RNeasy Mini Kit (QIAGEN).

30

[0134] Assays were performed on 8 cell lines obtained from 5 different human tumours which are correlated with MYCN expression, i.e.:

35

- as a neuroblastoma model use was made of the cell lines Kelly, IMR-32 (where the MYCN gene is amplified and superexpressed) and SKNBE2c and LAN1 (where the MYCN gene is amplified and superexpressed and the p53 gene is mutated);
- as a rhabdomyosarcoma model use was made of the cell line RH30, where the MYCN gene is amplified and superexpressed; ;
- as a Wilms' tumour model use was made of the cell line WiT49, where the MYCN gene is amplified and superexpressed;
- as a retinoblastoma model use was made of the cell line Y79, where the MYCN gene is amplified and superexpressed; and
- as a model of small cell lung cancer use was made of the cell line H69 where the MYCN gene is amplified and superexpressed.

40

[0135] As a control, use was made of the same cell lines as listed above treated with sterile water instead of oligonucleotides. Each RNA sample was quantified (in duplicate) using a NanoDrop ND-1000 spectrophotometer (NanoDrop Technologies).

45

[0136] The first strand of cDNA was produced using the cDNA Synthesis Kit for RT-PCR (Roche). For the cDNA synthesis reaction a total of 1 µg of RNA was used. For the real-time PCR, 10 ng of cDNA in a final volume of 20 µl was used with SYBR Green Master Mix 2X (Applied Biosystems) (3 identical experiments were conducted in triplicate). The sequences and the concentrations of the primer used to carry out the Real-Time PCRs are shown in Table 4. Two housekeeping genes were used as positive controls: GAPDH and beta-actin (ACTB).

[0137] The conditions of the QT-PCR reaction were: 10 min at 95 °C, 20 sec at 95 °C, and 30 sec at 60 °C, for 50 cycles.

55

Table 4

| Primer | Sequence | Concentration | SEQ ID NO |
|------------|---------------------|---------------|---------------|
| MYCN sense | CGACCACAAGGCCCTCAGT | 300 nM | SEQ ID NO: 37 |

(continued)

| Primer | Sequence | Concentration | SEQ ID NO |
|-----------------|------------------------|---------------|---------------|
| MYCN antisense | TGACCACGTCGATTCTTCCT | 300 nM | SEQ ID NO: 38 |
| ACTB sense | GAGCACAGAGCCTCGCCTTG | 300 nM | SEQ ID NO: 39 |
| ACTB antisense | ACCATCACGCCCTGGTGCCTG | 300 nM | SEQ ID NO: 40 |
| GAPDH sense | CCAATATGATTCCACCCATGGC | 300 nM | SEQ ID NO: 41 |
| GAPDH antisense | CTTGATTTGGAGGGATCTCGC | 300 nM | SEQ ID NO: 42 |

Treatments with oligonucleotides - Cellular proliferation assay.

[0138] For the purpose of assessing the gene modulation ability of the oligonucleotides of the present invention, the effect of their administration on cell viability was determined.

[0139] For this purpose 5×10^3 cells per well were seeded into 96-well cell culture plates (experiments conducted in triplicate) with 100 μ l of OPTI-MEM (GIBCO BRL) medium containing 4% FBS and 2mM of L-glutammine.

[0140] The PNA oligonucleotides were administered at different concentrations (1 μ M-2,5 μ M-5 μ M-10 μ M) in order to observe a dose-effect relationship.

[0141] As regards all the other oligonucleotides, the concentrations at which they administered are listed in the paragraph:

Treatments with Oligonucleotides - QT-PCR.

[0142] The viability of the treated cells was determined at 48, 72, 96 and 168 hours after treatment.

[0143] Cell viability was assessed by means of an ATP-Lite assay (Luminescence ATP Detection Assay System, PerkinElmer) and is reported as the ratio between the average signal of the treated wells compared to the average value of the wells of untreated control cells. The cells were processed following the instructions provided with the kit.

[0144] The assays were performed on the same cell lines used to determine the levels of the MYCN gene messenger and listed in the paragraph: Treatments with Oligonucleotides - QT-PCR.

RESULTS

[0145] As far as the PNA oligonucleotides are concerned, the results regarding their ability to inhibit the transcription of the MYCN gene and proliferation of Kelly cells are shown in Table 1. Table 6 shows the values (in percentage form) of the proliferating Kelly cells at the different concentrations of PNA analyzed.

Table 5

| SEQ ID NO | Sequence | Cell Prolif (%) 1 μ M 72h | Cell Prolif (%) 2,5 μ M 72h | Cell Prolif. (%) 5 μ M 72h | Cell Prolif. (%) 10 μ M 72h |
|--------------|------------------------------------|----------------------------------|------------------------------------|-----------------------------------|------------------------------------|
| SEQ ID NO: 1 | ATGCCGGGCATGAT <u>CT</u> | 89 | 66 | 50 | 24 |
| SEQ ID NO: 2 | <u>GGGTGGATGCGGGG</u> <u>GG</u> | 74 | 32 | 12 | 2 |
| SEQ ID NO: 3 | GATGCGGGGGGCTC <u>CT</u> | 78 | 41 | 19 | 3 |
| SEQ ID NO: 4 | GTCGGCGGGAGGTA <u>AG</u> | 78 | 48 | 21 | 3 |
| SEQ ID NO: 5 | GCTGGGTGGATGCG <u>GG</u> | 79 | 53 | 27 | 5 |

(continued)

| SEQ ID NO | Sequence | Cell Prolif (%) 1µM 72h | Cell Prolif (%) 2,5µM 72h | Cell Prolif. (%) 5µM 72h | Cell Prolif. (%) 10µM 72h |
|--------------------|------------------------------------|-------------------------|---------------------------|--------------------------|---------------------------|
| 5 SEQ ID NO: 6 | TGGACGCGCTGGGT <u>GG</u> | 82 | 59 | 35 | 13 |
| 10 SEQ ID NO: 7 | CGCGCTGGGTGGAT <u>GC</u> | 80 | 54 | 31 | 8 |
| 15 SEQ ID NO: 8 | GTCTGGACGCGCTG <u>GG</u> | 80 | 57 | 32 | 11 |
| 20 SEQ ID NO: 9 | CCCTGCAGTCGGCG <u>GG</u> | 86 | 64 | 42 | 20 |
| SEQ ID NO: 12 | TGTCTGGACGCGCT <u>GG</u> | 84 | 60 | 40 | 16 |

[0146] The results demonstrate that the inhibition effect of these PNAs on the translated protein of the target genes is highly selective and specific. Moreover, it was observed that the arrest of growth of the tumour cells used as a model (characterized by amplification of the MYCN gene) was directly followed by apoptosis following the administration of the antigenic PNAs.

[0147] In general, the PNA antigenic oligonucleotides having SEQ ID: 2-SEQ ID NO: 13 (containing one or more groups of Gs) have greater antigenic effectiveness (i.e. inhibition of MYCN mRNA and of the proliferation of tumour cells with MYCN expression) compared to PNA oligonucleotides devoid of groups of Gs (SEQ ID NO: 16-SEQ ID NO: 23).

[0148] In particular, the results shown in Table 1 and in Table 5 demonstrate that the sequences SEQ ID NO: 2-SEQ ID NO: 13 have greater antigenic effectiveness than the sequence SEQ ID NO: 1, the subject of patent EP 1618195.

[0149] Sequences SEQ ID NO: 7-SEQ ID NO: 12 (containing two groups of two or three consecutive Gs) show greater antigenic effectiveness than the sequences SEQ ID NO: 1, and SEQ ID NO: 13-SEQ ID NO: 15 (containing only one group of two or three consecutive Gs).

[0150] Sequences SEQ ID NO: 4-SEQ ID NO: 6 (containing three groups of two or three consecutive Gs) show greater antigenic effectiveness than the sequences SEQ ID NO: 7-SEQ ID NO: 12 (containing two groups of two or three consecutive Gs).

[0151] SEQ ID NO: 3, containing a group of six consecutive Gs, shows greater antigenic effectiveness than the sequences SEQ ID NO: 1 and sequences SEQ ID NO: 4-SEQ ID NO: 12 containing one or two or three groups of Gs (in which each group consists of at most two or three consecutive Gs).

[0152] Sequence SEQ ID NO: 2, containing three groups of consecutive Gs, which, in addition to two groups of two and three consecutive Gs, also comprises a group of six consecutive Gs, shows greater antigenic effectiveness than both SEQ ID NO: 3, containing only one group of six Gs, and sequences SEQ ID NO: 1 and sequences SEQ ID NO: 4-SEQ ID NO: 12 containing one or two or three groups of Gs (in which each group consists of at most two or three consecutive Gs).

[0153] Moreover, the oligonucleotides having SEQ ID NO 1-23 were administered in lines of fibroblastoid-type cells (NIH-3T3 and Phoenix) and the results are shown in Table 1 (last two columns).

[0154] The results clearly demonstrate that the oligonucleotides analyzed are not specifically effective against and are not toxic for these cells (i.e. cells that do not express the target gene, which in this case is MYCN).

[0155] In fact, no changes were observed in cellular proliferation in these two lines of non-tumoural fibroblasts, and this result means that the PNA oligonucleotides act with a specific effect of inhibiting the expression of MYCN, whereas they do not have a non-specific, toxic effect in cells that do not express MYCN.

[0156] Therefore, it may be deduced that the PNA oligonucleotides designed on the basis of the parameters described in the present invention act specifically and effectively on the target gene and therefore on the cells which express/over-express this gene.

[0157] The PNAs having SEQ ID NO: 1-12 were also tested on different cell lines which overexpress MYCN (Kelly, SKNBE2c, RH30, WiT49, WERI-Rb1 and H69). The results are shown in Table 6 and confirm the selectivity and specificity of the inhibition effect of the PNAs on the product of protein translation of the MYCN gene.

Table 6

| SEQ ID NO | SEQ ID NO: 1 | SEQ ID NO: 2 | SEQ ID NO: 3 | SEQ ID NO: 4 | SEQ ID NO: 5 | SEQ ID NO: 6 | SEQ ID NO: 7 | SEQ ID NO: 8 | SEQ ID NO: 9 | SEQ ID NO: 10 | SEQ ID NO: 11 | SEQ ID NO: 12 |
|-------------------------|--------------|--------------|--------------|--------------|--------------|--------------|--------------|--------------|--------------|---------------|---------------|---------------|
| Kelly %mRNA | 48 | 75 | 68 | 65 | 62 | 60 | 58 | 54 | 51 | 51 | 50 | 50 |
| Kelly %cell prol. | 66 | 32 | 41 | 48 | 53 | 54 | 57 | 59 | 60 | 64 | 64 | 65 |
| SKNBE2c%mRNA | 41 | 75 | 68 | 60 | 59 | 62 | 60 | 62 | 58 | 41 | 40 | 40 |
| SKNBE2c%cell prol. | 72 | 52 | 53 | 59 | 64 | 57 | 58 | 56 | 64 | 70 | 71 | 71 |
| RH30 %mRNA | 39 | 74 | 65 | 57 | 56 | 56 | 55 | 58 | 50 | 29 | 31 | 32 |
| RH30 %cell prol. | 76 | 56 | 59 | 62 | 64 | 64 | 63 | 60 | 71 | 80 | 78 | 77 |
| WiT49 %mRNA | 62 | 78 | 75 | 73 | 69 | 68 | 70 | 72 | 66 | 60 | 60 | 62 |
| WiT49 %cell prol. | 74 | 54 | 57 | 60 | 63 | 65 | 60 | 62 | 68 | 70 | 72 | 73 |
| WERI-Rb1 %mRNA | 31 | 46 | 43 | 41 | 38 | 37 | 38 | 40 | 36 | 30 | 30 | 30 |
| WERI-Rb1 %cell prol. | 82 | 79 | 78 | 79 | 80 | 80 | 79 | 78 | 79 | 85 | 84 | 84 |
| H69 %mRNA | 40 | 58 | 55 | 55 | 54 | 55 | 55 | 56 | 50 | 41 | 41 | 40 |

| | | | | | | | | | | | | |
|----------------------------------|----|----|----|----|----|----|----|----|----|----|----|----|
| H69 %cell prol. | 77 | 61 | 63 | 66 | 68 | 69 | 67 | 66 | 70 | 79 | 78 | 78 |
|----------------------------------|----|----|----|----|----|----|----|----|----|----|----|----|

[0158] The antigenic PNA oligonucleotide having SEQ ID NO: 5 was analyzed in a more detailed manner.

[0159] In particular, modified oligonucleotides, in which the groups of consecutive guanines present in the sequence were modified (the groups of consecutive guanines are underlined, whereas the modified nucleotides are in boldface type) so as to interrupt the consecutiveness of the guanines, were synthesized and analyzed in vitro on Kelly cells (which overexpress MYCN).

[0160] The results (summarized in Table 7) clearly demonstrate that the fact of the guanines being consecutive is of fundamental importance for the purposes of the activity of modulating the gene's expression. In fact, the PNAs having SEQ ID NO: 43, in which all the groups of consecutive guanines of the PNAs having SEQ ID NO: 5 were mutated, causes the inhibitory activity of the PNAs to be lost, whereas the mutation of one or two of the three groups of consecutive guanines considerably compromises, but does not completely suppress, the inhibitory activity of the molecule.

Table 7

| SEQ ID NO | SEQUENCE | mRNA Inhibition (%) (Kelly) | (Kelly) Cell Prolif. % |
|---------------|------------------------------------|-----------------------------|------------------------|
| SEQ ID NO: 5 | GCT <u>GGG</u> TGGATG <u>CGGG</u> | 62 | 53 |
| SEQ ID NO: 43 | GCTGAG <u>T</u> CGATGCGTG | 0 | 100 |
| SEQ ID NO: 44 | GCTGAGT <u>GG</u> ATGCGTG | 25 | 89 |
| SEQ ID NO: 45 | GCT <u>GGG</u> TCGATG <u>CGGG</u> | 47 | 69 |
| SEQ ID NO: 46 | GTT <u>GGG</u> TGGATG <u>T</u> GGG | 58 | 60 |

[0161] Chimeric oligonucleotides comprising DNA monomers and LNA monomers and having the MYCN gene as their target were tested in vitro on alveolar rhabdomyosarcoma cells (RH30). The results are summarized in Table 8 and demonstrate these oligonucleotides have an intense, specific antigenic activity.

Table 8

| SEQ ID NO | NAME | SEQUENCE | mRNA Inhibition (%) | Cell Prolif. (%) |
|---------------|--------------|-----------------------------|---------------------|------------------|
| SEQ ID NO: 24 | DNA-LNA 25-1 | <u>ATGCCGGGCATGA</u> TCT | 23 | 75 |
| SEQ ID NO: 25 | DNA-LNA 25-2 | <u>ATGCCGGGCATGA</u> TCT | 24 | 78 |

[0162] The antisense oligonucleotides designed and synthesized by the Applicant according to the parameters described in the present invention were analyzed in vitro on rhabdomyosarcoma cells (RH30). The results are summarized in the Table 9 and show that the oligonucleotides are capable of inhibiting the MYCN transcript in a specific and effective manner; moreover, they are also capable of selectively inhibiting tumour cell proliferation in a more effective manner than the standard antisense oligonucleotides presently available for the MYCN gene (Chung DH, Bioch Bioph Res Commun, 2006, 351(1):192-7).

[0163] In particular, the siRNA identified by the Applicant and directed against MYCN mRNA, described in Table 11, exert an antisense activity with an inhibition of MYCN mRNA ranging from a minimum of 70% to a maximum of 85%.

Table 9

| SEQ ID NO | | SEQUENCE SENSE | SEQUENCE ANTISENSE | mRNA Inhibition (%) | Cell Prolif (%) |
|---------------|--------------|--------------------------------|-------------------------|------------------------|--------------------|
| SEQ ID NO: 26 | siMYCN (795) | UGAAGAUGAU <u>GAAGAGGAA</u> | UUCCUCUUCA UCAUCUUCA | 82 | 28 |

(continued)

| SEQ ID NO | | SEQUENCE SENSE | SEQUENCE ANTISENSE | mRNA Inhibition (%) | Cell Prolif (%) |
|-----------|------------------|---|---|------------------------|--------------------|
| 5 | SEQ ID NO: 27 | siMYCN (799) <u>GAUGAUGAAG</u> <u>AGGAAGAUG</u> | CAUCUCCUC UUCAUCAUC | 70 | 43 |
| 10 | SEQ ID NO: 28 | siMYCN (801) <u>UGAUGAAGAG</u> <u>GAAGAUGAA</u> | UUCAUCUCC UCUUCAUCA | 78 | 35 |
| 15 | SEQ ID NO: 29 | siMYCN (808) <u>GAGGAAGAUG</u> <u>AAGAGGAAG</u> | CUUCCUCUUC AUCUUCUC | 85 | 28 |
| 20 | SEQ ID NO: 30 | siMYCN (810) <u>GGAAGAUGAA</u> <u>GAGGAAGAA</u> | UUCUUCUCU UCAUCUCC | 81 | 30 |
| 25 | SEQ ID NO: 31 | MYCN-PTO (1) as | <u>CGTGGAGCAG</u> CTCGGCAT | 34 | 73 |
| 30 | SEQ ID NO: 32 | MYCN-PTO (763) as | <u>CAGGGTGTCC</u> <u>TCTCCGGA</u> | 45 | 65 |
| 35 | SEQ ID NO: 33 | siRNA-2'-OMe-RNA 808 | <u>GAGGAAGAUG</u> <u>AAGAGGAAGT</u> T | 13 | 85 |
| 40 | SEQ ID NO: 34 | siRNA-2'F-RNA 808 | <u>CAGGAAGAUG</u> <u>AAGAGGAAGU</u> U | 59 | 52 |
| 45 | SEQ ID NO: 35 | siRNA-LNA 808 | <u>GAGGAAGAUG</u> <u>AAGAGGAAGT</u> T | 34 | 68 |
| 50 | SEQ ID NO: 36 | siRNA-ANA 808 | <u>CAGGAAGAUG</u> <u>AAGAGGAAGA</u> A | 69 | 35 |

[0164] For the purpose of verifying whether the oligonucleotides of the present invention are capable of lowering the concentrations of the chemotherapeutic agents presently used in the therapeutic protocols against cancer, the same were administered in concomitance with chemotherapy drugs.

[0165] Studies were conducted on the effect of associations between PNA and chemotherapy drugs (carboplatin, etoposide (VP16), cisplatin and vincristine) on different human and mouse neuroblastoma tumour cell lines (SMS-KAN, LAN 1, IMR-32, SMS-KCN, Kelly, NHO2A, SKNBE2c).

[0166] The therapeutic schedule used in the treatments carried out provided for the cells to be treated with PNA and then afterwards, at a pre-established time (it could be 6 or 12 hours) the chemotherapeutic agent was administered.

[0167] The results demonstrate that the association of these compounds with the oligonucleotides of the invention determines, at specific concentration ranges, a greater therapeutic effect than individual treatments with the same compounds.

[0168] In particular, it may be observed that the treatment - be it concomitant and simultaneous or at different time intervals (3 hours, 6 hours, 12 hours, 24 hours, 48 hours, 72 hours, etc.) - with a combination consisting of the oligonucleotides of the invention and other compounds with a pharmacological effect serves to enhance the desired therapeutic effect.

[0169] Therefore, combining one or more of the oligonucleotides identified by the Applicant with chemotherapy drugs

makes it possible to reduce the chemotherapy drug concentration by as much as 10 times compared to the present standard therapeutic concentration while still obtaining the same effect.

[0170] In particular, the results of this study demonstrate that SEQ ID NO: 1 has a synergic effect, in terms of inhibiting cell proliferation, when it is administered in combination with vincristine, or etoposide, or carboplatin, or cisplatin, compared to the effects obtained in treatments with the oligonucleotide or with the aforesaid chemotherapeutic agents on their own.

EXAMPLE 2

[0171] For the purpose of supporting the present invention, the oligonucleotides summarized in Table 10 were also synthesized. In particular, Table 10 shows the oligonucleotide sequences, the SEQ ID NO, the percentage value of the (G+C) content and the gene the synthesized oligonucleotides are directed against. The synthesis of the oligonucleotides was achieved as in example 1.

Table 10

| | SEQ ID NO | | % GC |
|----|---------------|---------------------------|------|
| 20 | SEQ ID NO: 66 | TCGGGAGCAGTGGGCA | 68.8 |
| | SEQ ID NO: 67 | GC <u>GGGTCGCGGGC</u> ACG | 87.5 |
| | SEQ ID NO: 68 | T <u>GGAGGTCGGCGCCG</u> G | 81.3 |
| | SEQ ID NO: 69 | TCGGC <u>GGGAGGTA</u> AGG | 68.8 |
| 25 | SEQ ID NO: 70 | CTCAGAGGCTT <u>GGCGG</u> | 68.8 |
| | SEQ ID NO: 71 | GC <u>GGCCGGCTAGGGT</u> G | 81.3 |
| | SEQ ID NO: 72 | CGGCC <u>GGCTAGGGT</u> GG | 81.3 |
| | SEQ ID NO: 73 | CGAC <u>GGCGGTGGCGG</u> | 87.5 |
| | SEQ ID NO: 74 | <u>GGACGGGGCGGT</u> GG | 81.3 |
| 30 | SEQ ID NO: 75 | GC <u>GGCGGCATGGGT</u> GC | 81.3 |
| | SEQ ID NO: 76 | <u>GGCGGCGGCATGGGT</u> G | 81.3 |
| | SEQ ID NO: 77 | GC <u>AGGAGAGGACGGT</u> A | 62.5 |
| | SEQ ID NO: 78 | C <u>AGGAGAGGACGGT</u> AC | 62.5 |
| 35 | SEQ ID NO: 79 | <u>GGCAGGAGAGGACGGT</u> | 68.8 |
| | SEQ ID NO: 80 | <u>GGATGGCGCACGCT</u> GG | 75.0 |
| | SEQ ID NO: 81 | <u>GGGAAGGATGGCGC</u> AC | 68.8 |
| | SEQ ID NO: 82 | CCAC <u>GGTGGTGGAGGA</u> | 68.8 |
| 40 | SEQ ID NO: 83 | AC <u>GGCAAGCGGC</u> GGGA | 75.0 |
| | SEQ ID NO: 84 | GGAC <u>GGCAAGCGGC</u> GG | 81.3 |
| 45 | | | |

[0172] The following, in particular, were synthesized: SEQ ID NO: 66-69 directed against MYCN, SEQ ID NO: 70-74 directed against MYC, SEQ ID NO: 75, 76 directed against BIRC5, SEQ ID NO: 77-79 directed against ALK, SEQ ID NO: 80-82 directed against BCL2 and SEQ ID NO: 83, 84 directed against PLK4.

[0173] The oligonucleotides were tested in vitro (at a concentration of 2.5 μ M) to determine their ability to inhibit the transcription of the gene they are directed against by measuring the levels of the gene messenger in cellular models in which the gene of interest is overexpressed. The methods used are the ones described in example 1.

[0174] In particular, SEQ ID NO: 66-69 (directed against MYCN) were tested in vitro in the Kelly and H69 cell lines; SEQ ID NO: 70-74 (directed against MYC) were tested in vitro in the H82 and RD cell lines, SEQ ID NO: 75, 76 (directed against BIRC5) were tested in vitro in the Kelly cell line, SEQ ID NO: 77-79 (directed against ALK) were tested in vitro in the Kelly cell line, SEQ ID NO: 80-82 (directed against BCL2) were tested in vitro in the Kelly cell line and SEQ ID NO: 83, 84 (directed against PLK4) were tested in vitro in the Kelly cell line.

[0175] The ability of the tested oligonucleotides to inhibit the messenger of the gene of interest was measured and

the proliferative capacity of the cells following administration of the oligonucleotides was also assessed. The results are summarized in Table 11.

[0176] The data show a potent inhibitory activity against the mRNA of the target gene and an inhibition of cell proliferation which rises with increases in the number of groups of Gs present in the sequence of the oligonucleotide.

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Table 11

| | SEQ ID NO | | mRNA inhibit. (%) | Cell Prolif. (%) |
|----------------------|------------------|--------------------------------|-------------------------|------------------------|
| 5 MCN Kelly | SEQ ID NO: 66 | TCGGGAGCAGTGGGC A | 57 | 60 |
| | SEQ ID NO: 67 | GC <u>GGGTCGCGGGCAC</u> G | 60 | 56 |
| | SEQ ID NO: 68 | TGGAGGT <u>CGGCCCGCG</u> G | 74 | 35 |
| | SEQ ID NO: 69 | TC <u>GGCGGGAGGTAAAG</u> G | 76 | 30 |
| 10 MCN H69 | SEQ ID NO: 66 | TCGGGAGCAGTGGGC A | 41 | 73 |
| | SEQ ID NO: 67 | GC <u>GGGTCGCGGGCAC</u> G | 42 | 71 |
| | SEQ ID NO: 68 | TGGAGGT <u>CGGCCCGCG</u> G | 62 | 49 |
| | SEQ ID NO: 69 | TC <u>GGCGGGAGGTAAAG</u> G | 67 | 45 |
| 15 MTC H62 | SEQ ID NO: 70 | CT <u>CAGAGGCTTGGCG</u> G | 43 | 79 |
| | SEQ ID NO: 71 | GC <u>GGGCCGGCTAGGGT</u> G | 46 | 75 |
| | SEQ ID NO: 72 | CG <u>GCAGGCTAGGTG</u> G | 52 | 64 |
| | SEQ ID NO: 73 | CGAC <u>GGCGGTGGCGG</u> G | 56 | 65 |
| 20 MTC H62 | SEQ ID NO: 74 | GGAC <u>GGGGCGGGTGG</u> A | 61 | 61 |
| | SEQ ID NO: 70 | CT <u>CAGAGGCTTGGCG</u> G | 34 | 73 |
| | SEQ ID NO: 71 | GC <u>GGGCCGGCTAGGGT</u> G | 42 | 65 |
| | SEQ ID NO: 72 | CG <u>GCAGGCTAGGTG</u> G | 58 | 53 |
| 25 MTC RD | SEQ ID NO: 73 | CGAC <u>GGCGGTGGCGG</u> G | 59 | 51 |
| | SEQ ID NO: 74 | GGAC <u>GGGGCGGGTGG</u> A | 64 | 47 |
| | SEQ ID NO: 75 | GC <u>GGCGGGCATGGGTG</u> C | 69 | 43 |
| | SEQ ID NO: 76 | GC <u>GGGGCGGGCATGGGT</u> G | 78 | 35 |
| 30 BIRCS Kelly | | | | |
| | | | | |

| | | | | | |
|----|---------------|------------------|----------------------|----|----|
| 5 | ALK Kelly | SEQ ID NO: 77 | GCAGGAGAGGACGGT A | 57 | 55 |
| | | SEQ ID NO: 78 | CAGGAGAGGACGGTA C | 68 | 46 |
| | | SEQ ID NO: 79 | GGCAGGAGAGGACGG T | 79 | 39 |
| 10 | BCL2 Kelly | SEQ ID NO: 80 | GGATGGCGCACGCTG G | 59 | 62 |
| | | SEQ ID NO: 81 | GGGAAGGATGGCGCA C | 62 | 58 |
| | | SEQ ID NO: 82 | CCACGGTGGTGGAGG A | 68 | 55 |
| 15 | PLK4 Kelly | SEQ ID NO: 83 | ACGGCAAGCGGG A | 62 | 59 |
| | | SEQ ID NO: 84 | GGACGGCAAGCGCG G | 74 | 49 |

25 [0177] Finally, the oligonucleotides having SEQ ID NO 66-84 were administered in cell lines of a fibroblastoid type (Phoenix and NIH-3T3). In these cells the oligonucleotides tested did not show to be toxic.

[0178] These results indicate that the PNA oligonucleotides act through a specific inhibition effect on the expression of the gene of interest, whereas they do not have any non-specific, toxic effect in cells which do not express the gene.

30 [0179] It can thus be deduced that the PNA oligonucleotides designed on the basis of the parameters described in the present invention act specifically and effectively on the target gene and therefore on the cells which express/over-express this gene.

SEQUENCE LISTING

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Venturelli, Leonardo
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Montemurro, Lucy

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Claims

1. A single strand anti-gene oligonucleotide complementary to the anti-sense DNA strand of a target gene, comprising 10 6-30 nucleotides, preferably 12-24 nucleotides, said oligonucleotide being **characterized by** a sequence comprising at least three groups of at least two consecutive guanines.
2. The oligonucleotide according to claim 1, wherein the groups of at least two consecutive guanines are continuous 15 with each other or spaced apart by at least one nucleotide, said nucleotide not being a guanine.
3. The oligonucleotide according to claim 1 or 2, wherein the groups of at least two consecutive guanines are at least 20 four, five or six in number.
4. The oligonucleotide according to any one of claims 1-3, wherein said oligonucleotide is conjugated with a carrier 25 sequence at the 3' and/or 5' end of said oligonucleotide, preferably a carrier selected from the group consisting of: SEQ ID NO: 47-56.
5. The oligonucleotide according to any one of claims 1-4, wherein said oligonucleotide is at least one molecule of 30 natural nucleic acid, preferably DNA or RNA, possibly chemically modified, or synthetic nucleic acid, preferably PNA, LNA or morpholino, possibly chemically modified, or a combination of said natural nucleic acid and said synthetic nucleic acid.
6. The oligonucleotide according to claim 5, wherein 1) said at least one DNA molecule comprises at least one LNA, 35 methylphosphonate-LNA, BNA, RNG, DNG, GNA, UNA, ENA, ANA, F-ANA, PNA, G-PNA or morpholino nucleotide; or comprises a 2'-O-Methyl, 2'-O-Methoxyethyl or 2'-fluoro RNA nucleotide; or 2) said at least one RNA molecule comprises at least one RNA nucleotide selected from among: a 2'-O-Methyl nucleotide, a 2'-O-Methoxyethyl nucleotide, a 2'-fluoro nucleotide; or at least one nucleotide of a nucleic acid selected from among: LNA, Methylphosphonate LNA, BNA, RNG, DNG, GNA, UNA, ENA, ANA, F-ANA, PNA, G-PNA or morpholino.
7. The oligonucleotide according to claim 6, wherein said at least one DNA molecule or said at least one RNA molecule 40 comprises at least one nucleotide modified at the level of the phosphodiester bond as a phosphorothioate bond.
8. The oligonucleotide according to claim 5, wherein said PNA has a modified backbone wherein the alpha carbon 45 has the side chain of arginine or lysine as a substituent.
9. The oligonucleotide according to any one of claims 1-8, wherein said oligonucleotide is directed against at least one gene responsible for a disease of genetic or viral origin or against at least one tumour gene, wherein said gene is 50 preferably selected from the group consisting of: at least one gene belonging to the MYC family, preferably MYC, MYCN or MYCL1, BIRC5, BCL2, PLK4, ALK, PKM2, CASP8 and RASSF1.
10. The oligonucleotide according to any one of claims 1-9, wherein said oligonucleotide is selected from among: SEQ 55 ID NO: 2, 4, 5, 6, 46, and SEQ ID NO: 68-84.
11. The oligonucleotide according to claim 10, wherein SEQ ID NO: 2, 4, 5, 6, 68 and 69 are directed against the MYCN gene.
12. The oligonucleotide according to claim 10, wherein SEQ ID NO: 70-74 are directed against the gene MYC; wherein 60 SEQ ID NO: 75, 76 are directed against the gene BIRC5; wherein SEQ ID NO: 77-79 are directed against the gene ALK; wherein SEQ ID NO: 80-82 are directed against the gene BCL2; wherein SEQ ID NO: 83, 84 are directed 65 against the gene PLK4.
13. A composition comprising at least one oligonucleotide according to any one of claims 1 to 12 and at least one 70 pharmacologically acceptable excipient.

14. The composition comprising at least one oligonucleotide according to any one of claims 1 to 12 in combination with a selected compound with pharmacological action, preferably in the group consisting of: NGF, somatostatin, retinoic acid, actinomycin D, asparaginase, bleomycin, busulphan, capecitabine, carboplatin, cyclophosphamide, cyclosporine, cisplatin, cytarabine, clorambucil, dacarbazine, daunorubicin, docetaxel, doxorubicin hydrochloride, epirubicin hydrochloride, etoposide, fludarabine phosphate, fluorouracil, gemcitabine, idarubicin hydrochloride, hydroxyurea, ifosfamide, irinotecan hydrochloride, melphalan, mercaptopurine, methotrexate, mitomycin, mitoxantrone, oxaliplatin, paclitaxel, procarbazine, raltitrexed, streptozocin, tegafur-uracil, temozolomide, thioguanine, thiotepa, topotecan, vinblastine, vincristine sulphate, vindesine and vinorelbine; wherein said combination is preferably: SEQ ID: NO 5 and carboplatin, or etoposide or cisplatin or vincristine.

15. An oligonucleotide according to any one of claims 1 to 12 or a composition according to claim 15 for use as a medicament.

16. The oligonucleotide or composition for use according to claim 15, for the treatment of a tumour wherein said tumour is an adult or a pediatric tumour, preferably selected from the group consisting in: neuroblastoma, retinoblastoma, medulloblastoma, ependymoma, pheochromocytoma, embryonal carcinoma, germ cell tumour, alveolar rhabdomyosarcoma, embryonal rhabdomyosarcoma, Wilms' tumour, clear cell sarcoma of the kidney, synovial sarcoma, hepatoblastoma, acute lymphoid leukaemia, chronic lymphoid leukaemia, acute lymphoblastic leukaemia, chronic lymphoblastic leukaemia, Burkitt's lymphoma, acute myeloid leukaemia, chronic myeloid leukaemia, acute megakaryoblastic leukaemia, B chronic lymphoid leukaemia, T-cell leukaemia, lymphomas, small cell lung cancer (microcytoma), lung adenocarcinoma, squamous cell lung carcinoma, typical and atypical primary lung cancer, large cell lung carcinoma, large-cell neuroendocrine lung carcinoma, glioblastoma, hepatocarcinoma, basal cell carcinoma, ovarian tumour, breast tumour and colon cancer.

25 Patentansprüche

1. Einzelstrang-Anti-Gen-Oligonukleotid, komplementär zum Antisense-DNA-Strang eines Zielgens, umfassend 6-30 Nukleotide, vorzugsweise 12-24 Nukleotide, wobei das Oligonukleotid durch eine Sequenz gekennzeichnet ist, die mindestens drei Gruppen von mindestens zwei aufeinanderfolgenden Guaninen umfasst.
2. Oligonukleotid nach Anspruch 1, wobei die Gruppen von mindestens zwei aufeinanderfolgenden Guaninen kontinuierlich zueinander sind oder durch mindestens ein Nukleotid beabstandet sind, wobei das Nukleotid kein Guanin ist.
3. Oligonukleotid nach Anspruch 1 oder 2, wobei die Gruppen von mindestens zwei aufeinanderfolgenden Guaninen mindestens vier, fünf oder sechs an der Zahl sind.
4. Oligonukleotid nach einem der Ansprüche 1 bis 3, wobei das Oligonukleotid mit einer Trägersequenz am 3'-und/oder 5'-Ende des Oligonukleotids konjugiert ist, vorzugsweise einem Träger, ausgewählt aus der Gruppe bestehend aus: SEQ ID NO: 47-56.
5. Oligonukleotid nach einem der Ansprüche 1 bis 4, wobei das Oligonukleotid mindestens ein Molekül natürlicher Nukleinsäure, vorzugsweise DNA oder RNA, gegebenenfalls chemisch modifizierter, oder synthetischer Nukleinsäure, vorzugsweise PNA, LNA oder Morpholin, gegebenenfalls chemisch modifiziert, oder eine Kombination der natürlichen Nukleinsäure und der synthetischen Nukleinsäure ist.
6. Oligonukleotid nach Anspruch 5, wobei 1) das mindestens eine DNA-Molekül mindestens ein LNA, Methylphosphonat-LNA, BNA, RNG, DNG, GNA, UNA, ENA, F-ANA, PNA, G-PNA oder Morpholin-Nukleotid umfasst; oder ein 2'-Methyl-, 2'-Methoxyethyl- oder 2'-Fluor-RNA-Nukleotid umfasst; oder 2) das mindestens eine RNA-Molekül mindestens ein RNA-Nukleotid umfasst, ausgewählt aus: einem 2'-Methylnukleotid, einem 2'-Methoxyethylnukleotid, einem 2'-Fluornukleotid; oder mindestens ein Nukleotid einer Nukleinsäure ausgewählt aus: LNA, Methylphosphonat LNA, BNA, RNG, DNG, GNA, UNA, ENA, F-ANA, PNA, G-PNA oder Morpholin.
7. Oligonukleotid nach Anspruch 6, wobei das mindestens eine DNA-Molekül oder das mindestens eine RNA-Molekül mindestens ein Nukleotid umfasst, das auf der Ebene der Phosphodiesterbindung als Phosphorothioatbindung modifiziert ist.
8. Oligonukleotid nach Anspruch 5, wobei das PNA ein modifiziertes Rückgrat aufweist, wobei der alpha-Kohlenstoff

die Seitenkette von Arginin oder Lysin als Substituenten aufweist.

9. Oligonukleotid nach einem der Ansprüche 1 bis 8, wobei das Oligonukleotid gegen mindestens ein Gen gerichtet ist, das für eine Erkrankung genetischen oder viralen Ursprungs verantwortlich ist oder gegen mindestens ein Tumor-Gen, wobei das Gen vorzugsweise aus der Gruppe ausgewählt ist, bestehend aus: mindestens einem Gen, das zur MYC-Familie, vorzugsweise MYC, MYCN oder MYCL1, BIRC5, BCL2, PLK4, ALK, PKM2, CASP8 und RASSF1 gehört.

10. Oligonukleotid nach einem der Ansprüche 1 bis 9, wobei das Oligonukleotid ausgewählt ist aus: SEQ ID NO: 2 ,4, 5, 6, 46, und SEQ ID NO: 68-84.

11. Oligonukleotid nach Anspruch 10, wobei SEQ ID NO: 2, 4, 5, 6, 68 und 69 gegen das MYCN-Gen gerichtet sind.

12. Oligonukleotid nach Anspruch 10, wobei SEQ ID NO: 70-74 gegen das Gen MYC gerichtet sind; wobei SEQ ID NO: 75, 76 gegen das Gen BIRC5 gerichtet sind; wobei SEQ ID NO: 77-79 gegen das Gen ALK gerichtet sind; wobei SEQ ID NO: 80-82 gegen das Gen BCL2 gerichtet sind; wobei SEQ ID NO: 83, 84 gegen das Gen PLK4 gerichtet sind.

13. Zusammensetzung, umfassend mindestens ein Oligonukleotid nach einem der Ansprüche 1 bis 12 und mindestens einen pharmakologisch annehmbaren Exzipienten.

14. Zusammensetzung umfassend mindestens ein Oligonukleotid nach einem der Ansprüche 1 bis 12 in Kombination mit einer ausgewählten Verbindung mit pharmakologischer Wirkung, vorzugsweise in der Gruppe bestehend aus: NGF, Somatostatin, Retinsäure, Actinomycin D, Asparaginase, Bleomycin, Busulphan-Capicitabin, Carboplatin, Cyclophosphamid, Ciclosporin, Cisplatin, Cytarabin, Clorambucil, Dacarbazine, Daunorubicin, Docetaxel, Doxorubicin-Hydrochlorid, Epirubicin-Hydrochlorid, Etoposid, Fludarabin-Phosphat, Fluorouracil, Gemcitabin, Idarubicin-Hydrochlorid, Hydroxyantrastostoff, Ifophosphamide, Irinotecan-Hydrochlorid, Melphalan, Mercaptopurin, Methotrexat, Mitomycin, Mitoxantron, Oxaliplatin, Paclitaxel, Procarbazine, Raltitrexed, Streptozocin, Tegafur-Uracil, Temozolomid, Thioguanin, Thiotepa, Topotecan, Vinblastin, Vincristinsulfat, Vindesin und Vinorelbine; wobei die Kombination vorzugsweise ist: SEQ ID: NO 5 und Carboplatin oder Etoposid oder Cisplatin oder Vincristine.

15. Oligonukleotid nach einem der Ansprüche 1 bis 12 oder Zusammensetzung nach Anspruch 15 zur Verwendung als Medikament.

16. Oligonukleotid oder Zusammensetzung zur Verwendung nach Anspruch 15 für die Behandlung eines Tumors, wobei der Tumor ein erwachsener oder ein Pediatrie-Tumor ist, vorzugsweise ausgewählt aus der Gruppe bestehend aus: Neuroblastom, Retinoblastom, Medulloblastom, Ependymom, Pheochromozytom, embryonalem Karzinom, Keimzell-tumor, alveolärem Ribdomyosarkom, embryonalem Ribdomiosarkom, Wilms-Tumor, klarem Zell-Sarkom der Niere, Synovial-Sarkom, Hepatoblastom, akuter lymphatischer Leukämie, chronischer lymphatischer Leukämie, akuter lymphoblastischer Leukämie, chronischer lymphoblastischer Leukämie, Burkitt-Lymphom, akuter myeloider Leukämie, chronischer myeloider Leukämie, akuter megakaryoblastischer Leukämie, B chronischer lymphatischer Leukämie, T-Zell-Leukämie, Lymphome, kleinzelligem Lungenkrebs (Mikrozytom), Lungen-Adenokarzinom, Plattenepithelkarzinom, typischem und atypischem primären Lungenkrebs, großzelligem Lungenkarzinom, großzelligem neuroendokrinem Lungenkarzinom, Glioblastom, Hepatokarzinom, Basalzellkarzinom, Ovarialtumor, Brusttumor und Darmkrebs.

Revendications

1. Oligonucléotide antigène à brin simple complémentaire au brin de l'ADN antisens d'un gène cible, comprenant 6-30 nucléotides, de préférence 12-24 nucléotides, ledit oligonucléotide étant **caractérisé par** une séquence comprenant au moins trois groupes d'au moins deux guanines consécutives.

2. Oligonucléotide selon la revendication 1, dans lequel les groupes d'au moins deux guanines consécutives sont continus les uns avec les autres ou espacés par un nucléotide, ledit nucléotide n'étant pas une guanine.

3. Oligonucléotide selon la revendication 1 ou 2, dans lequel les groupes d'au moins deux guanines consécutives sont au moins au nombre de quatre, cinq ou six.

4. Oligonucléotide selon l'une quelconque des revendications 1-3, dans lequel ledit oligonucléotide est conjugué à une séquence porteuse à l'extrémité 3' et/ou 5' dudit oligonucléotide, de préférence un porteur choisi parmi le groupe consistant en : SEQ ID N° : 47-56.

5. Oligonucléotide selon l'une quelconque des revendications 1-4, dans lequel ledit oligonucléotide est au moins une molécule d'acide nucléique naturel, de préférence de l'ADN ou de l'ARN, si possible modifiée chimiquement, ou un acide nucléique synthétique, de préférence de l'APN, LNA ou un morpholino, si possible modifié chimiquement, ou une combinaison dudit acide nucléique naturel et dudit acide nucléique synthétique.

10. 6. Oligonucléotide selon la revendication 5, dans lequel 1) ladite au moins une molécule d'ADN comprend au moins un LNA, méthylphosphonate-LNA, BNA, RNG, DNG, GNA, UNA, ENA, ANA, F-ANA, PNA, G-PNA ou un nucléotide-morpholino ; ou comprend un 2'0- Méthyle, 2'0-Méthoxyéthyle ou 2'-fluoro nucléotide ANR ; ou 2) ladite au moins une molécule d'ARN comprend au moins un nucléotide d'ARN sélectionné parmi : un 2'0-Méthyl nucléotide, un 2'0-Méthoxyéthyl nucléotide, un 2'-fluoro nucléotide ; ou au moins un nucléotide d'un acide nucléique sélectionné parmi : LNA, méthylphosphonate-LNA, BNA, RNG, DNG, GNA, UNA, ENA, ANA, F-ANA, PNA, G-PNA ou un morpholino.

15. 7. Oligonucléotide selon la revendication 6, dans lequel ladite au moins une molécule d'ADN ou ladite au moins une molécule d'ARN comprend au moins un nucléotide modifié au niveau de la liaison phosphodiester comme étant une liaison phosphorothioate.

20. 8. Oligonucléotide selon la revendication 5, dans lequel ledit PNA comporte une chaîne principale modifiée dans laquelle le carbone alpha comporte la chaîne latérale d'arginine ou de lysine en tant que substituant.

25. 9. Oligonucléotide selon l'une quelconque des revendications 1-8, dans lequel ledit oligonucléotide est dirigé contre au moins un gène responsable d'une maladie d'origine génétique ou virale ou contre au moins un gène tumoral, dans lequel ledit gène est de préférence choisi dans le groupe consistant en : au moins un gène appartenant à la famille MYC, de préférence MYC, MYCN ou MYCL1, BIRC5, BCL2, PLK4, ALK, PKM2, CASP8 et RASSF1.

30. 10. Oligonucléotide selon l'une quelconque des revendications 1-9, dans lequel ledit oligonucléotide est choisi parmi les : SEQ ID N : 2, 4, 5, 6, 46, et SEQ ID N° : 68-84.

35. 11. Oligonucléotide selon la revendication 10, dans lequel les SEQ ID N° : 2, 4, 5, 6, 68 et 69 sont dirigées contre le gène MYCN.

12. Oligonucléotide selon la revendication 10, dans lequel les SEQ ID N° : 70-74 sont dirigées contre le gène MYC ; dans lequel les SEQ ID N° : 75, 76 sont dirigées contre le gène BIRC5 ; dans lequel les SEQ ID N° : 77-79 sont dirigées contre le gène ALK ; dans lequel les SEQ ID N° : 80-82 sont dirigées contre le gène BCL2 ; dans lequel les SEQ ID N° : 83, 84 sont dirigées contre le gène PLK4.

40. 13. Composition comprenant au moins un oligonucléotide selon l'une quelconque des revendications de 1 à 12 et au moins un excipient pharmacologiquement acceptable.

14. Composition comprenant au moins un oligonucléotide selon l'une quelconque des revendications de 1 à 12 en combinaison avec un composant sélectionné ayant une action pharmacologique, de préférence dans le groupe consistant en : NGF, somatostatine, acide rétinoïque, actinomycine D, asparaginase, bléomycine, busulfan capé-citabine, carboplatine, cyclophosphamide, ciclosporine, cisplatine, cytarabine, chlorambucil, dacarbazine, daunorubicine, docétaxel, doxorubicine chlorhydrate, épirubicine chlorhydrate, étoposide, fludarabine phosphate, fluorouracile, gemcitabine, chlorhydrate d'idarubicine, hydroxycarbamide, isophosphamide, irinotécanchlorhydrate, melphalan, mercaptopurine, méthotrexate, mitomycine, mitoxantrone, oxaliplatin, paclitaxel, procarbazine, raltitrexed, streptozocine, tégarfur-uracil, témozolamide, thioguanine, thiotepla, topotécane, vinblastine, vincristine sulfate, vindésine et vinorelbine ; dans laquelle ladite combinaison est de préférence : SEQ ID : N° 5 et carboplatine, ou étoposide ou cisplatine ou vincristine.

50. 15. Oligonucléotide selon l'une quelconque des revendications de 1 à 12 ou composition selon la revendication 15, destinée à un usage médicamenteux.

55. 16. Oligonucléotide ou composition à utiliser selon la revendication 15, destiné(e) au traitement d'une tumeur, dans laquelle ladite tumeur est une tumeur typique de l'adulte ou une tumeur pédiatrique, de préférence choisie dans le

groupe consistant en : neuroblastome, rétinoblastome, médulloblastome, épendymome, phéochromocytome, carcinome embryonnaire, tumeur germinale, rhabdomyosarcome alvéolaire, rhabdomyosarcome embryonnaire, tumeur de Wilms, hypernéphrome, sarcome synovial, hépatoblastome, leucémie aiguë lymphoblastique, leucémie lymphoïde chronique, leucémie aiguë lymphoblastique, leucémie lymphoblastique chronique, lymphome de Burkitt, leucémie aiguë myéloïde, leucémie myéloïde chronique, Leucémie aiguë mégacaryoblastique, Leucémie lymphoïde chronique B, lymphome à cellules T, lymphomes, cancer pulmonaire à petites cellules (microcytome), adénocarcinome pulmonaire, carcinome épidermoïde pulmonaire, cancer primitif du poumon typique et atypique, cancer pulmonaire à grandes cellules, cancer du poumon neuroendocrinien à grandes cellules, glioblastome, carcinome hépatocellulaire, épithélioma cutané basocellulaire, tumeur ovarienne, tumeur au sein et cancer du côlon.

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Öligonukleotidok a génexpresszió befolyásolására, valamint őzek alkalmazása

SZABADALMI IGÉNYPONTOK

1. Egy megcélzott gén antiszensz DNS szálával komplementer, 6-30 nukleotídból, előnyösen 12-24 nukleotídból álló egyszálú anti-gén oligonukleotid, a szóban forgó oligonukleotídot olyan szekvencia jellemzi, amely legalább két egymást követő guanint tartalmazó csoportokból legalább hármat tartalmaz.

2. Az 1. igénypont szerinti oligonukleotid, amelyben a legalább két egymást követő guanint tartalmazó csoportok egymással összefüggnek, vagy legfeljebb egy nukleotid választja el őket, és a szóban forgó nukleotid nem guanin.

3. Az 1. vagy 2. igénypont szerinti oligonukleotid, a legalább két egymást követő guanint tartalmazó csoportok száma legalább négy, öt vagy hat.

4. Az 1-3. igénypontok bármelyike szerinti oligonukleotid, ahol a szóban forgó oligonukleotid a szóban forgó oligonukleotid a 3' és/vagy az 5' végén egy hordozó szekvenciához van konjugálva, előnyösen a 47-56. számú szekvenciák közül választható hordozóhoz.

5. Az 1-4. igénypontok bármelyike szerinti oligonukleotid, ahol a szóban forgó oligonukleotid legalább egy természetes nukleinsav, előnyösen DNS vagy RNS molekula, esetleg kémiailag módosítva, vagy egy szintetikus nukleinsav, előnyösen PNS, LNS vagy morfolino, esetleg kémiailag módosítva, vagy a szóban forgó természetes nukleinsav és a szóban forgó szintetikus nukleinsav kombinációja.

6. Az 5. igénypont szerinti oligonukleotid, amelyben 1) a szóban forgó oligonukleotid legalább egy LNS-t, metílfoszfonát-LNS-t, BNS-t, RNG-t, DNG-t, GNS-t, UNS-t, ENS-t, ANS-t, F-ANS-t, PNS-t, G-PNS-t vagy morfolinó nukleotidot tartalmaz, vagy egy 2'-O-metil, 2'-O-metoxietil, vagy 2'-fluoro RNS nukleotidot; vagy 2) a szóban forgó legalább egy RNS molekula tartalmaz legalább egy RNS nukleotidot, amely a következők közül választható ki: egy 2'-O-metil nukleotid, egy 2'-O-metoxietil nukleotid, egy 2'-fluor nukleotid; vagy legalább egy nukleotidot vagy nukleinsavat, amelyek az alábbiak közül választhatók: LNS, metílfoszfonát-LNS, BNS, RNG, DNG, GNS, UNS, ENS, ANS, F-ANS, PNS, G-PNS vagy morfolinó.

7. A 6. igénypont szerinti oligonukleotid, amelyben a szóban forgó legalább egy DNS molekula vagy a szóban forgó legalább egy RNS molekula egy, a foszfodiészter kötés szintjén módosított nukleotidot tartalmaz, foszforotíóát kötés formájában.



8. Az 5. igénypont szerinti oligonukleotid, amelyben a szóban forgó PNS-nek van egy módosított gerince, amelyben az alfa szénnek szubsztituensként arginin vagy lizin oldallánca van.

9. Az 1-8. igénypontok bármelyike szerinti oligonukleotid, ahol a szóban forgó oligonukleotid legalább egy genetikai vagy virális eredetű betegségért felelős gén, vagy legalább egy tumor gén ellen irányul, ahol a szóban forgó gént előnyösen a következő csoportból választhatjuk ki: legalább egy, a MYC családba tartozó gén, előnyösen MYC, MYCN vagy MYCL1, BIRC5, BCL2, PLK4, ALK, PKM2, CASP8 és RASSF1.

10. Az 1-9. igénypontok bármelyike szerinti oligonukleotid, ahol a szóban forgó oligonukleotidot a következők közül választjuk ki: 2., 4., 5., 6., 46. számú szekvencia, és a 68-84. számú szekvenciák.

11. A 10. igénypont szerinti oligonukleotid, ahol a 2., 4., 5., 6., 68. és 69. számú szekvencia az MYCN gén ellen irányul.

12. A 10. igénypont szerinti oligonukleotid, ahol 70-74. számú szekvencia az MYC gén ellen irányul; ahol a 75., 76. számú szekvencia a BIRC5 gén ellen irányul; ahol a 77-79. számú szekvencia az ALK gén ellen irányul; ahol a 80-82. számú szekvencia a BCL2 gén ellen irányul; ahol a 83., 84. számú szekvencia a PLK4 gén ellen irányul.

13. Készítmény, amely az 1-12. igénypontok bármelyike szerinti oligonukleotidok közül legalább egyet tartalmaz, valamint tartalmaz legalább egy gyógyászatilag elfogadható segédanyagot.

14. Készítmény, amely az 1-12. igénypontok bármelyike szerinti oligonukleotidok közül legalább egyet tartalmaz egy kiválasztott farmakológiai hatással rendelkező vegyülettel kombinálva, amely előnyösen a következő csoportból választható: NGF, szomatotropin, rétinasav, aktinomycin D, aszparagináz, bleomicin, buszulfán, kapocitabin, karboplatin, ciklofoszfamid, ciklosporin, ciszplatin, citarabin, klorambucil, dakarbazin, daunorubicin, docetaxel, doxorubicin hidroklorid, epirubicin hidroklorid, etopozid, fludarabin foszfát, fluor-uracil, gemcitabin, idarubicin hidroklorid, hidroxi-karbamid, ifoszfamid, irinotekán hidroklorid, melfalan, merkapto-purin, metotrexát, mitomicin, mitoxantron, oxaliplatin, paclitaxel, prokarbazin, raltitrexed, sztreptozocin, tegafur-uracil, temozolamid, tio-guanin, tiotepe, topotekán, vinblasztin, vinkrisztin-szulfát, vindezin és vinorelbín; ahol a szóban forgó kombináció előnyösen: 5. számú szekvencia és karboplatin, vagy etopozid, vagy ciszplatin vagy vinkrisztin.

15. Az 1-12. igénypontok bármelyike szerinti oligonukleotid, vagy a 15. igénypont szerinti készítmény, gyógyszerként történő alkalmazás céljára.

16. A 15. igénypont szerinti oligonukleotid vagy készítmény alkalmazása egy tumor kezelésére, ahol a szóban forgó tumor egy felnőttben vagy egy gyermekben levő tumor, és előnyösen a

következő csoportból választhatjuk ki: neuroblasztóma, retinoblasztóma, medulloblasztóma, ependimóma, pheochromocitóma, embrionális karcinóma, csírasejt tumor, alveolaris rhabdomyosarcoma, embrionális rhabdomyosarcoma, Wilms tumor, a vese fiszta sejt szarkómája, szinoviális szarkóma, hepatoblasztóma, akut limfoid leukémia, krónikus limfoid leukémia, akut limfoblasztos leukémia, krónikus limfoblasztos leukémia, Burkitt limfóma, akut mieloid leukémia, krónikus mieloid leukémia, akut megakarioblasztos leukémia, B krónikus limfoid leukémia, T-sejtes leukémia, limfómák, kissejtes tüdőrák (mikrocitóma), tüdő adenokarcinóma, pikkelyes sejtes tüdő karcinóma, tipikus és atípikus primer tüdőrák, nagysejtes tüdő karcinóma, nagysejtes neuroendokrin tüdő karcinóma, glioblasztóma, hepatokarcinóma, bazális sejtes karcinóma, petefészek tumor, emlőtumor és vastagbél rák.