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Bulletin De Paie Cameroun.pdf Oligonucleotides are biopolymers which are used to interfere with various cellular functions. In particular, they are used as antisense oligonucleotides for specifically inhibiting the expression of nucleic acids and thus the production of the corresponding proteins. Specifically, antisense oligonucleotides which are complementary to a region of the nucleic acids are hybridized to the nucleic acids and thus prevent the nucleic acids from being transcribed into corresponding proteins. An ideal antisense oligonucleotide comprises in its structure the same bases as the target nucleic acid, namely: a region of bases which is complementary to the target region and a region of bases which is not complementary to the target region. These regions are called the "overhanging region" and the "toe region", respectively. It is thus possible to influence the efficiency of antisense oligonucleotides by modifying the overhanging region with lipophilic or hydrophilic groups. It is desirable for antisense oligonucleotides to have a sufficient resistance to enzymatic hydrolysis in order to avoid unnecessary degradation of the antisense oligonucleotide before it is expressed. This resistance must not be accompanied by an increase of the size of the antisense oligonucleotide. On the contrary, the size of the antisense oligonucleotide is a parameter which influences the tissue penetration and thus the efficiency of the antisense oligonucleotide. An unsaturated carbon-carbon chain having at least 10 carbon atoms and an unsaturated carbon-carbon chain having at least 12 carbon atoms are usually used for bonding the hydrophobic group to the oligonucleotide. An oligonucleotide comprising such a hydrophobic group is more resistant to enzymatic hydrolysis than an oligonucleotide comprising a saturated carbon-carbon chain. However, its size is typically doubled compared to an oligonucleotide comprising an unsaturated carbon-carbon chain having less than 12 carbon atoms. Oligonucleotides comprising 12 carbon atoms have already been disclosed, for example, in the document WO 2004/064191. The use of polyalkylene glycol derivatives for increasing the stability and the tissue penetration of oligonucleotides is known. The document WO 2005/050581 describes oligonucleotides which are more stable and more efficiently transfected into cells. They comprise a group

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