FAMILIES OF NON-CROSS - HYBRIDIZING POLYNUCLEOTIDES FOR USE AS TAGS AND TAG COMPLEMENTS, MANUFACTURE AND USE THEREOF

FIELD OF THE INVENTION

This invention relates to families of oligonucleotide tags for use, for example, in sorting molecules. Members of a given family of tags can be distinguished one from the other by specific hybridization to their tag complements.

BACKGROUND OF THE INVENTION

Specific hybridization of oligonucleotides and their analogs is a fundamental process that is employed in a wide variety of research, medical, and industrial applications, including the identification of disease-related polynucleotides in diagnostic assays, screening for clones of novel target polynucleotides, identification of specific polynucleotides in blots of mixtures of polynucleotides, therapeutic blocking of inappropriately expressed genes and DNA sequencing. Sequence specific hybridization is critical in the development of high throughput multiplexed nucleic acid assays. As formats for these assays expand to encompass larger amounts of sequence information acquired through projects such as the Human Genome project, the challenge of sequence specific hybridization with high fidelity is becoming increasingly difficult to achieve.

In large part, the success of hybridization using oligonucleotides depends on minimizing the number of false positives and false negatives. Such problems have made the simultaneous use of multiple hybridization probes in a single experiment i.e. multiplexing, particularly in the analysis of multiple gene sequences on a gene microarray, very difficult. For example, in certain binding assays, a number of nucleic acid molecules are bound to a chip with the desire that a given "target" sequence will bind selectively to its complement attached to the chip. Approaches have been developed that involve the use of oligonucleotide tags attached to a solid support that can be used to specifically hybridize to the tag complements that are coupled to probe sequences. Chetverin *et al.* (WO 93/17126) uses sectioned, binary oligonucleotide arrays to sort and survey nucleic acids. These arrays have a constant nucleotide sequence attached to an adjacent variable nucleotide sequence, both bound to a solid support by a covalent linking moiety. These binary arrays have advantages compared with ordinary arrays in that they can be used to sort strands according to their terminal sequences so that each strand binds to a fixed location on an array. The design of the terminal sequences in this approach comprises the use of constant and variable

sequences. United States Patent Nos. 6,103,463 and 6,322,971 issued to Chetverin *et al.* on August 15, 2000 and November 27, 2001, respectively.

This concept of using molecular tags to sort a mixture of molecules is analogous to molecular tags developed for bacterial and yeast genetics (Hensel *et al.*, Science; 269, 400-403: 1995 and Schoemaker *et al.*, Nature Genetics; 14, 450-456: 1996). Here, a method termed "signature tagged" mutagenesis in which each mutant is tagged with a different DNA sequence is used to recover mutant genes from a complex mixture of approximately 10,000 bacterial colonies. In the tagging approach of Barany *et al.* (WO 9731256), known as the "zip chip", a family of nucleic acid molecules, the "zip-code addresses", each different from each other, are set out on a grid. Target molecules are attached to oligonucleotide sequences complementary to the "zipcode addresses," referred to as "zipcodes," which are used to specifically hybridize to the address locations on the grid. While the selection of these families of polynucleotide sequences used as addresses is critical for correct performance of the assay, the performance has not been described.

Working in a highly parallel hybridization environment requiring specific hybridization imposes very rigorous selection criteria for the design of families of oligonucleotides that are to be used. The success of these approaches is dependent on the specific hybridization of a probe and its complement. Problems arise as the family of nucleic acid molecules cross-hybridize or hybridize incorrectly to the target sequences. While it is common to obtain incorrect hybridization resulting in false positives or an inability to form hybrids resulting in false negatives, the frequency of such results must be minimized. In order to achieve this goal certain thermodynamic properties of forming nucleic acid hybrids must be considered. The temperature at which oligonucleotides form duplexes with their complementary sequences known as the T_m (the temperature at which 50% of the nucleic acid duplex is dissociated) varies according to a number of sequence dependent properties including the hydrogen bonding energies of the canonical pairs A-T and G-C (reflected in GC or base composition), stacking free energy and, to a lesser extent, nearest neighbour interactions. These energies vary widely among oligonucleotides that are typically used in hybridization assays. For example, hybridization of two probe sequences composed of 24 nucleotides, one with a 40% GC content and the other with a 60% GC content, with its complementary target under standard conditions theoretically may have a 10°C difference in melting temperature (Mueller *et al.*, Current Protocols in Mol. Biol.; 15, 5:1993). Problems in hybridization occur when the hybrids are allowed to form under hybridization conditions that include a single hybridization temperature that is not optimal for correct hybridization of all

oligonucleotide sequences of a set. Mismatch hybridization of non-complementary probes can occur forming duplexes with measurable mismatch stability (Santalucia *et al.*, Biochemistry; 38: 3468-77, 1999). Mismatching of duplexes in a particular set of oligonucleotides can occur under hybridization conditions where the mismatch results in a decrease in duplex stability that results in a higher T_m than the least stable correct duplex of that particular set. For example, if hybridization is carried out under conditions that favor the AT-rich perfect match duplex sequence, the possibility exists for hybridizing a GC-rich duplex sequence that contains a mismatched base having a melting temperature that is still above the correctly formed AT-rich duplex. Therefore, design of families of oligonucleotide sequences that can be used in multiplexed hybridization reactions must include consideration for the thermodynamic properties of oligonucleotides and duplex formation that will reduce or eliminate cross hybridization behavior within the designed oligonucleotide set.

The development of such families of tags has been attempted over the years with varying degrees of success. There are a number of different approaches for selecting sequences for use in multiplexed hybridization assays. The selection of sequences that can be used as zipcodes or tags in an addressable array has been described in the patent literature in an approach taken by Brenner and co-workers. United States Patent No. 5,654,413 describes a population of oligonucleotide tags (and corresponding tag complements) in which each oligonucleotide tag includes a plurality of subunits, each subunit consisting of an oligonucleotide having a length of from three to six nucleotides and each subunit being selected from a minimally cross hybridizing set, wherein a subunit of the set would have at least two mismatches with any other sequence of the set. Table II of the Brenner patent specification describes exemplary groups of 4mer subunits that are minimally cross hybridizing according to the aforementioned criteria. In the approach taken by Brenner, constructing non cross-hybridizing oligonucleotides, relies on the use of subunits that form a duplex having at least two mismatches with the complement of any other subunit of the same set. The ordering of subunits in the construction of oligonucleotide tags is not specifically defined.

Parameters used in the design of tags based on subunits are discussed in Barany *et al.* (WO 9731256). For example, in the design of polynucleotide sequences that are for example 24 nucleotides in length (24mer) derived from a set of four possible tetramers in which each 24mer "address" differs from its nearest 24mer neighbour by 3 tetramers. They discuss further that, if each tetramer differs from each other by at least two nucleotides, then each 24mer will differ from the next by at least six nucleotides. This is determined

without consideration for insertions or deletions when forming the alignment between any two sequences of the set. In this way a unique "zip code" sequence is generated. The zip code is ligated to a label in a target dependent manner, resulting in a unique "zip code" which is then allowed to hybridize to its address on the chip. To minimize cross-hybridization of a "zip code" to other "addresses", the hybridization reaction is carried out at temperatures of 75-80°C. Due to the high temperature conditions for hybridization, 24mers that have partial homology hybridize to a lesser extent than sequences with perfect complementarity and represent 'dead zones'. This approach of implementing stringent hybridization conditions for example, involving high temperature hybridization, is also practiced by Brenner *et. al.*

The current state of technology for designing non-cross hybridizing tags based on subunits does not provide sufficient guidance to construct a family of relatively large numbers of sequences with practical value in assays that require stringent non-cross hybridizing behavior.

A multiplex sequencing method has been described in United States Patent No. 4,942,124, which issued to Church on July 17, 1990. The method requires at least two vectors which differ from each other at a tag sequence. It is stated that a tag sequence in one vector will not hybridize under stringent hybridization conditions to a tag sequence (i.e., complementary probes do not cross-hybridize) in another vector. Exemplary stringent hybridization conditions are given as 42°C in 500-1000 mM sodium phosphate buffer. A set of 42 20-mer tag sequences, all of which lack G residues, is given in Figure 3 of the specification. Details of how the sequences were obtained are not provided, although Church states that initially 92 were chosen on the basis of their having sufficient sequence diversity to insure uniqueness.

So while it is possible for a person knowledgeable in the field to design a small number of non-cross hybridizing tags, it is difficult to design a larger number such tags. A co-pending application of the owner of this patent application describes such a set of 210 non-cross hybridizing tags that have a practical value. A method described in international patent application No. PCT/CA 01/00141 published under WO 01/59151 on August 16, 2001. Little guidance is provided, however, for the provision of a larger set, say 1000 or so, of non-cross hybridizing tags. Since having sets of approximately 1000 non-cross hybridizing tags, or more, would be of considerable practical value, it would be useful to develop such a set.

Thus, while it is desirable with such arrays to have, at once, a large number of address molecules, the address molecules should each be highly selective for its own complement sequence. While such an array provides the

advantage that the family of molecules making up the grid is entirely of design, and does not rely on sequences as they occur in nature, the provision of a family of molecules, which is sufficiently large and where each individual member is sufficiently selective for its complement over all the other zipcode molecules (i.e., where there is sufficiently low cross-hybridization, or cross-talk) continues to elude researchers.

SUMMARY OF INVENTION

A family of 1168 sequences was obtained using a computer algorithm to have desirable hybridization properties for use in nucleic acid detection assays. The sequence set of 1168 oligonucleotides was partially characterized in hybridization assays, demonstrating the ability of family members to correctly hybridize to their complementary sequences with minimal cross hybridization. These are the sequences having SEQ ID NOs:1 to 1168 of Table I.

Variant families of sequences (seen as tags or tag complements) of a family of sequences taken from Table I are also part of the invention. For the purposes of discussion, a family or set of oligonucleotides will often be described as a family of tag complements, but it will be understood that such a set could just easily be a family of tags.

A family of complements is obtained from a set of oligonucleotides based on a family of oligonucleotides such as those of Table I. To simplify discussion, providing a family of complements based on the oligonucleotides of Table I will be described.

Firstly, the groups of sequences based on the oligonucleotides of Table I can be represented as shown in Table IA.

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
1 1 1 2 2 3 2 3 1 1 1 3 1 2 2 3 2 2 2 3 2 3 2 1	1
3 2 2 1 3 1 3 2 2 1 1 2 2 3 2 1 2 2 2 3 1 2 3 1	2
1 2 3 2 2 1 1 1 3 2 1 1 3 2 3 2 2 3 1 1 1 2 3 2	3
2 3 1 2 3 2 2 1 3 1 1 3 2 1 2 1 2 2 3 2 3 1 1 2	4
2 2 2 3 2 3 2 1 3 1 1 2 1 2 3 2 3 2 2 3 2 2 1 1	5
1 2 1 1 3 2 3 2 1 1 3 2 3 1 1 1 2 1 1 3 1 1 3 1	6
1 1 3 1 3 2 1 2 2 2 3 2 2 3 2 3 1 3 2 2 1 1 1 2	7
3 2 3 2 2 2 1 2 3 2 2 1 2 1 2 3 2 3 1 1 3 2 2 2	8
1 1 1 3 1 3 1 1 2 1 3 1 1 2 1 2 3 2 3 2 1 1 3 2	9
2 1 2 3 1 1 1 3 1 3 2 3 1 3 1 2 1 1 2 3 2 2 2 1	10
1 2 3 1 3 1 1 1 2 1 2 3 2 2 1 3 1 1 2 3 2 3 1 2	11
2 2 1 3 2 2 3 2 2 3 1 2 3 2 2 2 1 3 2 1 3 2 2 2	12
3 2 1 1 1 3 1 3 2 1 2 1 1 3 2 2 2 3 1 2 3 1 2 1	13
1 1 1 3 2 1 1 3 1 1 2 3 1 2 3 2 1 1 2 1 1 3 2 3	14
3 2 1 3 1 1 1 2 1 3 2 2 2 1 2 2 3 1 2 3 1 2 2 3	15
2 3 2 1 1 3 2 3 1 1 1 2 1 3 2 3 1 3 2 2 1 2 2 2	16

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
1 1 1 2 1 3 1 2 3 1 2 1 2 1 1 3 2 3 1 3 1 1 2 3	17
1 2 1 1 3 2 2 1 2 1 1 3 2 3 2 2 1 2 3 2 3 1 3 2	18
2 1 2 1 3 1 2 1 1 1 3 1 3 1 2 3 1 2 2 2 3 2 2 3	19
1 3 1 3 2 2 3 1 3 1 1 2 3 2 1 2 1 3 2 1 2 2 1 2	20
1 1 3 2 1 3 2 2 2 3 2 1 1 3 1 1 2 3 1 2 2 3 2 1	21
2 2 1 2 3 1 1 1 2 2 3 1 3 2 3 1 1 3 1 2 2 3 1 2	22
3 2 1 2 1 2 3 2 1 1 1 2 2 3 2 2 1 2 3 2 2 3 1 3	23
3 1 1 2 2 3 2 1 2 1 1 1 3 2 1 2 2 1 3 1 2 3 2 3	24
2 1 3 1 2 3 1 3 1 2 2 1 1 3 2 3 2 2 1 2 2 2 3 1	25
3 2 2 1 1 3 2 2 2 3 2 2 2 1 2 3 2 1 2 1 3 1 1 3	26
3 1 3 2 1 2 2 1 3 2 1 1 1 3 2 3 1 2 1 2 3 1 2 1	27
3 2 3 1 1 2 3 1 2 2 2 1 3 2 1 1 1 2 3 1 2 2 3 1	28
3 1 2 2 3 1 1 3 2 2 1 2 1 3 1 1 1 2 3 1 2 2 1 3	29
1 3 2 3 1 2 1 1 1 2 3 2 2 1 3 2 2 3 1 1 2 2 3 2	30
2 1 2 1 2 1 3 2 1 1 1 2 3 2 2 2 3 2 3 2 3 2 2 3	31
2 2 1 1 3 2 3 2 2 1 3 2 2 1 2 2 2 3 2 2 3 2 1 3	32
3 2 1 3 2 1 1 2 1 2 3 1 1 3 2 3 1 3 1 1 2 1 2 1	33
2 1 3 2 3 2 1 2 1 3 1 1 2 3 2 1 3 1 2 2 2 1 3 2	34
2 2 3 2 1 3 1 2 2 1 3 1 2 3 2 3 2 2 2 3 2 1 1 1	35
2 1 3 2 1 2 1 3 1 3 2 1 3 1 3 1 2 3 1 2 1 2 2 2	36
1 2 2 3 2 3 1 1 1 3 1 1 1 3 1 3 1 1 3 1 1 1 2 2	37
2 3 2 3 1 3 1 1 2 2 1 1 3 1 2 2 1 1 3 1 1 2 3 2	38
1 2 1 2 2 1 3 2 2 1 1 3 1 1 1 3 1 1 3 1 3 2 2 3	39
2 2 3 2 1 3 2 2 3 1 3 1 1 1 2 1 2 3 2 1 3 2 2 2	40
2 1 3 1 3 2 2 3 2 2 1 1 1 3 1 3 2 3 2 1 1 1 2 1	41
3 2 2 1 2 3 1 2 3 2 3 2 1 2 1 1 3 2 1 1 2 1 2 3	42
2 2 2 3 2 2 1 3 1 1 2 3 1 3 1 1 3 1 2 2 2 1 2 3	43
1 3 2 1 2 1 3 2 2 2 1 1 1 3 1 1 3 2 1 3 2 1 3 1	44
3 2 3 1 3 1 2 1 2 1 3 1 2 2 2 1 3 1 1 1 3 2 1 1	45
2 2 3 2 2 2 1 2 1 3 2 3 1 1 3 2 3 1 1 2 1 3 2 1	46
1 1 3 2 1 1 3 2 1 3 2 1 1 2 1 3 2 3 2 3 2 2 1 1	47
1 2 2 2 3 2 3 1 3 2 2 1 2 3 1 1 1 3 1 2 1 1 3 1	48
3 1 1 1 3 2 1 3 1 3 1 1 2 1 1 1 3 1 2 1 1 3 1 1	49
1 2 2 2 1 1 3 1 2 2 3 2 2 1 1 3 1 3 2 1 3 1 1 3	50
3 2 2 2 1 1 1 3 1 2 2 3 2 1 1 3 1 1 2 3 2 3 2 1	51
2 2 2 3 2 3 1 1 3 1 2 3 1 1 3 2 1 2 2 2 3 2 1 2	52
2 3 2 3 2 2 2 1 3 1 1 2 2 2 1 3 2 1 2 3 2 3 2 1	53
3 1 2 1 1 2 3 1 2 2 1 2 1 3 1 1 1 3 2 3 2 2 2 3	54
3 2 2 1 2 2 2 3 2 1 1 3 2 2 1 1 3 1 2 1 3 2 1 3	55
1 3 2 2 2 1 2 2 3 1 1 1 3 1 3 2 2 2 3 1 1 2 1 3	56
2 2 3 2 3 2 2 2 1 2 2 3 2 3 2 1 3 2 2 2 1 1 1 3	57
1 2 2 3 2 3 1 3 1 1 3 1 2 1 2 3 1 1 1 3 2 2 1 2	58
2 3 1 3 1 1 2 3 2 1 1 1 3 1 1 2 3 2 2 2 1 2 2 3	59
1 2 3 2 3 1 1 1 3 2 2 1 2 3 1 2 3 2 2 1 1 2 2 3	60
3 2 2 2 1 3 2 1 2 2 1 3 2 2 3 2 2 1 1 3 1 2 2 3	61
3 1 2 2 3 1 2 1 2 2 2 3 1 1 2 3 2 2 2 3 2 2 2 3	62
2 3 1 1 2 2 3 1 1 1 3 2 3 2 1 1 2 3 2 2 3 2 1 2	63
3 1 2 2 3 2 1 2 2 3 2 2 3 1 3 1 1 2 1 3 1 1 2 1	64
1 1 1 2 2 2 3 1 3 1 2 2 2 3 2 3 1 2 1 3 1 3 2 1	65
3 2 1 1 2 2 1 3 1 2 2 2 3 2 2 2 3 2 2 3 2 2 3 2	66
3 2 2 2 3 2 1 2 2 3 2 2 1 3 2 3 1 1 2 1 2 1 3 2	67
1 2 3 2 1 3 2 1 3 2 1 3 1 2 3 2 2 2 1 2 3 1 1 2	68
2 3 2 2 2 1 1 1 3 1 2 3 1 2 2 3 1 1 3 1 1 1 2 3	69
2 3 2 3 1 2 1 1 2 3 1 2 3 2 2 1 2 2 2 3 2 3 2 1	70
1 2 1 3 2 2 3 2 3 1 3 1 1 2 2 2 3 2 1 1 2 2 1 3	71
1 2 1 3 1 2 3 2 1 1 3 1 3 1 1 1 2 2 3 2 3 1 1 1	72

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
1 3 1 2 2 1 1 3 1 3 1 1 3 2 2 1 1 2 1 3 1 3 2 1	73
3 1 1 3 2 1 1 1 2 2 3 2 3 1 1 2 3 1 1 1 3 1 1 1	74
1 1 2 3 2 1 1 3 1 1 1 3 1 1 3 1 2 2 3 2 2 3 2 1	75
2 2 2 3 1 2 2 2 1 2 3 2 3 2 2 1 2 3 2 2 3 1 3 2	76
3 2 1 2 2 3 1 3 1 1 1 2 2 2 3 1 1 3 1 1 2 3 1 1	77
3 1 1 2 2 3 2 1 2 3 1 1 1 2 3 1 1 2 2 3 2 1 1 3	78
2 1 2 2 3 2 1 3 1 1 3 2 1 1 1 3 2 2 1 3 1 1 3 2	79
2 2 2 1 2 3 2 1 1 2 3 1 2 1 1 3 2 3 2 1 3 2 2 3	80
1 2 1 2 1 3 2 2 3 1 1 1 2 2 3 2 3 1 2 1 3 2 3 2	81
1 2 1 1 3 1 1 1 2 2 1 3 1 3 1 3 2 2 3 2 1 1 1 3	82
3 1 1 2 2 3 2 3 1 1 1 2 3 2 3 1 2 2 3 1 2 1 2 1	83
1 1 1 2 1 1 3 2 1 3 2 2 2 1 1 2 3 1 3 1 3 1 1 3	84
3 1 2 2 1 1 1 3 1 1 3 2 1 1 3 2 3 1 1 2 3 2 2 2	85
2 1 2 3 2 3 2 3 2 2 3 2 2 2 1 3 2 3 2 2 1 2 2 1	86
3 1 3 2 2 1 2 1 2 3 2 1 3 2 2 1 3 1 3 2 2 1 2 1	87
3 1 1 1 3 1 1 1 3 1 1 3 2 3 2 2 1 1 3 2 2 1 1 1	88
2 1 3 2 1 2 2 1 3 2 1 1 3 2 1 2 3 2 3 1 2 2 3 2	89
2 2 3 2 3 2 3 1 2 2 3 1 1 2 1 2 2 3 2 3 1 1 1 2	90
1 2 3 2 3 1 1 1 3 1 3 2 2 1 1 3 2 3 1 2 2 1 1 1	91
3 1 2 2 3 1 1 2 3 1 2 2 3 1 3 1 2 1 2 3 2 1 1 1	92
1 1 3 1 2 3 1 2 1 3 2 2 1 1 3 2 3 2 1 1 3 2 2 1	93
2 1 3 2 2 3 2 2 1 2 2 3 1 3 1 1 2 2 2 1 3 1 1 3	94
2 2 2 1 2 1 3 2 3 1 1 2 2 1 2 3 1 3 2 3 1 1 1 3	95
3 1 2 1 3 1 2 2 2 1 3 1 1 2 3 1 1 2 2 1 1 3 2 3	96
2 2 2 3 1 1 3 1 1 3 1 3 1 2 2 2 3 1 1 1 2 2 3 1	97
1 2 3 1 1 2 1 1 3 1 3 2 2 3 1 2 1 1 1 2 3 2 3 1	98
2 3 2 2 2 1 2 3 2 1 3 2 3 2 1 3 1 2 2 3 1 1 2 2	99
2 2 2 1 1 3 2 3 1 3 2 2 1 2 1 3 1 1 3 2 1 3 2 1	100
3 1 2 2 2 1 2 3 2 3 2 2 2 3 1 1 3 2 2 1 1 3 1 2	101
2 1 3 2 2 1 3 1 3 1 1 1 3 2 3 1 2 1 1 1 3 2 2 1	102
3 2 1 1 2 3 1 2 1 1 2 3 1 1 3 2 3 2 1 2 1 2 1 3	103
1 1 2 3 1 1 3 2 3 2 2 1 3 2 1 2 1 3 1 2 1 3 2 1	104
2 1 1 1 2 2 3 1 3 2 2 2 3 2 2 2 3 1 2 2 3 2 1 3	105
2 1 1 2 3 1 1 3 1 1 2 1 1 3 2 1 2 3 1 3 2 3 2 2	106
1 1 1 2 3 2 1 1 2 1 3 2 3 2 2 3 2 2 1 3 2 2 1 3	107
1 3 1 3 2 2 1 3 2 3 1 1 1 2 3 2 2 3 2 2 1 1 1 2	108
3 1 1 1 2 1 3 1 1 1 2 3 2 1 2 2 3 2 2 2 3 2 3 1	109
1 3 2 2 1 2 1 1 3 2 2 2 3 2 3 1 3 1 1 2 2 1 1 3	110
3 1 3 2 2 2 1 2 1 3 2 2 1 3 1 1 2 1 2 3 2 2 3 2	111
1 3 1 3 2 2 1 2 2 1 3 1 1 3 1 1 3 1 2 2 2 1 1 3	112
3 1 3 2 2 1 1 2 3 1 1 1 2 1 1 3 2 1 2 2 2 3 2 3	113
1 2 3 1 2 3 1 1 2 1 3 2 2 3 1 1 3 2 1 2 1 2 1 3	114
1 2 1 3 1 2 1 2 3 1 3 1 2 3 1 1 1 3 2 2 1 3 2 1	115
2 1 2 3 2 1 1 1 3 1 1 1 3 2 3 1 1 1 3 1 1 3 1 1	116
2 3 1 1 2 3 2 1 3 1 1 1 2 3 1 1 2 3 2 2 3 1 1 1	117
1 1 2 2 3 1 1 2 1 3 2 3 2 3 2 3 1 3 2 2 2 1 1 2	118
1 3 1 2 1 2 2 3 2 2 2 3 1 2 2 1 1 2 3 1 1 3 1 3	119
1 1 1 3 2 2 3 2 1 1 1 3 2 2 3 1 1 3 1 2 1 1 1 3	120
3 2 2 1 1 3 1 3 1 2 2 1 2 3 1 3 1 2 3 2 1 2 2 1	121
1 3 1 1 3 1 2 1 2 1 1 3 1 1 3 1 2 2 3 1 1 2 2 3	122
3 2 1 3 1 1 1 2 2 2 3 1 1 2 2 3 1 2 3 2 3 1 1 1	123
1 1 3 1 3 2 1 3 1 2 2 3 1 2 1 1 3 2 1 2 1 2 3 1	124
2 3 1 2 1 2 1 3 2 1 3 2 3 1 1 3 1 1 1 2 1 1 3 2	125
1 3 1 2 1 1 2 3 1 2 3 1 3 1 1 1 2 3 1 1 3 1 2 1	126
1 2 3 2 3 1 1 1 3 2 1 2 2 2 3 2 3 1 2 1 2 1 3 2	127
1 1 2 1 1 3 1 3 1 1 2 2 3 1 2 1 2 3 1 1 3 1 2 3	128

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
2 1 1 3 2 3 2 1 2 2 2 1 3 2 1 3 1 1 2 3 1 1 3 2	129
2 1 2 3 2 2 1 3 1 2 2 2 3 2 2 3 1 3 1 2 2 3 1 2	130
1 3 2 2 2 3 2 1 2 3 1 1 3 1 3 1 2 1 3 2 1 2 2 2	131
3 1 3 1 1 1 2 3 2 2 1 2 3 2 1 2 2 2 1 3 2 1 3 2	132
2 1 2 3 2 3 1 3 1 1 2 3 2 3 2 2 2 3 1 2 2 2 1 1	133
3 2 1 2 3 2 2 2 3 2 2 2 1 2 1 3 1 1 2 3 2 1 2 3	134
3 1 3 2 1 2 1 2 1 3 1 1 3 1 1 1 3 1 1 1 2 2 2 3	135
1 2 3 1 3 2 3 1 1 3 2 1 1 1 2 3 2 1 3 2 2 1 2 2	136
2 2 1 1 3 1 1 3 2 3 1 3 2 2 1 2 2 3 2 3 1 2 1 2	137
1 2 3 1 1 1 2 3 1 3 1 1 2 1 2 2 3 2 2 3 2 2 2 3	138
3 1 2 2 1 1 2 3 1 2 2 1 2 3 2 3 1 1 2 2 3 1 2 3	139
3 1 1 1 2 3 2 2 1 1 1 3 1 2 1 2 3 1 1 1 3 2 1 3	140
2 1 2 2 3 2 2 3 1 2 2 2 3 1 2 1 2 2 1 3 2 3 2 3	141
2 2 2 1 2 3 2 2 2 3 2 3 2 1 2 3 2 1 1 3 2 1 3 2	142
1 1 2 2 3 1 1 1 3 1 1 2 2 3 2 3 2 3 1 1 2 2 3 1	143
2 3 1 3 2 2 2 3 1 1 2 2 2 3 2 2 2 3 1 3 2 1 1 2	144
3 1 2 3 2 1 2 1 1 2 3 1 2 3 2 3 2 3 2 1 1 1 2 2	145
1 2 3 2 3 1 3 1 3 1 1 3 1 1 2 2 2 3 2 2 2 1 2 2	146
3 2 3 1 2 1 1 1 3 2 1 2 2 3 2 2 3 1 2 1 3 1 1 1	147
3 1 1 3 2 1 3 1 1 2 1 3 1 1 1 3 2 2 1 1 2 1 3 1	148
2 2 3 2 3 2 1 3 2 2 1 1 3 1 3 2 2 3 2 2 2 1 1 2	149
2 1 3 2 1 3 2 1 1 3 2 2 3 2 2 1 3 1 1 2 1 3 2 2	150
1 1 2 2 2 3 1 1 3 2 1 2 1 1 2 3 1 1 2 3 2 3 2 3	151
2 1 3 1 1 1 2 2 3 2 1 3 2 1 2 2 2 3 1 3 1 3 1 1	152
2 3 2 1 2 1 2 3 2 2 1 1 2 3 1 3 1 2 3 2 2 3 2 1	153
2 1 2 2 2 3 1 2 1 1 3 1 3 1 1 2 3 1 1 3 1 1 3 2	154
2 2 3 1 1 2 1 3 2 3 2 1 1 2 3 1 1 2 1 2 3 1 2 3	155
3 2 1 3 2 2 2 3 2 3 1 1 2 1 3 1 1 2 2 1 3 2 2 2	156
1 1 1 3 1 2 3 1 2 2 3 2 1 1 2 2 2 3 2 3 2 3 1 1	157
3 1 1 3 1 2 2 3 2 2 3 1 3 2 2 1 1 2 1 3 1 2 1 1	158
1 3 1 2 2 1 2 3 2 1 3 2 3 1 2 3 2 1 1 1 2 3 2 2	159
3 1 1 2 2 2 1 3 1 2 3 2 1 3 1 2 1 2 3 1 1 2 3 2	160
3 1 2 1 3 1 1 3 2 3 2 1 2 2 1 1 3 2 1 1 3 2 2 1	161
2 1 2 3 1 1 2 2 1 2 3 1 3 1 1 3 1 1 2 1 3 1 3 2	162
2 2 2 3 2 2 1 2 3 1 1 3 2 3 1 2 2 2 3 2 2 2 3 2	163
3 2 1 1 1 3 1 2 2 3 2 3 2 2 1 2 1 2 3 1 1 1 2 3	164
2 2 3 2 3 1 2 1 3 2 1 3 2 2 1 3 1 2 1 2 2 2 3 2	165
3 1 1 2 2 1 1 3 1 2 1 1 1 3 1 1 3 1 3 1 1 3 2 1	166
3 1 2 2 3 2 1 3 1 1 2 3 1 1 2 2 2 3 2 1 3 2 1 2	167
1 1 1 2 1 1 3 1 3 1 3 1 3 1 1 2 3 1 2 2 2 1 3 2	168
1 1 2 2 1 2 3 2 3 1 1 2 1 3 1 2 2 3 2 2 3 1 1 3	169
2 2 1 1 3 1 2 2 2 1 2 3 2 3 1 2 1 3 2 1 3 1 3 2	170
2 2 1 1 1 3 1 2 1 3 2 3 2 2 2 3 2 2 3 2 3 2 2 1	171
2 1 2 2 3 1 2 2 2 1 2 3 1 1 3 1 3 2 1 2 1 3 2 3	172
1 1 1 2 2 2 3 1 2 3 1 3 2 1 3 2 2 2 1 1 3 1 3 1	173
1 2 1 1 1 3 2 2 3 2 2 2 3 1 2 3 2 2 2 3 1 1 2 3	174
3 1 2 2 3 2 3 1 2 3 1 1 2 1 1 2 3 2 2 1 2 2 3 1	175
3 1 2 3 1 1 3 1 1 1 2 1 2 3 1 2 1 2 3 1 1 2 1 3	176
2 2 1 1 1 3 2 2 1 2 2 3 1 1 3 2 3 1 1 3 2 2 3 1	177
2 2 3 2 1 1 3 1 1 1 2 1 3 1 3 1 2 2 2 3 2 3 2 2	178
3 1 3 1 2 2 3 1 3 2 2 2 1 1 3 2 1 2 2 1 3 1 2 2	179
1 3 2 3 1 2 1 1 2 1 3 1 1 2 3 1 2 1 1 1 2 3 2 3	180
3 1 2 1 1 2 1 3 2 3 1 1 2 2 2 3 1 3 2 2 3 2 1 2	181
1 3 1 2 1 2 2 2 3 2 1 3 2 1 3 1 1 1 3 2 1 2 3 2	182
3 2 2 1 2 3 1 1 2 3 2 2 3 1 1 2 2 2 3 1 1 2 3 2	183
1 2 3 1 1 1 3 1 2 2 2 1 3 2 2 3 2 3 1 3 1 2 1 2	184

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
1 1 1 2 1 3 1 3 1 1 3 2 2 1 2 3 1 2 3 2 3 1 2 1	185
2 2 1 3 2 3 1 3 1 1 1 2 3 2 2 2 1 1 2 3 2 3 1 2	186
2 3 1 1 3 1 1 2 1 2 3 2 3 1 1 1 2 2 1 3 2 2 2 3	187
3 2 2 2 3 1 2 1 3 2 2 2 1 1 2 3 1 3 2 1 2 2 3 1	188
3 2 2 3 2 1 1 3 2 1 1 2 3 1 2 1 1 1 3 2 1 2 3 1	189
2 1 1 3 1 3 2 1 3 2 1 1 2 2 3 2 2 3 2 2 2 1 3 1	190
2 2 2 3 1 3 1 3 1 3 2 1 2 3 2 1 2 3 1 2 2 1 2 2	191
1 2 2 3 1 2 2 3 2 3 1 1 2 2 1 3 1 2 1 3 1 1 3 1	192
3 1 2 2 1 3 2 1 2 2 2 1 3 2 1 3 2 1 1 2 1 3 1 3	193
2 1 2 3 2 1 2 2 1 3 1 3 1 2 1 2 2 3 1 1 1 3 2 3	194
2 1 2 3 2 3 1 1 1 3 2 1 1 2 3 1 2 1 1 1 2 3 1 3	195
3 2 1 1 2 2 1 3 2 1 1 2 3 1 2 2 2 3 1 1 2 3 1 3	196
3 2 2 2 1 2 2 3 2 1 1 1 3 1 2 3 2 1 1 3 2 3 1 1	197
2 1 3 2 1 3 1 1 2 2 3 2 2 3 2 2 1 1 1 3 1 1 2 3	198
2 1 2 2 3 2 2 1 3 2 2 1 2 3 2 1 3 2 3 2 3 2 1 1	199
3 1 3 2 3 1 1 1 3 2 2 1 2 1 2 3 1 1 1 3 2 1 2 1	200
1 2 1 2 1 3 1 1 3 2 2 3 1 2 3 1 3 2 2 2 1 2 3 1	201
2 2 2 1 3 1 1 3 2 1 1 3 1 1 2 1 1 3 2 3 1 3 2 1	202
2 3 2 3 2 1 2 1 1 3 1 2 1 2 2 2 3 2 1 1 3 1 1 3	203
2 1 3 1 1 3 1 3 2 2 3 2 1 2 2 3 2 2 1 2 1 1 3 2	204
3 2 3 2 2 1 2 2 1 3 2 2 2 1 1 3 2 2 1 3 1 3 2 1	205
1 1 2 1 2 1 3 2 3 1 2 3 2 3 1 1 1 2 2 3 1 1 2 3	206
2 2 1 3 1 3 1 1 2 1 3 1 3 2 3 1 2 2 1 2 1 3 2 2	207
3 1 1 3 2 3 1 3 2 2 1 1 2 3 1 2 2 2 3 2 1 1 1 2	208
1 1 2 3 2 1 1 1 3 2 1 1 1 3 1 1 1 3 2 3 1 2 3 1	209
3 2 2 1 3 2 2 1 2 3 1 2 3 1 1 2 1 2 2 3 2 3 2 1	210
1 1 1 2 3 1 3 2 2 1 3 1 3 2 1 3 1 1 2 2 1 2 3 2	211
3 1 2 1 2 1 3 1 1 3 1 2 2 1 3 2 2 1 3 2 3 1 2 1	212
1 2 1 3 2 2 2 3 2 2 3 1 3 1 2 2 2 1 2 3 1 3 2 1	213
2 1 3 1 1 2 1 3 2 2 1 3 2 1 3 2 1 1 3 1 3 2 1 2	214
3 1 1 2 2 2 3 2 1 2 2 3 2 3 1 1 3 2 2 2 1 3 2 1	215
3 2 1 3 2 1 1 3 1 1 3 1 3 1 1 2 2 1 3 1 2 2 1 1	216
1 1 2 3 2 3 2 2 1 2 3 2 1 2 3 2 1 1 1 2 1 3 2 3	217
3 1 1 2 2 1 3 2 2 1 3 1 3 2 1 1 1 2 2 3 2 2 2 3	218
3 1 1 1 2 2 3 1 1 3 1 2 1 3 2 1 1 3 1 1 1 2 3 1	219
3 2 3 2 1 2 2 1 2 3 2 3 1 2 2 2 1 2 3 1 2 1 3 1	220
2 1 2 2 1 2 3 1 3 1 1 1 3 2 2 3 1 1 2 1 3 2 1 3	221
2 1 2 3 2 1 2 2 3 2 1 2 2 3 1 3 2 1 3 1 2 3 1 1	222
3 2 3 1 2 2 3 1 1 2 1 3 2 1 3 1 2 2 3 2 2 2 1 1	223
1 3 2 1 1 3 2 2 3 2 2 2 3 1 2 2 3 1 1 1 2 2 2 3	224
3 1 1 3 2 2 2 3 1 2 2 2 1 1 3 2 2 2 1 1 3 1 1 3	225
3 1 3 1 1 3 1 2 1 1 1 2 3 1 2 1 2 2 3 2 2 1 2 3	226
1 2 3 1 2 3 1 3 2 2 3 2 2 1 1 2 1 3 2 2 1 3 2 2	227
2 1 2 3 1 2 1 2 2 2 3 1 1 3 1 3 2 3 2 2 1 1 3 1	228
3 1 3 1 2 3 1 2 2 1 1 1 3 2 3 1 2 2 2 1 2 3 1 1	229
1 2 1 3 2 2 1 1 3 1 3 2 3 1 2 3 1 3 1 1 2 1 1 1	230
2 2 2 1 2 2 3 2 2 1 3 1 2 1 1 1 3 1 3 2 2 3 1 3	231
1 3 1 1 2 1 2 2 3 1 2 1 3 2 2 3 1 1 3 2 2 3 1 1	232
2 1 3 2 3 2 1 1 1 3 2 3 2 1 3 1 2 2 3 2 1 1 1 2	233
1 3 2 1 3 2 3 1 2 1 2 3 1 2 2 2 3 1 1 2 1 2 2 3	234
2 3 2 1 2 2 3 1 1 2 2 1 3 1 1 2 1 3 2 3 1 3 1 1	235
2 3 1 2 1 2 3 1 3 1 2 1 3 1 1 3 2 2 2 1 1 2 3 2	236
3 1 1 3 1 1 3 2 1 1 3 2 1 2 1 1 1 3 2 1 1 1 2 3	237
2 2 2 1 1 3 2 3 2 3 1 2 1 1 3 1 1 1 3 1 2 1 3 1	238
2 1 2 2 3 2 2 3 1 1 2 3 2 3 2 2 2 1 1 1 3 1 3 1	239
3 1 1 2 1 1 2 3 1 2 3 1 3 1 2 3 1 2 2 1 2 2 3 1	240

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
2 1 3 1 3 1 1 1 3 1 3 1 3 1 1 2 2 3 2 1 2 2 1 1	241
1 2 3 2 1 2 1 1 2 3 1 3 1 2 1 2 3 2 2 2 3 2 3 1	242
1 1 2 1 3 1 2 1 1 3 1 2 2 3 1 2 2 3 2 3 2 2 2 3	243
2 2 2 3 1 2 3 1 2 1 1 2 1 3 1 1 3 1 3 1 1 2 3 1	244
1 3 1 2 3 1 1 2 1 1 3 2 2 3 2 3 1 1 2 3 2 2 2 1	245
1 3 1 2 3 1 1 1 3 1 1 1 3 2 3 2 1 3 1 1 2 1 2 2	246
2 3 2 2 1 1 1 2 3 2 1 2 3 2 1 3 2 1 1 2 2 3 1 3	247
2 1 3 2 1 3 2 3 2 3 1 1 3 2 2 1 2 2 2 3 2 2 1 2	248
1 3 2 3 1 1 2 3 2 2 2 3 2 1 1 1 3 1 3 2 2 2 1 1	249
3 1 2 1 1 1 2 3 1 3 1 1 2 2 3 1 3 2 1 1 2 2 3 2	250
2 3 1 2 3 1 3 1 1 1 2 2 3 2 2 2 1 1 3 2 3 2 2 2	251
1 1 1 2 1 1 3 2 1 3 2 3 2 3 1 3 2 1 1 2 1 3 2 1	252
2 1 2 3 1 1 1 2 1 2 3 2 3 1 2 1 3 2 1 1 3 1 3 1	253
1 2 2 3 2 1 1 3 1 3 2 3 1 2 2 1 2 1 3 1 2 3 1 2	254
1 3 1 3 2 1 1 3 1 1 2 3 1 1 1 3 1 3 1 2 1 1 2 1	255
2 1 1 3 2 1 1 3 2 1 3 1 2 3 2 2 1 1 1 3 1 3 1 2	256
1 1 1 2 1 3 1 1 1 3 1 1 2 2 3 2 1 3 1 3 2 1 3 2	257
1 2 1 3 1 2 2 2 1 1 3 2 3 1 1 3 1 3 1 3 2 2 1 2	258
3 1 1 2 3 2 2 2 3 2 1 1 1 2 3 2 1 2 1 3 1 2 1 3	259
1 1 1 2 1 3 1 1 2 3 1 3 2 1 3 2 3 1 1 1 2 1 2 3	260
2 2 3 1 1 2 2 1 2 3 2 1 3 1 3 1 1 1 3 2 1 1 1 3	261
2 1 3 2 1 1 1 2 2 3 1 3 1 3 2 1 3 2 2 3 1 1 2 2	262
2 3 2 1 1 1 3 2 3 2 2 2 1 2 1 3 2 3 2 3 2 1 1 2	263
1 2 1 2 3 1 2 2 2 3 1 3 1 2 3 1 3 1 1 2 3 2 1 1	264
1 1 2 1 2 2 3 1 2 1 2 3 2 3 2 2 3 2 3 1 1 3 2 1	265
1 3 2 3 1 3 1 2 2 1 2 3 1 3 2 1 2 2 3 1 2 2 2 1	266
2 2 3 2 1 2 2 2 1 3 1 2 1 3 2 3 1 3 1 2 2 1 2 3	267
1 2 1 3 1 1 1 2 3 1 1 1 3 1 2 1 3 1 2 1 3 1 1 3	268
3 1 2 2 3 2 1 2 1 2 3 2 1 1 1 3 2 1 3 2 2 2 1 3	269
2 1 2 3 1 1 2 3 2 2 1 2 2 3 2 3 2 3 2 2 3 1 2 2	270
3 1 2 1 2 2 1 3 2 1 3 1 3 2 1 1 3 2 1 2 1 2 2 3	271
2 3 1 3 1 2 3 1 1 2 2 2 3 2 3 2 2 1 2 3 1 2 1 2	272
2 1 2 3 1 1 2 3 1 1 3 2 1 1 1 3 1 3 1 2 3 2 1 1	273
3 1 3 2 3 1 1 2 2 2 3 2 2 3 2 1 1 2 2 2 3 2 2 2	274
1 3 1 1 1 2 2 3 2 1 3 1 3 2 2 1 1 2 2 3 2 3 2 1	275
3 2 3 2 2 1 1 2 3 1 1 1 3 2 2 3 2 3 1 1 2 1 1 2	276
2 3 2 3 1 2 2 2 3 2 2 1 1 3 1 1 3 1 2 2 1 1 2 3	277
1 3 2 1 3 2 1 2 2 3 2 1 1 1 3 2 1 2 1 1 1 3 1 3	278
2 3 1 2 2 3 2 2 3 2 1 2 1 3 2 2 1 2 2 3 2 3 2 1	279
3 1 2 2 3 2 1 3 2 2 2 1 1 2 3 2 2 1 1 3 1 1 2 3	280
1 2 3 1 1 1 2 1 1 3 1 1 1 2 2 3 1 3 2 1 3 1 3 1	281
2 1 2 3 1 2 3 1 2 1 2 2 2 3 2 2 3 2 1 2 3 2 3 2	282
2 2 2 1 3 1 3 2 2 2 3 1 2 2 1 3 2 1 2 3 2 2 2 3	283
1 1 2 1 1 3 1 3 1 2 2 3 2 3 1 2 3 1 3 1 1 1 2 1	284
1 1 2 3 1 1 2 1 3 1 1 2 1 3 1 3 1 1 2 3 2 1 3 1	285
3 2 1 3 2 1 3 2 1 1 2 2 2 3 1 1 2 3 2 2 2 3 1 1	286
1 3 2 3 1 3 2 1 1 2 2 3 1 2 2 3 1 2 2 3 2 2 1 1	287
3 1 1 2 1 1 2 3 2 2 2 1 3 2 3 2 3 2 2 2 3 1 1 1	288
1 2 1 2 3 1 1 1 3 2 1 3 1 3 1 1 1 3 2 3 2 2 1 2	289
2 3 1 3 2 2 1 2 2 3 2 1 2 2 2 1 3 2 2 2 3 1 1 3	290
2 1 3 2 2 3 1 3 2 2 2 1 1 1 3 2 2 3 1 1 1 3 1 1	291
2 1 1 1 3 1 3 2 3 1 2 3 2 1 1 1 2 1 3 1 1 3 2 2	292
2 3 2 1 3 2 3 2 2 2 1 3 1 3 2 1 1 3 2 2 1 2 2 1	293
1 3 1 3 1 2 2 1 1 2 3 2 3 2 2 3 1 1 1 3 1 2 2 1	294
3 2 1 1 2 1 1 3 2 2 3 2 3 1 1 1 3 1 1 3 1 2 2 1	295
3 1 3 1 2 3 2 2 1 2 1 3 1 2 1 1 2 3 1 1 1 3 1 1	296

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
2 2 2 1 3 2 2 3 1 2 2 3 2 2 3 1 1 2 1 3 1 3 2 1	297
1 2 2 1 2 2 3 1 1 1 3 2 1 3 1 2 3 2 2 1 3 1 2 3	298
2 2 2 1 2 3 2 3 2 3 1 2 2 3 1 3 2 3 2 2 2 1 1 2	299
2 1 2 2 2 1 3 2 2 1 3 1 2 1 3 1 2 1 3 1 3 1 3 2	300
1 2 3 2 3 2 2 2 1 2 3 2 3 1 1 1 3 1 2 2 2 3 2 1	301
1 2 1 3 2 1 1 2 2 1 3 1 1 3 1 3 1 1 3 1 1 2 3 2	302
2 1 2 3 1 3 2 3 1 2 2 1 3 1 1 2 2 3 2 1 2 2 2 3	303
2 2 1 1 2 3 2 1 2 2 3 2 2 2 1 1 1 3 1 3 2 3 2 3	304
1 2 1 3 1 3 1 1 2 2 1 1 3 1 1 2 2 3 2 2 2 3 1 3	305
3 2 2 1 2 1 1 3 2 1 3 1 1 1 2 3 2 1 2 1 3 1 1 3	306
1 3 2 1 1 2 2 1 3 2 2 2 3 1 1 1 2 3 2 3 2 1 3 2	307
3 1 1 1 3 1 2 2 1 2 3 1 2 2 3 2 1 1 1 3 2 3 1 2	308
3 2 1 1 3 1 2 2 1 3 1 1 3 2 2 1 1 2 3 1 1 3 1 1	309
3 1 3 1 1 2 3 2 2 3 1 1 2 1 1 3 1 1 3 2 1 1 2 2	310
2 2 1 1 3 1 3 2 3 2 2 2 3 1 1 2 1 3 2 3 2 2 2 1	311
1 2 1 1 1 3 1 1 1 3 1 3 2 1 2 3 1 3 1 2 2 1 2 3	312
1 3 2 2 1 2 2 3 1 2 2 3 1 1 3 1 2 3 1 3 1 1 1 2	313
3 2 2 2 3 2 3 2 2 2 3 2 1 2 1 1 3 2 2 3 2 2 1 1	314
2 2 3 2 1 2 3 2 3 1 3 2 2 2 1 3 1 2 2 1 1 2 3 1	315
2 1 3 2 2 1 1 1 3 2 1 2 1 3 2 2 3 2 2 2 3 1 3 2	316
1 1 1 2 2 2 3 2 3 2 2 3 1 3 1 2 2 2 3 2 1 2 1 3	317
2 1 2 2 1 3 2 3 2 2 1 2 3 1 2 1 1 1 3 1 3 1 1 3	318
2 1 2 1 1 3 1 1 3 2 1 1 2 2 2 3 1 3 1 1 3 1 3 2	319
2 1 1 3 2 2 3 1 3 1 2 3 2 2 2 3 2 2 2 3 1 2 1 1	320
3 2 3 2 1 3 1 2 2 2 1 2 3 1 1 2 2 3 1 3 2 1 1 2	321
2 1 2 1 3 1 3 1 1 3 2 3 2 2 2 1 3 2 2 3 2 1 2 1	322
1 2 1 1 1 3 1 1 3 1 1 2 1 3 2 2 3 2 2 3 2 3 2 1	323
1 3 1 2 2 3 1 1 1 2 1 3 1 2 2 1 3 1 1 1 3 2 2 3	324
3 2 2 3 2 2 1 2 1 1 3 1 1 1 2 1 3 2 2 2 3 2 2 3	325
1 3 1 1 1 2 1 3 1 3 2 1 1 3 1 3 2 3 2 2 2 1 1 1	326
1 3 1 3 1 2 1 3 2 1 3 2 1 1 1 2 1 3 2 2 1 2 2 3	327
1 1 1 2 3 1 2 2 3 2 3 2 1 1 3 2 2 1 2 3 2 1 2 3	328
1 1 3 1 1 3 2 1 1 3 1 3 1 3 1 1 1 2 2 2 3 1 1 2	329
3 2 3 2 3 2 1 2 2 2 1 3 2 2 3 1 2 1 1 2 2 3 1 2	330
1 2 2 3 2 2 3 2 2 3 2 2 3 1 3 1 1 1 2 3 2 1 2 2	331
1 3 1 2 1 1 3 2 2 1 1 1 3 2 1 1 1 3 1 3 1 1 2 3	332
2 1 3 2 2 3 1 1 3 2 2 1 3 2 2 2 1 1 3 2 3 2 2 1	333
1 3 2 1 1 3 1 1 2 3 2 1 1 2 1 2 3 1 2 3 1 2 1 3	334
1 2 3 1 3 1 2 2 3 1 1 1 3 1 2 2 2 1 2 3 1 1 2 3	335
2 3 1 2 2 3 1 1 2 2 1 3 1 3 1 3 1 1 2 3 2 1 2 1	336
1 3 2 2 1 3 2 1 1 3 1 3 1 1 2 1 2 1 3 2 3 1 1 2	337
1 2 2 1 1 3 1 2 2 3 2 1 2 1 3 2 2 1 3 2 3 1 2 3	338
3 1 3 1 2 1 1 1 3 1 1 2 2 3 1 1 1 2 1 3 1 1 3 1	339
1 3 1 3 2 1 1 1 2 3 2 2 1 1 3 1 1 1 3 1 1 3 2 2	340
1 1 1 3 2 2 2 3 2 2 1 2 3 2 3 2 3 1 1 3 1 1 2 2	341
1 2 2 3 2 3 2 2 2 1 1 3 1 1 1 2 1 2 3 1 2 3 1 3	342
2 1 2 2 3 1 1 1 2 3 1 3 1 2 3 2 1 2 3 2 1 3 2 2	343
1 2 2 2 3 2 3 2 3 1 2 3 2 2 2 3 1 1 1 2 1 2 3 1	344
2 1 1 3 1 2 1 1 2 1 3 2 3 1 3 1 3 1 1 1 2 2 3 1	345
1 2 2 2 1 2 3 1 2 2 1 3 2 3 2 1 1 3 2 3 2 2 3 2	346
3 1 2 2 1 1 3 1 1 2 1 1 1 3 2 3 2 3 1 1 3 1 1 2	347
3 2 1 1 2 2 3 1 2 3 1 1 3 1 3 2 2 1 3 2 2 2 1 2	348
2 3 2 3 2 2 1 2 3 2 2 1 2 1 1 3 1 1 3 2 3 1 2 1	349
1 3 1 3 1 1 1 2 2 3 1 1 2 2 2 1 3 1 1 1 2 3 2 3	350
2 2 1 2 2 3 1 1 2 3 2 3 1 3 1 1 1 3 2 1 2 2 2 3	351
2 3 2 2 1 1 2 3 1 3 1 1 3 1 2 1 1 2 3 1 2 1 3 2	352

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
3 1 1 1 3 2 1 2 2 2 3 2 2 3 1 2 2 1 2 2 3 2 2 3	353
2 1 3 2 2 2 1 2 3 2 1 3 2 2 1 1 2 2 3 2 2 3 1 3	354
3 2 2 3 1 1 1 3 1 2 1 3 2 2 2 3 1 2 1 2 3 2 1 2	355
2 2 1 3 1 1 3 1 2 1 3 1 2 2 1 2 2 3 1 3 1 1 1 3	356
1 1 2 1 1 2 3 2 2 3 2 3 1 1 1 2 1 3 1 2 3 2 3 1	357
1 3 2 1 1 3 1 1 1 3 2 2 2 1 3 2 2 2 1 3 2 2 1 3	358
2 1 3 2 2 2 1 1 2 3 1 3 1 2 3 2 2 2 3 1 2 1 2 3	359
2 2 1 1 1 3 1 2 3 2 2 1 1 1 3 1 1 2 3 1 3 2 3 1	360
1 1 1 3 2 3 2 3 2 1 2 1 2 3 2 2 1 3 1 1 1 3 2 1	361
1 2 2 1 1 3 2 2 1 2 3 2 3 2 2 2 1 2 3 2 3 2 2 3	362
2 2 2 3 1 1 3 1 1 3 2 3 2 2 2 3 2 1 2 2 1 2 3 2	363
2 3 2 2 1 1 3 1 1 3 2 2 2 1 3 2 2 1 1 1 3 2 2 3	364
2 2 2 1 1 3 2 1 2 1 1 3 1 2 2 3 2 3 2 3 1 3 1 2	365
1 3 1 2 1 2 2 2 3 1 2 1 3 1 2 1 3 1 1 3 1 1 1 3	366
1 2 2 2 1 3 1 3 2 2 3 2 1 1 3 1 1 3 1 2 1 2 2 3	367
3 1 3 1 1 1 2 2 3 2 1 1 2 2 3 2 2 1 3 1 3 2 1 2	368
3 1 1 3 2 1 2 1 2 3 2 2 1 1 3 1 2 3 2 1 1 2 1 3	369
1 1 2 1 2 2 3 1 1 3 1 2 3 2 1 3 2 3 1 3 2 2 1 2	370
3 1 3 2 2 2 1 3 1 1 1 2 3 1 2 1 1 1 3 1 1 2 2 3	371
2 1 1 3 1 1 1 2 3 1 3 2 2 1 2 1 2 3 2 2 3 1 3 1	372
2 2 3 1 2 1 2 1 1 3 1 1 3 2 2 3 2 3 1 2 1 1 3 2	373
1 1 3 2 3 2 2 2 1 1 2 3 2 1 1 3 1 3 1 1 2 3 1 1	374
3 2 2 3 2 3 1 3 1 1 2 2 1 3 1 1 1 2 1 3 2 1 2 1	375
2 2 2 1 3 2 2 2 3 1 2 3 2 3 2 2 2 1 2 3 1 3 1 2	376
3 2 1 1 2 2 3 1 1 1 3 2 1 2 3 1 3 2 1 3 2 1 1 2	377
2 1 3 2 2 3 1 1 2 1 1 3 1 2 2 3 1 3 1 3 1 1 1 2	378
2 2 1 1 3 2 3 1 1 3 2 3 2 2 3 2 2 2 1 2 2 3 1 1	379
1 2 2 3 1 2 2 2 3 2 2 3 1 1 1 2 1 1 3 2 3 2 2 3	380
2 3 1 1 2 2 3 2 2 3 1 2 1 1 3 2 2 1 2 3 1 1 3 1	381
3 2 2 2 3 2 2 1 2 2 3 1 3 2 1 1 3 2 2 3 1 1 2 2	382
2 3 1 2 2 2 1 3 2 1 2 3 2 1 2 2 1 3 1 3 2 2 3 1	383
2 1 1 1 2 1 3 1 3 1 2 3 1 3 1 1 2 1 1 3 1 1 1 3	384
1 3 1 1 2 3 2 2 1 2 1 2 3 2 1 3 1 3 1 1 1 2 2 3	385
1 2 2 2 1 2 3 2 1 3 2 2 3 1 3 1 3 2 3 1 2 1 1 1	386
3 2 1 1 1 3 1 2 1 3 2 2 2 3 1 3 2 1 1 2 2 2 3 1	387
3 1 1 1 2 1 3 2 1 2 1 1 2 3 2 2 1 1 3 2 3 1 3 1	388
1 2 2 3 2 1 2 1 2 2 3 2 3 2 2 3 1 1 3 1 1 1 3 2	389
3 1 3 2 2 1 1 3 2 3 2 1 1 1 2 3 1 1 1 2 3 2 1 1	390
1 2 1 3 1 2 2 3 2 3 2 3 1 1 1 3 1 1 1 3 1 1 2 2	391
2 2 1 1 2 1 3 1 1 3 2 2 2 3 2 1 3 2 1 2 3 1 2 3	392
2 2 3 2 1 2 3 2 3 1 3 1 1 2 1 1 1 3 2 2 2 1 3 2	393
3 2 3 1 2 2 1 3 1 2 1 2 3 1 2 3 1 2 1 2 3 1 1 2	394
2 3 1 1 3 1 1 3 1 1 2 2 2 1 3 1 2 2 2 3 2 1 1 3	395
2 3 2 1 2 3 1 2 2 1 2 2 3 1 2 2 1 3 2 3 2 3 2 2	396
2 3 2 3 1 1 1 3 1 3 1 1 2 3 1 2 1 3 1 2 1 2 2 2	397
1 1 2 2 3 1 1 1 2 3 1 3 2 3 2 3 2 2 2 1 1 3 1 1	398
1 2 2 1 2 1 3 1 3 2 2 1 3 2 2 2 1 3 1 1 2 3 1 3	399
1 1 1 3 1 2 1 3 1 1 1 2 2 3 1 3 2 3 2 1 2 3 1 2	400
3 2 1 3 2 2 2 3 2 2 1 1 2 3 2 2 3 2 1 2 1 1 2 3	401
1 3 1 3 1 2 1 2 2 1 3 1 1 2 3 2 1 1 3 1 1 2 1 3	402
1 3 1 1 3 2 2 2 3 1 1 1 2 1 2 3 1 2 1 3 1 1 2 3	403
2 1 3 1 1 2 3 2 1 1 1 3 2 2 2 1 3 2 1 2 1 3 1 3	404
1 3 2 1 3 1 2 3 2 1 2 3 2 2 1 1 2 3 2 3 1 1 2 1	405
2 3 1 1 1 3 2 3 1 1 1 2 1 2 3 1 1 1 2 3 2 2 3 2	406
1 2 1 3 2 1 2 1 2 2 3 1 3 2 2 2 3 2 1 2 3 1 1 3	407
3 1 1 3 1 1 1 2 3 2 2 2 3 2 1 3 1 1 2 1 1 3 2 1	408

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
1 1 2 3 1 3 2 1 2 2 3 1 1 3 1 1 1 2 3 2 1 2 1 3	409
3 2 3 1 2 1 3 1 1 2 2 2 3 2 3 2 2 2 1 1 2 3 1 1	410
2 3 2 1 3 2 1 2 3 1 1 3 1 1 2 1 1 2 3 1 1 1 2 3	411
1 2 1 3 1 1 3 2 2 1 1 2 3 1 2 1 1 2 2 3 2 3 2 3	412
3 2 3 1 2 2 3 2 1 1 3 2 1 1 3 2 1 1 1 3 1 2 1 1	413
2 1 2 3 2 1 3 2 2 2 3 2 3 2 2 1 2 2 2 3 1 1 3 1	414
2 3 1 3 2 1 1 3 2 2 2 3 2 1 2 3 2 2 2 1 1 3 2 1	415
2 1 1 1 2 3 2 1 2 3 1 3 2 3 2 3 2 1 1 1 3 1 1 1	416
3 2 1 1 3 1 3 2 1 2 2 3 1 1 1 2 2 1 3 2 1 1 3 1	417
3 2 2 3 1 3 2 3 2 1 1 1 3 1 2 2 1 2 2 3 1 2 1 1	418
1 3 2 1 2 3 1 3 2 2 1 2 2 1 3 1 2 1 1 1 3 2 3 1	419
1 2 2 2 3 2 2 1 2 1 3 1 3 2 2 3 2 3 2 2 3 2 1 2	420
2 1 1 2 2 1 3 2 1 3 2 3 2 3 2 2 3 1 1 1 2 2 2 3	421
2 3 2 1 2 2 3 1 3 1 2 2 3 2 2 1 2 2 3 2 1 2 2 3	422
3 2 2 1 2 2 1 3 1 1 3 1 3 1 2 1 1 2 2 3 1 3 2 2	423
2 2 3 1 3 2 2 3 2 3 1 2 2 1 1 3 2 1 3 2 1 2 1 2	424
3 1 2 1 3 2 1 2 1 1 2 3 1 2 2 3 1 1 3 2 1 1 2 3	425
3 2 3 1 1 1 3 1 2 1 2 2 2 3 1 3 1 3 1 2 1 1 1 2	426
1 3 2 2 1 2 3 1 2 2 2 3 1 1 3 1 1 1 2 2 3 2 2 3	427
3 2 1 1 3 2 1 2 2 2 3 1 1 2 2 2 3 1 2 3 1 3 2 2	428
2 1 1 2 1 3 2 3 2 2 1 2 1 1 3 2 3 1 1 1 3 1 3 2	429
1 1 1 2 3 1 1 2 2 3 1 2 3 2 3 2 1 2 1 2 3 1 1 3	430
1 3 1 1 1 3 2 3 1 3 2 2 3 2 2 1 1 3 2 1 2 2 2 1	431
2 2 2 1 2 3 2 3 2 3 1 1 2 2 3 2 3 2 1 2 1 2 1 3	432
3 2 1 1 2 1 2 3 1 2 1 3 1 1 1 2 3 2 1 1 1 3 1 3	433
3 1 3 1 1 2 2 3 2 2 2 1 1 1 3 1 2 1 3 2 2 3 2 1	434
3 1 1 2 2 2 3 2 2 1 1 3 1 1 2 3 1 3 2 2 2 3 1 2	435
1 2 1 3 2 3 1 2 3 1 2 2 1 1 1 3 1 3 1 1 2 2 2 3	436
1 2 1 3 1 2 3 2 2 2 1 3 2 2 3 1 3 1 2 2 1 2 2 3	437
1 1 3 1 3 2 3 2 1 1 1 2 1 3 1 1 1 3 2 3 1 2 1 2	438
2 3 2 3 2 1 2 2 3 1 2 2 3 2 2 3 1 3 1 2 1 1 1 2	439
2 1 3 2 1 2 1 3 2 3 1 3 1 1 1 3 1 3 2 2 1 1 1 2	440
1 1 1 3 1 2 1 1 3 1 1 1 3 1 3 1 2 3 1 2 3 2 2 2	441
3 1 1 3 2 2 1 2 2 3 1 1 1 2 1 3 1 3 1 1 3 2 1 2	442
1 2 3 2 1 2 3 2 1 2 1 3 1 1 1 3 1 3 2 1 1 1 2 3	443
3 1 2 3 2 2 2 3 2 1 1 1 3 1 2 2 3 1 1 1 2 2 3 1	444
1 1 2 2 2 1 3 1 3 1 3 2 1 2 2 2 3 2 3 2 2 3 2 1	445
1 1 2 2 2 3 2 2 2 3 1 1 1 3 1 1 1 3 2 1 1 3 2 3	446
1 1 1 3 1 3 2 1 3 2 3 2 2 1 2 2 3 2 2 1 3 1 2 1	447
3 2 1 2 3 2 2 3 2 1 2 1 2 3 2 2 3 2 2 3 1 2 1 2	448
3 2 1 3 1 1 2 2 2 3 2 2 3 1 3 2 1 2 2 2 3 2 1 1	449
1 2 3 1 1 2 2 2 1 3 2 2 1 3 2 3 2 1 1 3 1 1 1 3	450
1 2 3 1 2 1 1 3 1 1 1 2 3 2 2 3 1 2 3 1 1 3 2 1	451
2 2 3 1 2 3 1 2 3 1 1 3 1 2 1 1 2 3 2 1 3 1 2 1	452
1 3 1 2 3 1 2 1 2 3 1 2 1 2 1 3 1 2 2 1 3 1 2 3	453
2 2 3 1 1 1 3 2 2 1 3 1 1 1 3 1 2 1 3 1 2 3 2 2	454
3 2 2 2 1 1 2 3 2 2 1 3 2 2 1 3 1 1 1 3 2 1 1 3	455
3 1 3 1 2 2 2 1 1 3 2 2 2 3 1 1 3 2 3 1 1 1 2 1	456
2 2 2 3 2 2 1 3 2 1 3 2 2 3 2 2 1 2 1 1 3 1 3 1	457
2 1 2 3 1 3 1 1 2 1 3 2 2 2 3 2 2 1 3 2 3 1 1 2	458
2 2 3 1 1 1 3 2 2 2 1 1 1 3 1 1 3 1 3 1 2 1 1 3	459
1 1 3 2 3 1 3 2 2 3 1 1 1 2 3 1 1 1 2 1 2 3 2 2	460
3 2 2 1 3 1 1 1 2 3 1 1 1 2 3 1 3 2 1 3 2 2 1 2	461
2 1 1 3 2 1 2 2 3 2 1 2 2 2 3 2 3 2 3 2 3 2 1 2	462
2 3 2 1 2 2 1 3 2 1 1 1 3 1 1 3 1 3 1 3 1 1 2 1	463
3 1 3 1 1 3 1 3 1 1 1 2 1 1 3 2 2 3 1 1 1 2 1 1	464

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
3 2 1 1 1 3 2 1 3 1 1 1 2 1 3 1 1 2 2 3 1 3 2 2	465
3 2 3 2 3 2 2 1 2 2 2 3 2 2 2 3 2 1 1 1 3 2 1 2	466
2 2 2 3 1 2 3 2 1 2 3 1 1 2 1 2 1 3 2 1 2 3 1 3	467
1 1 3 1 2 2 3 2 3 2 3 1 1 2 1 3 2 2 3 1 1 1 2 2	468
2 1 2 1 1 1 3 2 2 2 3 1 1 3 1 2 3 1 3 2 3 1 2 1	469
1 3 1 2 1 1 1 3 1 3 1 2 2 2 1 1 3 1 2 3 2 1 2 3	470
3 1 1 3 1 1 2 2 1 1 3 2 2 3 1 3 1 1 2 2 1 1 3 1	471
2 1 3 1 3 1 1 1 2 2 2 3 1 2 1 1 1 3 1 1 1 3 1 3	472
1 1 1 3 2 2 2 1 2 3 1 1 3 2 2 1 2 2 3 1 3 2 1 3	473
1 1 1 2 1 3 2 3 2 1 1 3 2 1 1 1 3 1 3 1 2 3 1 2	474
2 1 2 3 1 2 3 1 2 2 2 1 3 2 2 1 2 1 1 3 1 3 2 3	475
2 1 3 1 2 1 1 1 2 3 2 2 1 2 3 1 2 3 1 3 2 1 1 3	476
1 3 1 2 2 3 1 2 2 3 2 3 1 2 3 1 2 2 2 3 2 1 2 1	477
2 2 1 1 3 1 1 3 1 1 2 2 3 2 1 2 1 2 3 1 3 1 3 2	478
3 2 1 3 1 1 2 3 2 2 2 1 3 1 3 2 2 3 1 1 2 1 2 1	479
3 1 3 1 1 1 2 1 3 2 1 1 3 1 1 3 2 1 1 1 2 1 3 1	480
1 2 2 3 1 1 3 2 2 3 2 2 1 2 3 2 3 1 1 3 1 2 2 2	481
2 1 1 1 2 3 2 2 3 2 3 2 1 3 1 3 2 1 1 2 2 1 3 1	482
1 1 1 2 1 1 3 1 3 2 2 2 3 1 3 1 1 3 2 2 3 2 2 2	483
1 3 2 2 3 2 1 1 2 1 1 3 1 1 3 2 3 1 2 2 2 1 1 3	484
3 2 2 1 3 1 1 2 3 2 1 2 1 2 1 3 1 3 2 2 1 3 1 2	485
2 2 3 1 2 1 2 2 3 1 1 1 3 1 3 1 1 1 3 2 2 1 2 3	486
2 2 1 1 1 3 1 3 1 3 1 1 1 2 3 2 2 2 3 1 2 2 1 3	487
2 3 2 3 1 1 2 2 2 3 1 3 2 1 2 2 1 3 2 1 1 3 1 1	488
2 1 1 2 2 2 3 1 1 2 3 2 3 1 1 1 3 2 2 3 2 2 1 3	489
1 2 3 2 3 2 2 2 3 1 1 1 3 1 2 3 1 2 3 1 2 2 2 1	490
1 1 3 2 2 1 2 3 2 2 3 1 2 1 2 2 3 1 3 2 3 1 1 1	491
2 1 3 1 2 1 1 1 3 1 1 3 1 2 1 3 1 3 1 2 2 2 1 3	492
3 1 2 3 1 1 2 3 2 1 3 1 2 1 2 1 2 3 2 1 1 2 3 1	493
3 1 1 3 1 1 2 1 3 2 2 2 1 2 3 2 1 1 1 2 3 1 2 3	494
3 2 1 3 2 1 2 1 2 1 3 2 2 1 1 1 3 1 2 3 1 3 2 2	495
3 2 2 1 2 2 2 3 2 3 2 1 2 3 1 2 2 1 2 3 1 2 2 3	496
1 3 1 3 1 2 2 1 3 1 1 1 2 2 3 1 3 1 3 1 1 2 2 1	497
3 2 1 2 3 1 2 1 3 1 3 2 2 2 1 2 1 3 2 3 1 2 1 1	498
3 2 2 1 3 1 1 1 3 1 1 2 3 1 1 1 2 2 3 1 1 3 2 1	499
1 1 3 1 1 2 3 1 3 1 1 2 1 2 1 3 1 3 1 2 3 1 1 2	500
1 1 1 3 1 3 1 1 2 1 3 2 3 2 2 2 1 1 3 1 1 3 1 2	501
3 1 2 3 2 3 2 2 1 2 2 3 1 2 1 3 1 1 1 2 2 1 3 1	502
2 1 3 1 3 2 2 1 2 1 3 1 3 1 2 1 2 2 3 2 1 2 3 1	503
3 1 3 1 3 2 2 3 1 1 2 1 1 3 2 2 1 1 1 3 1 2 1 2	504
1 3 1 2 1 2 3 1 1 1 2 1 3 1 2 2 3 2 2 1 3 1 3 1	505
3 1 3 2 3 1 1 2 1 3 1 1 1 3 1 2 1 2 3 2 2 1 1 2	506
1 1 1 3 1 3 1 2 1 2 2 3 1 1 3 1 3 1 1 2 1 1 1 3	507
3 2 2 1 2 1 3 1 1 2 1 1 3 2 2 3 2 1 1 1 3 2 3 2	508
2 3 1 2 1 3 2 1 2 3 1 2 1 1 2 3 2 3 2 2 2 1 2 3	509
2 2 2 3 2 2 3 2 2 1 1 3 2 1 2 3 2 3 1 2 2 2 1 3	510
2 1 1 1 3 2 3 2 2 3 2 3 2 2 1 1 1 3 1 2 2 1 1 3	511
2 3 2 3 2 2 2 3 1 2 2 3 1 2 2 1 1 2 3 2 2 1 2 3	512
1 2 2 1 1 2 3 1 1 2 3 1 3 2 3 2 2 3 2 1 1 2 3 2	513
2 1 3 1 2 3 2 2 2 3 2 3 1 3 2 2 2 3 1 2 1 2 2 1	514
3 1 1 2 3 1 1 2 1 3 2 1 1 2 1 3 1 2 3 1 2 2 2 3	515
1 1 2 1 3 2 3 2 3 2 2 3 2 2 1 2 1 2 3 1 2 2 1 3	516
2 1 3 1 2 2 1 3 1 1 3 1 2 3 2 2 3 2 3 2 1 2 2 1	517
1 1 2 3 2 3 2 3 2 3 2 2 1 1 1 2 3 1 1 2 2 2 3 2	518
3 1 1 2 2 1 1 3 2 1 2 1 2 3 1 3 2 3 2 1 3 1 1 1	519
2 2 1 2 2 3 2 3 2 3 2 1 1 3 2 1 3 2 3 2 1 1 1 2	520

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
3 2 1 3 2 1 1 1 3 1 3 1 1 2 2 3 2 2 2 1 3 2 1 2	521
1 1 3 2 2 2 3 2 1 1 3 1 1 3 2 1 3 2 2 3 1 1 2 1	522
1 3 2 2 1 2 1 3 2 1 2 1 3 2 1 3 2 1 2 1 3 1 3 1	523
3 1 1 1 3 1 1 1 2 3 2 3 2 1 2 1 3 2 2 2 1 1 2 3	524
2 2 3 2 3 1 3 2 1 1 2 3 1 1 2 3 1 2 3 2 1 2 2 1	525
3 2 1 3 1 3 2 2 3 2 1 1 1 2 1 3 1 3 1 1 2 1 1 1	526
1 2 2 1 1 2 3 2 1 3 1 2 2 3 2 1 1 3 1 3 1 2 1 3	527
2 2 1 3 2 3 2 3 2 2 2 3 2 1 3 1 2 1 3 1 1 2 2 1	528
1 3 1 3 1 3 2 2 2 3 2 3 2 1 2 1 2 3 2 1 2 1 1 1	529
2 2 1 1 3 2 2 2 1 3 2 3 1 3 1 2 2 2 3 2 2 1 1 3	530
1 2 3 1 1 3 2 2 2 1 2 2 3 1 1 2 1 3 2 1 3 2 3 1	531
1 2 1 2 2 2 3 2 3 2 2 3 2 1 2 3 2 2 2 3 2 3 1 1	532
1 1 1 3 2 3 2 2 2 1 2 1 3 1 1 3 1 2 2 2 3 1 2 3	533
1 1 3 1 3 1 2 1 2 3 1 2 2 2 3 2 2 1 3 2 2 3 2 1	534
1 1 3 1 1 3 1 1 1 2 3 1 3 2 3 1 2 1 1 2 3 2 1 1	535
2 1 3 2 3 2 2 2 3 1 2 1 2 3 2 2 1 1 3 1 1 3 2 2	536
3 2 1 3 1 1 1 3 2 3 1 2 1 3 1 2 2 1 3 2 1 1 2 1	537
3 1 2 1 1 1 2 3 2 2 1 1 3 2 2 1 3 2 1 2 3 1 2 3	538
1 3 1 2 2 1 3 1 1 3 1 1 2 2 3 2 2 2 1 3 1 1 2 3	539
1 2 1 2 2 2 3 1 3 1 1 3 2 3 2 3 1 1 1 2 3 1 1 2	540
2 3 1 3 2 1 1 1 2 1 3 2 2 2 1 2 3 1 3 2 1 3 2 1	541
2 2 1 3 1 3 1 3 2 1 3 1 2 1 1 1 3 1 2 2 2 3 1 2	542
1 2 2 3 2 2 2 1 1 3 2 2 3 2 2 3 1 2 1 1 3 1 2 3	543
3 2 2 3 2 1 1 1 3 2 2 1 1 1 3 2 3 2 3 1 1 2 2 2	544
1 2 1 3 1 2 2 3 2 3 2 3 2 2 2 3 2 2 1 2 1 3 2 1	545
3 2 1 1 3 2 2 1 2 2 3 1 3 1 1 2 3 1 2 1 1 2 1 3	546
2 1 3 1 2 2 1 3 2 2 3 1 2 1 1 3 2 3 2 3 2 1 1 2	547
1 1 1 2 3 2 1 1 1 2 3 1 1 3 1 3 2 3 2 2 2 3 2 2	548
3 1 2 1 3 1 1 3 1 1 1 2 3 2 1 2 1 2 1 3 2 3 1 2	549
2 1 2 1 3 1 3 2 3 2 1 2 3 2 2 1 2 3 1 2 1 1 1 3	550
2 1 2 3 1 1 3 2 3 1 2 1 1 3 1 2 3 1 1 3 1 1 2 2	551
2 3 2 2 3 1 3 1 1 2 1 3 2 1 1 3 1 3 1 1 2 2 2 1	552
2 1 3 1 2 1 1 2 3 2 3 1 1 3 2 1 1 2 1 1 3 2 3 1	553
3 2 1 2 2 1 2 3 1 2 3 1 2 1 3 2 1 3 2 1 1 3 2 1	554
1 3 1 2 1 2 3 1 2 2 2 1 3 2 1 2 2 3 1 1 2 3 2 3	555
1 1 2 2 1 1 3 2 2 2 3 2 1 3 1 3 2 3 1 2 2 2 3 1	556
1 1 3 1 1 1 2 1 3 1 2 3 2 1 3 2 1 1 3 1 2 3 2 2	557
2 2 3 1 3 1 1 3 2 2 3 2 2 3 2 1 1 2 1 1 3 1 1 2	558
1 3 2 3 2 3 1 1 1 2 1 3 2 3 1 1 1 3 2 2 2 1 1 1	559
2 2 2 1 2 3 2 1 3 2 1 3 1 2 2 2 1 2 3 2 3 1 1 3	560
1 2 2 1 1 2 3 1 3 1 1 1 2 2 1 3 2 3 2 3 2 2 1 3	561
1 2 3 2 2 1 1 2 1 3 2 3 1 2 1 3 2 1 1 1 3 2 3 1	562
2 1 2 3 2 2 3 1 2 1 1 1 2 3 1 2 2 1 2 3 1 3 2 3	563
2 2 1 2 2 1 3 1 3 2 2 3 2 3 2 3 2 3 1 2 1 2 1 2	564
2 3 2 2 3 2 2 1 2 3 1 2 2 3 1 3 2 2 1 3 1 1 2 1	565
1 1 2 2 2 3 1 3 2 2 1 1 3 1 1 3 1 1 3 2 3 2 1 1	566
1 1 1 3 1 2 1 1 1 3 2 2 1 1 3 2 3 2 2 2 3 2 1 3	567
2 3 2 2 3 1 3 1 2 3 1 2 1 2 2 3 2 1 2 1 1 3 2 2	568
2 1 1 1 2 1 3 2 3 1 1 2 3 1 3 2 2 1 2 1 3 1 3 2	569
1 2 1 3 1 2 3 2 2 1 2 3 1 2 1 3 2 2 1 3 2 2 1 3	570
3 2 2 1 1 3 2 3 1 1 3 1 2 1 2 3 2 1 2 2 3 2 2 1	571
2 1 1 3 1 1 1 3 2 1 1 1 3 2 2 2 3 2 1 3 1 2 3 2	572
1 1 3 1 3 1 1 1 3 2 2 2 3 1 2 2 3 1 1 2 1 1 1 3	573
1 2 1 2 2 1 3 1 2 3 2 3 1 3 2 2 1 2 1 2 3 2 3 2	574
1 3 2 2 2 3 1 3 2 2 2 1 3 2 1 2 2 3 2 3 1 1 2 1	575
1 2 3 2 2 1 1 1 2 3 1 3 1 3 1 2 2 3 2 3 2 1 2 1	576

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
2 1 1 1 2 3 2 2 3 2 3 1 2 2 1 2 2 3 2 3 1 3 1 2	577
2 1 1 3 1 1 2 2 3 1 1 3 2 1 1 3 1 3 2 2 1 2 2 3	578
1 3 1 3 1 2 1 3 1 1 2 2 1 1 3 2 2 2 3 2 2 3 1 2	579
3 1 1 3 1 1 2 3 2 2 1 1 3 1 1 1 2 1 2 3 2 1 1 3	580
2 1 2 2 2 3 2 3 1 2 2 1 1 3 1 1 3 2 2 3 1 3 1 1	581
1 3 2 2 1 3 1 1 2 2 2 3 2 3 2 1 3 2 1 3 1 1 2 2	582
1 1 3 2 2 2 1 2 2 3 2 2 3 1 2 3 2 2 3 2 1 2 2 3	583
3 1 1 2 3 1 3 2 2 2 1 1 3 1 3 2 2 2 1 2 1 3 2 1	584
1 3 2 3 1 1 3 1 2 2 3 2 1 2 3 2 1 3 2 1 2 1 1 1	585
1 3 2 2 3 1 1 1 2 3 1 3 2 1 2 2 1 1 3 2 1 1 2 3	586
1 2 3 2 3 2 2 1 2 2 2 3 1 3 1 2 3 1 3 2 1 1 2 2	587
1 1 1 2 1 3 2 3 2 2 3 2 2 3 1 1 3 2 2 3 2 2 1 2	588
3 2 1 3 1 3 1 1 1 3 1 2 1 2 1 2 3 2 1 3 2 2 2 1	589
3 1 3 1 3 2 1 2 2 2 3 1 2 3 1 1 2 3 1 2 2 1 2 1	590
3 1 3 2 1 2 1 1 3 2 2 2 1 3 2 3 2 1 2 1 2 2 3 1	591
1 2 1 1 2 3 2 3 1 2 2 1 2 2 3 1 2 2 3 1 3 1 3 1	592
2 2 1 3 2 2 3 2 2 1 2 3 2 3 1 3 1 3 2 1 1 2 1 1	593
1 1 1 2 3 1 3 2 1 2 1 2 2 3 1 1 2 2 3 2 3 1 2 3	594
1 1 2 2 1 3 1 1 3 2 1 1 3 2 1 3 1 3 2 2 2 1 1 3	595
2 3 2 1 1 3 2 2 2 1 1 1 3 2 1 1 3 1 1 1 2 3 2 3	596
3 1 1 1 2 3 1 2 1 1 3 2 2 3 1 2 1 2 1 1 3 1 1 3	597
1 1 2 3 1 3 2 1 3 2 2 2 3 2 1 2 2 2 3 1 3 2 2 2	598
1 3 2 3 1 1 2 3 2 1 1 3 1 2 2 1 2 3 2 1 2 2 2 3	599
3 2 1 1 2 2 3 1 1 2 2 3 1 1 1 3 1 2 1 1 3 2 3 2	600
2 1 2 3 2 2 2 1 1 3 2 1 3 2 3 1 1 1 2 1 3 1 3 2	601
3 2 1 2 2 3 1 1 1 2 2 3 1 1 2 2 1 3 1 1 3 2 1 3	602
1 1 2 1 2 3 2 1 1 2 3 2 1 3 2 2 3 1 1 1 3 2 3 1	603
2 3 1 1 2 1 2 2 3 1 3 1 1 2 2 1 2 3 1 3 1 3 2 2	604
2 1 3 2 3 2 1 1 1 2 3 1 2 3 1 1 3 1 1 1 3 2 1 2	605
3 2 1 2 3 2 3 2 1 1 1 3 1 1 1 2 2 2 3 1 2 3 2 1	606
1 1 2 2 3 2 2 2 3 1 1 1 3 2 2 2 3 2 2 3 1 3 1 1	607
1 1 2 2 3 2 2 2 3 1 3 2 1 3 2 1 2 2 1 3 2 1 3 2	608
2 1 1 2 2 3 1 3 2 2 2 3 1 1 2 1 1 3 1 3 1 3 2 2	609
2 3 2 2 3 1 2 2 3 2 1 1 3 2 3 2 2 2 1 2 2 3 2 2	610
3 1 1 1 2 2 2 3 2 3 1 3 2 1 2 3 2 1 2 2 2 3 1 1	611
2 1 1 3 1 1 2 3 1 1 2 3 2 3 1 1 3 2 3 1 1 2 1 2	612
2 1 1 2 3 2 3 1 1 3 2 2 2 3 2 3 1 1 1 3 1 2 1 2	613
2 2 3 2 1 2 1 2 3 1 1 1 3 2 1 1 3 1 1 3 1 1 3 2	614
2 1 3 1 3 1 3 1 1 3 1 1 3 1 1 1 2 1 1 3 1 1 2 1	615
1 2 2 2 3 1 1 1 2 3 2 2 1 1 2 3 1 3 1 3 1 3 1 2	616
2 2 3 2 3 2 3 2 1 2 1 2 1 3 2 1 2 2 1 3 1 1 2 3	617
1 2 2 3 2 2 1 3 2 1 2 2 3 1 2 3 2 3 1 1 3 2 2 1	618
2 3 2 2 2 3 2 1 2 2 2 3 1 1 2 3 1 1 1 2 3 1 1 3	619
2 3 2 2 1 3 1 2 2 3 2 3 2 2 1 1 1 2 3 2 1 3 2 2	620
2 1 2 1 3 1 3 2 1 2 2 3 2 1 2 1 3 1 3 1 3 1 1 1	621
1 1 1 2 1 3 2 1 1 3 1 1 2 3 2 1 3 2 2 3 2 2 3 1	622
2 3 1 3 2 3 2 3 1 2 2 2 1 2 3 1 2 2 1 1 3 2 2 1	623
1 3 1 1 2 2 2 3 2 2 3 2 1 3 2 3 2 2 1 2 3 1 2 2	624
3 1 2 2 3 1 1 3 1 1 1 3 1 1 1 2 1 3 2 2 2 3 1 1	625
3 1 2 1 1 2 1 3 1 3 1 1 2 1 3 2 1 3 1 3 2 2 1 1	626
3 1 2 2 3 1 1 1 2 2 2 3 2 1 3 2 2 1 2 1 3 2 3 1	627
3 1 2 2 2 1 1 3 1 1 3 1 2 3 1 1 2 1 1 2 3 2 1 3	628
2 2 2 3 1 3 1 3 1 1 1 3 2 1 3 1 1 2 1 1 3 1 2 1	629
3 1 2 2 1 1 3 1 3 2 1 1 1 2 3 1 3 2 1 2 1 1 3 1	630
2 2 2 3 1 2 1 3 1 1 2 2 3 1 1 1 2 2 2 3 1 3 1 3	631
2 3 1 1 3 1 1 3 1 3 2 3 2 2 1 2 1 1 3 1 2 2 2 1	632

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
3 2 3 1 1 1 2 3 1 2 2 2 1 3 1 3 2 1 1 2 1 1 3 2	633
1 1 1 2 1 1 3 1 1 2 1 3 1 3 1 3 1 3 1 1 1 3 2 2	634
3 2 2 3 2 1 1 1 3 2 1 1 2 1 3 1 3 1 1 1 2 2 1 3	635
1 3 2 3 1 2 2 2 1 3 1 2 2 1 2 3 2 3 1 2 3 1 2 2	636
1 3 1 3 2 1 2 1 3 2 2 2 1 3 1 2 2 2 1 2 3 2 1 3	637
1 2 3 1 2 2 1 3 1 2 1 3 2 3 1 1 1 2 2 3 2 2 1 3	638
1 2 3 1 1 1 2 3 2 1 2 2 1 3 2 2 2 1 3 1 3 2 2 3	639
1 2 1 2 2 3 1 3 2 3 1 3 1 3 2 2 1 2 2 3 2 2 1 1	640
1 3 1 2 3 2 3 2 1 2 2 3 1 1 2 2 1 1 3 1 1 3 2 2	641
2 1 1 2 3 2 3 2 2 3 1 2 1 3 1 1 2 1 3 1 3 1 2 1	642
1 1 1 2 2 1 3 2 2 3 1 1 1 3 2 1 2 3 1 3 1 1 1 3	643
2 2 2 1 3 1 3 2 2 3 1 1 3 1 1 1 2 3 2 2 1 1 2 3	644
3 1 2 2 3 2 2 3 1 2 2 1 2 2 3 1 2 3 1 1 2 2 2 3	645
2 3 2 2 3 2 2 3 2 2 3 1 1 2 2 3 1 1 3 1 1 2 2 1	646
1 2 2 1 1 3 2 1 1 3 1 1 2 2 3 1 3 1 3 2 2 2 3 1	647
3 2 1 2 3 2 2 3 2 1 1 2 3 2 1 2 2 1 1 3 1 1 1 3	648
2 1 3 2 2 3 2 3 1 2 2 2 1 2 3 2 1 1 2 3 1 2 2 3	649
2 3 1 2 1 1 2 3 1 1 1 3 2 2 2 1 2 1 3 1 3 1 3 1	650
3 2 1 1 3 1 2 2 3 2 2 2 3 2 1 2 3 1 2 1 1 3 1 2	651
2 2 3 1 1 2 2 1 1 3 1 3 2 1 1 3 1 2 3 2 2 2 1 3	652
1 1 3 2 3 2 2 2 3 2 2 2 1 3 1 3 2 1 1 1 3 1 2 1	653
1 3 1 3 1 3 1 2 1 1 1 3 2 1 2 1 3 1 1 3 2 2 1 1	654
1 2 2 1 2 3 1 1 2 1 3 2 2 1 3 1 1 1 3 1 3 1 3 2	655
2 2 3 2 2 3 1 2 1 2 2 1 3 1 3 1 1 2 3 2 3 2 2 2	656
2 2 2 1 2 2 3 1 3 1 3 2 2 2 3 2 2 1 2 2 2 3 2 3	657
1 3 2 3 2 2 1 1 3 1 1 3 2 2 3 1 2 2 1 2 2 3 1 2	658
3 1 3 1 1 1 2 3 1 2 2 3 1 1 2 3 2 2 3 1 2 1 1 2	659
3 1 2 1 1 3 2 1 2 2 1 3 2 1 2 3 1 3 2 3 2 1 1 2	660
2 2 2 3 1 2 2 2 1 1 3 1 3 2 3 2 2 3 1 1 2 3 2 1	661
1 1 3 2 2 1 3 2 1 1 1 2 1 3 1 3 2 1 3 1 1 1 3 1	662
3 2 1 1 1 3 2 1 2 3 1 1 2 1 2 3 2 3 1 1 1 2 3 2	663
2 1 1 2 1 1 3 2 3 2 3 2 2 3 2 3 1 1 2 3 2 1 1 2	664
1 1 1 3 1 2 2 2 1 3 1 3 2 1 3 1 1 1 3 1 3 1 1 2	665
2 2 1 3 2 2 2 3 1 3 2 2 3 1 1 1 2 1 3 2 1 1 1 3	666
2 1 1 2 1 3 2 1 2 3 1 3 2 1 1 3 1 2 2 3 1 1 1 3	667
3 1 1 3 1 2 2 1 3 1 2 2 3 1 2 3 2 2 1 3 2 2 1 1	668
2 1 1 1 3 1 3 1 3 1 1 3 2 2 1 3 2 1 1 2 1 3 1 1	669
2 1 1 3 2 1 2 3 1 3 1 1 1 2 3 1 2 3 2 3 2 2 1 2	670
3 1 3 2 2 2 3 2 2 2 3 2 2 1 3 2 2 1 2 2 3 1 2 1	671
1 1 3 2 1 1 1 3 1 1 1 2 3 2 2 1 1 3 1 3 2 1 3 2	672
1 2 3 1 3 1 1 2 2 3 2 2 3 2 2 3 1 1 1 2 3 2 1 1	673
2 2 1 3 1 2 2 1 3 1 3 2 1 3 2 1 3 1 1 3 1 1 2 1	674
2 1 3 2 3 1 2 3 1 1 3 1 1 3 2 2 1 3 1 1 1 2 2 1	675
2 1 1 2 3 2 1 3 2 1 1 2 3 2 3 1 2 3 1 2 1 1 3 2	676
2 2 3 1 3 1 1 1 3 1 1 2 1 1 3 2 1 3 2 3 2 1 1 1	677
2 1 1 2 3 1 3 2 3 1 3 1 2 2 1 2 1 3 1 2 2 3 2 2	678
3 2 1 2 1 1 3 1 1 1 2 3 2 3 2 3 2 2 2 3 1 2 2 1	679
3 2 3 1 1 2 3 2 3 2 2 1 1 2 3 1 1 3 1 2 1 2 1 2	680
3 1 1 1 3 2 2 1 2 2 1 3 2 1 3 2 2 1 1 1 3 1 2 3	681
2 1 3 1 1 2 2 3 2 3 2 2 2 3 1 2 1 1 3 2 3 1 2 1	682
2 3 1 2 2 2 1 3 1 2 2 3 1 3 1 3 2 2 1 1 1 2 3 1	683
1 2 2 1 2 2 3 1 3 2 2 2 3 1 1 2 3 2 2 3 1 2 1 3	684
1 2 1 3 2 1 3 2 2 1 2 3 2 2 2 3 1 2 2 2 1 3 2 3	685
1 2 1 3 1 1 3 1 1 3 1 1 2 1 1 1 3 2 2 1 3 1 3 1	686
3 1 2 3 2 2 3 1 1 1 3 2 1 1 2 3 1 1 2 2 2 3 2 1	687
3 1 3 1 2 2 3 1 2 1 3 2 1 3 1 1 1 2 3 1 2 1 1 1	688

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
2 3 1 3 1 3 1 1 2 1 1 1 3 2 1 2 3 1 1 2 2 2 3 1	689
2 1 2 1 1 1 3 1 2 3 1 2 3 2 3 1 1 2 2 1 3 2 1 3	690
2 2 1 2 3 2 1 1 3 1 1 2 3 2 2 2 3 1 3 1 3 1 1 1	691
1 3 2 1 1 1 2 3 1 2 3 1 1 2 3 1 2 1 2 3 1 2 3 1	692
3 1 1 1 2 2 2 3 2 3 2 2 1 1 1 3 2 2 3 1 1 2 3 1	693
3 1 2 3 1 1 2 3 1 2 2 3 2 3 2 2 2 1 1 3 2 1 2 1	694
3 1 1 1 2 1 1 3 2 3 1 3 1 3 2 2 1 1 2 3 1 1 1 2	695
2 3 2 2 3 1 1 1 2 1 3 2 2 1 2 2 1 3 2 2 2 3 2 3	696
2 2 2 3 1 3 1 3 2 1 2 1 2 2 3 1 2 1 2 3 1 3 1 1	697
1 2 2 3 2 3 2 3 2 1 1 1 3 2 1 1 3 1 2 2 2 1 1 3	698
2 1 2 1 3 2 2 2 3 1 1 3 2 3 2 3 1 2 3 2 1 2 2 2	699
3 2 3 1 1 3 2 2 1 2 1 3 2 3 2 1 2 1 1 1 3 1 1 2	700
3 2 1 2 3 2 2 3 1 1 2 1 3 2 1 1 1 2 1 3 1 2 2 3	701
2 2 1 3 1 1 1 3 2 3 2 3 1 2 2 2 3 2 3 2 1 2 2 2	702
2 2 2 1 3 2 1 1 2 1 2 3 2 1 1 3 1 3 1 2 3 2 3 1	703
1 3 2 1 2 3 2 1 2 1 3 1 2 3 1 2 3 2 2 2 3 2 2 2	704
1 2 2 2 1 1 3 2 1 1 1 3 2 3 2 1 3 1 3 1 2 1 1 3	705
1 2 2 2 3 2 3 2 2 3 1 1 2 2 3 2 1 1 1 3 2 3 1 1	706
1 2 3 2 2 1 2 2 1 3 1 2 2 3 2 3 1 2 3 1 1 2 3 1	707
2 1 3 2 1 3 2 1 3 1 1 2 1 2 3 1 1 1 2 2 1 3 1 3	708
2 2 2 1 1 2 3 1 3 1 1 3 1 3 2 2 1 3 1 3 2 1 2 1	709
1 1 1 3 2 2 2 1 3 2 1 3 1 3 2 3 2 1 2 3 2 1 1 1	710
1 2 1 2 1 2 3 1 2 1 3 2 1 3 1 3 2 1 3 1 2 2 1 3	711
2 3 1 3 1 1 3 2 2 1 1 2 2 3 2 1 2 1 3 1 2 2 3 1	712
2 1 2 1 3 1 3 1 2 3 2 2 1 2 1 2 3 1 1 3 2 2 3 2	713
1 1 1 2 2 2 3 2 2 1 1 3 2 2 3 2 2 3 2 2 3 2 2 3	714
2 2 3 2 2 3 1 1 3 1 2 3 1 1 1 3 2 1 3 1 1 2 2 1	715
1 1 3 1 3 1 2 1 1 3 2 1 3 2 3 2 2 2 1 2 3 2 2 2	716
1 1 2 1 1 3 1 1 3 1 1 3 2 3 1 1 1 3 1 2 2 3 1 2	717
2 1 1 3 2 2 1 1 1 3 2 2 3 1 2 3 1 2 2 3 1 2 1 3	718
1 2 1 2 1 1 3 1 2 1 1 3 1 3 2 3 2 1 1 3 2 3 1 2	719
3 2 2 1 1 1 2 3 2 2 3 2 2 3 2 2 2 1 1 3 2 3 1 2	720
3 1 3 2 2 1 1 3 2 2 1 2 2 1 3 2 2 1 1 3 1 1 3 2	721
2 1 2 2 1 3 1 3 2 2 2 3 1 3 1 1 2 1 1 3 2 1 3 2	722
2 1 1 2 3 2 2 3 2 2 1 2 3 2 3 2 2 1 3 1 2 3 2 2	723
3 1 1 1 3 2 2 3 1 2 1 3 1 1 2 3 2 1 1 2 3 2 2 2	724
2 3 1 2 1 3 1 2 3 1 1 2 2 3 1 2 2 3 1 2 2 1 3 2	725
1 2 3 1 2 1 3 1 3 2 1 1 1 3 1 1 2 1 1 3 2 2 3 2	726
1 3 2 1 1 3 2 3 2 2 1 3 1 2 1 3 2 1 2 2 3 1 1 2	727
1 2 3 2 1 3 1 2 2 1 1 1 3 2 1 3 2 3 2 1 2 3 2 2	728
2 2 1 2 2 3 1 2 1 1 2 3 1 3 1 3 1 3 2 2 1 1 1 3	729
1 2 2 2 3 2 2 1 2 3 1 2 1 1 1 2 3 2 3 2 1 3 2 3	730
2 2 3 1 1 3 1 1 1 2 1 1 3 1 3 2 1 1 2 1 1 3 1 3	731
2 3 2 3 2 1 1 2 1 1 3 2 1 3 2 1 1 3 1 2 2 1 3 1	732
1 2 3 1 1 1 3 2 2 1 3 1 3 2 2 2 1 2 3 1 2 1 1 3	733
1 2 2 1 3 2 2 1 1 3 1 3 1 3 2 2 2 3 2 1 3 1 2 2	734
2 3 2 1 3 2 1 2 2 3 2 1 2 3 1 2 2 1 1 1 3 2 3 2	735
1 3 2 2 3 1 2 1 1 1 3 1 1 3 1 1 3 1 3 2 1 2 1 2	736
3 2 1 1 2 3 1 3 1 2 1 1 1 3 1 3 1 3 1 2 1 1 2 2	737
2 3 2 3 2 2 3 1 1 3 1 2 1 1 1 3 2 2 2 1 2 3 1 2	738
1 1 3 1 1 3 1 3 2 1 3 2 2 1 3 1 1 2 2 3 1 2 2 1	739
3 1 1 2 3 1 1 3 1 2 3 1 1 3 2 2 2 3 2 2 1 1 2 1	740
1 1 1 2 2 3 2 2 3 1 3 1 2 1 1 3 1 2 1 3 2 3 1 2	741
2 3 1 2 2 3 2 2 2 1 1 2 3 1 2 3 2 3 2 3 1 2 2 1	742
1 2 3 1 1 3 2 1 2 2 3 2 2 3 1 3 2 3 1 2 2 2 1 1	743
3 2 3 2 1 1 1 2 3 2 2 2 3 1 3 1 2 3 2 1 2 1 2 2	744

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
1 1 2 2 3 1 2 3 1 3 2 2 2 1 1 1 3 1 3 2 2 3 1 2	745
2 2 2 3 2 3 2 1 1 2 1 2 3 1 2 2 3 1 3 1 3 2 2 2	746
3 2 1 3 2 1 3 1 2 3 1 2 2 1 1 3 1 1 3 1 2 1 1 1	747
2 2 2 1 1 2 3 2 3 1 1 1 2 2 2 3 2 2 3 2 3 1 3 2	748
3 2 1 1 1 3 1 1 2 2 1 3 1 2 1 1 1 3 1 3 2 3 1 2	749
1 1 2 1 3 2 2 1 1 3 2 2 2 1 1 3 1 3 2 2 3 2 3 2	750
3 2 3 2 3 1 2 3 2 2 2 1 2 1 3 1 2 2 2 3 2 2 1 2	751
3 2 1 2 1 3 2 3 2 3 1 2 2 1 3 1 2 2 2 3 2 1 1 1	752
3 2 2 3 2 1 1 3 1 1 1 3 1 2 1 2 3 2 1 1 3 1 1 1	753
1 2 1 2 2 1 3 1 2 2 3 2 1 1 1 3 1 3 1 3 2 3 1 1	754
3 1 3 2 3 1 2 1 2 2 3 1 1 1 2 2 1 3 1 2 2 3 2 1	755
2 1 1 3 1 1 3 2 2 1 1 1 3 1 1 3 1 3 1 3 2 2 2 1	756
3 1 2 3 2 2 1 3 1 2 1 1 1 3 2 2 2 1 1 3 2 1 3 2	757
3 2 3 1 2 2 3 2 1 2 3 1 3 1 1 1 2 3 2 2 1 1 1 2	758
2 3 1 2 2 1 2 2 3 2 1 1 3 1 1 1 3 1 2 2 3 1 3 1	759
1 1 3 1 1 2 2 3 2 3 2 1 1 3 2 2 2 1 2 3 1 3 2 1	760
2 2 3 2 1 2 2 2 1 3 1 1 3 1 2 2 2 3 2 1 3 1 2 3	761
2 1 2 1 2 3 2 2 2 3 2 3 2 1 1 3 1 1 3 1 1 1 2 3	762
3 1 2 1 1 2 3 2 3 2 3 1 1 2 2 2 3 2 3 1 1 2 1 1	763
2 2 1 3 1 1 1 2 3 2 3 1 3 1 2 2 2 1 1 3 1 3 2 2	764
1 3 2 3 2 1 3 1 1 2 2 2 3 2 1 2 2 2 2 1 3 2 2 3 2	765
2 1 3 2 2 1 1 3 1 2 1 3 1 3 2 1 1 1 2 3 1 2 1 3	766
3 1 1 3 2 3 1 2 1 2 2 3 2 1 1 1 2 2 3 1 2 1 1 3	767
3 2 1 1 2 2 3 2 3 2 2 1 3 1 2 2 2 1 1 3 1 1 3 2	768
2 3 1 2 1 2 2 2 3 2 3 1 1 2 2 3 1 2 1 3 2 1 2 3	769
1 1 3 2 1 1 1 3 1 3 1 2 1 2 1 3 2 2 1 1 3 2 2 3	770
1 2 2 1 3 2 2 1 1 3 2 2 1 2 2 2 3 2 3 1 3 2 3 1	771
1 3 1 2 3 1 1 3 2 1 3 2 2 2 1 2 3 1 1 2 2 1 3 1	772
2 3 1 3 2 2 1 3 2 2 1 1 3 2 3 1 2 1 3 2 2 1 1 1	773
2 2 1 2 2 3 2 1 3 1 2 2 2 1 3 1 3 1 1 3 1 2 3 1	774
2 1 2 2 2 3 2 3 2 2 2 3 2 2 3 1 2 2 1 3 1 2 1 3	775
3 2 1 2 1 1 2 3 2 3 2 3 2 3 1 1 1 3 2 2 1 2 1 1	776
2 1 2 1 2 3 2 2 3 1 3 2 1 2 1 1 1 3 1 3 1 3 1 1	777
2 2 1 3 2 2 1 3 2 2 2 1 1 1 3 1 2 2 3 2 3 1 3 2	778
2 2 2 1 3 1 1 2 1 1 3 2 3 1 2 3 2 3 1 2 3 1 1 1	779
1 3 1 3 2 1 1 2 3 2 3 2 1 1 1 2 1 3 2 2 1 3 1 2	780
2 3 2 3 1 2 1 1 1 3 1 3 1 1 1 2 2 1 3 2 2 3 2 2	781
3 1 1 2 2 2 1 3 2 3 1 1 2 3 2 2 2 3 1 3 1 2 2 1	782
2 3 2 3 1 2 3 2 3 2 1 1 3 2 1 2 1 2 3 1 1 1 2 2	783
2 2 3 2 3 1 1 2 3 1 2 2 1 1 2 3 1 1 2 1 3 1 1 3	784
1 1 2 3 2 2 3 2 2 2 1 3 1 2 2 3 1 3 1 1 1 3 2 2	785
1 3 1 2 2 3 2 3 2 2 1 3 2 1 2 2 1 3 2 1 2 1 1 3	786
2 2 3 1 2 3 2 1 2 2 1 3 1 1 1 3 2 2 2 1 2 3 2 3	787
2 1 2 3 1 2 2 3 2 3 2 3 2 2 1 3 1 3 1 1 2 2 2 1	788
2 1 3 2 3 2 1 3 1 2 1 2 2 2 3 1 2 1 3 2 2 1 2 3	789
1 3 2 2 2 1 1 2 3 2 3 2 2 2 1 3 2 2 3 2 2 1 2 3	790
2 3 2 3 2 1 1 1 3 2 1 3 1 1 1 3 2 1 1 1 3 1 2 2	791
3 2 2 1 2 3 1 2 1 2 1 3 2 3 1 3 2 2 3 2 2 1 2 2	792
2 2 2 3 1 2 2 3 1 1 2 3 2 2 1 1 2 1 3 2 3 2 3 2	793
1 3 1 3 2 1 2 2 1 3 2 1 3 2 2 1 2 2 3 2 1 1 3 2	794
2 1 1 3 2 1 3 1 1 1 3 1 1 3 1 1 3 1 2 1 2 2 2 3	795
1 3 1 1 1 3 1 3 1 1 2 2 1 2 3 2 1 1 2 3 1 1 1 3	796
2 2 1 3 1 2 2 2 3 2 2 1 3 2 3 2 3 1 2 2 2 1 1 3	797
3 1 2 3 1 2 2 1 1 3 1 2 1 2 1 3 1 3 1 2 1 3 2 2	798
1 2 1 2 2 2 3 1 3 2 3 1 2 2 1 1 3 1 3 2 1 1 2 3	799
2 3 2 1 2 2 3 2 3 1 3 2 2 1 1 3 2 1 2 1 1 3 2 2	800

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
1 1 2 2 2 1 3 2 1 3 1 1 1 3 2 3 2 2 3 2 3 2 2 2	801
3 2 2 1 3 1 1 3 1 2 2 1 1 3 2 2 3 1 1 2 1 1 2 3	802
2 1 1 1 3 2 1 2 3 2 3 1 3 1 2 3 1 2 2 2 1 2 3 2	803
2 3 1 1 1 2 3 1 2 2 1 1 1 3 1 2 3 1 1 3 1 2 3 1	804
2 2 1 2 2 1 3 1 2 3 2 2 3 1 3 2 3 2 2 2 3 2 2 2	805
2 1 3 2 3 2 2 2 1 1 1 3 1 3 2 1 3 2 1 2 1 2 3 1	806
1 3 2 2 1 2 1 1 3 2 1 1 1 2 3 1 2 3 2 2 3 1 2 3	807
2 2 1 1 3 1 3 1 3 1 1 1 2 1 1 3 2 3 2 1 2 2 3 2	808
3 1 2 1 2 2 3 1 1 1 2 3 2 3 2 1 1 1 2 3 1 2 3 1	809
1 2 3 1 2 3 1 1 2 2 1 1 3 1 1 1 3 1 1 1 3 1 3 1	810
3 1 1 2 1 3 2 2 2 3 1 2 2 2 3 2 3 2 2 2 3 2 2 1	811
1 3 2 2 3 2 2 2 1 3 1 2 2 2 3 1 2 2 2 3 2 1 1 3	812
3 2 1 2 3 1 3 1 2 2 2 3 1 2 1 2 1 1 3 1 2 2 1 3	813
2 2 2 1 2 1 3 2 3 1 3 2 1 2 1 3 2 3 1 2 3 1 2 2	814
2 1 2 1 2 3 2 3 1 1 3 1 2 1 2 1 1 3 2 3 2 2 2 3	815
1 2 2 3 1 2 1 3 1 2 3 1 2 1 3 2 1 1 2 2 3 1 3 2	816
2 3 1 2 1 3 1 2 3 2 3 1 1 3 1 1 2 2 2 3 1 2 2 2	817
3 1 1 3 1 2 1 2 2 3 1 1 1 3 1 1 2 2 2 3 1 2 3 2	818
3 1 2 3 2 2 2 1 3 2 3 2 1 3 1 2 1 2 1 3 1 2 2 2	819
3 1 1 2 1 2 2 3 1 3 2 2 1 2 1 1 3 2 1 3 2 1 3 2	820
1 3 2 3 1 3 2 1 1 3 2 1 1 2 1 3 1 1 1 3 1 2 2 2	821
3 2 1 3 1 1 2 1 1 3 2 1 1 2 2 2 3 2 3 1 3 1 2 1	822
3 1 3 2 2 1 2 2 2 3 1 3 1 2 2 2 1 3 1 2 3 2 2 2	823
3 1 1 1 2 3 1 2 3 1 2 2 3 1 1 2 2 2 1 3 1 3 1 2	824
1 1 1 2 1 3 2 3 2 3 1 3 1 1 2 1 3 2 2 1 1 3 2 1	825
1 2 3 2 3 2 2 1 1 3 2 2 3 2 1 3 1 1 3 1 1 2 1 1	826
1 2 1 1 2 3 1 3 2 2 1 1 2 1 3 2 3 2 1 1 3 1 1 3	827
1 2 1 1 3 1 3 1 2 3 2 2 2 1 1 3 2 2 1 3 1 1 1 3	828
2 3 2 2 1 3 2 3 2 2 1 3 1 1 1 2 1 2 3 1 1 1 3 1	829
2 2 2 1 3 1 1 3 1 2 2 3 2 2 1 3 1 2 1 1 3 2 2 3	830
3 2 3 2 1 1 2 3 2 1 2 1 1 3 1 2 1 3 2 2 1 1 3 2	831
2 1 2 2 1 3 1 3 1 3 1 1 1 2 2 3 2 1 3 1 3 1 2 2	832
2 1 3 2 3 1 3 1 2 1 1 1 3 2 1 1 1 3 2 2 2 1 2 3	833
2 2 3 2 3 1 1 1 3 2 2 1 1 3 2 1 1 3 2 2 1 3 2 2	834
1 1 1 3 2 3 2 1 1 3 2 2 3 1 1 3 1 1 2 1 2 2 3 1	835
3 1 1 2 1 3 1 3 2 3 2 2 1 2 2 2 3 1 1 1 2 1 3 1	836
2 1 2 1 1 3 1 3 1 3 1 3 1 2 1 1 3 2 1 1 2 1 1 3	837
2 3 1 3 2 3 1 1 1 2 2 3 1 2 1 3 1 3 2 1 1 1 2 2	838
3 1 2 3 1 1 2 1 1 3 2 2 2 1 1 3 2 3 1 3 1 1 1 2	839
3 2 3 2 3 1 2 1 2 3 2 2 2 1 2 2 3 1 2 2 1 1 3 2	840
2 1 1 1 3 2 3 1 3 2 3 2 1 1 1 2 3 1 2 1 1 2 3 1	841
3 2 1 3 1 3 2 2 2 3 1 2 2 2 3 1 1 1 3 1 1 2 1 2	842
3 1 1 2 1 2 2 3 2 2 1 2 3 2 2 2 3 2 2 1 2 3 1 3	843
3 2 3 2 1 1 2 1 1 3 1 2 3 2 1 2 2 3 2 2 3 2 2 2	844
2 1 1 1 2 2 3 1 2 2 3 2 3 1 3 2 2 3 1 1 3 1 1 2	845
2 3 1 3 1 2 1 3 2 2 1 2 1 3 2 2 1 1 3 2 2 2 1 3	846
1 3 2 2 2 3 2 2 1 1 3 1 2 2 1 2 3 2 1 3 1 1 1 3	847
3 1 1 2 3 2 3 2 1 3 1 1 2 1 1 3 1 3 1 2 2 1 1 1	848
3 2 1 2 2 1 2 3 1 1 1 3 1 1 3 2 2 3 2 2 3 2 2 2	849
3 2 3 2 2 1 2 1 3 1 1 3 2 2 1 1 1 2 3 2 2 1 1 3	850
2 2 1 1 3 1 3 2 1 3 2 3 1 1 2 1 2 3 1 2 1 3 2 1	851
1 1 2 3 2 2 1 2 1 1 3 1 2 3 1 3 1 3 2 2 2 1 3 2	852
1 2 1 2 1 1 3 1 2 2 2 3 1 2 3 2 1 3 2 3 2 1 3 2	853
2 1 2 3 2 2 2 3 2 2 3 2 2 3 2 2 1 1 3 2 2 2 3 1	854
3 1 2 1 3 2 2 2 1 3 2 1 2 1 3 1 1 3 1 2 1 1 1 3	855
3 2 2 3 1 1 2 1 2 1 3 1 3 1 2 1 3 2 1 1 1 2 1 3	856

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
1 3 1 3 1 1 3 1 2 2 2 1 3 2 1 1 3 1 1 2 3 1 2 1	857
2 3 1 1 2 3 1 3 1 1 1 3 1 2 1 2 2 3 1 3 2 1 2 2	858
2 3 1 1 3 1 2 2 1 2 1 3 2 1 3 2 2 3 2 1 2 1 3 1	859
3 1 2 2 1 3 2 1 3 2 1 2 2 3 1 1 3 1 2 2 1 2 3 2	860
2 3 1 1 1 2 3 2 3 2 1 2 2 2 3 2 1 2 3 2 2 2 1 3	861
1 2 2 1 1 1 3 2 2 3 1 2 1 2 3 1 1 1 3 1 1 3 2 3	862
1 1 2 3 2 1 3 1 3 1 2 2 3 2 1 3 2 3 1 1 2 1 2 2	863
2 2 1 2 2 2 3 2 2 3 1 3 2 3 2 1 1 1 2 3 2 3 1 2	864
1 2 3 2 1 1 2 2 3 2 3 1 1 2 1 1 2 3 2 1 2 3 2 3	865
3 1 2 2 2 3 2 1 2 1 3 1 3 1 2 2 1 3 2 1 1 3 2 1	866
1 1 2 1 2 2 3 2 2 3 2 2 2 1 3 1 3 1 1 1 3 1 1 3	867
1 2 3 1 2 3 1 2 3 2 1 2 2 2 3 2 1 1 1 3 1 3 2 1	868
1 1 2 3 2 1 2 2 2 3 2 3 2 3 1 2 2 3 2 3 2 1 1 1	869
1 3 2 3 2 2 1 2 3 1 1 3 1 1 2 1 3 2 1 1 3 1 1 2	870
3 2 2 1 2 3 2 1 3 1 3 1 2 3 1 1 1 3 1 1 1 2 2 1	871
3 2 2 2 3 2 1 2 2 1 3 1 2 1 1 1 2 3 1 3 2 2 3 2	872
2 3 1 2 2 2 1 2 3 1 3 1 2 2 1 1 3 1 3 1 1 1 3 1	873
2 2 2 3 2 3 2 3 2 2 1 2 2 3 2 1 1 2 2 3 1 3 1 2	874
3 1 2 3 2 3 2 3 1 2 1 2 3 1 2 2 1 1 1 3 1 1 1 2	875
1 3 1 2 2 1 2 1 3 1 2 2 2 3 2 1 3 1 3 1 1 1 3 2	876
3 1 1 3 1 3 2 1 2 3 2 1 1 2 1 3 2 1 2 2 3 2 1 2	877
2 2 2 3 2 1 1 2 3 2 2 3 2 2 3 1 3 2 2 2 1 1 3 1	878
1 3 2 1 1 1 2 1 3 2 1 3 2 1 2 3 1 1 2 1 1 3 1 3	879
3 1 1 2 3 2 2 3 1 1 2 2 3 1 1 1 2 1 2 3 1 3 2 2	880
1 3 2 1 3 2 2 1 1 2 2 3 1 2 1 3 2 1 1 3 2 2 2 3	881
1 3 2 3 2 1 1 1 3 1 1 1 2 3 1 1 2 3 1 1 2 1 1 3	882
2 3 2 2 1 3 1 2 1 2 2 2 3 2 3 1 1 1 2 3 2 3 1 1	883
2 3 2 1 2 3 2 2 3 1 3 2 2 2 3 1 1 2 2 3 2 2 1 2	884
2 3 1 3 2 3 1 1 2 2 1 3 2 2 1 2 3 2 2 3 2 2 1 2	885
3 1 1 3 1 1 1 3 1 1 1 2 3 1 3 1 1 1 3 1 2 2 1 2	886
2 2 1 1 3 2 1 1 3 2 2 3 2 3 2 2 3 1 2 1 2 2 1 3	887
1 2 3 1 2 3 2 3 2 2 2 3 1 2 2 2 3 1 1 2 2 3 1 1	888
1 1 3 2 1 1 3 2 3 1 1 1 2 2 3 2 2 3 2 2 2 3 1 1	889
1 2 3 1 1 3 2 3 2 1 1 1 3 2 2 2 3 1 1 1 3 1 1 1	890
1 3 1 3 1 3 2 1 1 3 1 2 1 1 2 2 3 2 1 2 1 3 2 1	891
2 2 2 1 2 3 1 3 1 2 1 3 1 2 3 1 1 1 2 1 1 3 2 3	892
1 3 1 1 1 2 2 1 3 2 1 3 2 1 1 2 3 1 2 2 2 3 2 3	893
3 1 2 2 2 3 1 3 1 2 2 3 1 1 2 3 1 3 1 1 2 1 2 1	894
3 1 2 2 1 3 1 1 1 3 1 2 3 1 1 2 1 1 1 3 1 2 3 1	895
2 1 3 1 2 1 3 1 1 1 3 2 1 2 1 2 3 2 2 3 2 1 3 2	896
3 1 1 3 1 2 1 3 2 1 1 1 3 2 1 1 1 3 2 1 1 3 2 2	897
1 1 1 2 3 2 3 2 3 2 2 2 1 3 2 1 3 2 2 3 2 1 1 1	898
2 2 3 2 2 3 1 1 3 2 1 1 3 1 3 1 2 3 1 1 2 1 1 1	899
2 1 2 2 2 3 1 3 1 3 1 1 1 3 1 1 1 3 1 3 2 2 2 1	900
2 1 2 2 2 1 3 2 3 1 2 3 1 1 2 2 2 3 2 3 1 2 3 2	901
2 2 1 2 1 3 2 3 1 2 3 1 2 3 1 2 1 1 3 2 2 3 1 2	902
2 1 1 1 3 1 2 1 1 2 2 3 2 1 3 1 1 1 3 2 1 3 2 3	903
3 2 2 2 1 3 2 1 2 2 3 1 2 1 2 2 3 2 3 2 3 2 1 1	904
3 2 3 2 2 3 2 3 1 1 2 1 1 3 1 2 2 3 1 1 1 2 1 2	905
1 1 1 3 1 1 1 3 2 1 2 1 1 1 3 2 3 1 3 1 2 1 3 1	906
2 1 2 2 2 3 2 1 1 3 1 1 3 2 3 2 1 3 1 2 1 2 2 3	907
2 1 3 1 1 3 1 2 3 1 1 1 2 2 3 2 3 1 2 2 2 3 2 2	908
1 2 1 1 2 1 3 2 1 1 3 2 3 1 1 2 3 1 2 3 1 3 1 2	909
1 1 2 3 2 3 1 1 2 1 1 3 1 2 1 1 1 3 2 3 2 3 2 1	910
1 2 2 3 1 1 3 1 2 1 1 1 3 1 2 3 2 2 3 2 2 2 1 3	911
2 3 1 1 1 2 1 3 1 1 3 2 3 1 3 1 2 2 1 2 1 3 2 1	912

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
1 3 2 2 1 2 2 3 2 3 1 1 1 3 1 3 2 2 2 1 2 3 1 2	913
1 1 1 2 1 3 2 1 3 2 3 1 2 1 3 1 3 1 1 3 1 2 2 2	914
1 3 2 3 2 1 2 3 1 1 3 2 3 2 1 1 2 1 1 3 1 2 2 1	915
2 3 1 2 2 1 1 3 1 2 2 3 2 3 2 1 3 2 3 2 2 1 2 1	916
1 3 2 2 2 1 2 3 1 2 1 2 2 2 3 2 1 3 1 2 3 1 3 2	917
2 1 2 3 2 3 2 1 2 3 1 1 3 1 2 2 1 2 1 3 2 2 1 3	918
3 1 1 1 2 2 3 2 2 3 2 1 2 1 3 1 3 2 3 1 2 1 2 1	919
2 1 3 1 1 1 2 1 3 2 2 2 1 1 3 2 1 2 1 3 2 3 2 3	920
2 3 1 2 2 2 1 3 1 2 3 2 2 2 1 2 2 3 2 3 1 3 1 1	921
1 1 3 2 2 3 1 2 1 2 2 2 3 2 2 3 2 2 1 3 1 2 3 1	922
2 3 1 2 3 2 3 1 2 1 1 2 3 1 3 1 1 2 1 1 1 3 1 1	923
1 1 1 3 2 2 2 1 3 2 2 2 3 2 1 2 2 1 3 2 1 3 1 3	924
1 3 2 2 2 3 1 2 3 2 3 1 2 1 3 2 1 1 1 2 1 3 1 1	925
1 1 3 2 3 2 2 1 2 2 3 1 1 2 3 2 3 1 2 3 2 2 1 2	926
1 1 1 2 2 3 1 1 3 2 3 2 3 1 2 1 1 2 3 2 2 2 3 2	927
3 2 2 2 1 3 2 3 1 2 2 1 1 1 3 1 2 1 3 1 2 2 1 3	928
1 2 1 1 3 2 3 2 1 2 1 1 3 1 3 1 1 3 2 3 2 2 1 1	929
1 2 3 1 1 2 2 2 3 2 2 2 3 2 3 1 2 3 1 1 3 2 2 1	930
1 1 1 3 1 1 2 2 3 1 3 1 1 1 2 3 1 1 1 3 2 2 1 3	931
1 3 2 3 2 1 1 3 1 3 2 1 2 1 1 1 3 2 1 2 2 2 3 1	932
3 1 1 2 2 1 1 3 1 2 2 3 2 2 1 2 1 2 3 2 3 1 3 2	933
2 1 2 3 1 1 1 3 2 3 2 2 3 2 2 2 1 1 3 2 1 1 3 1	934
2 1 1 1 3 2 1 1 1 2 3 2 2 1 2 3 2 3 1 3 1 3 1 1	935
1 1 1 3 1 2 1 2 2 3 1 2 2 3 1 3 1 2 1 3 1 3 2 2	936
1 1 3 2 3 1 2 1 2 3 1 1 2 1 2 3 2 3 1 3 1 1 1 2	937
1 1 1 2 1 3 1 3 2 2 2 3 2 2 1 1 2 3 2 1 1 3 2 3	938
3 1 2 2 2 1 3 1 2 3 1 3 2 2 1 1 3 1 1 2 2 2 1 3	939
2 2 3 2 1 1 1 2 3 1 3 2 3 2 3 1 1 2 1 2 2 3 2 1	940
1 3 2 1 3 2 3 2 1 2 2 2 3 1 3 1 2 1 1 2 1 3 1 1	941
2 3 1 3 2 2 1 1 1 3 1 3 2 2 3 2 2 3 1 2 1 2 2 2	942
1 1 1 3 1 3 2 3 2 1 2 2 1 3 1 1 1 2 1 3 2 2 2 3	943
3 2 2 2 1 3 2 2 1 2 2 2 3 1 2 3 1 3 1 2 1 1 2 3	944
1 1 3 2 3 2 1 1 1 2 3 1 1 2 1 1 1 3 1 3 2 2 3 2	945
1 1 2 1 1 1 3 2 3 1 3 2 1 3 1 1 3 2 3 2 1 1 2 2	946
2 1 2 2 3 1 3 2 2 2 3 2 3 2 1 1 1 3 1 1 3 1 2 1	947
2 2 2 1 2 1 3 2 2 3 2 2 3 2 3 2 2 3 1 1 1 3 2 2	948
1 2 3 1 1 1 2 1 2 3 1 2 2 3 2 3 2 2 2 3 2 2 3 2	949
1 1 1 3 1 3 1 2 3 2 1 1 1 3 2 3 1 3 2 2 1 2 2 1	950
2 2 3 1 1 3 1 1 1 3 2 2 1 3 1 2 3 1 2 3 1 1 2 2	951
1 2 3 2 2 1 2 2 2 3 2 2 2 1 3 2 2 2 3 2 3 2 3 1	952
1 1 1 2 1 2 3 1 1 2 2 2 3 1 1 3 1 3 1 1 3 2 3 1	953
3 1 2 2 1 3 1 2 1 2 1 3 1 1 2 1 2 2 3 1 1 3 1 3	954
2 2 1 3 1 1 2 1 1 3 1 3 1 1 1 2 3 2 1 2 3 2 3 2	955
2 2 2 1 2 3 1 1 1 3 1 3 1 1 3 2 3 2 1 2 2 1 2 3	956
3 2 1 1 3 2 1 2 2 1 1 3 2 3 2 3 1 2 2 2 1 3 2 1	957
1 2 1 1 1 3 1 3 1 1 3 2 1 1 1 3 1 3 2 1 1 1 3 2	958
1 2 2 3 2 2 1 1 2 2 3 1 1 3 2 3 2 1 2 3 1 1 1 3	959
2 1 2 1 2 1 3 2 2 3 1 3 2 2 3 1 3 2 1 1 3 1 2 2	960
2 1 3 1 2 3 1 3 1 2 1 2 1 2 3 1 1 1 3 1 2 1 3 2	961
1 2 1 1 3 1 1 3 1 2 3 1 2 2 2 3 2 3 2 1 1 1 2 3	962
2 2 1 3 2 1 1 2 1 1 3 1 1 1 3 1 2 3 1 1 3 1 3 1	963
3 1 2 2 2 3 2 3 1 3 2 1 1 1 3 2 1 1 1 2 1 3 1 1	964
1 1 1 2 1 3 1 2 3 2 1 3 1 1 2 2 2 3 2 3 2 3 2 2	965
3 1 1 1 2 2 1 3 2 3 2 2 2 3 2 3 2 3 2 1 2 2 1 2	966
1 2 2 2 3 1 3 2 1 2 3 1 2 1 3 1 1 3 1 2 2 3 2 2	967
1 2 1 3 1 3 2 2 3 1 1 3 2 1 2 3 2 1 1 1 3 2 2 2	968

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
2 1 1 2 2 2 3 2 3 1 1 2 3 2 2 3 2 2 1 2 2 3 2 3	969
2 2 1 3 2 2 2 1 2 3 1 3 1 3 2 3 1 3 1 2 2 2 1 1	970
3 2 2 3 2 2 1 3 1 3 2 3 2 2 2 1 2 3 1 1 1 2 2 2	971
2 2 2 1 2 2 3 2 3 1 2 3 2 3 1 1 1 2 1 1 3 1 3 1	972
3 2 1 1 3 2 1 1 2 1 2 3 1 2 1 3 2 3 1 2 2 1 1 3	973
2 3 1 3 1 2 3 1 2 3 2 1 2 1 2 3 2 1 3 2 1 1 2 1	974
1 1 2 2 3 1 3 1 1 1 3 1 3 1 2 1 1 1 2 3 1 2 1 3	975
2 2 2 3 1 1 3 2 3 1 2 3 2 2 1 3 1 1 2 3 2 2 2 1	976
1 3 2 2 3 2 2 3 2 3 2 1 1 2 2 3 2 2 1 3 2 1 1 1	977
1 2 1 3 2 3 1 3 1 1 3 2 3 1 2 1 1 3 1 2 1 2 2 2	978
3 2 3 2 3 1 2 1 1 3 2 1 1 2 2 3 1 3 2 2 1 1 1 2	979
2 1 3 2 2 1 2 2 3 2 2 2 3 2 3 1 1 2 2 2 3 1 3 1	980
1 2 1 3 2 2 3 1 1 2 1 3 2 1 1 2 2 2 3 1 1 3 1 3	981
1 2 3 2 2 2 3 2 3 2 2 2 3 1 1 2 1 3 1 3 1 1 2 1	982
2 3 1 2 1 1 1 3 1 2 1 2 3 1 3 1 3 1 2 2 3 2 1 1	983
2 1 1 1 3 1 2 3 1 3 1 2 3 2 2 3 2 2 1 1 1 3 2 2	984
1 1 3 2 3 1 1 1 2 2 2 3 2 1 1 3 1 1 2 2 1 3 2 3	985
3 1 1 1 2 3 1 3 1 3 2 2 1 2 2 3 1 2 1 3 2 2 2 1	986
2 2 2 3 2 1 1 1 2 3 1 3 1 2 1 2 1 3 2 3 2 2 1 3	987
3 2 2 1 1 2 2 3 2 3 1 2 1 2 2 2 3 1 2 2 1 3 2 3	988
1 3 1 3 2 3 2 2 3 1 2 1 1 1 3 1 2 3 2 2 2 1 2 1	989
1 1 2 2 3 2 3 1 3 1 1 1 2 2 3 1 2 1 1 3 1 1 3 1	990
2 2 1 1 1 3 1 3 1 1 2 2 3 1 3 1 1 3 1 3 1 1 1 2	991
2 2 3 2 2 1 3 1 1 3 1 1 2 2 3 1 1 2 3 2 1 2 3 2	992
1 3 2 2 1 1 3 1 2 1 2 3 2 3 2 3 1 2 3 2 2 2 1 1	993
2 3 1 3 2 2 1 2 3 2 2 3 2 1 1 2 1 3 1 1 1 2 2 3	994
2 2 1 3 1 2 1 1 3 2 2 2 1 3 1 3 1 2 2 3 1 3 1 1	995
1 2 3 1 3 2 1 1 2 1 1 3 1 3 2 1 2 2 2 3 1 1 3 2	996
2 3 2 2 2 1 1 3 2 3 2 1 1 2 3 1 2 2 2 3 2 2 1 3	997
2 2 3 1 1 3 1 1 3 1 2 2 3 2 2 1 2 2 3 2 2 3 1 1	998
2 1 2 1 3 1 1 1 3 1 2 2 1 1 1 3 1 3 2 3 1 1 2 3	999
2 1 1 1 2 2 3 2 2 1 3 1 1 1 2 2 2 3 1 3 2 3 2 3	1000
1 2 2 3 2 2 1 3 2 3 2 3 2 2 1 2 2 3 1 2 2 1 2 3	1001
3 1 3 1 1 2 2 1 2 3 2 3 2 3 1 1 2 1 2 1 3 1 1 1	1002
2 2 3 1 2 2 3 1 2 1 1 1 3 2 1 1 1 3 1 3 2 3 2 1	1003
3 2 3 2 3 2 1 1 1 2 2 3 1 1 2 1 2 3 2 2 1 1 2 3	1004
1 1 1 3 2 1 1 1 3 1 1 1 3 1 1 3 2 2 2 3 1 1 1 3	1005
2 2 2 1 3 2 2 3 1 1 3 1 1 2 1 3 1 1 1 3 1 1 1 3	1006
3 2 3 2 1 1 2 1 1 3 1 3 2 3 1 1 2 1 3 2 1 1 2 2	1007
2 1 2 2 3 1 1 1 2 1 1 3 1 3 1 3 1 2 2 2 3 2 3 1	1008
1 2 3 1 3 1 1 1 3 1 1 3 1 1 3 2 2 1 1 3 1 2 2 2	1009
1 1 3 1 3 2 3 1 3 2 1 2 1 2 2 3 2 2 1 1 1 3 1 1	1010
2 2 2 3 2 1 1 1 3 2 3 1 2 3 1 2 3 2 1 1 3 1 2 1	1011
3 1 2 3 2 2 1 2 3 2 3 1 2 3 1 1 1 2 1 2 3 2 1 2	1012
3 2 1 3 1 1 2 1 1 1 3 2 3 2 2 1 1 1 3 2 3 2 2 1	1013
1 1 1 3 1 3 2 1 2 3 2 3 2 3 2 1 2 3 1 2 1 2 2 2	1014
1 1 1 3 1 2 1 1 3 1 3 2 2 1 3 2 1 1 1 2 2 3 2 3	1015
1 1 3 1 1 2 2 1 3 1 3 1 1 2 1 1 3 2 3 2 3 1 2 1	1016
3 1 2 1 1 3 1 1 1 3 2 3 1 1 1 2 3 2 1 1 1 2 2 3	1017
3 1 2 3 1 1 1 3 1 2 3 2 2 2 1 1 1 3 2 2 2 3 2 2	1018
1 3 2 3 2 1 1 3 2 1 1 2 1 1 3 2 2 2 3 1 3 1 1 1	1019
3 2 2 3 1 3 1 1 2 2 1 3 1 1 2 2 2 3 1 2 1 1 1 3	1020
2 2 1 1 3 1 1 1 2 2 2 3 2 1 2 3 2 3 2 2 3 2 2 3	1021
1 3 1 1 3 1 2 2 2 1 3 1 2 3 1 1 1 2 3 1 3 2 2 2	1022
2 1 1 3 2 2 2 3 1 3 1 2 1 1 1 3 1 2 3 1 2 1 2 3	1023
2 3 1 3 1 2 1 3 2 2 2 3 2 1 1 2 1 2 3 2 2 2 3 2	1024

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
1 3 2 2 2 3 1 1 1 2 2 3 2 1 1 3 2 2 2 3 1 2 3 1	1025
2 1 3 1 1 2 2 3 1 2 2 1 1 2 3 1 2 3 1 3 2 1 3 2	1026
1 3 1 3 1 2 2 2 3 2 1 1 2 1 1 3 2 1 2 2 3 1 1 3	1027
1 2 1 1 2 3 1 2 3 2 1 1 2 3 2 1 1 3 2 1 3 2 3 2	1028
2 3 1 1 1 2 2 2 3 1 2 3 1 3 1 3 1 2 1 2 3 2 2 1	1029
2 3 2 3 2 1 1 1 3 2 1 2 1 3 2 2 2 1 2 3 2 2 1 3	1030
2 3 1 1 2 1 1 3 2 3 1 1 1 2 1 3 1 1 2 3 1 1 2 3	1031
1 1 1 3 1 1 1 3 1 2 2 3 2 1 1 2 1 1 3 2 1 3 1 3	1032
1 1 2 3 1 1 1 2 1 3 2 3 2 2 1 1 1 2 3 1 3 2 3 2	1033
3 2 1 3 1 2 1 1 1 3 1 2 3 2 3 1 1 2 2 1 2 3 1 2	1034
3 1 2 1 3 2 1 2 1 2 3 2 3 2 3 2 1 2 2 2 3 2 2 2	1035
1 2 3 2 2 2 3 2 1 3 1 1 1 2 3 2 2 2 3 1 1 3 1 2	1036
1 1 1 2 2 2 3 2 1 3 1 3 1 3 1 1 1 2 2 2 3 2 2 3	1037
2 1 3 1 1 2 1 1 3 1 2 2 1 3 2 1 1 3 2 3 2 1 3 1	1038
2 3 1 2 2 2 1 3 1 3 1 1 1 2 1 2 3 1 3 2 1 3 1 1	1039
1 1 2 1 3 1 3 2 1 2 3 2 2 3 2 2 2 1 2 3 1 3 1 1	1040
3 1 2 3 1 2 3 1 1 3 1 3 2 2 2 1 2 2 3 2 1 1 1 2	1041
1 1 3 2 1 1 1 3 1 1 3 1 1 3 1 1 1 2 3 2 3 2 2 1	1042
2 2 3 1 1 3 1 1 2 2 1 1 3 2 3 2 2 2 1 3 2 3 2 1	1043
1 3 1 1 1 3 1 1 2 3 2 2 3 1 2 2 2 1 2 3 1 2 3 2	1044
3 1 2 2 1 1 1 3 1 3 1 2 3 2 2 3 1 2 2 3 1 1 1 2	1045
1 1 2 3 1 2 1 1 2 2 3 2 2 3 1 3 1 3 1 3 2 1 1 2	1046
3 2 2 2 3 2 2 3 1 1 1 3 2 3 2 1 1 1 3 2 1 2 1 2	1047
2 3 1 3 2 2 1 2 1 2 3 1 3 1 1 1 3 2 3 2 1 1 2 2	1048
2 2 3 2 3 1 3 1 1 1 3 1 1 3 2 1 2 1 2 1 3 1 1 2	1049
3 2 1 1 3 2 2 2 1 3 1 3 2 2 1 2 1 3 1 3 2 2 2 1	1050
3 1 2 1 3 1 2 1 3 1 2 1 1 3 2 2 1 1 2 2 3 1 1 3	1051
1 3 1 3 1 2 3 1 2 2 3 2 2 2 1 2 3 2 1 2 2 1 2 3	1052
1 1 1 3 2 2 1 1 3 1 1 1 2 2 3 2 1 3 2 3 1 2 1 3	1053
2 2 2 3 1 2 1 2 2 3 2 2 2 3 2 3 1 3 2 3 2 1 2 1	1054
1 2 2 2 3 2 1 3 1 1 1 3 2 2 3 2 2 1 2 3 1 3 2 2	1055
3 1 2 2 2 3 1 3 2 1 1 3 2 2 2 1 2 1 3 1 2 3 1 1	1056
1 1 3 1 2 1 1 1 3 2 3 1 3 2 2 3 1 2 2 2 1 3 1 2	1057
3 1 2 1 2 2 3 2 1 1 3 1 2 1 2 3 2 2 3 2 1 1 1 3	1058
3 2 1 1 3 1 3 2 3 2 1 2 2 3 2 1 1 3 2 2 1 1 2 2	1059
3 2 3 2 3 1 2 2 1 3 2 1 1 2 3 1 1 3 2 1 2 2 2 1	1060
3 2 1 1 3 1 1 1 3 1 2 2 1 1 3 2 3 2 2 1 3 2 1 1	1061
1 3 2 1 3 1 1 1 3 2 2 3 1 1 1 2 2 3 1 2 2 1 2 3	1062
2 1 1 3 1 3 1 1 3 2 2 3 1 3 2 1 1 2 3 2 1 2 2 2	1063
3 2 2 1 1 3 1 1 1 2 1 3 2 1 3 1 2 1 1 3 2 3 1 1	1064
2 1 1 3 2 1 1 1 2 2 3 1 1 1 3 2 3 2 1 2 1 3 2 3	1065
1 1 3 1 2 3 2 1 2 3 2 2 2 1 2 2 3 2 2 3 2 3 2 1	1066
1 2 2 2 1 3 1 1 2 1 2 1 3 2 3 1 1 3 1 3 1 2 1 3	1067
3 2 2 1 2 3 1 1 1 3 1 3 2 1 2 3 2 3 2 2 1 1 1 2	1068
2 1 2 2 1 2 3 2 3 1 1 3 1 1 3 1 1 2 3 1 2 2 1 3	1069
2 1 1 2 1 1 3 2 2 3 1 1 3 1 3 1 1 2 2 3 2 2 3 2	1070
2 3 1 2 3 2 2 2 3 1 2 3 2 1 1 2 2 3 2 2 1 1 1 3	1071
3 2 3 1 1 1 3 1 2 2 2 3 1 3 2 2 2 3 2 1 2 1 1 2	1072
1 3 1 3 1 1 2 1 2 1 3 1 2 2 3 1 3 1 2 2 2 3 2 2	1073
2 2 2 3 1 3 1 2 3 2 3 1 2 3 1 2 1 1 1 3 2 2 1 1	1074
3 2 2 3 2 1 1 1 2 2 3 2 1 3 2 1 1 1 3 1 1 3 2 1	1075
3 2 3 2 2 1 2 3 1 2 3 2 2 3 2 2 2 3 2 1 2 2 1 2	1076
1 2 2 1 2 2 3 2 3 2 1 3 1 2 3 2 1 2 2 1 1 3 1 3	1077
3 2 2 1 3 1 1 1 3 1 2 2 2 1 3 1 1 3 2 2 1 3 2 2	1078
2 2 3 2 3 2 1 2 2 1 1 3 1 3 1 3 2 3 1 1 1 2 1 2	1079
3 2 2 2 1 1 3 1 2 1 3 1 1 1 3 1 3 2 3 1 2 2 2 1	1080

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
1 1 2 3 1 3 1 1 1 2 1 3 1 2 1 3 2 2 1 2 2 3 2 3	1081
2 3 1 1 2 2 3 1 1 2 1 1 3 1 1 2 2 2 3 2 2 3 2 3	1082
1 1 2 1 1 3 1 2 2 3 1 1 2 2 1 3 2 3 1 3 2 1 1 3	1083
1 1 2 3 2 2 2 3 1 3 1 3 1 2 2 2 1 3 2 1 1 1 3 1	1084
1 3 2 2 2 1 3 1 1 2 1 3 1 1 1 2 3 2 3 2 2 2 3 1	1085
2 1 2 1 1 3 2 1 1 3 2 3 2 2 1 1 3 1 2 2 2 3 1 3	1086
3 2 1 3 2 3 1 1 2 1 1 3 2 2 1 3 2 3 2 2 1 1 2 1	1087
1 1 3 2 3 2 3 2 2 1 1 1 3 2 1 1 1 2 3 2 1 3 1 2	1088
1 3 1 3 1 2 3 2 2 2 1 2 3 2 2 3 2 3 1 1 2 2 1 1	1089
1 3 2 2 3 1 1 2 1 2 2 3 1 2 3 1 2 1 1 3 1 1 3 1	1090
2 3 1 1 2 3 2 3 1 3 1 2 3 2 2 2 1 3 1 1 2 1 1 2	1091
1 1 2 1 1 2 3 1 2 3 2 1 1 3 2 2 2 3 1 3 2 2 2 3	1092
1 1 1 3 1 3 2 3 1 1 2 1 3 1 1 1 2 1 1 3 1 3 1 1	1093
1 1 2 1 1 1 3 2 2 1 2 2 3 1 3 1 3 1 3 2 2 2 1 3	1094
1 3 2 1 3 2 3 2 2 3 2 1 3 2 2 2 1 3 2 1 2 1 2 1	1095
3 2 1 1 3 1 1 2 3 2 1 2 2 1 3 1 2 1 2 2 2 3 2 3	1096
3 1 2 1 1 1 2 3 2 2 2 3 1 2 1 1 1 3 2 1 3 2 2 3	1097
1 2 1 3 2 1 2 3 2 1 2 3 2 3 2 3 1 1 3 1 2 2 2 1	1098
1 2 3 1 1 2 3 2 1 3 1 3 2 3 1 2 2 1 3 2 2 2 1 1	1099
3 2 1 3 2 1 2 2 2 1 3 2 3 1 2 3 2 1 1 3 1 1 2 1	1100
1 3 1 1 2 2 3 2 1 2 2 3 1 1 3 1 1 3 1 1 2 1 2 3	1101
2 2 2 1 2 1 3 1 1 2 2 3 1 3 1 3 1 1 3 2 2 1 1 3	1102
1 1 1 3 2 1 3 2 1 3 1 3 1 2 2 2 3 1 3 1 1 2 2 1	1103
2 2 2 1 1 1 3 1 1 1 3 2 1 2 2 3 2 1 1 3 1 3 2 3	1104
1 1 1 2 2 3 1 3 1 1 1 3 2 3 1 1 2 3 1 1 3 2 2 2	1105
1 1 3 1 1 1 2 1 1 3 2 1 2 3 1 2 1 3 2 1 3 2 1 3	1106
1 2 2 2 3 1 1 2 2 3 2 1 2 2 3 2 1 3 2 2 2 3 2 3	1107
1 1 3 1 3 1 1 2 1 1 2 3 2 1 3 1 3 1 2 1 2 1 1 3	1108
2 3 2 3 2 1 1 2 1 3 2 2 3 2 2 1 1 2 3 1 3 2 1 1	1109
2 1 2 1 3 2 2 3 2 1 3 2 2 2 1 3 1 2 3 1 1 2 3 2	1110
1 2 2 3 2 3 2 2 1 3 1 1 2 3 1 2 3 2 2 1 1 2 1 3	1111
3 2 2 2 3 2 1 2 1 3 2 1 2 2 2 3 1 2 2 3 1 2 3 2	1112
1 3 1 3 2 1 1 1 3 2 1 2 3 1 3 2 2 1 2 3 1 1 2 1	1113
3 1 1 1 3 2 2 2 1 1 3 2 3 1 2 3 2 1 2 1 2 2 3 2	1114
2 2 1 1 1 2 3 1 2 1 1 1 3 1 3 2 1 3 2 3 1 1 3 2	1115
2 2 1 1 1 2 3 2 3 2 3 1 3 1 1 3 1 2 3 1 1 2 1 1	1116
1 2 2 2 3 2 1 2 1 1 1 3 2 3 1 1 3 1 1 3 1 3 1 1	1117
2 3 1 2 2 1 3 2 1 2 2 2 3 2 3 1 1 3 1 3 1 2 2 2	1118
2 2 2 3 1 1 2 3 1 1 1 2 2 3 1 2 3 1 2 1 3 1 2 3	1119
1 3 1 3 2 1 1 3 1 2 2 1 1 3 1 1 2 1 1 3 1 1 1 3	1120
1 2 2 3 1 1 2 2 3 1 3 1 1 3 2 3 1 1 3 2 1 1 1 2	1121
2 2 2 1 3 1 3 1 1 3 2 1 2 2 3 2 2 2 3 1 1 1 3 1	1122
2 1 1 1 3 2 3 1 1 1 3 1 2 2 2 3 1 1 1 2 3 1 2 3	1123
3 1 1 1 3 2 2 1 3 1 3 1 1 1 2 3 2 1 3 1 1 1 2 2	1124
3 2 3 1 1 2 1 1 2 3 1 1 3 1 1 3 2 2 1 2 3 2 2 1	1125
2 2 3 2 3 1 1 2 1 1 1 3 2 1 3 1 2 3 2 3 2 2 1 2	1126
2 2 1 2 1 2 3 1 2 1 2 3 1 3 2 2 2 3 2 3 2 2 3 1	1127
2 2 3 1 2 2 2 3 2 3 2 3 1 3 2 1 2 2 1 3 2 2 1 2	1128
1 1 1 3 2 3 1 2 2 1 1 3 2 2 1 3 2 2 2 3 1 3 1 2	1129
2 2 3 2 1 2 2 2 3 2 1 2 1 1 2 3 2 2 3 1 1 3 1 3	1130
3 2 2 2 3 1 1 1 2 2 1 3 2 3 2 3 1 3 1 1 1 2 1 2	1131
1 1 2 3 2 2 3 1 3 1 2 2 3 1 2 1 1 2 3 2 2 3 1 1	1132
2 1 3 2 1 3 2 1 3 2 1 2 2 3 2 2 3 2 1 1 2 1 1 3	1133
3 2 2 3 2 1 1 2 2 2 3 1 3 2 3 2 2 1 3 2 2 1 2 2	1134
2 3 1 1 2 1 2 3 1 2 1 3 2 2 1 3 2 1 1 2 2 3 2 3	1135
2 3 1 2 1 3 2 1 2 3 2 2 2 3 2 3 1 2 2 1 1 1 3 1	1136

Table IA: Numeric sequences corresponding to nucleotide base patterns of a set of oligonucleotides

Numeric Pattern	Sequence Identifier
3 1 2 3 2 1 2 1 1 1 3 1 3 2 1 2 3 2 2 1 2 1 1 3	1137
1 3 2 3 1 3 1 2 2 2 1 3 1 1 3 1 2 3 2 2 1 2 2 1	1138
1 2 3 1 3 1 1 2 2 2 3 2 2 1 1 1 3 1 3 1 1 1 3 2	1139
1 1 1 3 1 1 2 2 1 3 2 1 2 3 1 2 1 3 1 2 3 1 3 1	1140
2 1 3 1 3 2 2 3 2 1 2 1 3 2 2 2 1 2 1 3 2 2 3 1	1141
3 2 1 3 1 1 2 3 1 2 2 3 2 2 2 1 3 1 1 3 1 2 2 2	1142
3 2 2 2 1 2 3 2 2 2 3 1 3 1 1 3 1 3 2 2 1 2 2 2	1143
2 1 3 1 1 3 2 2 2 3 1 1 1 3 2 2 1 2 2 3 1 2 2 3	1144
3 1 2 3 1 1 3 1 3 2 1 2 2 2 3 2 2 1 2 1 2 3 2 1	1145
3 1 2 3 1 1 2 1 2 1 3 2 1 1 3 2 1 2 2 3 1 3 2 1	1146
2 1 3 2 3 1 2 3 1 1 1 2 2 2 3 1 3 1 2 1 3 1 2 1	1147
3 1 1 1 3 1 1 1 2 2 3 1 1 3 1 3 2 2 2 3 1 2 1 2	1148
1 2 2 2 3 1 3 2 1 2 2 2 3 2 3 2 1 2 2 3 1 1 2 3	1149
1 2 3 1 3 2 2 3 1 1 1 2 2 2 3 1 1 3 2 1 2 2 3 2	1150
2 2 1 1 2 1 3 2 3 1 3 1 3 1 3 2 1 2 1 2 3 2 1 1	1151
1 2 2 1 1 3 1 3 1 3 2 3 1 3 2 1 1 1 2 3 2 1 1 1	1152
1 1 3 1 1 2 1 3 1 2 3 1 3 1 2 2 1 3 1 1 1 2 1 3	1153
1 3 2 2 2 1 1 1 3 1 3 2 2 1 3 1 1 2 2 3 1 1 1 3	1154
3 2 1 1 3 1 2 2 2 3 2 2 3 1 1 2 1 1 1 3 1 1 3 1	1155
1 3 1 3 1 1 1 3 1 1 3 2 2 1 1 1 3 2 3 1 2 1 2 2	1156
2 1 1 2 1 3 1 3 1 1 3 1 3 1 2 3 2 1 2 3 1 1 2 1	1157
2 2 1 2 2 1 3 2 3 1 2 1 1 3 2 3 1 1 3 2 2 2 1 3	1158
1 2 1 1 2 3 2 1 1 1 3 1 2 3 1 3 2 2 2 1 2 3 1 3	1159
2 2 3 1 2 2 2 3 1 3 1 3 2 2 3 1 2 1 1 3 1 2 2 2	1160
1 2 3 1 2 2 1 2 2 3 2 3 2 3 2 1 3 1 1 2 2 1 3 1	1161
2 1 2 1 1 1 3 1 2 1 2 1 3 2 1 3 1 2 3 1 2 3 2 3	1162
2 2 2 1 3 2 2 3 1 3 1 2 3 1 1 3 2 2 1 2 2 1 3 1	1163
1 2 2 3 1 1 2 2 3 1 2 1 2 1 3 2 3 2 1 1 1 3 2 3	1164
3 1 1 3 1 1 1 3 1 2 2 1 2 2 3 2 1 2 2 3 1 3 2 2	1165
1 2 2 3 1 3 2 3 2 1 3 2 3 1 2 2 2 1 3 1 1 1 2 1	1166
1 1 2 1 1 1 3 2 3 2 2 2 1 1 3 1 3 2 1 3 1 3 2 1	1167
3 2 1 3 1 3 1 2 1 1 2 2 3 1 2 3 2 3 2 1 1 2 2 2	1168

In Table IA, each of the numerals 1 to 3 (numeric identifiers) represents a nucleotide base and the pattern of numerals 1 to 3 of the sequences in the above list corresponds to the pattern of nucleotide bases present in the oligonucleotides of Table I, which oligonucleotides have been found to be non-cross-hybridizing, as described further in the detailed examples. Each nucleotide base is selected from the group of nucleotide bases consisting of A, C, G, and T/U. A particularly preferred embodiment of the invention, in which a specific base is assigned to each numeric identifier is shown in Table I, below.

In one broad aspect, the invention is a composition comprising molecules for use as tags or tag complements wherein each molecule comprises an oligonucleotide selected from a set of oligonucleotides based on a group of sequences as specified by numeric identifiers set out on Table IA wherein each of 1 to 3 is a nucleotide base

selected to be different from the others of 1 to 3 with the *proviso* that up to three nucleotide bases of each sequence can be substituted with any nucleotide base provided that:

for any pair of sequences of the set:

M1 ≤ 16, M2 ≤ 13, M3 ≤ 20, M4 ≤ 16, and M5 ≤ 19, where:

M1 is the maximum number of matches for any alignment in which there are no internal indels;

M2 is the maximum length of a block of matches for any alignment;

M3 is the maximum number of matches for any alignment having a maximum score;

M4 is the maximum sum of the lengths of the longest two blocks of matches for any alignment of maximum score; and

M5 is the maximum sum of the lengths of all the blocks of matches having a length of at least 3, for any alignment of maximum score;

wherein:

the score of an alignment is determined according to the equation (A x *m*) - (B x *mm*) - (C x (*og* + *eg*)) - (D x *eg*)), wherein:

for each of (i) to (iv):

(i) *m* = 6, *mm* = 6, *og* = 0 and *eg* = 6,

(ii) *m* = 6, *mm* = 6, *og* = 5 and *eg* = 1,

(iii) *m* = 6, *mm* = 2, *og* = 5 and *eg* = 1, and

(iv) *m* = 6, *mm* = 6, *og* = 6 and *eg* = 0,

A is the total number of matched pairs of bases in the alignment;

B is the total number of internal mismatched pairs in the alignment;

C is the total number of internal gaps in the alignment; and

D is the total number of internal indels in the alignment minus the total number of internal gaps in the alignment; and

wherein the maximum score is determined separately for each of (i), (ii), (iii) and (iv).

An explanation of the meaning of the parameters set out above is given in the section describing detailed embodiments.

In another broad aspect, the invention is a composition containing molecules for use as tags or tag complements wherein each molecule comprises an oligonucleotide selected from a set of oligonucleotides based on a group of sequences as set out in Table IA wherein each of 1 to 3 is a nucleotide base selected to be different from the others of 1 to 3 with the *proviso* that up to three nucleotide bases of each sequence can be substituted with any nucleotide base provided that:

for any pair of sequences of the set:

M1 ≤ 19, M2 ≤ 17, M3 ≤ 21, M4 ≤ 18, and M5 ≤ 20, where:

M1 is the maximum number of matches for any alignment in which there are no internal indels;

M2 is the maximum length of a block of matches for any alignment;

M3 is the maximum number of matches for any alignment having a maximum score;

M4 is the maximum sum of the lengths of the longest two blocks of matches for any alignment of maximum score; and

M5 is the maximum sum of the lengths of all the blocks of matches having a length of at least 3, for any alignment of maximum score;

wherein

the score of an alignment is determined according to the equation (A x *m*) - (B x *mm*) - (C x (*og* + *eg*)) - (D x *eg*)), wherein:

for each of (i) to (iv):

(i) *m* = 6, *mm* = 6, *og* = 0 and *eg* = 6,

(ii) *m* = 6, *mm* = 6, *og* = 5 and *eg* = 1,

(iii) *m* = 6, *mm* = 2, *og* = 5 and *eg* = 1, and

(iv) *m* = 6, *mm* = 6, *og* = 6 and *eg* = 0,

A is the total number of matched pairs of bases in the alignment;

B is the total number of internal mismatched pairs in the alignment;

C is the total number of internal gaps in the alignment; and

D is the total number of internal indels in the alignment minus the total number of internal gaps in the alignment; and

wherein the maximum score is determined separately for each of (i), (ii), (iii) and (iv).

In another broad aspect, the invention is a composition comprising molecules for use as tags or tag complements wherein each molecule comprises an oligonucleotide selected from a set of oligonucleotides based on a group of sequences set out in Table IA wherein each of 1 to 3 is a nucleotide base selected to be different from the others of 1 to 3 with the *proviso* that up to three nucleotide bases of each sequence can be substituted with any nucleotide base provided that:

for any pair of sequences of the set:

M1 ≤ 19, M2 ≤ 17, M3 ≤ 21, M4 ≤ 18, and M5 ≤ 20, where:

M1 is the maximum number of matches for any alignment in which there are no internal indels;

M2 is the maximum length of a block of matches for any alignment;

M3 is the maximum number of matches for any alignment having a maximum

score;

M4 is the maximum sum of the lengths of the longest two blocks of matches for any alignment of maximum score; and

M5 is the maximum sum of the lengths of all the blocks of matches having a length of at least 3, for any alignment of maximum score, wherein:

the score of an alignment is determined according to the equation 3A - B - 3C - D, wherein:

A is the total number of matched pairs of bases in the alignment;

B is the total number of internal mismatched pairs in the alignment;

C is the total number of internal gaps in the alignment; and

D is the total number of internal indels in the alignment minus the total number of internal gaps in the alignment.

In preferred aspects, the invention provides a composition in which, for the group of 24mer sequences in which 1 = A, 2 = T and 3 = G, under a defined set of conditions in which the maximum degree of hybridization between a sequence and any complement of a different sequence of the group of 24mer sequences does not exceed 30% of the degree of hybridization between said sequence and its complement, for all said oligonucleotides of the composition, the maximum degree of hybridization between an oligonucleotide and a complement of any other oligonucleotide of the composition does not exceed 50% of the degree of hybridization of the oligonucleotide and its complement.

More preferably, the maximum degree of hybridization between a sequence and any complement of a different sequence does not exceed 30% of the degree of hybridization between said sequence and its complement, the degree of hybridization between each sequence and its complement varies by a factor of between 1 and up to 10, more preferably between 1 and up to 9, more preferably between 1 and up to 8, more preferably between 1 and up to 7, more preferably between 1 and up to 6, and more preferably between 1 and up to 5.

It is also preferred that the maximum degree of hybridization between a sequence and any complement of a different sequence does not exceed 25%, more preferably does not exceed 20%, more preferably does not exceed 15%, more preferably does not exceed 10%, more preferably does not exceed 5%.

Even more preferably, the above-referenced defined set of conditions results in a level of hybridization that is the same as the

level of hybridization obtained when hybridization conditions include 0.2 M NaCl, 0.1 M Tris, 0.08% Triton X-100, pH 8.0 at 37°C.

In the composition, the defined set of conditions can include the group of 24mer sequences being covalently linked to beads.

In a particular preferred aspect, for the group of 24mers the maximum degree of hybridization between a sequence and any complement of a different sequence does not exceed 15% of the degree of hybridization between said sequence and its complement and the degree of hybridization between each sequence and its complement varies by a factor of between 1 and up to 9, and for all oligonucleotides of the set, the maximum degree of hybridization between an oligonucleotide and a complement of any other oligonucleotide of the set does not exceed 20% of the degree of hybridization of the oligonucleotide and its complement.

It is possible that each 1 is one of A, T/U, G and C; each 2 is one of A, T/U, G and C; and each 3 is one of A, T/U, G and C; and each of 1, 2 and 3 is selected so as to be different from all of the others of 1, 2 and 3. More preferably, 1 is A or T/U, 2 is A or T/U and 3 is G or C. Even more preferably, 1 is A, 2 is T/U, and 3 is G.

In certain preferred composition, each of the oligonucleotides is from twenty-two to twenty-six bases in length, or from twenty-three to twenty-five, and preferably, each oligonucleotide is of the same length as every other said oligonucleotide.

In a particularly preferred embodiment, each oligonucleotide is twenty-four bases in length.

It is preferred that no oligonucleotide contains more than four contiguous bases that are identical to each other.

It is also preferred that the number of G's in each oligonucleotide does not exceed L/4 where L is the number of bases in said sequence.

For reasons described below, the number of G's in each said oligonucleotide is preferred not to vary from the average number of G's in all of the oligonucleotides by more than one. Even more preferably, the number of G's in each said oligonucleotide is the same as every other said oligonucleotide. In the embodiment disclosed below in which oligonucleotides were tested, the sequence of each was twenty-four bases in length and each oligonucleotide contained 6 G's.

It is also preferred that, for each nucleotide, there is at most six bases other than G between every pair of neighboring pairs of G's.

Also, it is preferred that, at the 5'-end of each oligonucleotide at least one of the first, second, third, fourth, fifth, sixth and

seventh bases of the sequence of the oligonculeotide is a G. Similarly, it is preferred, at the 3'-end of each oligonucleotide that at least one of the first, second, third, fourth, fifth, sixth and seventh bases of the sequence of the oligonucleotide is a G.

It is possible to have sequence compositions that include one hundred and sixty said molecules, or that include one hundred and seventy said molecules, or that include one hundred and eighty said molecules, or that include one hundred and ninety said molecules, or that include two hundred said molecules, or that include two hundred and twenty said molecules, or that include two hundred and forty said molecules, or that include two hundred and sixty said molecules, or that include two hundred and eighty said molecules, or that include three hundred said molecules, or that include four hundred said molecules, or that include five hundred said molecules, or that include six hundred said molecules, or that include seven hundred said molecules, or that include eight hundred said molecules, or that include nine hundred said molecules, or that include one thousand said molecules.

It is possible, in certain applications, for each molecule to be linked to a solid phase support so as to be distinguishable from a mixture containing other of the molecules by hybridization to its complement. Such a molecule can be linked to a defined location on a solid phase support such that the defined location for each molecule is different than the defined location for different others of the molecules.

In certain embodiments, each solid phase support is a microparticle and each said molecule is covalently linked to a different microparticle than each other different said molecule.

In another broad aspect, the invention is a composition comprising a set of 150 molecules for use as tags or tag complements wherein each molecule includes an oligonucleotide having a sequence of at least sixteen nucleotide bases wherein for any pair of sequences of the set:

$M1 \leq 19/24 \times L1$, $M2 \leq 17/24 \times L1$, $M3 \leq 21/24 \times L1$, $M4 \leq 18/24 \times L1$, $M5 \leq 20/24 \times L1$, where L1 is the length of the shortest sequence of the pair, where:

M1 is the maximum number of matches for any alignment of the pair of sequences in which there are no internal indels;

M2 is the maximum length of a block of matches for any alignment of the pair of sequences;

M3 is the maximum number of matches for any alignment of the pair of sequences having a maximum score;

M4 is the maximum sum of the lengths of the longest two blocks of matches for any alignment of the pair of sequences of maximum score; and

M5 is the maximum sum of the lengths of all the blocks of matches having a length of at least 3, for any alignment of the pair of sequences of maximum score, wherein:

the score of an alignment is determined according to the equation (A x *m*) - (B x *mm*) - (C x (*og* + *eg*)) - (D x *eg*)), wherein:

for each of (i) to (iv):

(i) *m* = 6, *mm* = 6, *og* = 0 and *eg* = 6,

(ii) *m* = 6, *mm* = 6, *og* = 5 and *eg* = 1,

(iii) *m* = 6, *mm* = 2, *og* = 5 and *eg* = 1, and

(iv) *m* = 6, *mm* = 6, *og* = 6 and *eg* = 0,

A is the total number of matched pairs of bases in the alignment;

B is the total number of internal mismatched pairs in the alignment;

C is the total number of internal gaps in the alignment; and

D is the total number of internal indels in the alignment minus the total number of internal gaps in the alignment; and

wherein the maximum score is determined separately for each of (i), (ii), (iii) and (iv).

In yet another broad aspect, the invention is a composition that includes a set of 150 molecules for use as tags or tag complements wherein each molecule has an oligonucleotide having a sequence of at least sixteen nucleotide bases wherein for any pair of sequences of the set:

M1 ≤ 19, M2 ≤ 17, M3 ≤ 21, M4 ≤ 18, and M5 ≤ 20, where:

M1 is the maximum number of matches for any alignment of the pair of sequences in which there are no internal indels;

M2 is the maximum length of a block of matches for any alignment of the pair of sequences;

M3 is the maximum number of matches for any alignment of the pair of sequences having a maximum score;

M4 is the maximum sum of the lengths of the longest two blocks of matches for any alignment of the pair of sequences of maximum score; and

M5 is the maximum sum of the lengths of all the blocks of matches having a length of at least 3, for any alignment of the pair of sequences of maximum score, wherein:

the score of a said alignment is determined according to the equation

3A - B - 3C - D, wherein:

A is the total number of matched pairs of bases in the alignment;

B is the total number of internal mismatched pairs in the alignment;

C is the total number of internal gaps in the alignment; and

D is the total number of internal indels in the alignment minus the total number of internal gaps in the alignment.

In certain embodiments of the invention, each sequence of a composition has up to fifty bases. More preferably, however, each sequence is between sixteen and forty bases in length, or between sixteen and thirty-five bases in length, or between eighteen and thirty bases in length, or between twenty and twenty-eight bases in length, or between twenty-one and twenty-seven bases in length, or between twenty-two and twenty-six bases in length.

Often, each sequence is of the same length as every other said sequence. In particular embodiments disclosed herein, each sequence is twenty-four bases in length.

Again, it can be preferred that no sequence contains more than four contiguous bases that are identical to each other, etc., as described above.

In certain preferred embodiments, the composition is such that, under a defined set of conditions, the maximum degree of hybridization between an oligonucleotide and any complement of a different oligonucleotide of the composition does not exceed about 30% of the degree of hybridization between said oligonucleotide and its complement, more preferably 20%, more preferably 15%, more preferably 10%, more preferably 6%.

Preferably, the set of conditions results in a level of hybridization that is the same as the level of hybridization obtained when hybridization conditions include 0.2 M NaCl, 0.1 M Tris, 0.08% Triton X-100, pH 8.0 at 37°C, and the oligonucleotides are covalently linked to microparticles. Of course it is possible that these specific conditions be used for determining the level of hybridization.

It is also preferred that under such a defined set of conditions, the degree of hybridization between each oligonucleotide and its complement varies by a factor of between 1 and up to 8, more preferably up to 7, more preferably up to 6, more preferably up to 5. In a particular disclosed embodiment, the observed variance in the degree of hybridization was a factor of only 5.3, i.e., the degree of

hybridization between each oligonucleotide and its complement varied by a factor of between 1 and 5.6.

In certain preferred embodiments, under the defined set of conditions, the maximum degree of hybridization between a said oligonucleotide and any complement of a different oligonucleotide of the composition does not exceed about 15%, more preferably 10%, more preferably 6%.

In one preferred embodiment, the set of conditions results in a level of hybridization that is the same as the level of hybridization obtained when hybridization conditions include 0.2 M NaCl, 0.1 M Tris, 0.08% Triton X-100, pH 8.0 at 37°C, and the oligonucleotides are covalently linked to microparticles.

Also, under the defined set of conditions, it is preferred that the degree of hybridization between each oligonucleotide and its complement varies by a factor of between 1 and up to 8, more preferably up to 7, more preferably up to 6, more preferably up to 5.

Any composition of the invention can include one hundred and sixty of the oligonucleotide molecules, or one hundred and seventy of the oligonucleotide molecules, or one hundred and eighty of the oligonucleotide molecules, or one hundred and ninety of the oligonucleotide molecules, or two hundred of the oligonucleotide molecules, or two hundred and twenty of the oligonucleotide molecules, or two hundred and forty of the oligonucleotide molecules, or two hundred and sixty of the oligonucleotide molecules, or two hundred and eighty of the oligonucleotide molecules, or three hundred of the oligonucleotide molecules, or four hundred of the oligonucleotide molecules, or five hundred of the oligonucleotide molecules, or six hundred of the oligonucleotide molecules, or seven hundred of the oligonucleotide molecules, or eight hundred of the oligonucleotide molecules, or nine hundred of the oligonucleotide molecules, or one thousand or more of the oligonucleotide molecules.

A composition of the invention can be a family of tags, or it can be a family of tag complements.

An oligonucleotide molecule belonging to a family of molecules of the invention can have incorporated thereinto one more analogues of nucleotide bases, preference being given those that undergo normal Watson-Crick base pairing.

Another aspect of the present invention there is provided, a kit for sorting and identifying polynucleotides, the kit comprising one or more solid phase supports each having one or more spatially discrete regions, each such

region having a uniform population of substantially identical tag complements covalently attached and the tag complements each being selected from the set of oligonucleotides as defined above.

The one or more solid phase supports can be a planar substrate in which the one or more spatially discrete regions is a plurality of spatially addressable regions.

The tag complements can also be coupled to microparticles. Microparticles preferably each have a diameter in the range of from 5 to 40 μm.

Such a kit preferably includes microparticles that are spectrophotometrically unique, and therefore distinguisable from each other according to conventional laboratory techniques. Of course for such kits to work, each type of microparticle would generally have only one tag complement associated with it, and usually there would be a different oligonucleotide tag complement associated with (attached to) each type of microparticle.

The invention includes methods of using families of oligonucleotides of the invention.

A further aspect of the present invention, there is provided, a method of analyzing a biological sample comprising a biological sequence for the presence of a mutation or polymorphism at a locus of the nucleic acid, the method comprising:

(A) amplifying the nucleic acid molecule in the presence of a first primer having a 5′-sequence having the sequence of a tag complementary to the sequence of a tag complement belonging to a family of tag complements as defined above to form an amplified molecule with a 5′-end with a sequence complementary to the sequence of the tag;

(B) extending the amplified molecule in the presence of a polymerase and a second primer having 5′-end complementary the 3′-end of the amplified sequence, with the 3′-end of the second primer extending to immediately adjacent said locus, in the presence of a plurality of nucleoside triphosphate derivatives each of which is: (i) capable of incorporation during transciption by the polymerase onto the 3′-end of a growing nucleotide strand; (ii) causes termination of polymerization; and (iii) capable of differential detection, one from the other, wherein there is a said derivative complementary to each possible nucleotide present at said locus of the amplified sequence;

(C) specifically hybridizing the second primer to a tag complement having the tag complement sequence of (A); and

(D) detecting the nucleotide derivative incorporated into the second primer in (B) so as to identify the base located at the locus of the nucleic

acid.

Another aspect of the present invention, there is provided, a method of analyzing a biological sample comprising a plurality of nucleic acid molecules is analyzed for the presence of a mutation or polymorphism at a locus of each nucleic acid molecule, for each nucleic acid molecule, the method comprising:

(A) amplifying the nucleic acid molecule in the presence of a first primer having a 5'-sequence having the sequence of a tag complementary to the sequence of a tag complement belonging to a family of tag complements as defined above to form an amplified molecule with a 5'-end with a sequence complementary to the sequence of the tag;

(B) extending the amplified molecule in the presence of a polymerase and a second primer having 5'-end complementary the 3'-end of the amplified sequence, the 3'-end of the second primer extending to immediately adjacent said locus, in the presence of a plurality of nucleoside triphosphate derivatives each of which is: (i) capable of incorporation during transciption by the polymerase onto the 3'-end of a growing nucleotide strand; (ii) causes termination of polymerization; and (iii) capable of differential detection, one from the other, wherein there is a said derivative complementary to each possible nucleotide present at said locus of the amplified molecule;

(C) specifically hybridizing the second primer to a tag complement having the tag complement sequence of (A); and

(D) detecting the nucleotide derivative incorporated into the second primer in (B) so as to identify the base located at the locus of the nucleic acid;

wherein each tag of (A) is unique for each nucleic acid molecule and steps (A) and (B) are carried out with said nucleic molecules in the presence of each other.

Another aspect of the present invention, there is provided a method of analyzing a biological sample comprising a plurality of double stranded complementary nucleic acid molecules for the presence of a mutation or polymorphism at a locus of each nucleic acid molecule, for each nucleic acid molecule. The method includes steps of:

(A) amplifying the double stranded molecule in the presence of a pair of first primers, each primer having an identical 5'-sequence having the sequence of a tag complementary to the sequence of a tag complement belonging to a family of tag complements as defined above to form amplified molecules with 5'-ends with a sequence complementary to the sequence of the tag;

(B) extending the amplified molecules in the presence of a polymerase and a

pair of second primers each second primer having a 5'-end complementary a 3'-end of the amplified sequence, the 3'-end of each said second primer extending to immediately adjacent said locus, in the presence of a plurality of nucleoside triphosphate derivatives each of which is: (i) capable of incorporation during transciption by the polymerase onto the 3'-end of a growing nucleotide strand; (ii) causes termination of polymerization; and (iii) capable of differential detection, one from the other;

(C) specifically hybridizing each of the second primers to a tag complement having the tag complement sequence of (A); and

(D) detecting the nucleotide derivative incorporated into the second primers in (B) so as to identify the base located at said locus;

wherein the sequence of each tag of (A) is unique for each nucleic acid molecule and steps (A) and (B) are carried out with said nucleic molecules in the presence of each other.

In yet another aspect, the invention is a method of analyzing a biological sample containing a plurality of nucleic acid molecules for the presence of a mutation or polymorphism at a locus of each nucleic acid molecule, for each nucleic acid molecule, the method comprising:

(a) hybridizing the molecule and a primer, the primer having a 5'-sequence having the sequence of a tag complementary to the sequence of a tag complement belonging to a family of tag complements as defined above and a 3'-end extending to immediately adjacent the locus;

(b) enzymatically extending the 3'-end of the primer in the presence of a plurality of nucleoside triphosphate derivatives each of which is: (i) capable of enzymatic incorporation onto the 3'-end of a growing nucleotide strand; (ii) causes termination of said extension; and (iii) capable of differential detection, one from the other, wherein there is a said derivative complementary to each possible nucleotide present at said locus;

(c) specifically hybridizing the extended primer formed in step (b) to a tag complement having the tag complement sequence of (a); and

(d) detecting the nucleotide derivative incorporated into the primer in step (b) so as to identify the base located at the locus of the nucleic acid molecule;

wherein each tag of (a) is unique for each nucleic acid molecule and steps (a) and (b) are carried out with said nucleic molecules in the presence of each other.

The derivative can be a dideoxy nucleoside triphosphate.

Each respective complement can be attached as a uniform population of substantially identical complements in spacially discrete regions on one or more solid phase support(s).

Each tag complement can include a label, each such label being different for respective complements, and step (d) can include detecting the presence of the different labels for respective hybridization complexes of bound tags and tag complements.

Another aspect of the invention includes a method of determining the presence of a target suspected of being contained in a mixture, the method comprising the steps of:

(i) labelling the target with a first label;

(ii) providing a first detection moiety capable of specific binding to the target and including a first tag;

(iii) exposing a sample of the mixture to the detection moiety under conditions suitable to permit (or cause) said specific binding of the molecule and target;

(iv) providing a family of suitable tag complements as defined above wherein the family contains a first tag complement having a sequence complementary to that of the first tag;

(v) exposing the sample to the family of tag complements under conditions suitable to permit (or cause) specific hybridization of the first tag and its tag complement;

(vi) determining whether a said first detection moiety hybridized to a first said tag complement is bound to a said labelled target in order to determine the presence or absence of said target in the mixture.

Preferably , the first tag complement is linked to a solid support at a specific location of the support and step (vi) includes detecting the presence of the first label at said specified location.

Also, the first tag complement can include a second label and step (vi) includes detecting the presence of the first and second labels in a hybridized complex of the moiety and the first tag complement.

Further, the target can be selected from the group consisting of organic molecules, antigens, proteins, polypeptides, antibodies and nucleic acids. The target can be an antigen and the first molecule can be an antibody specific for that antigen.

The antigen is usually a polypeptide or protein and the labelling step can include conjugation of fluorescent molecules, digoxigenin, biotinylation and the like.

The target can be a nucleic acid and the labelling step can include incorporation of fluorescent molecules, radiolabelled nucleotide, digoxigenin, biotinylation and the like.

DETAILED DESCRIPTION OF THE INVENTION

FIGURES

Reference is made to the attached figures in which,

Figure 1 illustrates generally the steps followed to obtain a family of sequences of the present invention;

Figure 2 shows the intensity of the signal (MFI) for each perfectly matched sequence (probe sequences indicated in Table I) and its complement (target, at 50 fmol) obtained as described in Example 1;

Figure 3 is a three dimensional representation showing cross-hybridization observed for the sequences of Figure 2 as described in Example 1. The results shown in Figure 2 are reproduced along the diagonal of the drawing; and

Figure 4 is illustrative of results obtained for an individual target (SEQ ID NO:90, target No. 90) when exposed to the 100 probes of Example 1. The MFI for each bead is plotted.

DETAILED EMBODIMENTS

The invention provides a method for sorting complex mixtures of molecules by the use of families of oligonucleotide sequence tags. The families of oligonucleotide sequence tags are designed so as to provide minimal cross hybridization during the sorting process. Thus any sequence within a family of sequences will not significantly cross-hybridize with any other sequence derived from that family under appropriate hybridization conditions known by those skilled in the art. The invention is particularly useful in highly parallel processing of analytes.

Families of Oligonucleotide Sequence Tags

The present invention includes a family of 24mer polynucleotides that have been demonstrated to be minimally cross-hybridizing with each other. This family of polynucleotides is thus useful as a family of tags, and their complements as tag complements.

In order to be considered for inclusion into the family, a sequence had to satisfy a certain number of rules regarding its composition. For example, repetitive regions that present potential hybridization problems such as four or more of a similar base (e.g., AAAA or TTTT) or pairs of Gs were forbidden. Another rule is that each sequence contains exactly six Gs and no Cs, in order

to have sequences that are more or less isothermal. Also required for a 24mer to be included is that there must be at most six bases between every neighboring pair of Gs. Another way of putting this is that there are at most six non-Gs between any two consecutive Gs. Also, each G nearest the 5'-end (resp. 3'-end) of its oligonucleotide (the left-hand (resp. right-hand) side as written in Table I) was required to occupy one of the first to seventh positions (counting the 5'-terminal (resp. 3'-terminal) position as the first position.)

The process used to design families of sequences that do not exhibit cross-hybridization behavior is illustrated generally in Figure 1). Depending on the application for which these families of sequences will be used, various rules are designed. A certain number of rules can specify constraints for sequence composition (such as the ones described in the previous paragraph). The other rules are used to judge whether two sequences are too similar. Based on these rules, a computer program can derive families of sequences that exhibit minimal or no cross-hybridization behavior. The exact method used by the computer program is not crucial since various computer programs can derive similar families based on these rules. Such a program is for example described in international patent application No. PCT/CA 01/00141 published under WO 01/59151 on August 16, 2001. Other programs can use different methods, such as the ones summarized below.

A first method of generating a maximum number of minimally cross-hybridizing polynucleotide sequences starts with any number of non-cross-hybridizing sequences, for example just one sequence, and increases the family as follows. A certain number of sequences is generated and compared to the sequences already in the family. The generated sequences that exhibit too much similarity with sequences already in the family are dropped. Among the "candidate sequences" that remain, one sequence is selected and added to the family. The other candidate sequences are then compared to the selected sequence, and the ones that show too much similarity are dropped. A new sequence is selected from the remaining candidate sequences, if any, and added to the family, and so on until there are no candidate sequences left. At this stage, the process can be repeated (generating a certain number of sequences and comparing them to the sequences in the family, etc.) as often as desired. The family obtained at the end of this method contains only minimally cross-hybridizing sequences.

A second method of generating a maximum number of minimally cross-hybridizing polynucleotide sequences starts with a fixed-size family of polynucleotide sequences. The sequences of this family can be generated randomly or designed by some other method. Many sequences in this family may

not be compatible with each other, because they show too much similarity and are not minimally cross-hybridizing. Therefore, some sequences need to be replaced by new ones, with less similarity. One way to achieve this consists of repeatedly replacing a sequence of the family by the best (that is, lowest similarity) sequence among a certain number of (for example, randomly generated) sequences that are not part of the family. This process can be repeated until the family of sequences shows minimal similarity, hence minimal cross-hybridizing, or until a set number of replacements has occurred. If, at the end of the process, some sequences do not obey the similarity rules that have been set, they can be taken out of the family, thus providing a somewhat smaller family that only contains minimally cross-hybridizing sequences. Some additional rules can be added to this method in order to make it more efficient, such as rules to determine which sequence will be replaced.

Such methods have been used to obtain the 1168 non-cross-hybridizing tags of Table I that are the subject of this patent application.

One embodiment of the invention is a composition comprising molecules for use as tags or tag complements wherein each molecule comprises an oligonucleotide selected from a set of oligonucleotides based on the group of sequences set out in Table IA, wherein each of the numeric identifiers 1 to 3 (see the Table) is a nucleotide base selected to be different from the others of 1 to 3. According to this embodiment, several different families of specific sets of oligonucleotide sequences are described, depending upon the assignment of bases made to the numeric identifiers 1 to 3.

The sequences contained in Table I have a mathematical relationship to each other, described as follows.

Let S and T be two DNA sequences of lengths s and t respectively. While the term "alignment" of nucleotide sequences is widely used in the field of biotechnology, in the context of this invention the term has a specific meaning illustrated here. An *alignment* of S and T is a $2xp$ matrix A (with $p \geq s$ and $p \geq t$) such that the first (or second) row of A contains the characters of S (or T respectively) in order, interspersed with p-s (or p-t respectively) spaces. It assumed that no column of the alignment matrix contains two spaces, i.e., that any alignment in which a column contains two spaces is ignored and not considered here. The columns containing the same base in both rows are called *matches*, while the columns containing different bases are called *mismatches*. Each column of an alignment containing a space in its first row is called an *insertion* and each colmun containing a space in its second row is called a *deletion* while a column of the alignment containing a space in either

row is called an *indel*. Insertions and deletions within a sequence are represented by the character '-'. A *gap* is a continuous sequence of spaces in one of the rows (that is neither immediately preceded nor immediately followed by another space in the same row), and the *length of a gap* is the number of spaces in that gap. An *internal gap* is one in which its first space is preceded by a base and its last space is followed by a base and an *internal indel* is an indel belonging to an internal gap. Finally, a *block* is a continuous sequence of matches (that is neither immediately preceded nor immediately followed by another match), and the *length of a block* is the number of matches in that block. In order to illustrate these definitions, two sequences S = TGATCGTAGCTACGCCGCG (of length s = 19; SEQ ID NO:1169) and T = CGTACGATTGCAACGT (of length t = 16; SEQ ID NO:1170) are considered. Exemplary alignment R_1 of S and T (with p = 23) is:

Alignment R_1:

-	-	-	-	T	G	A	T	C	G	T	A	G	C	T	A	C	G	C	C	G	C	G
C	G	T	A	C	G	A	T	-	-	T	-	G	C	A	A	C	G	T	-	-	-	-
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23

Columns 1 to 4, 9, 10, 12 and 20 to 23 are indels, columns 6, 7, 8, 11, 13, 14, 16, 17 and 18 are matches, and columns 5, 15 and 19 are mismatches. Columns 9 and 10 form a gap of length 2, while columns 16 to 18 form a block of length 3. Columns 9, 10 and 12 are internal indels.

A score is assigned to the alignment A of two sequences by assigning weights to each of matches, mismatches and gaps as follows:

- the reward for a match m,
- the penalty for a mismatch mm,
- the penalty for opening a gap og,
- the penalty for extending a gap eg.

Once these values are set, a score to each column of the alignment is assigned according to the following rules:

1. assign 0 to each column preceding the first match and to each column following the last match.
2. for each of the remaining columns, assign m if it is a match, $-mm$ if it is a mismatch, $-og-eg$ if it is the first indel

of a gap, *-eg* if it is an indel but not the first indel of a gap.

The *score of the alignment* A is the sum of the scores of its columns. An *alignment* is said to be *of maximum score* if no other alignment of the same two sequences has a higher score (with the same values of *m*, *mm*, *og* and *eg*). A person knowledgeable in the field will recognize this method of scoring an alignment as scoring a local (as opposed to global) alignment with affine gap penalties (that is, gap penalties that can distinguish between the first indel of a gap and the other indels). It will be appreciated that the total number of indels that open a gap is the same as the total number of gaps and that an internal indel is not one of those assigned a 0 in rule (1) above. It will also be noted that foregoing rule (1) assigns a 0 for non-internal mismatches. An internal mismatch is a mismatch that is preceded and followed (not necessarily immediately) by a match.

As an illustration, if the values of *m*, *mm*, *og* and *eg* are set to 3, 1, 2 and 1 respectively, alignment R_1 has a score of 19, determined as shown below:

Scoring of Alignment R_1

-	-	-	-	T	G	A	T	C	G	T	A	G	C	T	A	C	G	C	C	G	C	G
C	G	T	A	C	G	A	T	-	-	T	-	G	C	A	A	C	G	T	-	-	-	-
0	0	0	0	0	3	3	3	-3	-1	3	-3	3	3	-1	3	3	3	0	0	0	0	0

Note that for two given sequences S and T, there are numerous alignments. There are often several alignments of maximum score.

Based on these alignments, five sequence similarity measures are defined as follows. For two sequences S and T, and weights {*m*, *mm*, *og*, *eg* }:

- *M1* is the maximum number of matches over all alignments free of internal indels;
- *M2* is the maximum length of a block over all alignments;
- *M3* is the maximum number of matches over all alignments of maximum score;
- *M4* is the maximum sum of the lengths of the longest two blocks over all alignments of maximum score;

- *M5* is the maximum sum of the lengths of all the blocks of length at least 3, over all alignments of maximum score.

Notice that, by definition, the following inequalities between these similarity measures are obtained: $M4 \leq M3$ and $M5 \leq M3$. Also, in order to determine *M2* it is sufficient to determine the maximum length of a block over all alignments free of internal indels. For two given sequences, the values of *M3* to *M5* can vary depending on the values of the weights {*m*, *mm*, *og*, *eg* }, but not *M1* and *M2*.

For weights {3, 1, 2, 1}, the illustrated alignment is not a maximum score alignment of the two example sequences. But for weights {6, 6, 0, 6} it is; hence this alignment shows that for these two example sequences, and weights {6, 6, 0, 6}, $M2 \geq 3$, $M3 \geq 9$, $M4 \geq 6$ and $M5 \geq 6$. In order to determine the exact values of *M1* to *M5*, all the necessary alignments need to be considered. *M1* and *M2* can be found by looking at the *s+t-1* alignments free of internal indels, where *s* and *t* are the lengths of the two sequences considered. Mathematical tools known as dynamic programming can be implemented on a computer and used to determine *M3* to *M5* in a very quick way. Using a computer program to do these calculations, it was determined that:

- with the weights {6, 6, 0, 6}, $M1 = 8$, $M2 = 4$, $M3 = 10$, $M4 = 6$ and $M5 = 6$;
- with the weights {3, 1, 2, 1}, $M1 = 8$, $M2 = 4$, $M3 = 10$, $M4 = 6$ and $M5 = 4$.

According to the preferred embodiment of this invention, two sequences S and T each of length 24 are *too similar* if at least one of the following happens:

- $M1 > 16$ or
- $M2 \leq 13$ or
- $M3 > 20$ or
- $M4 > 16$ or
- $M5 > 19$

when using either weights {6, 6, 0, 6}, or {6, 6, 5, 1}, or {6, 2, 5, 1}, or {6, 6, 6, 0}. In other words, the five similarity measures between S and T are determined for each of the above four sets of weights, and checked against these thresholds (for a total of 20 tests).

The above thresholds of 16, 13, 20, 16 and 19, and the above sets of weights, were used to obtain the sequences listed in Table I. Additional sequences can thus be added to those of Table I as long as the above alignment rules are obeyed for all sequences.

It is also possible to alter thresholds *M1*, *M2*, etc., while remaining within the scope of this invention. It is thus possible to substitute or add sequences to those of Table I, or more generally to those of Table IA to obtain other sets of sequences that would also exhibit reasonably low cross-hybridization. More specifically, a set of 24mer sequences in which there are no two sequences that are too similar, where too similar is defined as:

- *M1* > *19* or
- *M2* > *17* or
- *M3* > *21* or
- *M4* > *18* or
- *M5* > *20*

when using either weights {6, 6, 0, 6}, or {6, 6, 5, 1}, or {6, 2, 5, 1}, or {6, 6, 6, 0}, would also exhibit low cross-hybridization. Reducing any of the threshold values provides sets of sequences with even lower cross-hybridization. Alternatively, 'too similar' can also be defined as:

- *M1* > *19* or
- *M2* > *17* or
- *M3* > *21* or
- *M4* > *18* or
- *M5* > *20*

when using either weights {3, 1, 2, 1}. Alternatively, other combinations of weights will lead to sets of sequences with low cross-hybridization.

Notice that using weights {6, 6, 0, 6} is equivalent to using weights {1, 1, 0, 1}, or weights {2, 2, 0, 2}, … (that is, for any two sequences, the values of M1 to M5 are exactly the same whether weights {6, 6, 0, 6} or {1, 1, 0, 1} or {2, 2, 0, 2} or any other multiple of {1, 1, 0, 1} is used).

When dealing with sequences of length other than 24, or sequences of various lengths, the definition of similarity can be adjusted. Such adjustments are obvious to the persons skilled in the art. For example, when comparing a sequence of length L1 with a sequence of length L2 (with L1<L2), they can be considered as too similar when

M1 > 19/24 x L1

M2 > 17/24 x L1

M3 > 21/24 x L1

M4 > 18/24 x L1

$M5 > 20/24 \times L1$

when using either weights {6, 6, 0, 6}, or {6, 6, 5, 1}, or {6, 2, 5, 1} or {6, 6, 6, 0}.

Polynucleotide sequences can be composed of a subset of natural bases most preferably A, T and G. Sequences that are deficient in one base possess useful characteristics, for example, in reducing potential secondary structure formation or reduced potential for cross hybridization with nucleic acids in nature. Also, it is preferable to have tag sequences that behave isothermally. This can be achieved for example by maintaining a constant base composition for all sequences such as six Gs and eighteen As or Ts for each sequence. Additional sets of sequences can be designed by extrapolating on the original family of non-cross-hybridizing sequences by simple methods known to those skilled in the art.

In order to validate the sequence set, a subset of sequences from the family of 1168 sequence tags was selected and characterized, in terms of the ability of these sequences to form specific duplex structures with their complementary sequences, and the potential for cross-hybridization within the sequence set. See Example 1, below. The subset of 100 sequences was randomly selected, and analyzed using the *Luminex*^{100} LabMAP™ platform. The 100 sequences were chemically immobilized onto the set of 100 different *Luminex* microsphere populations, such that each specific sequence was coupled to one spectrally distinct microsphere population. The pool of 100 microsphere-immobilized probes was then hybridized with each of the 100 corresponding complementary sequences. Each sequence was examined individually for its specific hybridization with its complementary sequence, as well as for its non-specific hybridization with the other 99 sequences present in the reaction. This analysis demonstrated the propensity of each sequence to hybridize only to its complement (perfect match), and not to cross-hybridize appreciably with any of the other oligonucleotides present in the hybridization reaction.

It is within the capability of a person skilled in the art, given the family of sequences of Table I, to modify the sequences, or add other sequences while largely retaining the property of minimal cross-hybridization which the polynucleotides of Table I have been demonstrated to have.

There are 1168 polynucleotide sequences given in Table I. Since all 1168 of this family of polynucleotides can work with each other as a minimally cross-hybridizing set, then any plurality of polynucleotides that is a subset of the 1168 can also act as a minimally cross-hybridizing set of polynucleotides. An application in which, for example, 30 molecules are to be

sorted using a family of polynucleotide tags and tag complements could thus use any group of 30 sequences shown in Table I. This is not to say that some subsets may be found in a practical sense to be more preferred than others. For example, it may be found that a particular subset is more tolerant of a wider variety of conditions under which hybridization is conducted before the degree of cross-hybridization becomes unacceptable.

It may be desirable to use polynucleotides that are shorter in length than the 24 bases of those in Table I. A family of subsequences (i.e., subframes of the sequences illustrated) based on those contained in Table I having as few as 10 bases per sequence could be chosen, so long as the subsequences are chosen to retain homological properties between any two of the sequences of the family important to their non cross-hybridization.

The selection of sequences using this approach would be amenable to a computerized process. Thus for example, a string of 10 contiguous bases of the first 24mer of Table I could be selected: AAATTGTGAAAGATTGTTTGTGTA (SEQ ID NO:1).

The same string of contiguous bases from the second 24mer could then be selected and compared for similarity against the first chosen sequence: GTTAGAGTTAATTGTATTTGATGA (SEQ ID NO:2). A systematic pairwise comparison could then be carried out to determine if the similarity requirements are violated. If the pair of sequences does not violate any set property, a 10mer subsequence can be selected from the third 24mer sequence of Table I, and compared to each of the first two 10mer sequences (in a pairwise fashion to determine its compatibility therewith, etc. In this way a family of 10mer sequences may be developed.

It is within the scope of this invention, to obtain families of sequences containing 11mer, 12mer, 13mer, 14mer, 15mer, 16mer, 17mer, 18mer, 19mer, 20mer, 21mer, 22mer and 23mer sequences by analogy to that shown for 10mer sequences.

It may be desirable to have a family of sequences in which there are sequences greater in length than the 24mer sequences shown in Table I. It is within the capability of a person skilled in the art, given the family of sequences shown in Table I, to obtain such a family of sequences. One possible approach would be to insert into each sequence at one or more locations a nucleotide, non-natural base or analogue such that the longer sequence should not have greater similarity than any two of the original non-cross-hybridizing sequences of Table I and the addition of extra bases to the tag sequences should not result in a major change in the thermodynamic properties of the tag sequences of that set for example the GC content must be maintained between 10%-40% with a variance from the average of 20%. This

method of inserting bases could be used to obtain, for example, a family of sequences up to 40 bases long.

Given a particular family of sequences that can be used as a family of tags (or tag complements), e.g., those of Table I, a skilled person will readily recognize variant families that work equally as well.

Again taking the sequences of Table I for example, every T could be converted to an A and vice versa and no significant change in the cross-hybridization properties would be expected to be observed. This would also be true if every G were converted to a C.

Also, all of the sequences of a family could be taken to be constructed in the 5'-3' direction, as is the convention, or all of the constructions of sequences could be in the opposition direction (3'-5').

There are additional modifications that can be carried out. For example, C has not been used in the family of sequences. Substitution of C in place of one or more G's of a particular sequence would yield a sequence that is at least as low in homology with every other sequence of the family as was the particular sequence chosen for modification. It is thus possible to substitute C in place of one or more G's in any of the sequences shown in Table I. Analogously, substituting of C in place of one or more A's is possible, or substituting C in place of one or T's is possible.

It is preferred that the sequences of a given family are of the same, or roughly the same length. Preferably, all the sequences of a family of sequences of this invention have a length that is within five bases of the base-length of the average of the family. More preferably, all sequences are within four bases of the average base-length. Even more preferably, all or almost all sequences are within three bases of the average base-length of the family. Better still, all or almost all sequences have a length that is within two of the base-length of the average of the family, and even better still, within one of the base-length of the average of the family.

It is also possible for a person skilled in the art to derive sets of sequences from the family of sequences described in this specification and remove sequences that would be expected to have undesirable hybridization properties.

Methods For Synthesis Of Oligonucleotide Families

Preferably oligonucleotide sequences of the invention are synthesized directly by standard phosphoramidite synthesis approaches and the like (Caruthers et al, Methods in Enzymology; 154, 287-313: 1987; Lipshutz et al, Nature Genet.; 21, 20-24: 1999; Fodor et al, Science; 251, 763-773: 1991). Alternative chemistries involving non

natural bases such as peptide nucleic acids or modified nucleosides that offer advantages in duplex stability may also be used (Hacia et al; Nucleic Acids Res ;27: 4034-4039, 1999; Nguyen et al, Nucleic Acids Res.;27, 1492-1498: 1999; Weiler et al, Nucleic Acids Res.; 25, 2792-2799:1997). It is also possible to synthesize the oligonucleotide sequences of this invention with alternate nucleotide backbones such as phosphorothioate or phosphoroamidate nucleotides. Methods involving synthesis through the addition of blocks of sequence in a stepwise manner may also be employed (Lyttle et al, Biotechniques, 19: 274-280 (1995). Synthesis may be carried out directly on the substrate to be used as a solid phase support for the application or the oligonucleotide can be cleaved from the support for use in solution or coupling to a second support.

Solid Phase Supports

There are several different solid phase supports that can be used with the invention. They include but are not limited to slides, plates, chips, membranes, beads, microparticles and the like. The solid phase supports can also vary in the materials that they are composed of including plastic, glass, silicon, nylon, polystyrene, silica gel, latex and the like. The surface of the support is coated with the complementary tag sequences by any conventional means of attachment.

In preferred embodiments, the family of tag complement sequences is derivatized to allow binding to a solid support. Many methods of derivatizing a nucleic acid for binding to a solid support are known in the art (Hermanson G., Bioconjugate Techniques; Acad. Press: 1996). The sequence tag may be bound to a solid support through covalent or non-covalent bonds (Iannone et al, Cytometry; 39: 131-140, 2000; Matson et al, Anal. Biochem.; 224: 110-106, 1995; Proudnikov et al, Anal Biochem; 259: 34-41, 1998; Zammatteo et al, Analytical Biochemistry; 280:143-150, 2000). The sequence tag can be conveniently derivatized for binding to a solid support by incorporating modified nucleic acids in the terminal 5′ or 3′ locations.

A variety of moieties useful for binding to a solid support (e.g., biotin, antibodies, and the like), and methods for attaching them to nucleic acids, are known in the art. For example, an amine-modified nucleic acid base (available from, eg., Glen Research) may be attached to a solid support (for example, Covalink-NH, a polystyrene surface grafted with secondary amino groups, available from Nunc) through a bifunctional crosslinker (e.g., bis(sulfosuccinimidyl suberate), available from Pierce). Additional spacing

moieties can be added to reduce steric hindrance between the capture moiety and the surface of the solid support.

Attaching Tags to Analytes for Sorting

A family of oligonucleotide tag sequences can be conjugated to a population of analytes most preferably polynucleotide sequences in several different ways including but not limited to direct chemical synthesis, chemical coupling, ligation, amplification, and the like. Sequence tags that have been synthesized with primer sequences can be used for enzymatic extension of the primer on the target for example in PCR amplification.

Detection of Single Nucleotide Polymorphisms Using Primer Extension

There are a number of areas of genetic analysis where families of non-cross-hybridizing sequences can be applied including disease diagnosis, single nucleotide polymorphism analysis, genotyping, expression analysis and the like. One such approach for genetic analysis, referred to as the primer extension method (also known as Genetic Bit Analysis (Nikiforov et al, Nucleic Acids Res.; 22, 4167-4175: 1994; Head et al Nucleic Acids Res.; 25, 5065-5071: 1997)), is an extremely accurate method for identification of the nucleotide located at a specific polymorphic site within genomic DNA. In standard primer extension reactions, a portion of genomic DNA containing a defined polymorphic site is amplified by PCR using primers that flank the polymorphic site. In order to identify which nucleotide is present at the polymorphic site, a third primer is synthesized such that the polymorphic position is located immediately 3′ to the primer. A primer extension reaction is set up containing the amplified DNA, the primer for extension, up to 4 dideoxynucleoside triphosphates (each labeled with a different fluorescent dye) and a DNA polymerase such as the Klenow subunit of DNA Polymerase 1. The use of dideoxy nucleotides ensures that a single base is added to the 3′ end of the primer, a site corresponding to the polymorphic site. In this way the identity of the nucleotide present at a specific polymorphic site can be determined by the identity of the fluorescent dye-labeled nucleotide that is incorporated in each reaction. One major drawback to this approach is its low throughput. Each primer extension reaction is carried out independently in a separate tube.

Universal sequences can be used to enhance the throughput of primer extension assay as follows. A region of genomic DNA containing multiple polymorphic sites is amplified by PCR. Alternatively, several genomic regions containing one or more polymorphic sites each are amplified together in a multiplexed PCR reaction. The primer extension reaction

is carried out as described above except that the primers used are chimeric, each containing a unique universal tag at the 5′ end and the sequence for extension at the 3′ end. In this way, each gene-specific sequence would be associated with a specific universal sequence. The chimeric primers would be hybridized to the amplified DNA and primer extension is carried out as described above. This would result in a mixed pool of extended primers, each with a specific fluorescent dye characteristic of the incorporated nucleotide. Following the primer extension reaction, the mixed extension reactions are hybridized to an array containing probes that are reverse complements of the universal sequences on the primers. This would segregate the products of a number of primer extension reactions into discrete spots. The fluorescent dye present at each spot would then identify the nucleotide incorporated at each specific location. A number of additional methods for the detection of single nucleotide polymorphisms, including but not limited to, allele specific polymerase chain reaction (ASPCR), allele specific primer extension (ASPE) and oligonucleotide ligation assay (OLA) can be performed by someone skilled in the art in combination with the tag sequences described herein.

Kits Using Families Of Tag Sequences

The families of non cross-hybridizing sequences may be provided in kits for use in for example genetic analysis. Such kits include at least one set of non-cross-hybridizing sequences in solution or on a solid support. Preferably the sequences are attached to microparticles and are provided with buffers and reagents that are appropriate for the application. Reagents may include enzymes, nucleotides, fluorescent labels and the like that would be required for specific applications. Instructions for correct use of the kit for a given application will be provided.

EXAMPLES

EXAMPLE 1 - Cross Talk Behavior of Sequence on Beads

A group of 100 sequences, randomly selected from Table I, was tested for feasibility for use as a family of minimally cross-hybridizing oligonucleotides. The 100 sequences selected are separately indicated in Table I along with the numbers assigned to the sequences in the tests.

The tests were conducted using the Luminex LabMAP™ platform available from Luminex Corporation, Austin, Texas, U.S.A. The one hundred sequences, used as probes, were synthesized as oligonucleotides by Integrated DNA Technologies (IDT, Coralville, Iowa, U.S.A.). Each probe included a C_6

aminolink group coupled to the 5'-end of the oligonucleotide through a C_{12} ethylene glycol spacer. The C_6 aminolink molecule is a six carbon spacer containing an amine group that can be used for attaching the oligonucleotide to a solid support. One hundred oligonucleotide targets (probe complements), the sequence of each being the reverse complement of the 100 probe sequences, were also synthesized by IDT. Each target was labelled at its 5'-end with biotin. All oligonucleotides were purified using standard desalting procedures, and were reconstituted to a concentration of approximately 200 µM in sterile, distilled water for use. Oligonucleotide concentrations were determined spectrophotometrically using extinction coefficients provided by the supplier.

Each probe was coupled by its amino linking group to a carboxylated fluorescent microsphere of the LapMAP system according to the *Luminex*^{100} protocol. The microsphere, or bead, for each probe sequence has unique, or spectrally distinct, light absorption characteristics which permits each probe to be distinguished from the other probes. Stock bead pellets were dispersed by sonication and then vortexing. For each bead population, five million microspheres (400 µL) were removed from the stock tube using barrier tips and added to a 1.5 mL Eppendorf tube (USA Scientific). The microspheres were then centrifuged, the supernatant was removed, and beads were resuspended in 25 µL of 0.2 M MES (2-(N-morpholino)ethane sulfonic acid) (Sigma), pH 4.5, followed by vortexing and sonication. One nmol of each probe (in a 25 µL volume) was added to its corresponding bead population. A volume of 2.5 µL of EDC cross-linker (1-ethyl-3-(3-dimethylaminopropyl) carbodiimide hydrochloride (Pierce), prepared immediately before use by adding 1.0 mL of sterile ddH_2O to 10 mg of EDC powder, was added to each microsphere population. Bead mixes were then incubated for 30 minutes at room temperature in the dark with periodic vortexing. A second 2.5 µL aliquot of freshly prepared EDC solution was then added followed by an additional 30 minute incubation in the dark. Following the second EDC incubation, 1.0 mL of 0.02% Tween-20 (BioShop) was added to each bead mix and vortexed. The microspheres were centrifuged, the supernatant was removed, and the beads were resuspended in 1.0 mL of 0.1% sodium dodecyl sulfate (Sigma). The beads were centrifuged again and the supernatant removed. The coupled beads were resuspended in 100 µL of 0.1 M MES pH 4.5. Bead concentrations were then determined by diluting each preparation 100-fold in ddH_2O and enumerating using a

Neubauer BrightLine Hemacytometer. Coupled beads were stored as individual populations at 8°C protected from light.

The relative oligonucleotide probe density on each bead population was assessed by Terminal Deoxynucleotidyl Transferase (TdT) end-labelling with biotin-ddUTPs. TdT was used to label the 3'-ends of single-stranded DNA with a labeled ddNTP. Briefly, 180 µL of the pool of 100 bead populations (equivalent to about 4000 of each bead type) to be used for hybridizations was pipetted into an Eppendorf tube and centrifuged. The supernatant was removed, and the beads were washed in 1x TdT buffer. The beads were then incubated with a labelling reaction mixture, which consisted of 5x TdT buffer, 25mM $CoCl_2$, and 1000 pmol of biotin-16-ddUTP (all reagents were purchased from Roche). The total reaction volume was brought up to 85.5 µL with sterile, distilled H_2O, and the samples were incubated in the dark for 1 hour at 37°C. A second aliquot of enzyme was added, followed by a second 1 hour incubation. Samples were run in duplicate, as was the negative control, which contained all components except the TdT. In order to remove unincorporated biotin-ddUTP, the beads were washed 3 times with 200 µL of hybridization buffer, and the beads were resuspended in 50 µL of hybridization buffer following the final wash. The biotin label was detected spectrophotometrically using SA-PE (streptavidin-phycoerythrin conjugate). The streptavidin binds to biotin and the phycoerythrin is spectrally distinct from the probe beads. The 10mg/mL stock of SA-PE was diluted 100-fold in hybridization buffer, and 15 µL of the diluted SA-PE was added directly to each reaction and incubated for 15 minutes at 37°Celsius. The reactions were analyzed on the *Luminex*^{100} LabMAP. Acquisition parameters were set to measure 100 events per bead using a sample volume of 50 µL.

The results obtained are shown in Figure 2. As can be seen the Mean Fluorescent Intensity (MFI) of the beads varies from 840.3 to 3834.9, a range of 4.56-fold. Assuming that the labelling reactions are complete for all of the oligonucleotides, this illustrates the signal intensity that would be obtained for each type of bead at this concentration if the target (i.e., labelled complement) was bound to the probe sequence to the full extent possible.

The cross-hybridization of targets to probes was evaluated as follows. 100 oligonucleotide probes linked to 100 different bead populations, as described above, were combined to generate a master bead mix, enabling multiplexed reactions to be carried out. The pool of

microsphere-immobilized probes was then hybridized individually with each biotinylated target. Thus, each target was examined individually for its specific hybridization with its complementary bead-immobilized sequence, as well as for its non-specific hybridization with the other 99 bead-immobilized universal sequences present in the reaction. For each hybridization reaction, 25 µL bead mix (containing about 2500 of each bead population in hybridization buffer) was added to each well of a 96-well Thermowell PCR plate and equilibrated at 37°C. Each target was diluted to a final concentration of 0.002 fmol/µL in hybridization buffer, and 25 µL (50 fmol) was added to each well, giving a final reaction volume of 50 µL. Hybridization buffer consisted of 0.2 M NaCl, 0.1 M Tris, 0.08% Triton X-100, pH 8.0 and hybridizations were performed at 37°C for 30 minutes. Each target was analyzed in triplicate and six background samples (i.e. no target) were included in each plate. A SA-PE conjugate was used as a reporter, as described above. The 10 mg/mL stock of SA-PE was diluted 100-fold in hybridization buffer, and 15 µL of the diluted SA-PE was added directly to each reaction, without removal of unbound target, and incubated for 15 minutes at 37°C. Finally, an additional 35 µL of hybridization buffer was added to each well, resulting in a final volume of 100 µL per well prior to analysis on the *Luminex*^{100} LabMAP. Acquisition parameters were set to measure 100 events per bead using a sample volume of 80 µL.

The percent hybridization was calculated for any event in which the NET MFI was at least 3 times the zero target background. In other words, a calculation was made for any sample where $(MFI_{sample}-MFI_{zero\ target\ background})/MFI_{zero\ target\ background} \geq 3$.

The net median fluorescent intensity ($MFI_{sample}-MFI_{zero\ target\ background}$) generated for all of the 10,000 possible target/probe combinations was calculated. As there are 100 probes and 100 targets, there are 100 x 100 = 10,0000 possible different interactions possible of which 100 are the result of perfect hybridizations. The remaining 9900 result from hybridization of a target with a mismatched probe. A cross-hybridization event is then defined as a non-specific event whose net median fluorescent intensity exceeds 3 times the zero target background. In other words, a cross-talk calculation is only be made for any sample where $(MFI_{sample}-MFI_{zero\ target\ background})/MFI_{zero\ target\ background} \geq 3$. Cross-hybridization events were quantified by expressing the value of the cross-hybridization signal as a percentage of the perfect match hybridization signal with the same probe.

The results obtained are illustrated in Figure 3. The ability of each target to be specifically recognized by its matching probe is shown. Of the possible 9900 non-specific hybridization events that could have occurred when the 100 targets were each exposed to the pool of 100 probes, 6 events were observed. Of these 6 events, the highest non-specific event generated a signal equivalent to 5.3% of the signal observed for the perfectly matched pair (i.e. specific hybridization event).

Each of the 100 targets was thus examined individually for specific hybridization with its complement sequence as incorporated onto a microsphere, as well as for non-specific hybridization with the complements of the other 99 target sequences. Representative hybridization results for target (complement of probe 90, Table I) are shown in Figure 4. Probe 90 was found to hybridize only to its perfectly-matched target. No cross-hybridization with any of the other 99 targets was observed.

The foregoing results demonstrate the possibility of incorporating the 1168 sequences of Table I, or any subset thereof, into a multiplexed system with the expectation that most if not all sequences can be distinguished from the others by hybridization. That is, it is possible to distinguish each target from the other targets by hybridization of the target with its precise complement and minimal hybridization with complements of the other targets.

EXAMPLE 2 - Tag sequences used in sorting polynucleotides

The family of non cross hybridizing sequence tags or a subset thereof can be attached to oligonucleotide probe sequences during synthesis and used to generate amplified probe sequences. In order to test the feasibility of PCR amplification with non cross hybridizing sequence tags and subsequently addressing each respective sequence to its appropriate location on two-dimensional or bead arrays, the following experiment was devised. A 24mer tag sequence can be connected in a 5'-3' specific manner to a p53 exon specific sequence (20mer reverse primer). The connecting p53 sequence represents the inverse complement of the nucleotide gene sequence. To facilitate the subsequent generation of single stranded DNA post-amplification the tag-Reverse primer can be synthesized with a phosphate modification (PO_4) on the 5'-end. A second PCR primer can also be generated for each desired exon, represented by the Forward (5'-3') amplification primer. In this instance the Forward primer can be labeled with a 5'-biotin modification to allow detection with Cy3-avidin or equivalent.

A practical example of the aforementioned description is as follows: For exon 1 of the human p53 tumor suppressor gene sequence the following tag-Reverse primer (SEQ ID NO:1171) can be generated:

222087 222063

5'-PO4-*ATGTTAAAGTAAGTGTTGAAATGT* -TCCAGGGAAGCGTGTCACCGTCGT-3'

Tag Sequence # 3 Exon 1 Reverse

The numbering above the Exon-1 reverse primer represents the genomic nucleotide positions of the indicated bases.

The corresponding Exon-1 Forward primer sequence (SEQ ID NO:1172) is as follows:

221873 221896

5'-Biotin-TCATGGCGACTGTCCAGCTTTGTG-3'

In combination these primers will amplify a product of 214 bp plus a 24 bp tag extension yielding a total size of 238 bp.

Once amplified, the PCR product can be purified using a QIAquick PCR purification kit and the resulting DNA can be quantified. To generate single stranded DNA, the DNA is subjected to λ-exonuclease digestion thereby resulting in the exposure of a single stranded sequence (anti-tag) complementary to the tag-sequence covalently attached to the solid phase array. The resulting product is heated to 95°C for 5 minutes and then directly applied to the array at a concentration of 10-50 nM. Following hybridization and concurrent sorting, the tag-Exon 1 sequences are visualized using Cy3-streptavidin. In addition to direct visualization of the biotinylated product, the product itself can now act as a substrate for further analysis of the amplified region, such as SNP detection and haplotype determination.

DEFINITIONS

Non-cross-hybridization: Describes the absence of hybridization between two sequences that are not perfect complements of each other.

Cross-hybridization: The hydrogen bonding of a single-stranded DNA sequence that is partially but not entirely complementary to a single-stranded substrate.

Homology or Similarity: How closely related two or more separate strands of DNA are to each other, based on their base sequences.

Analogue: The symbols A, G, T/U, C take on their usual meaning in the art here. In the case of T and U, a person skilled in the art would understand that these are equivalent to each other with respect to the inter-strand hydrogen-bond (Watson-Crick) binding properties at work in the context of this invention. The two bases are thus interchangeable and hence the designation of T/U. A chemical, which resembles a nucleotide base is an analogue thereof. A base that does not normally appear in DNA but can substitute for the ones, which do, despite minor differences in structure. Analogues particularly useful in this invention are of the naturally occurring bases can be inserted in their respective places where desired. Such an analogue is any non-natural base, such as peptide nucleic acids and the like that undergoes normal Watson-Crick pairing in the same way as the naturally occurring nucleotide base to which it corresponds.

Complement: The opposite or "mirror" image of a DNA sequence. A complementary DNA sequence has an "A" for every "T" and a "C" for every "G". Two complementary strands of single stranded DNA, for example a tag sequence and its complement, will join to form a double-stranded molecule.

Complementary DNA (cDNA): DNA that is synthesized from a *messenger RNA* template; the single- stranded form is often used as a *probe* in *physical mapping*.

Oligonucleotide: Refers to a short nucleotide polymer whereby the nucleotides may be natural nucleotide bases or analogues thereof.

Tag: Refers to an oligonucleotide that can be used for specifically sorting analytes with at least one other oligonucleotide that when used together do not cross hybridize.

Table I

Sequence	SEQ ID NO:	No. in Ex 1
A A A T T G T G A A A G A T T G T T T G T G T A	1	1
G T T A G A G T T A A T T G T A T T T G A T G A	2	-
A T G T T A A A G T A A G T G T T T G A A A T G T	3	-
T G A T G T T A G A A G T A T A T T G T G A A T	4	-
T T T G T G T A G A A T A T G T G T T T G T T A A	5	-

Table I

Sequence	SEQ ID NO:	No. in Ex 1
ATAAGTGTAAGTGAAATAAGAAGA	6	-
AAGAGTATTTGTTGTGAGTTAAAAT	7	-
GTGTTTATGTTATATGTGAAGTTT	8	-
AAAGAGAA'TAGAATATGTGTAAGT	9	-
TATGAAAGAGTGAGATAATGTTTA	10	-
ATGAGAAATATGTTAGAATGTGAT	11	-
TTAGTTGTTGATGTTTAGTAGTTT	12	-
GTAAAGAGTATAAGTTTGATGATA	13	-
AAAGTAAGAATGATGTAATAAGTG	14	-
GTAGAAATAGTTTATTGATGATTG	15	-
TGTAAGTGAAATAGTGAGTTATTT	16	2
AAATAGATGATATAAGTGAGAATG	17	-
ATAAGTTATAAGTGTTATGTGAGT	18	-
TATAGATAAAGAGATGATTTGTTG	19	-
AGAGTTGAGAATGTATAGTATTAT	20	-
AAGTAGTTTGTAAGAATGATTGTA	21	-
TTATGAAATTGAGTGAAGATTGAT	22	-
GTATATGTAAAATTGTTATGTTGAG	23	-
GAATTGTATAAAGTATTAGATGTG	24	4
TAGATGAGATTAAGTGTTATTTGA	25	-
GTTAAGTTTGTTTATGTATAGAAG	26	-
GAGTATTAGTAAAGTGATATGATA	27	-
GTGAATGATTTAGTAAATGATTGA	28	-
GATTGAAGTTATAGAAATGATTAG	29	-
AGTGATAAATGTTAGTTGAATTGT	30	-
TATATAGTAAATGTTTGTGTGTTG	31	-
TTAAGTGTTAGTTATTTGTTGTAG	32	-
GTAGTAATATGAAGTGAGAATATA	33	-
TAGTGTATAGAATGTAGATTTAGT	34	-
TTGTAGATTAGATGTGTTTGTAAA	35	-
TAGTATAGAGTAGAGATGATATTT	36	-
ATTGTGAAAGAAAGAGAAGAAATT	37	7
TGTGAGAATTAAGATTAAGAATGT	38	-
ATATTAGTTAAGAAAGAAGAGTTG	39	-
TTGTAGTTGAGAAATATGTAGTTT	40	-
TAGAGTTGTTAAAGAGTGTAAATA	41	-
GTTATGATGTGTATAAGTAATATG	42	-
TTTGTTAGAATGAGAAGATTTATG	43	10
AGTATAGTTTAAAGAAGTAGTAGA	44	-
GTGAG_ATATAGATTTAGAAAGTAA	45	-
TTGTTTATAGTGAAGTGAATAGTA	46	-
AAGTAAGTAGTAATAGTGTGTTAA	47	-
ATTTGTGAGTTATGAAAGATAAGA	48	-
GAAAGTAGAGA.ATAAAGATAAGAA	49	-
ATTTAAGATTGTTAAGAGTAGAAG	50	-
GTTTAAAGA,TTGTAAGAATGTGTA	51	-
TTTGTGAAGATGAAGTATTTGTAT	52	-
TGTGTTTAGAATTTAGTATGTGTA	53	-
GATAATGATTATAGAAAGTGTTTG	54	-
GTTATTTGTAAGTTAAGATAGTAG	55	-
AGTTTATTGAAAGAGTTTGAATAG	56	-
TTGTGTTTATTGTGTAGTTTAAAG	57	-
ATTGTGAGAAGATATGAAAGTTAT	58	-
TGAGAATGTAAAGAATGTTTATTG	59	13
ATGTGAAAGTTATGATGTTAATTG	60	-
GTTTAGTATTAGTTGTTAAGATTG	61	-
GATTGATATTTGAATGTTTGTTTG	62	14

Table I

Sequence	SEQ ID NO:	No. in Ex 1
TGAATTGAAAGTGTAATGTTGTAT	63	-
GATTGTATTGTTGAGAATAGAATA	64	-
AAATTTGAGATTTGTGATAGAGTA	65	-
GTAATTAGATTTGTTTGTTGTTGT	66	-
GTTTGTATTGTTAGTGAATATAGT	67	-
ATGTAGTAGTAGATGTTTATGAAT	68	-
TGTTTAAAGATGATTGAAGAAATG	69	-
TGTGATAATGATGTTATTTGTGTA	70	-
ATAGTTGTGAGAATTTGTAATTAG	71	-
ATAGATGTAAGAGAAATTGTGAAA	72	-
AGATTAAGAGAAGTTAATAGAGTA	73	-
GAAGTAAATTGTGAATGAAAGAAA	74	-
AATGTAAGAAAGAAGATTGTTGTA	75	-
TTTGATTTATGTGTTATGTTGAGT	76	-
GTATTGAGAAATTTGAAGAATGAA	77	-
GAATTGTATGAAATGAATTGTAAG	78	-
TATTGTAGAAGTAAAGTTAGAAGT	79	-
TTTATGTAATGATAAGTGTAGTTG	80	-
ATATAGTTGAAATTGTGATAGTGT	81	-
ATAAGAAATTAGAGAGTTGTAAAG	82	-
GAATTGTGAAATGTGATTGATATA	83	-
AAATAAGTAGTTTAATGAGAGAAG	84	-
GATTAAAGAAGTAAGTGAATGTTT	85	-
TATGTGTGTTGTTTAGTGTTATTA	86	-
GAGTTATATGTAGTTAGAGTTATA	87	-
GAAAGAAAGAAGTGTTAAGTTAAA	88	-
TAGTATTAGTAAGTATGTGATTGT	89	-
TTGTGTGATTGAAATATTGTGAAAT	90	-
ATGTGAAAGAGTTAAGTGATTAAA	91	-
GATTGAATGATTGAGATATGTAAA	92	-
AAGATGATAGTTAAGTGTAAGTTA	93	17
TAGTTGTTATTGAGAATTTAGAAG	94	-
TTTATAGTGAATTATGAGTGAAAG	95	-
GATAGATTTAGAATGAATTAAGTG	96	18
TTTGAAGAAGAGATTTGAAATTGA	97	-
ATGAATAAGAGTTGATAAATGTGA	98	-
TGTTTATGTAGTGTAGATTGAATT	99	-
TTTAAGTGAGTTATAGAAGTAGTA	100	19
GATTTATGTGTTTGAAGTTAAGAT	101	-
TAGTT-AGAGAAAAGTGATAAAGTTA	102	-
GTAATGATAATGAAGTGTATATAG	103	-
AATGAAGTGTTAGTATAGATAGTA	104	-
TAAATTGAGTTTGTTTGATTGTAG	105	-
TAATGAAGAATAAGTATGAGTGTT	106	-
AAATGTAATAGTGTTGTTAGTTAG	107	-
AGAGTTAGTGAAATGTTGTTAAAT	108	-
GAAATAGAAATGTATTGTTTGTGA	109	-
AGTTATAAGTTTGTGAGAATTAAG	110	-
GAGTTTATAGTTAGAATATGTTGT	111	-
AGAGTTATTAGAAGAAGATTTAAG	112	-
GAGTTAATGAAATAAGTATTTGTG	113	-
ATGATGAATAGTTGAAGTATATAG	114	-
ATAGATATGAGATGAAAGTTAGTA	115	-
TATGTAAAGAAAGTGAAAGAAGAA	116	-
TGAATGTAGAAATGAATGTTGAAA	117	-
AATTGAATAGTGTGTGAGTTTAAT	118	-
AGATATTGTTTGATTAATGAAGAG	119	-

Table I

Sequence	SEQ ID NO:	No. in Ex 1
AAAGTTGTAAAGTTGAAGATAAAG	120	-
GTTAAGAGATTATGAGATGTATTA	121	-
AGAAGATATAAGAAGATTGAATTG	122	-
GTAGAAATTTGAATTGATGTGAAA	123	-
AAGAGTAGATTGATAAGTATATGA	124	-
TGATATAGTAGTGAAGAAATAAGT	125	22
AGATAATGATGAGAAATGAAGATA	126	-
ATGTGAAAGTATTTGTGATATAGT	127	-
AATAAGAGAATTGATATGAAGATG	128	23
TAAGTGTATTTAGTAGAATGAAGT	129	-
TATGTTAGATTTGTTGAGATTGAT	130	-
AGTTTGTATGAAGAGATAGTATTT	131	-
GAGAAATGTTATGTATTTAGTAGT	132	-
TATGTGAGAATGTGTTTGATTTAA	133	-
GTATGTTTGTTTATAGAATGTATG	134	-
GAGTATATAGAAGAAAGAAATTTG	135	-
ATGAGTGAAGTAAATGTAGTTATT	136	-
TTAAGAAGTGAGTTATTGTGATAT	137	-
ATGAAATGAGAATATTGTTGTTTG	138	-
GATTAATGATTATGTGAATTGATG	139	-
GAAATGTTAAAGATATGAAAGTAG	140	-
TATTGTTGATTTGATATTAGTGTG	141	-
TTTATGTTTGTGTATGTAAGTAGT	142	-
AATTGAAAGAATTGTGTGAATTGA	143	-
TGAGTTTGAATTTGTTTGAGTAAT	144	-
GATGTATAATGATGTGTGTAAATT	145	-
ATGTGAGAGAAGAATTTGTTTATT	146	-
GTGATAAAGTATTGTTGATAGAAA	147	-
GAAGTAGAATAGAAAGTTAATAGA	148	-
TTGTGTAGTTAAGAGTTGTTTAAT	149	24
TAGTAGTAAGTTGTTAGAATAGTT	150	-
AATTTGAAGTATAATGAATGTGTG	151	-
TAGAAATTGTAGTATTTGAGAGAA	152	-
TGTATATGTTAATGAGATGTTGTA	153	25
TATTTGATAAGAGAATGAAGAAGT	154	26
TTGAATAGTGTAATGAATATGATG	155	-
GTAGTTTGTGAATAGAATTAGTTT	156	-
AAAGATGATTGTAATTTGTGTGAA	157	-
GAAGATTGTTGAGTTAATAGATAA	158	-
AGATTATGTAGTGATGTAAATGTT	159	-
GAATTTAGATGTAGATATGAATGT	160	-
GATAGAAGTGTATTAAGTAAGTTA	161	-
TATGAATTATGAGAAGAATAGAGT	162	-
TTTGTTATGAAGTGATTTGTTTGT	163	-
GTAAAGATTGTGTTATATGAAATG	164	-
TTGTGATAGTAGTTAGATATTTGT	165	28
GAATTAAGATAAAGAAGAGAAGTA	166	-
GATTGTAGAATGAATTTGTAGTAT	167	-
AAATAAGAGAGAGAATGATTTAGT	168	-
AATTATGTGAATAGATTGTTGAAG	169	-
TTAAGATTTATGTGATAGTAGAGT	170	-
TTAAAGATAGTGTTTGTTGTGTTA	171	-
TATTGATTTATGAAGAGTATAGTG	172	-
AAATTTGATGAGTAGTTTAAGAGA	173	-
ATAAAGTTGTTTGATGTTTGAATG	174	-
GATTGTGATGAATAATGTTATTGA	175	-
GATGAAGAAATATGATATGAATAG	176	-

Table I

Sequence	SEQ ID NO:	No. in Ex 1
TTAAAGTTATTGAAGTGAAGTTGA	177	-
TTGTAAGAAATAGAGATTTGTGTT	178	-
GAGATTGAGTTTAAGTATTAGATT	179	-
AGTGATAATAGAATGATAAATGTG	180	-
GATAATAGTGAATTTGAGTTGTAT	181	-
AGATATTTGTAGTAGAAAGTATGT	182	-
GTTATGAATGTTGAATTTGAATGT	183	-
ATGAAAGATTTAGTTGTGAGATAT	184	30
AAATAGAGAAGTTATGATGTGATA	185	-
TTAGTGAGAAATGTTTAATGTGAT	186	-
TGAAGAATATGTGAAATTAGTTTG	187	-
GTTTGATAGTTTAATGAGTATTGA	188	-
GTTGTAAGTAATGATAAAGTATGA	189	-
TAAGAGTAGTAATTGTTGTTTAGA	190	-
TTTGAGAGAGTATGTATGATTATT	191	-
ATTGATTGTGAATTAGATAGAAGA	192	-
GATTAGTATTTAGTAGTAATAGAG	193	31
TATGTATTAGAGATATTGAAAGTG	194	-
TATGTGAAAAGTAATGATAAATGAG	195	-
GTAATTAGTAATGATTTGAAATGAG	196	-
GTTTATTGTAAAGATGTAAGTGAA	197	-
TAGTAGAATTGTTGTTAAAGAATG	198	32
TATTGTTAGTTATGTAGTGTGTAA	199	-
GAGTGAAAAGTTATATGAAAGTATA	200	-
ATATAGAAGTTGATGAGTTTATGA	201	-
TTTAGAAGTAAGAATAAGTGAGTA	202	-
TGTGTATAAGATATTTGTAAGAAG	203	-
TAGAAGAGTTGTATTGTTATAAGT	204	-
GTGTTATTAGTTTTAAAGTTAGAGTA	205	-
AATATAGTGATGTGTGAAAATTGAAATG	206	-
TTAGAGAAATAGAGAGTGATTATAGTT	207	-
GAAGTGAGTTAATGATTTGTAAAAT	208	-
AATGTAAAAGTAAAAGAAAGTGATGA	209	33
GTTAGTTATGATGAATATTTGTGTA	210	34
AAATGAGTTAGAGTAGAATTATGT	211	-
GATATAGAAGATTAGTTAGTGATA	212	-
ATAGTTTGTTGAGATTTATGAGTA	213	-
TAGAATAGTTAGTAGTAAGAGTAT	214	-
GAATTTGTATTGTGAAGTTTAGTA	215	-
GTAGTAAGAAGAGAATTAGATTAA	216	-
AATGTGTTATGTATGTAAATAGTG	217	-
GAATTAGTTAGAGTAAATTGTTTG	218	-
GAAATTGAAGATAGTAAGAAATGA	219	-
GTGTATTATGTGATTTATGATAGA	220	-
TATTATGAGAAAGTTGAATAGTAG	221	35
TATGTATTGTATTGAGTAGATGAA	222	-
GTGATTGAAATAGTAGATTGTTTAA	223	36
AGTAAGTTGTTTGATTGAAATTTG	224	-
GAAGTTTGATTTAAGTTTAAGAAG	225	-
GAGAAGATAAATGATATTGTTATG	226	-
ATGATGAGTTGTTAATAGTTAGTT	227	-
TATGATATTTGAAGAGTGTTAAGA	228	-
GAGATGATTAAAGTGATTTATGAA	229	-
ATAGTTAAGAGTGATGAGAATAAA	230	-
TTTATTGTTAGATAAAGAGTTGAG	231	-
AGAATATTGATAGTTGAAGTTGAA	232	-
TAGTGTAAAGTGTAGATTGTAAAT	233	-

Table I

Sequence	SEQ ID NO:	No. in Ex 1
AGTAGTGATATGATTTGAATATTG	234	-
TGTATTGAATTAGAATAGTGAGAA	235	-
TGATATGAGATAGAAGTTTAATGT	236	-
GAAGAAGTAAGTATAAAGTAAATG	237	-
TTTAAGTGTGATAAGAAAGATAGA	238	-
TATTGTTGAATGTGTTTAAAGAGA	239	38
GAATAATGATGAGATGATTATTGA	240	-
TAGAGAAAGAGAGAATTGTATTAA	241	39
ATGTATAATGAGATATGTTTGTGA	242	-
AATAGATAAGATTGATTGTGTTTG	243	40
TTTGATGATAATAGAAGAGAATGA	244	-
AGATGAATAAGTTGTGAATGTTTA	245	-
AGATGAAAAGAAAAGTGTAGAATATT	246	-
TGTTAAATGTATGTAGTAATTGAG	247	41
TAGTAGTGTGAAAGTTATTTGTTAT	248	-
AGTGAATGTTTTGTAAAAGAGTTTAA	249	-
GATAAATGAGAATTGAGTAATTGT	250	-
TGATGAGAAAATTGTTTAAGTGTTT	251	-
AAATAAGTAGTGTGAGTAATAGTA	252	-
TATGAAATATATGTGATAGTAAGAGA	253	-
ATTGTAAGAGTGATTATAGATGAT	254	-
AGAGTAAGAAATGAAAAGAGATAATA	255	-
TAAGTAAGTAGATGTTAAAAGAGAT	256	-
AAATAGAAAAGAATTGTAGAGTAGT	257	-
ATAGATTTAAGTGAAGAGAGTTATT	258	42
GAATGTTTGTAAATGTATAGATAG	259	43
AAATAGAATGAGTAGTGAAAATATG	260	-
TTGAATTATGTAGAGAAAAGTAAAAG	261	-
TAGTAAAATTGAGAGTAGTTGAATT	262	-
TGTAAAAGTGTTTATAGTGTGTAAAT	263	-
ATATGATTTGAGATGAGAAATGTAAA	264	-
AATATTGATATGTGTTGTGAAGTA	265	-
AGTGAGATTATGAGTATTGATTTA	266	44
TTGTATTTAGATAGTGAGATTATTG	267	-
ATAGAAATGAAAAGATAGATAGAAAG	268	-
GATTGTATATGTAAAAGTAGTTTAG	269	-
TATGAATGTTATTGTGTGTTGATT	270	45
GATATTAGTAGAGTAAGTATATTTG	271	-
TGAGATGAAATTTGTGTTATGATAT	272	-
TATGAATGAAAGTAAAAGAGATGTAAA	273	-
GAGTGAAATTTGTTGTAATTTGTTTT	274	-
AGAAATTGTAGAGTTAATTGTGTA	275	-
GTGTTAATGAAAAGTTGTGAATAAAT	276	-
TGTGATTTGTTAAGAAGATTAATG	277	-
AGTAGTATTGTAAAAGTATAAAGAG	278	-
TGATTGTTGTATAGTTATTGTGTA	279	-
GATTGTAGTTTAATGTTAAGAATG	280	-
ATGAAATAAGAAATTGAGTAGAGA	281	-
TATGATGATATTTGTTGTATGTGT	282	-
TTTAGAGTTTGATTAGTATGTTTG	283	-
AATAAGAGATTGTGATGAGAAATA	284	-
AATGAATAGAATAGAGAATGTAGA	285	-
GTAGTAGTAATTTGAATGTTTGAA	286	47
AGTGAGTAATTGATTGATTGTTAA	287	-
GAATAATGTTTAGTGTGTTTGAAA	288	-
ATATGAAAAGTAGAGAAAGTGTTAT	289	-
TGAGTTATTGTATTTAGTTTGAAG	290	-

Table I

Sequence	SEQ ID NO:	No. in Ex 1
TAGTTGAGTTTAAAGTTGAAAGAA	291	-
TAAAGAGTGATGTAAATAGAAGTT	292	-
TGTAGTGTTTAGAGTAAGTTATTA	293	-
AGAGATTAATGTGTTGAAAGATTA	294	-
GTAATAAGTTGTGAAAGAAGATTA	295	-
GAGATGTTATAGATAATGAAAGAA	296	-
TTTAGTTGATTGTTGAATAGAGTA	297	-
ATTATTGAAAGTAGATGTTAGATG	298	-
TTTATGTGTGATTGAGTGTTTAAT	299	-
TATTTAGTTAGATAGATAGAGAGT	300	-
ATGTGTTTATGTGAAAGATTTGTA	301	-
ATAGTAATTAGAAGAGAAGAATGT	302	-
TATGAGTGATTAGAATTGTATTTG	303	-
TTAATGTATTGTTTAAAGAGTGTG	304	-
ATAGAGAATTAAGAATTGTTTGAG	305	-
GTTATAAGTAGAAATGTATAGAAG	306	-
AGTAATTAGTTTGAAATGTGTAGT	307	-
GAAAGATTATGATTGTAAAGTGAT	308	-
GTAAGATTAGAAGTTAATGAAGAA	309	48
GAGAATGTTGAATAAGAAGTAATT	310	-
TTAAGAGTGTTTGAATAGTGTTTA	311	-
ATAAAGAAAGAGTATGAGATTATG	312	-
AGTTATTGATTGAAGATGAGAAAT	313	-
GTTTGTGTTTGTATAAGTTGTTAA	314	50
TTGTATGTGAGTTTAGATTAATGA	315	-
TAGTTAAAGTATAGTTGTTTGAGT	316	-
AAATTTGTGTTGAGATTTGTATAG	317	-
TATTAGTGTTATGATAAAGAGAAG	318	-
TATAAGAAGTAATTTGAGAAGAGT	319	-
TAAGTTGAGATGTTTGTTTGATAA	320	-
GTGTAGATTTATGAATTGAGTAAT	321	-
TATAGAGAAGTGTTTAGTTGTATA	322	-
ATAAAGAAGAATAGTTGTTGTGTA	323	-
AGATTGAAATAGATTAGAAAGTTG	324	-
GTTGTTATAAGAAATAGTTTGTTG	325	-
AGAAATAGAGTAAGAGTGTTTAAA	326	-
AGAGATAGTAGTAAATAGTTATTG	327	-
AAATGATTGTGTAAGTTATGTATG	328	-
AAGAAAGTAAGAGAGAAATTTGAAT	329	-
GTGTGTATTTAGTTGATAATTGAT	330	-
ATTGTTGTTGTTGAGAAATGTATT	331	-
AGATAAGTTAAAGTAAAGAGAATG	332	-
TAGTTGAAGTTAGTTTAAGTGTTA	333	-
AGTAAGAATGTAATATGATGATAG	334	-
ATGAGATTGAAAGATTTATGAATG	335	-
TGATTGAATTAGAGAGAATGTATA	336	-
AGTTAGTAAGAGAATATAGTGAAT	337	-
ATTAAGATTGTATAGTTAGTGATG	338	-
GAGATAAAGAATTGAAATAGAAGA	339	-
AGAGTAAATGTTAAGAAAGAAGTT	340	-
AAAGTTTGTTATGTGTGAAGAATT	341	-
ATTGTGTTTAAGAAATATGATGAG	342	-
TATTGAAATGAGATGTATGTAGTT	343	-
ATTTGTGTGATGTTTGAAATATGA	344	-
TAAGATAATAGTGAGAGAAATTGA	345	-
ATTTATGATTAGTGTAAGTGTTGT	346	-
GATTAAGAATAAAGTGTGAAGAAT	347	-

Table I

Sequence	SEQ ID NO:	No. in Ex 1
GTAATTGATGAAGAGTTAGTTTAT	348	-
TGTGTTATGTTATAAGAAGTGATA	349	-
AGAGAAATTGAATTTAGAAATGTG	350	-
TTATTGAATGTGAGAAAGTATTTG	351	-
TGTTAATGAGAAGATAATGATAGT	352	-
GAAAGTATTTGTTGATTATTGTTG	353	-
TAGTTTATGTAGTTAATTGTTGAG	354	-
GTTGAAAGATAGTTTGATATGTAT	355	-
TTAGAAGATAGATTATTGAGAAAG	356	-
AATAATGTTGTGAAATAGATGTGA	357	56
AGTAAGAAAGTTTAGTTTAGTTAG	358	-
TAGTTTAATGAGATGTTTGATATG	359	-
TTAAAGATGTTAAAGAATGAGTGA	360	-
AAAGTGTGTATATGTTAGAAAGTA	361	-
ATTAAGTTATGTGTTTATGTGTTG	362	-
TTTGAAGAAGTGTTTGTATTATGT	363	-
TGTTAAGAAGTTTAGTTAAAGTTG	364	-
TTTAAGTATAAGATTGTGTGAGAT	365	-
AGATATTTGATAGATAGAAGAAAG	366	-
ATTTAGAGTTGTAAGAAGATATTG	367	-
GAGAAATTGTAATTGTTAGAGTAT	368	-
GAAGTATATGTTAAGATGTAATAG	369	-
AATATTGAAGATGTAGTGAGTTAT	370	-
GAGTTTAGAAATGATAAAGAATTG	371	-
TAAGAAATGAGTTATATGTTGAGA	372	60
TTGATATAAGAAGTTGTGATAAGT	373	-
AAGTGTTTAATGTAAGAGAAATGAA	374	61
GTTGTGAGAATTAGAAATAGTATA	375	-
TTTAGTTTGATGTGTTTATGAGAT	376	-
GTAATTGAAAGTATGAGTAGTAAT	377	-
TAGTTGAATAAGATTGAGAGAAAT	378	-
TTAAGTGAAGTGTTGTTTATTGAA	379	-
ATTGATTTGTTGAAATAAGTGTTG	380	-
TGAATTGTTGATAAGTTATGAAGA	381	-
GTTTGTTATTGAGTAAGTTGAATT	382	-
TGATTTAGTATGTATTAGAGTTGA	383	-
TAAATAGAGATGAGAATAAGAAAG	384	-
AGAATGTTATATGTAGAGAAATTG	385	-
ATTTATGTAGTTGAGAGTGATAAA	386	-
GTAAA-GATAGTTTGAGTAATTTGA	387	-
GAAATAGTATAATGTTAAGTGAGA	388	-
ATTGTATATTGTGTTGAAGAAAGT	389	-
GAGTTAAGTGTAAATGAAATGTAA	390	-
ATAGATTGTGTGAAAGAAAGAATT	391	-
TTAATAGAAGTTTGTAGTATGATG	392	-
TTGTATGTGAGAATAAAGTTTAGT	393	-
GTGATTAGATATGATGATATGAAT	394	-
TGAAGAAGAATTTAGATTTGTAAG	395	-
TGTATGATTATTGATTAGTGTGTT	396	-
TGTGAAAGAGAATGATAGATATTT	397	-
AATTGAAATGAGTGTGTTTAAGAA	398	-
ATTATAGAGTTAGTTTAGAATGAG	399	-
AAAGATAGAAATTGAGTGTATGAT	400	-
GTAGTTTGTTAATGTTGTATAATG	401	-
AGAGATATTAGAATGTAAGAATAG	402	64
AGAAGTTTGAAATATGATAGAATG	403	-
TAGAATGTAAAGTTTAGTATAGAG	404	-

Table I

Sequence	SEQ ID NO:	No. in Ex 1
AGTAGATGTATGTTAATGTGAATA	405	-
TGAAAGTGAAATATGAAATGTTGT	406	-
ATAGTATATTGAGTTTGTATGAAG	407	-
GAAGAAATGTTTGTAGAATAAGTA	408	-
AATGAGTATTGAAGAAATGTATAG	409	-
GTGATAGAATTTGTGTTTAATGAA	410	66
TGTAGTATGAAGAATAATGAAATG	411	-
ATAGAAGTTAATGATAATTGTGTG	412	-
GTGATTGTAAGTAAGTAAAGATAA	413	-
TATGTAGTTTGTGTTATTTGAAGA	414	-
TGAGTAAGTTTGTATGTTTAAGTA	415	67
TAAATGTATGAGTGTGTAAAGAAA	416	-
GTAAGAGTATTGAAATTAGTAAGA	417	68
GTTGAGTGTAAAGATTATTGATAA	418	-
AGTATGAGTTATTAGATAAAGTGA	419	-
ATTTGTTATAGAGTTGTGTTGTAT	420	-
TAATTAGTAGTGTGTTGAAATTTG	421	-
TGTATTGAGATTGTTATTGTATTG	422	-
GTTATTAGAAGAGATAATTGAGTT	423	-
TTGAGTTGTGATTAAGTAGTATAT	424	-
GATAGTATAATGATTGAAGTAATG	425	-
GTGAAAGATATTTGAGAGATAAAT	426	-
AGTTATGATTTGAAGAAATTGTTG	427	-
GTAAGTATTTGAATTTGATGAGTT	428	-
TAATAGTGTTATAAGTGAAAGAGT	429	-
AAATGAATTGATGTGTATATGAAG	430	-
AGAAAGTGAGTTGTTAAGTATTTA	431	-
TTTATGTGTGAATTGTGTATATAG	432	-
GTAATATGATAGAAAATGTAAAGAG	433	-
GAGAATTGTTTAAAGATAGTTGTA	434	-
GAATTTGTTAAGAATGAGTTTGAT	435	-
ATAGTGATGATTAAAGAGAATTTG	436	-
ATAGATGTTTAGTTGAGATTATTG	437	-
AAGAGTGTAAATAGAAAGTGATAT	438	-
TGTGTATTGATTGTTGAGATAAAT	439	-
TAGTATAGTGAGAAAGAGTTAAAT	440	-
AAAGATAAGAAAGAGATGATGTTT	441	-
GAAGTTATTGAAATAGAGAAGTAT	442	-
ATGTATGTATAGAAAGAGTAAATG	443	-
GATGTTGTAAAGATTGAAATTGA	444	-
AATTTAGAGAGTATTTGTGTTGTA	445	-
AATTTGTTTGAAAGAAAGTAAGTG	446	-
AAAGAGTAGTGTTATTGTTAGATA	447	-
GTATGTTGTATATGTTGTTGATAT	448	-
GTAGAATTTGTTGAGTATTTGTAA	449	-
ATGAATTTAGTTAGTGTAAGAAAG	450	-
ATGATAAGAAATGTTGATGAAGTA	451	-
TTGATGATGAAGATAATGTAGATA	452	-
AGATGATATGATATAGATTAGATG	453	-
TTGAAAGTTAGAAAGATAGATGTT	454	-
GTTTAATGTTAGTTAGAAAGTAAG	455	-
GAGATTTAAGTTTGAAGTGAAATA	456	-
TTTGTTAGTAGTTGTTATAAGAGA	457	-
TATGAGAATAGTTTGTTAGTGAAT	458	-
TTGAAAGTTTAAAGAAGAGATAAG	459	-
AAGTGAGTTTGAAAATGAAATATGTT	460	-
GTTAGAAATGAAATGAGTAGTTAT	461	-

Table I.

Sequence	SEQ ID NO:	No. in Ex 1
TAAGTATTGTATTTGTGTGTGTAT	462	-
TGTATTAGTAAAGAAGAGAGAATA	463	-
GAGAAGAGAAATAAGTTGAAATAA	464	-
GTAAAGTAGAAATAGAATTGAGTT	465	-
GTGTGTTATTTGTTTGTAAAGTAT	466	69
TTTGATGTATGAATATAGTATGAG	467	-
AAGATTGTGTGAATAGTTGAAATT	468	-
TATAAAGTTTGAAGATGAGTGATA	469	-
AGATAAAGAGATTTAAGATGTATG	470	71
GAAGAATTAAGTTGAGAATTAAGA	471	-
TAGAGAAATTTGATAAAGAAAGAG	472	-
AAAGTTTATGAAGTTATTGAGTAG	473	-
AAATAGTGTAAGTAAAGAGATGAT	474	-
TATGATGATTTAGTTATAAGAGTG	475	-
TAGATAAAATGTTATGATGAGTAAG	476	-
AGATTGATTGTGATGATTTGTATA	477	-
TTAAGAAGAATTGTATATGAGAGT	478	73
GTAGAATGTTTAGAGTTGAAATATA	479	-
GAGAAATAGTAAGAAGTAAATAGA	480	-
ATTGAAGTTGTTATGTGAAGATTT	481	-
TAAATGTTGTGTAGAGTAATTAGA	482	-
AAATAAGAGTTTGAGAAGTTGTTT	483	-
AGTTGTAATAAGAAGTGATTTAAG	484	-
GTTAGAATGTATATAGAGTTAGAT	485	74
TTGATATTGAAAGAGAAAGTTATG	486	-
TTAAAGAGAGAAATGTTTGATTAG	487	-
TGTGAATTTGAGTATTAGTAAGAA	488	-
TAATTTGAATGTGAAAAGTTGTTAG	489	-
ATGTGTTTGAAAAGATGATGATTTA	490	-
AAGTTATGTTGATATTGAGTGAAA	491	-
TAGATAAAGAAGATAGAGATTTAG	492	-
GATGAATGTAGATATATGTAATGA	493	-
GAAGAATAGTTTATGTAAATGATG	494	-
GTAGTATATAGTTAAAGATGAGTT	495	-
GTTATTTGTGTATGATTATGATTG	496	-
AGAGATTAGAAATTGAGAGAATTA	497	-
GTATGATAGAGTTTATAGTGATAA	498	-
GTTAGAAAGAATGAAATTGAAGTA	499	-
AAGAATGAGAATATAGAGATGAAT	500	-
AAAGAGAATAGTGTTTAAGAAGAT	501	-
GATGTGTTATTGATAGAAATTAGA	502	-
TAGAGTTATAGAGATATTGTATGA	503	-
GAGAGTTGAATAAGTTAAAGATAT	504	-
AGATATGAAATAGATTGTTAGAGA	505	-
GAGTGAATAGAAAGATATGTTAAT	506	-
AAAGAGATATTGAAGAGAATAAAG	507	-
GTTATAGAATAAGTTGTAAAGTGT	508	-
TGATAGTATGATAATGTGTTTATG	509	-
TTTGTTGTTAAGTATGTGATTTAG	510	77
TAAAGTGTTGTGTTAAAGATTAAG	511	-
TGTGTTTGATTGATTAATGTTATG	512	-
ATTAATGAATGAGTGTTGTAATGT	513	-
TAGATGTTTGTGAGTTTGATATTA	514	-
GAATGAATAGTAATAGATGATTTG	515	-
AATAGTGTGTTGTTATATGATTAG	516	-
TAGATTAGAAGATGTTGTGTATTA	517	-
AATGTGTGTGTTAAATGAATTTGT	518	-

Table I

Sequence	SEQ ID NO:	No. in Ex 1
GAATTAAGTATATGAGTGTAGAAA	519	-
TTATTGTGTGTAAGTAGTGTAAAAT	520	-
GTAGTAAAGAGAAATTGTTTAGTAT	521	80
AAGTTTGTAAGAAGTAGTTGAATA	522	-
AGTTATAGTATAGTAGTATAGAGA	523	-
GAAAGAAATGTGTATAGTTTAATG	524	-
TTGTGAGTAATGAATGATGTATTA	525	-
GTAGAGTTGTAAATAGAGAATAAA	526	-
ATTAATGTAGATTGTAAGAGATAG	527	-
TTAGTGTGTTTGTAGATAGAATTA	528	-
AGAGAGTTTGTGTATATGTATAAA	529	81
TTAAGTTTAGTGAGATTTGTTAAG	530	-
ATGAAGTTTATTGAATAGTAGTGA	531	-
ATATTTGTGTTTGTATGTTTGTGAA	532	-
AAAGTGTTTATAGAAGATTTGATG	533	-
AAGAGATATGATTTGTTAGTTGTA	534	-
AAGAAGAAATGAGTGATAATGTAA	535	-
TAGTGTTTGATATGTTAAGAAGTT	536	-
GTAGAAAGTGATAGATTAGTAATA	537	-
GATAAATGTTAAGTTAGTATGATG	538	-
AGATTAGAAGAATTGTTTAGAATG	539	-
ATATTTGAGAAGTGTGAAATGAAT	540	-
TGAGTAAATAGTTTATGAGTAGTA	541	-
TTAGAGAGTAGATAAAGATTTGAT	542	-
ATTGTTTAAGTTGTTGATAAGATG	543	-
GTTGTAAAGTTAAAGTGTGAATTT	544	-
ATAGATTGTGTGTTTGTTATAGTA	545	-
GTAAGTTATTGAGAATGATAATAG	546	-
TAGATTAGTTGATAAGTGTGTAAT	547	83
AAATGTAAATGAAGAGTGTTTGTT	548	-
GATAGAAGAAATGTATATAGTGAT	549	-
TATAGAGTGTATGTTATGATAAAG	550	-
TATGAAGTGATAAGATGAAGAATT	551	-
TGTTGAGAATAGTAAGAGAATTTA	552	-
TAGATAATGTGAAGTAATAAGTGA	553	84
GTATTATGATGATAGTAGTAAGTA	554	-
AGATATGATTTAGTATTGAATGTG	555	-
AATTAAGTTTGTAGAGTGATTTGA	556	-
AAGAAATAGATGTAGTAAGATGTT	557	-
TTGAGAAGTTGTTGTAATAAGAAT	558	-
AGTGTGAAAATAGTGAAAGTTTAAA	559	-
TTTATGTAGTAGATTTATGTGAAG	560	-
ATTAATGAGAAATTAGTGTGTTAG	561	-
ATGTTAATAGTGATAGTAAAGTGA	562	-
TATGTTGATAAATGATTATGAGTG	563	-
TTATTAGAGTTGTGTGTGATATAT	564	-
TGTTGTTATGATTGAGTTAGAATA	565	-
AATTTGAGTTAAGAAGAAGTGTAA	566	-
AAAGATAAAGTTAAGTGTTTGTAG	567	88
TGTTGAGATGATATTGTATAAGTT	568	-
TAAATAGTGAATGAGTTATAGAGT	569	-
ATAGATGTTATGATAGTTAGTTAG	570	-
GTTAAGTGAAGATATGTATTGTTA	571	-
TAAGAAAGTAAAGTTTGTAGATGT	572	-
AAGAGAAAGTTTGATTGAATAAAG	573	-
ATATTAGATGTGAGTTATATGTGT	574	-
AGTTTGAGTTTAGTATTGTGAATA	575	-

Table I

Sequence	SEQ ID NO:	No. in Ex 1
ATGTTAAATGAGAGATTGTGTATA	576	-
TAAATGTTGTGATTATTGTGAGAT	577	-
TAAGAATTGAAGTAAGAGTTATTG	578	-
AGAGATAGAATTAAGTTTGTTGAT	579	-
GAAGAATGTTAAGAAATATGTAAG	580	-
TATTTGTGATTAAGAAGTTGAGAA	581	-
AGTTAGAATTTGTGTAGTAGAATT	582	-
AAGTTTATTGTTGATGTTGTATTG	583	-
GAATGAGTTTAAGAGTTTATAGTA	584	-
AGTGAAGATTGTATGTAGTATAAA	585	-
AGTTGAAATGAGTATTAAGTAATG	586	-
ATGTGTTATTTGAGATGAGTAATT	587	-
AAATAGTGTTGTTGAAGTTGTTAT	588	-
GTAGAGAAAGATATATGTAGTTTA	589	-
GAGAGTATTTGATGAATGATTATA	590	-
GAGTATAAGTTTAGTGTATATTGA	591	-
ATAATGTGATTATTGATTGAGAGA	592	-
TTAGTTGTTATGTGAGAGTAATAA	593	-
AAATGAGTATATTGAATTGTGATG	594	-
AATTAGAAGTAAGTAGAGTTTAAG	595	3
TGTAAGTTTAAAGTAAGAAATGTG	596	5
GAAATGATAAGTTGATATAAGAAG	597	-
AATGAGTAGTTTGTATTTGAGTTT	598	-
AGTGAATGTAAGATTATGTATTTG	599	6
GTAATTGAATTGAAAGATAAGTGT	600	8
TATGTTTAAGTAGTGAAATAGAGT	601	-
GTATTGAAATTGAATTAGAAGTAG	602	-
AATATGTAATGTAGTTGAAAGTGA	603	-
TGAATATTGAGAATTATGAGAGTT	604	-
TAGTGTAAATGATGAAGAAAGTAT	605	-
GTATGTGTAAAGAAATTTGATGTA	606	-
AATTGTTTGAAAGTTTGTTGAGAA	607	-
AATTGTTTGAGTAGTATTAGTAGT	608	-
TAATTGAGTTTGAATAAGAGAGTT	609	-
TGTTGATTGTAAGTGTTTATTGTT	610	-
GAAATTTGTGAGTATGTATTTGAA	611	-
TAAGAATGAATGTGAAGTGAATAT	612	-
TAATGTGAAGTTTGTGAAAGATAT	613	-
TTGTATATGAAAGTAAGAAGAAGT	614	-
TAGAGAGAAGAAGAAATAAGAATA	615	-
ATTTGAAATGTTAATGAGAGAGAT	616	-
TTGTGTGTATATAGTATTAGAATG	617	-
ATTGTTAGTATTGATGTGAAGTTA	618	-
TGTTTGTATTTGAATGAAATGAAG	619	-
TGTTAGATTGTGTTAAATGTAGTT	620	-
TATAGAGTATTGTATAGAGAGAAA	621	-
AAATAGTAAGAATGTAGTTGTTGA	622	-
TGAGTGTGATTTATGATTAAGTTA	623	-
AGAATTTGTTGTAGTGTTATGATT	624	-
GATTGAAGAAAGAAATAGTTTGAA	625	-
GATAATAGAGAATAGTAGAGTTAA	626	-
GATTGAAATTTGTAGTTATAGTGA	627	-
GATTTAAGAAGATGAATAATGTAG	628	-
TTTGAGAGAAAGTAGAATAAGATA	629	-
GATTAAGAGTAAATGAGTATAAGA	630	-
TTTGATAGAATTGAAATTTGAGAG	631	-
TGAAGAAGAGTGTTATAAGATTTA	632	-

Table I

Sequence	SEQ ID NO:	No. in Ex 1
GTGAAATGATTTAGAGTAATAAGT	633	-
AAATAAGAATAGAGAGAGAAAGTT	634	9
GTTGTAAAGTAATAGAGAAATTAG	635	-
AGTGATTTAGATTATGTGATGATT	636	-
AGAGTATAGTTTAGATTTATGTAG	637	-
ATGATTAGATAGTGAAATTGTTAG	638	-
ATGAAATGTATTAGTTTAGAGTTG	639	-
ATATTGAGTGAGAGTTATTGTTAA	640	-
AGATGTGTATTGAATTAAGAAGTT	641	-
TAATGTGTTTGATAGAATAGAGATA	642	-
AAATTAGTTGAAAGTATGAGAAAG	643	11
TTTAGAGTTGAAGAAATGTTAATG	644	-
GATTGTTGATTATTGATGAATTTG	645	-
TGTTGTTGTTGAATTGAAGAATTA	646	-
ATTAAGTAAGAATTGAGAGTTTGA	647	12
GTATGTTGTAATGTATTAAGAAAG	648	15
TAGTTGTGATTTATGTAATGATTG	649	-
TGATAATGAAAGTTTATAGAGAGA	650	-
GTAAGATTGTTTGTATGATAAGAT	651	-
TTGAATTAAGAGTAAGATGTTTAG	652	-
AAGTGTTTGTTTAGAGTAAAGATA	653	-
AGAGAGATAAAGTATAGAAGTTAA	654	-
ATTATGAATAGTTAGAAAGAGAGT	655	-
TTGTTGATATTAGAGAATGTGTTT	656	-
TTTATTGAGAGTTTGTTATTTGTG	657	-
AGTGTTAAGAAGTTGATTATTGAT	658	-
GAGAAATGATTGAATGTTGATAAT	659	-
GATAAGTATTAGTATGAGTGTAAT	660	-
TTTGATTTAAGAGTGTTGAATGTA	661	16
AAGTTAGTAAATAGAGTAGAAAGA	662	-
GTAAAGTATGAATATGTGAAATGT	663	-
TAATAAGTGTGTTGTGAATGTAAT	664	-
AAAGATTTAGAGTAGAAAGAGAAT	665	-
TTAGTTTGAGTTGAAATAGTAAAG	666	-
TAATAGTATGAGTAAGATTGAAAG	667	-
GAAGATTAGATTGATGTTAGTTAA	668	-
TAAAGAGAGAAGTTAGTAATAGAA	669	-
TAAGTATGAGAAAATGATGTGTTAT	670	-
GAGTTTGTTTGTTAGTTATTGATA	671	-
AAGTAAAGAAATGTTAAGAGTAGT	672	-
ATGAGAATTGTTGTTGAAATGTAA	673	-
TTAGATTAGAGTAGTAGAAGAATA	674	-
TAGTGATGAAGAAGTTAGAAATTA	675	-
TAATGTAGTAATGTGATGATAAGT	676	-
TTGAGAAAGAATAAGTAGTGTAAA	677	-
TAATGAGTGAGATTATAGATTGTT	678	-
GTATAAGAAATGTGTGTTTGATTA	679	-
GTGAATGTGTTAATGAAGATATAT	680	-
GAAAGTTATTAGTAGTTAAAGATG	681	-
TAGAATTGTGTTTGATAAGTGATA	682	-
TGATTTAGATTGAGAGTTAAAATGA	683	-
ATTATTGAGTTTGAATGTTGATAG	684	-
ATAGTAGTTATGTTTGATTTAGTG	685	-
ATAGAAGAAGAATAAAGTTAGAGA	686	-
GATGTTGAAAGTAATGAATTTGTA	687	-
GAGATTGATAGTAGAAATGATAAA	688	-
TGAGAGAATAAAGTATGAATTTGA	689	-

Table I

Sequence	SEQ ID NO:	No. in Ex 1
TATAAAGATGATGTGAATTAGTAG	690	-
TTATGTAAGAATGTTTGAGAGAAA	691	-
AGTAAATGATGAATGATATGATGA	692	-
GAAATTTGTGTTAAAGTTGAATGA	693	-
GATGAATGATTGTGTTTAAGTATA	694	-
GAAATAAGTGAGAGTTAATGAAAT	695	-
TGTTGAAATAGTTATTAGTTTGTG	696	-
TTTGAGAGTATATTGATATGAGAA	697	-
ATTGTGTGTAAAGTAAGATTTAAG	698	-
TATAGTTTGAAGTGTGATGTATTT	699	-
GTGAAGTTATAGTGTATAAAGAAT	700	-
GTATGTTGAATAGTAAAATAGATTG	701	-
TTAGAAAGTGTGATTTGTGTATTT	702	-
TTTAGTAATATGTAAGAGATGTGA	703	-
AGTATGTATAGATGATGTTTGTTT	704	-
ATTTAAGTAAAGTGTAGAGATAAG	705	20
ATTTGTGTTGAATTGTAAAGTGAA	706	-
ATGTTATTAGATTGTGATGAATGA	707	-
TAGTAGTAGAATATGAAATTAGAG	708	-
TTTAATGAGAAGAGTTAGAGTATA	709	-
AAAGTTTAGTAGAGTGTATGTAAA	710	-
ATATATGATAGTAGAGTAGATTAG	711	-
TGAGAAGTTAATTGTATAGATTGA	712	-
TATAGAGATGTTATATGAAGTTGT	713	-
AAATTTGTTAAGTTGTTGTTGTTG	714	-
TTGTTGAAGATGAAAAGTAGAATTA	715	-
AAGAGATAAGTAGTGTTTATGTTT	716	-
AATAAGAAGAAGTGAAAGATTGAT	717	-
TAAGTTAAAAGTTGATGATTGATAG	718	-
ATATAAGATAAGAGTGTAAGTGAT	719	-
GTTAAATGTTGTTGTTTAAGTGAT	720	-
GAGTTAAGTTATTAGTTAAGAAGT	721	-
TATTAGAGTTTGAGAATAAGTAGT	722	21
TAATGTTGTTATGTGTTAGATGTT	723	-
GAAAGTTGATAGAATGTAATGTTT	724	-
TGATAGATGAATTGATTGATTAGT	725	-
ATGATAGAGTAAAGAATAAGTTGT	726	-
AGTAAGTGTTAGATAGTATTGAAT	727	27
ATGTAGATTAAAGTAGTGTATGTT	728	-
TTATT-GATAATGAGAGAGTTAAAG	729	-
ATTTGTTATGATAAATGTGTAGTG	730	29
TTGAAGAAATAAGAGTAATAAGAG	731	-
TGTGTAATAAGTAGTAAGATTAGA	732	-
ATGAAAGTTAGAGTTTATGATAAG	733	37
ATTAGTTAAGAGAGTTTGTAGATT	734	-
TGTAGTATTGTATGATTAAAGTGT	735	-
AGTTGATAAAGAAGAAGAGTATAT	736	-
GTAATGAGATAAAGAGAGATAATT	737	-
TGTGTTGAAGATAAAGTTTATGAT	738	-
AAGAAGAGTAGTTAGAATTGATTA	739	-
GAATGAAGATGAAGTTTGTTAATA	740	-
AAATTGTTGAGATAAGATAGTGAT	741	-
TGATTGTTTAATGATGTGTGATTA	742	-
ATGAAGTATTGTTGAGTGATTTAA	743	-
GTGTAAATGTTTGAGATGTATATT	744	46
AATTGATGAGTTTAAAGAGTTGAT	745	-
TTTGTGTAATATGATTGAGAGTTT	746	-

Table I

Sequence	SEQ ID NO:	No. in Ex 1
GTAGTAGATGATTAAGAAGATAAA	747	-
TTTAATGTGAAATTTGTTGTGAGT	748	-
GTAAAGAATTAGATAAAGAGTGAT	749	-
AATAGTTAAGTTTAAGAGTTGTGT	750	-
GTGTGATGTTTATAGATTTGTTAT	751	-
GTATAGTGTGATTAGATTTGTAAA	752	49
GTTGTAAGAAAGATATGTAAGAAA	753	-
ATATTAGATTGTAAAGAGAGTGAA	754	-
GAGTGATATTGAAATTAGATTGTA	755	-
TAAGAAGTTAAAGAAGAGAGTTTA	756	-
GATGTTAGATAAAGTTTAAGTAGT	757	-
GTGATTGTATGAGAAATGTTAAAAT	758	-
TGATTATTGTAAGAAAGATTGAGA	759	-
AAGAATTGTGTAAGTTTATGAGTA	760	-
TTGTATTTAGAAGATTTGTAGATG	761	-
TATATGTTTGTGTAAGAAGAAATG	762	-
GATAATGTGTGAATTTGTGAATAA	763	-
TTAGAAATGTGAGATTTAAGAGTT	764	-
AGTGTAGAATTTGTATTTAGTTGT	765	-
TAGTTAAGATAGAGTAAATGATAG	766	-
GAAGTGATATTGTAAATTGATAAG	767	-
GTAATTGTGTTAGATTTAAGAAGT	768	-
TGATATTTGTGAATTGATAGTATG	769	-
AAGTAAAGAGATATAGTTAAGTTG	770	-
ATTAGTTAAGTTATTTGTGAGTGA	771	-
AGATGAAGTAGTTTATGAATTAGA	772	-
TGAGTTAGTTAAGTGATAGTTAAA	773	-
TTATTGTAGATTTAGAGAAGATGA	774	-
TATTTGTGTTTGTTGATTAGATAG	775	-
GTATAATGTGTGTGAAAGTTATAA	776	-
TATATGTTGAGTATAAAGAGAGAA	777	-
TTAGTTAGTTTAAAGATTGTGAGT	778	-
TTTAGAATAAGTGATGTGATGAAA	779	-
AGAGTAATGTGTAAATAGTTAGAT	780	-
TGTGATAAAGAGAAATTAGTTGTT	781	-
GAATTTAGTGAATGTTTGAGATTA	782	-
TGTGATGTGTAAGTATATGAAATT	783	-
TTGTGAATGATTAATGAATAGAAG	784	51
AATGTTGTTTAGATTGAGAAAGTT	785	-
AGATTGTGTTAGTATTAGTATAAG	786	-
TTGATGTATTAGAAAGTTTATGTG	787	-
TATGATTGTGTGTTAGAGAATTTA	788	-
TAGTGTAGATATTTGATAGTTATG	789	52
AGTTTAATGTGTTTAGTTGTTATG	790	-
TGTGTAAAGTAGAAAGTAAAGATT	791	-
GTTATGATATAGTGAGTTGTTATT	792	53
TTTGATTGAATGTTAATAGTGTGT	793	-
AGAGTATTAGTAGTTATTGTAAGT	794	54
TAAGTAGAAAGAAGAAGATATTTG	795	-
AGAAAGAGAATTATGTAATGAAAG	796	-
TTAGATTTGTTAGTGTGATTTAAG	797	-
GATGATTAAGATATAGAGATAGTT	798	-
ATATTTGAGTGATTAAGAGTAATG	799	-
TGTATTGTGAGTTAAGTATAAGTT	800	-
AATTTAGTAGAAAGTGTTGTGTTT	801	-
GTTAGAAGATTAAGTTGAATAATG	802	-
TAAAGTATGTGAGATGATTTATGT	803	-

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Sequence	SEQ ID NO:	No. in Ex 1
TGAAATGATTAAAGATGAAGATGA	804	-
TTATTAGATGTTGAGTGTTTGTTT	805	-
TAGTGTTTAAAGAGTAGTATATGA	806	-
AGTTATAAGTAAATGATGTTGATG	807	-
TTAAGAGAGAAATAAGTGTATTGT	808	-
GATATTGAAATGTGTAAATGATGA	809	-
ATGATGAATTAAGAAAGAAAGAGA	810	-
GAATAGTTTGATTTGTGTTTGTTA	811	-
AGTTGTTTAGATTTGATTTGTAAG	812	-
GTATGAGATTTGATATAAGATTAG	813	-
TTTATAGTGAGTATAGTGATGATT	814	-
TATATGTGAAGATATAAGTGTTTG	815	-
ATTGATAGATGATAGTAATTGAGT	816	-
TGATAGATGTGAAGAATTTGATTT	817	-
GAAGATATTGAAAGAATTTGATGT	818	55
GATGTTTAGTGTAGATATAGATTT	819	-
GAATATTGAGTTATAAGTAGTAGT	820	-
AGTGAGTAAGTAATAGAAAGATTT	821	-
GTAGAATAAGTAATTTGTGAGATA	822	-
GAGTTATTTGAGATTTAGATGTTT	823	-
GAAATGATGATTGAATTTAGAGAT	824	-
AAATAGTGTGAGAATAGTTAAGTA	825	-
ATGTGTTAAGTTGTAGAAGAATAA	826	-
ATAATGAGTTAATAGTGTAAGAAG	827	-
ATAAGAGATGTTTAAGTTAGAAAG	828	-
TGTTAGTGTTAGAAATATGAAAGA	829	-
TTTAGAAGATTGTTAGATAAGTTG	830	-
GTGTAATGTATAAGATAGTTAAGT	831	-
TATTAGAGAGAAATTGTAGAGATT	832	57
TAGTGAGATAAAGTAAAGTTTATG	833	-
TTGTGAAAGTTAAGTAAGTTAGTT	834	-
AAAGTGTAAGTTGAAGAATATTGA	835	-
GAATAGAGTGTTATTTGAAATAGA	836	-
TATAAGAGAGAGATAAGTAATAAG	837	-
TGAGTGAAATTGATAGAGTAAATT	838	-
GATGAATAAGTTTAAGTGAGAAAT	839	-
GTGTGATATGTTTATTGATTAAGT	840	-
TAAAGTGAGTGTAAATGATAATGA	841	-
GTAGAGTTTGATTTGAAAGAATAT	842	-
GAATATTGTTATGTTTGTTATGAG	843	-
GTGTAATAAGATGTATTGTTGTTT	844	-
TAAATTGATTGTGAGTTGAAGAAT	845	-
TGAGATAGTTATAGTTAAGTTTAG	846	-
AGTTTGTTAAGATTATGTAGAAAG	847	-
GAATGTGTAGAATAAGAGATTAAA	848	-
GTATTATGAAAGAAGTTGTTGTTT	849	-
GTGTTATAGAAGTTAAATGTTAAG	850	58
TTAAGAGTAGTGAATATGATAGTA	851	-
AATGTTATAAGATGAGAGTTTAGT	852	-
ATATAAGATTTGATGTAGTGTAGT	853	-
TATGTTTGTTGTTGTTAAGTTTGA	854	-
GATAGTTTAGTATAGAAGATAAAG	855	-
GTTGAATATAGAGATAGTAAATAG	856	-
AGAGAAGATTTAGTAAGAATGATA	857	-
TGAATGAGAAAGATATTGAGTATT	858	-
TGAAGATTATAGTAGTTGTATAGA	859	-
GATTAGTAGTATTGAAGATTATGT	860	-

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Sequence	SEQ ID NO:	No. in Ex 1
TGAAATGTGTATTTGTATGTTTAG	861	59
ATTAAAGTTGATATGAAAGAAGTG	862	-
AATGTAGAGATTGTAGTGAATATT	863	62
TTATTTGTTGAGTGTAAATGTGAT	864	-
ATGTAATTGTGAATAATGTATGTG	865	63
GATTTGTATAGAGATTAGTAAGTA	866	-
AATATTGTTGTTTAGAGAAAGAAG	867	-
ATGATGATGTATTTGTAAAGAGTA	868	-
AATGTATTTGTGTGATTGTGTAAA	869	-
AGTGTTATGAAGAATAGTAAGAAT	870	-
GTTATGTAGAGATGAAAGAAATTA	871	65
GTTTGTATTAGATAAATGAGTTGT	872	-
TGATTTATGAGATTAAGAGAAAGA	873	-
TTTGTGTGTTATTGTAATTGAGAT	874	70
GATGTGTGATATGATTAAAGAAAT	875	-
AGATTATAGATTTGTAGAGAAAGT	876	-
GAAGAGTATGTAATAGTATTGTAT	877	-
TTTGTAATGTTGTTGAGTTTAAGA	878	-
AGTAAATAGTAGTATGAATAAGAG	879	-
GAATGTTGAATTGAAATATGAGTT	880	-
AGTAGTTAATTGATAGTAAGTTTG	881	-
AGTGTAAAGAAATGAATGAATAAG	882	-
TGTTAGATATTTGTGAAATGTGAA	883	-
TGTATGTTGAGTTTGAATTGTTAT	884	-
TGAGTGAATTAGTTATGTTGTTAT	885	-
GAAGAAAGAAATGAGAAAGATTAT	886	-
TTAAGTAAGTTGTGTTGATATTAG	887	-
ATGATGTGTTTGATTTGAATTGAA	888	72
AAGTAAGTGAAATTGTTGTTTGAA	889	-
ATGAAGTGTAAAGTTTGAAAGAAA	890	-
AGAGAGTAAGATAATTGTATAGTA	891	-
TTTATGAGATAGATGAAATAAGTG	892	-
AGAAATTAGTAGTAATGATTTGTG	893	-
GATTTGAGATTGAATGAGAATATA	894	-
GATTAGAAAGATGAATAAAGATGA	895	-
TAGATAGAAAGTATATGTTGTAGT	896	-
GAAGATAGTAAAGTAAAGTAAGTT	897	-
AAATGTGTGTTTAGTAGTTGTAAA	898	75
TTGTTGAAGTAAGAGATGAATAAA	899	-
TATTTGAGAGAAAGAAAGAGTTTA	900	-
TATTTAGTGATGAATTTGTGATGT	901	-
TTATAGTGATGATGATAAGTTGAT	902	-
TAAAGATAATTGTAGAAAGTAGTG	903	-
GTTTAGTATTGATATTGTGTGTAA	904	-
GTGTTGTGAATAAGATTGAAATAT	905	-
AAAGAAAGTATAAAGTGAGATAGA	906	-
TATTTGTAAGAAGTGTAGATATTG	907	-
TAGAAGATGAAATTGTGATTTGTT	908	-
ATAATAGTAAGTGAATGATGAGAT	909	-
AATGTGAATAAGATAAAGTGTGTA	910	-
ATTGAAGATAAAGATGTTGTTTAG	911	-
TGAAATAGAAGTGAGATTATAGTA	912	76
AGTTATTGTGAAAGAGTTTATGAT	913	-
AAATAGTAGTGATAGAGAAGATTT	914	-
AGTGTATGAAGTGTAATAAGATTA	915	-
TGATTAAGATTGTGTAGTGTTATA	916	-
AGTTTATGATATTTGTAGATGAGT	917	-

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Sequence	SEQ ID NO:	No. in Ex 1
TATGTGTATGAAGATTATAGTTAG	918	78
GAAATTGTTGTATAGAGTGATATA	919	-
TAGAAATAGTTTAAGTATAGTGTG	920	-
TGATTTAGATGTTTATTGTGAGAA	921	-
AAGTTGATATTTGTTGTTAGATGA	922	-
TGATGTGATAATGAGAATAAAGAA	923	79
AAAGTTTAGTTTGTATTAGTAGAG	924	-
AGTTTGATGTGATAGTAAATAGAA	925	-
AAGTGTTATTGAATGTGATGTTAT	926	-
AAATTGAAGTGTGATAATGTTTGT	927	-
GTTTAGTGATTAAAGATAGATTAG	928	82
ATAAGTGTATAAGAGAAGTGTTAA	929	-
ATGAATTTGTTTGTGATGAAGTTA	930	-
AAAGAATTGAGAAATGAAAGTTAG	931	-
AGTGTAAGAGTATAAAGTATTTGA	932	-
GAATTAAGATTGTTATATGTGAGT	933	-
TATGAAAGTGTTGTTTAAGTAAGA	934	-
TAAAGTAAATGTTATGTGAGAGAA	935	-
AAAGATATTGATTGAGATAGAGTT	936	-
AAGTGATATGAATATGTGAGAAAT	937	-
AAATAGAGTTTGTTAATGTAAGTG	938	-
GATTTAGATGAGTTAAGAATTTAG	939	-
TTGTAAATGAGTGTGAATATTGTA	940	-
AGTAGTGTATTTGAGATAATAGAA	941	-
TGAGTTAAAGAGTTGTTGATATTT	942	-
AAAGAGTGTATTAGAAATAGTTTG	943	-
GTTTAGTTATTTGATGAGATAATG	944	-
AAGTGTAAATGAATAAAGAGTTGT	945	-
AATAAAGTGAGTAGAAGTGTAATT	946	-
TATTGAGTTTGTGTAAAGAAGATA	947	-
TTTATAGTTGTTGTGTTGAAAGTT	948	-
ATGAAATATGATTGTGTTTGTTGT	949	-
AAAGAGATGTAAAGTGAGTTATTA	950	-
TTGAAGAAAGTTAGATGATGAATT	951	-
ATGTTATTTGTTTAGTTTGTGTGA	952	-
AAATATGAATTTGAAGAGAAAGTGA	953	-
GATTAGATATAGAATATTGAAGAG	954	-
TTAGAATAAGAGAAATGTATGTGT	955	-
TTTATGAAAGAGAAGTGTATTATG	956	-
GTAAGTATTAAGTGTGATTTAGTA	957	-
ATAAAGAGAAGTAAAGAGTAAAGT	958	-
ATTGTTAATTGAAGTGTATGAAAG	959	-
TATATAGTTGAGTTGAGTAAGATT	960	-
TAGATGAGATATATGAAAGATAGT	961	-
ATAAGAAGATGATTTGTGTAAATG	962	-
TTAGTAATAAGAAAGATGAAGAGA	963	-
GATTTGTGAGTAAAGTAAATAGAA	964	-
AAATAGATGTAGAATTTGTGTGTT	965	-
GAAATTAGTGTTTGTGTGTATTAT	966	-
ATTTGAGTATGATAGAAGATTGTT	967	-
ATAGAGTTGAAGTATGTAAAGTTT	968	-
TAATTTGTGAATGTTGTTATTGTG	969	-
TTAGTTTATGAGAGTGAGATTTAA	970	-
GTTGTTAGAGTGTTTATGAAATTT	971	-
TTTATTGTGATGTGAAATAAGAGA	972	-
GTAAGTAATATGATAGTGATTAAG	973	-
TGAGATGATGTATATGTAGTAATA	974	-

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Sequence	SEQ ID NO:	No. in Ex 1
AATTGAGAAAGAGATAAATGATAG	975	85
TTTGAAGTGATGTTAGAATGTTTA	976	-
AGTTGTTGTGTAATTGTTAGTAAA	977	-
ATAGTGAGAAGTGATAAGATATTT	978	-
GTGTGATAAGTAATTGAGTTAAAT	979	-
TAGTTATTGTTTGTGAATTTGAGA	980	-
ATAGTTGAATAGTAATTTGAAGAG	981	-
ATGTTTGTGTTTGAATAGAGAATA	982	-
TGATAAAGATATGAGAGATTGTAA	983	-
TAAAGATGAGATGTTGTTAAAGTT	984	-
AAGTGAAATTTGTAAGAATTAGTG	985	-
GAAATGAGAGTTATTGATAGTTTA	986	-
TTTGTAAATGAGATATAGTGTTAG	987	-
GTTAATTGTGATATTTGATTAGTG	988	-
AGAGTGTTGATAAAGATGTTTATA	989	-
AATTGTGAGAAATTGATAAGAAGA	990	-
TTAAAGAGAATTGAGAAGAGAAAT	991	-
TTGTTAGAAGAATTGAATGTATGT	992	-
AGTTAAGATATGTGTGATGTTTAA	993	-
TGAGTTATGTTGTAATAGAAATTG	994	-
TTAGATAAGTTTAGAGATTGAGAA	995	-
ATGAGTAATAAGAGTATTTGAAGT	996	-
TGTTTAAGTGTAATGATTTGTTAG	997	-
TTGAAGAAGATTGTTATTGTTGAA	998	-
TATAGAAAGATTAAAGAGTGAATG	999	-
TAAATTGTTAGAAATTTGAGTGTG	1000	-
ATTGTTAGTGTGTTATTGATTATG	1001	-
GAGAATTATGTGTGAATATAGAAA	1002	-
TTGATTGATAAAGTAAAGAGTGTA	1003	-
GTGTGTAAATTGAATATGTTAATG	1004	-
AAAGTAAAGAAAGAAGTTTGAAAG	1005	-
TTTAGTTGAAGAATAGAAAGAAAG	1006	-
GTGTAATAAGAGTGAATAGTAATT	1007	-
TATTGAAATAAGAGAGATTTGTGA	1008	-
ATGAGAAAGAAGAAGTTAAGATTT	1009	-
AAGAGTGAGTATATTGTTAAAGAA	1010	-
TTTGTAAAGTGATGATGTAAGATA	1011	-
GATGTTATGTGATGAAATATGTAT	1012	-
GTAGAATAAAGTGTTAAAGTGTTA	1013	-
AAAGAGTATGTGTGTATGATATTT	1014	-
AAAGATAAGAGTTAGTAAATTGTG	1015	-
AAGAATTAGAGAATAAGTGTGATA	1016	-
GATAAGAAAGTGAAATGTAAATTG	1017	86
GATGAAAGATGTTTAAAGTTTGTT	1018	-
AGTGTAAGTAATAAGTTTGAGAAA	1019	-
GTTGAGAATTAGAATTTGATAAAG	1020	87
TTAAGAAATTTGTATGTGTTGTTG	1021	-
AGAAGATTTAGATGAAATGAGTTT	1022	-
TAAGTTTGAGATAAAGATGATATG	1023	-
TGAGATAGTTTGTAATATGTTTGT	1024	-
AGTTTGAAAATTGTAAGTTTGATGA	1025	-
TAGAATTGATTAATGATGAGTAGT	1026	-
AGAGATTTGTAATAAGTATTGAAG	1027	-
ATAATGATGTAATGTAAGTAGTGT	1028	-
TGAAATTTGATGAGAGATATGTTA	1029	-
TGTGTAAAGTATAGTTTATGTTAG	1030	-
TGAATAAGTGAAATAGAATGAATG	1031	-

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Sequence	SEQ ID NO:	No. in Ex 1
AAAGAAAGATTGTAATAAGTAGAG	1032	-
AATGAAATAGTGTTAAAATGAGTGT	1033	89
GTAGATAAAGATGTGAATTATGAT	1034	-
GATAGTATATGTGTGTATTTGTTT	1035	-
ATGTTTGTAGAAATGTTTGAAGAT	1036	-
AAATTTGTAGAGAGAAATTTGTTG	1037	-
TAGAATAAGATTAGTAAGTGTAGA	1038	-
TGATTTAGAGAAATATGAGTAGAA	1039	-
AATAGAGTATGTTGTTTATGAGAA	1040	-
GATGATGAAGAGTTTATTGTAAAAT	1041	-
AAGTAAAAGAAGAAGAAATGTGTTA	1042	-
TTGAAGAATTAAGTGTTTAGTGTA	1043	-
AGAAAGAATGTTGATTTATGATGT	1044	-
GATTAAAAGAGATGTTGATTGAAAAT	1045	-
AATGATAATTGTTGAGAGAGTAATT	1046	-
GTTTGTTGAAAGTGTAAAGTATAT	1047	90
TGAGTTATATGAGAAAGTGTAATT	1048	-
TTGTGAGAAAGAAGTATATAGAAT	1049	-
GTAAGTTTAGAGTTATAGAGTTTA	1050	-
GATAGATAGATAAGTTAATTGAAG	1051	-
AGAGATGATTGTTTATGTATTATG	1052	-
AAAGTTAAGAAATTGTAGTGATAG	1053	-
TTTGATATTGTTTGTGAGTGTATA	1054	-
ATTTGTAGAAAGTTGTTATGAGTT	1055	-
GATTTGAGTAAGTTTATAGATGAA	1056	-
AAGATAAAGTGAGTTGATTTAGAT	1057	-
GATATTGTAAGATATGTTGTAAAAG	1058	-
GTAAGAGTGTATTGTAAGTTAATT	1059	-
GTGTGATTAGTAATGAAGTATTTA	1060	91
GTAAGAAAGATTAAGTGTTAGTAA	1061	-
AGTAGAAAGTTGAAATTGATTATG	1062	92
TAAGAGAAGTTGAGTAATGTATTT	1063	-
GTTAAGAAATAGTAGATAAGTGAA	1064	-
TAAGTAAATTGAAAGTGTATAGTG	1065	-
AAGATGTATGTTTATTGTTGTGTA	1066	-
ATTTAGAATATAGTGAAGAGATAG	1067	-
GTTATGAAAGAGTATGTGTTAAAAT	1068	93
TATTATGTGAAGAAGAAATGATTAG	1069	-
TAATAAGTTGAAGAGAATTGTTGT	1070	-
TGATGTTTGATGTAATTGTTAAAG	1071	-
GTGAAAGATTTGAGTTTGTATAAT	1072	-
AGAGAATATAGATTGAGATTTGTT	1073	-
TTTGAGATGTGATGATAAAGTTAA	1074	-
GTTGTAAATTGTAGTAAAGAAGTA	1075	94
GTGTTATGATGTTGTTTGTATTAT	1076	-
ATTATTGTGTAGATGTATTAAGAG	1077	-
GTTAGAAAGATTTAGAAGTTAGTT	1078	-
TTGTGTATTAAGAGAGTGAAATAT	1079	-
GTTTAAGATAGAAAGAGTGATTTA	1080	-
AATGAGAAAATAGATAGTTATTGTG	1081	-
TGAATTGAAATAAGAATTTGTTGTG	1082	95
AATAAGATTGAATTAGTGAGTAAG	1083	-
AATGTTTGAGAGATTTAGTAAAGA	1084	-
AGTTTAGAAATAGAAATGTGTTTGA	1085	-
TATAAGTAAGTGTTAAGATTTGAG	1086	-
GTAGTGAATAAGTTAGTGTTAATA	1087	-
AAGTGTGTTAAAGTAAATGTAGAT	1088	-

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Sequence	SEQ ID NO:	No. in Ex 1
AGAGATGTTTATGTTGTGAATTAA	1089	-
AGTTGAATATTGATGATAAGAAGA	1090	-
TGAATGTGAGATGTTTAGAATAAT	1091	-
AATAATGATGTAAGTTTGAGTTTG	1092	-
AAAGAGTGAATAGAAATAAGAGAA	1093	-
AATAAAGTTATTGAGAGAGTTTAG	1094	-
AGTAGTGTTGTAGTTTAGTATATA	1095	-
GTAAGAATGTATTAGATATTTGTG	1096	-
GATAAATGTTTGATAAAGTAGTTG	1097	-
ATAGTATGTATGTGTGAAGATTTA	1098	-
ATGAATGTAGAGTGATTAGTTTAA	1099	-
GTAGTATTTAGTGATGTAAGAATA	1100	-
AGAATTGTATTGAAGAAGAATATG	1101	-
TTTATAGAATTGAGAGAAGTTAAG	1102	-
AAAGTAGTAGAGATTTGAGAATTA	1103	-
TTTAAAGAAAGTATTGTAAGAGTG	1104	-
AAATTGAGAAAAGTGAATGAAGTTT	1105	-
AAGAAATAAGTATGATAGTAGTAG	1106	-
ATTTGAATTGTATTGTAGTTTGTG	1107	-
AAGAGAATAATGTAGAGATATAAG	1108	-
TGTGTAATAGTTGTTAATGAGTAA	1109	-
TATAGTTGTAGTTTAGATGAATGT	1110	-
ATTGTGTTAGAATGATGTTAATAG	1111	-
GTTTGTATAGTATTTGATTGATGT	1112	-
AGAGTAAAGTATGAGTTATGAATA	1113	-
GAAAGTTTAAGTGATGTATATTGT	1114	96
TTAAATGATAAAGAGTAGTGAAGT	1115	-
TTAAATGTGTGAGAAGATGAATAA	1116	-
ATTTGTATAAAGTGAAGAAGAGAA	1117	97
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Table I

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ATTGAGTGTAGTGATTTAGAAATA	1166	-
AATAAAGTGTTTAAGAGTAGAGTA	1167	-
GTAGAGATAATTGATGTGTAATTT	1168	-

All references referred to in this specification are incorporated herein by reference.

The scope of protection sought for the invention described herein is defined by the appended claims. It will also be understood that any elements recited above or in the claims, can be combined with the elements of any claim. In particular, elements of a dependent claim can be combined with any element of a claim from which it depends, or with any other compatible element of the invention.

This application claims priority from United States Provisional Patent Application Nos. 60/263,710 and 60/303,799, filed January 25, 2001 and July 10, 2001. Both of these documents are incorporated herein by reference.

CLAIMS

1. A composition comprising molecules for use as tags or tag complements wherein each molecule comprises an oligonucleotide selected from a set of oligonucleotides based on a following group of sequences,

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wherein each of 1 to 3 is a nucleotide base selected to be different from the others of 1 to 3 with the *proviso* that up to three nucleotide bases of each sequence can be substituted with any nucleotide base provided that:

for any pair of sequences of the set:

M1 ≤ 16, M2 ≤ 13, M3 ≤ 20, M4 ≤ 16, and M5 ≤ 19, where:

M1 is the maximum number of matches for any alignment in which there are no internal indels;

M2 is the maximum length of a block of matches for any alignment;

M3 is the maximum number of matches for any alignment having a maximum score;

M4 is the maximum sum of the lengths of the longest two blocks of matches for any alignment of maximum score; and

M5 is the maximum sum of the lengths of all the blocks of matches

having a length of at least 3, for any alignment of maximum score;
wherein:

the score of an alignment is determined according to the equation (A x *m*) - (B x *mm*) - (C x (*og* + *eg*)) - (D x *eg*)), wherein:

for each of (i) to (iv):

(i) *m* = 6, *mm* = 6, *og* = 0 and *eg* = 6,

(ii) *m* = 6, *mm* = 6, *og* = 5 and *eg* = 1,

(iii) *m* = 6, *mm* = 2, *og* = 5 and *eg* = 1, and

(iv) *m* = 6, *mm* = 6, *og* = 6 and *eg* = 0,

A is the total number of matched pairs of bases in the alignment;

B is the total number of internal mismatched pairs in the alignment;

C is the total number of internal gaps in the alignment; and

D is the total number of internal indels in the alignment minus the total number of internal gaps in the alignment; and

wherein the maximum score is determined separately for each of (i), (ii), (iii) and (iv).

2. A composition comprising molecules for use as tags or tag complements wherein each molecule comprises an oligonucleotide selected from a set of oligonucleotides based on a group of sequences having the numeric pattern of sequences set out in claim 1, wherein each of 1 to 3 is a nucleotide base selected to be different from the others of 1 to 3 with the *proviso* that up to three nucleotide bases of each sequence can be substituted with any nucleotide base provided that:

for any pair of sequences of the set:

M1 ≤ 19, M2 ≤ 17, M3 ≤ 21, M4 ≤ 18, and M5 ≤ 20, where:

M1 is the maximum number of matches for any alignment in which there are no internal indels;

M2 is the maximum length of a block of matches for any alignment;

M3 is the maximum number of matches for any alignment having a maximum score;

M4 is the maximum sum of the lengths of the longest two blocks of matches for any alignment of maximum score; and

M5 is the maximum sum of the lengths of all the blocks of matches having a length of at least 3, for any alignment of maximum score;

wherein

the score of an alignment is determined according to the equation (A x *m*) - (B x *mm*) - (C x (*og* + *eg*)) - (D x *eg*)), wherein:

for each of (i) to (iv):

(i) $m = 6$, $mm = 6$, $og = 0$ and $eg = 6$,

(ii) $m = 6$, $mm = 6$, $og = 5$ and $eg = 1$,

(iii) $m = 6$, $mm = 2$, $og = 5$ and $eg = 1$, and

(iv) $m = 6$, $mm = 6$, $og = 6$ and $eg = 0$,

A is the total number of matched pairs of bases in the alignment;

B is the total number of internal mismatched pairs in the alignment;

C is the total number of internal gaps in the alignment; and

D is the total number of internal indels in the alignment minus the total number of internal gaps in the alignment; and

wherein the maximum score is determined separately for each of (i), (ii), (iii) and (iv).

3. A composition comprising molecules for use as tags or tag complements wherein each molecule comprises an oligonucleotide selected from a set of oligonucleotides based on a group of sequences having the numeric patterns set out in claim 1, wherein each of 1 to 3 is a nucleotide base selected to be different from the others of 1 to 3 with the *proviso* that up to three nucleotide bases of each sequence can be substituted with any nucleotide base provided that:

for any pair of sequences of the set:

M1 ≤ 19, M2 ≤ 17, M3 ≤ 21, M4 ≤ 18, and M5 ≤ 20, where:

M1 is the maximum number of matches for any alignment in which there are no internal indels;

M2 is the maximum length of a block of matches for any alignment;

M3 is the maximum number of matches for any alignment having a maximum score;

M4 is the maximum sum of the lengths of the longest two blocks of matches for any alignment of maximum score; and

M5 is the maximum sum of the lengths of all the blocks of matches having a length of at least 3, for any alignment of maximum score, wherein:

the score of an alignment is determined according to the equation 3A - B - 3C - D, wherein:

A is the total number of matched pairs of bases in the alignment;

B is the total number of internal mismatched pairs in the alignment;

C is the total number of internal gaps in the alignment; and

D is the total number of internal indels in the alignment minus the total number of internal gaps in the alignment; and

4. A composition according to any preceding claim, wherein for the group of 24mer sequences in which 1 = A, 2 = T and 3 = G, under a defined set of conditions in which the maximum degree of hybridization between a sequence and any complement of a different sequence of the group of 24mer sequences does not exceed 30% of the degree of hybridization between said sequence and its complement, for all said oligonucleotides of the composition, the maximum degree of hybridization between an oligonucleotide and a complement of any other oligonucleotide of the composition does not exceed 50% of the degree of hybridization of the oligonucleotide and its complement.

5. The composition of claim 4, wherein said maximum degree of hybridization between a sequence and any complement of a different sequence does not exceed 30% of the degree of hybridization between said sequence and its complement, the degree of hybridization between each sequence and its complement varies by a factor of between 1 and up to 10, more preferably between 1 and up to 9, more preferably between 1 and up to 8, more preferably between 1 and up to 7, more preferably between 1 and up to 6, and more preferably between 1 and up to 5.

6. The composition of claim 4 or 5, wherein the maximum degree of hybridization between a sequence and any complement of a different sequence does not exceed 25%, more preferably does not exceed 20%, more preferably does not exceed 15%, more preferably does not exceed 10%, more preferably does not exceed 5%.

7. The composition of any of claims 4 to 6, wherein said defined set of conditions results in a level of hybridization that is the same as the level of hybridization obtained when hybridization conditions include 0.2 M NaCl, 0.1 M Tris, 0.08% Triton X-100, pH 8.0 at 37°C.

8. The composition of any of claims 4 to 7, said defined set of conditions includes the group of 24mer sequences being covalently linked to beads.

9. The composition of claim 4 wherein, for the group of 24mers the maximum degree of hybridization between a sequence and any complement of a different sequence does not exceed 15% of the degree of hybridization between said sequence and its complement and the degree of hybridization

between each sequence and its complement varies by a factor of between 1 and up to 9, and for all oligonucleotides of the set, the maximum degree of hybridization between an oligonucleotide and a complement of any other oligonucleotide of the set does not exceed 20% of the degree of hybridization of the oligonucleotide and its complement.

10. The composition of any preceding claim wherein: each 1 is one of A, T/U, G and C; each 2 is one of A, T/U, G and C; and each 3 is one of A, T/U, G and C; and each of 1, 2 and 3 is selected so as to be different from all of the others of 1, 2 and 3.

11. The composition of claim 10, wherein 1 is A or T/U, 2 is A or T/U and 3 is G or C.

12. The composition of claim 11, wherein 1 is A, 2 is T/U and 3 is G.

13. The composition of any preceding claim, wherein each said oligonucleotide is from twenty-two to twenty-six bases in length, or from twenty-three to twenty-five.

14. The composition of any of claims 1 to 13, wherein each said oligonucleotide is of the same length as every other said oligonucleotide.

15. The composition of claim 14, wherein each said oligonucleotide is twenty-four bases in length.

16. The composition of any of claims 1 to 13 wherein no said oligonucleotide contains more than four contiguous bases that are identical to each other.

17. The composition of any of claims 1 to 12 and 16 wherein the number of G's in each said oligonucleotide does not exceed L/4 where L is the number of bases in said sequence.

18. The composition of any of claims 1 to 14 and 16 and 17, wherein the number of G's in each said oligonucleotide does not vary from the average number of G's in all of the said oligonucleotides by more than one.

19. The composition of any claims 1 to 14 and 16 to 18, wherein the number of G's in each said oligonucleotide is the same as every other said oligonucleotide.

20. The composition of claim 19, wherein each said oligonucleotide is twenty-four bases in length and each said oligonucleotide contains 6 G's.

21. The composition of any of claims 1 to 12 and claims 16 to 20, wherein, for each said nucleotide, there is at most six bases other than G between every pair of neighboring pairs of G's.

22. The composition of any of claims 1 to 12 and claims 16 to 21, wherein at the 5'-end of each said oligonucleotide at least one of the first, second, third, fourth, fifth, sixth and seventh bases of the sequence of the oligonculeotide is a G.

23. The composition of any of claims 1 to 12 and claims 16 to 21, wherein at the 3'-end of each said oligonucleotide at least one of the first, second, third, fourth, fifth, sixth and seventh bases of the sequence of the oligonucleotide is a G.

24. The composition of claim 22, wherein at the 3'-end of each said oligonucleotide at least one of the first, second, third, fourth, fifth, sixth and seventh bases of the sequence of the oligonucleotide is a G.

25. The composition of any of claims 1 to 24, comprising ten, or twenty, or thirty, or forty, or fifty, or sixty, or seventy, or eighty, or ninety, or one hundred, or one hundred and ten, or one hundred and twenty, or one hundred and thirty, or one hundred and forty, or one hundred and fity, or one hundred and sixty said molecules, or comprising one hundred and seventy said molecules, or comprising one hundred and eighty said molecules, or comprising one hundred and ninety said molecules, or comprising two hundred said molecules, or comprising two hundred and twenty said molecules, or comprising two hundred and forty said molecules, or comprising two hundred and sixty said molecules, or comprising two hundred and eighty said molecules, or comprising three hundred said molecules, or comprising four hundred said molecules, or comprising five hundred said molecules, or comprising six hundred said molecules, or comprising seven hundred said molecules, or comprising

eight hundred said molecules, or comprising nine hundred said molecules, or comprising one thousand said molecules, or comprising eleven hundred said molecules.

26. A composition of any of claims 1 to 25, wherein each said molecule is linked to a solid phase support so as to be distinguishable from a mixture of other said molecules by hybridization to its complement.

27. The composition of claim 26, wherein each molecule is linked to a defined location on a said solid phase support, the defined location for each said molecule being different than the defined location for different other said molecules.

28. The composition of claim 27, wherein each said solid phase support is a microparticle and each said molecule is covalently linked to a different microparticle than each other different said molecule.

29. A composition comprising a set of 150 molecules for use as tags or tag complements wherein each molecule comprises an oligonucleotide having a sequence of at least sixteen nucleotide bases wherein for any pair of sequences of the set:

M1 ≤ 19/24 x L1, M2 ≤ 17/24 x L1, M3 ≤ 21/24 x L1, M4 ≤ 18/24 x L1, M5 ≤ 20/24 x L1, where L1 is the length of the shortest sequence of the pair, where:

M1 is the maximum number of matches for any alignment of the pair of sequences in which there are no internal indels;

M2 is the maximum length of a block of matches for any alignment of the pair of sequences;

M3 is the maximum number of matches for any alignment of the pair of sequences having a maximum score;

M4 is the maximum sum of the lengths of the longest two blocks of matches for any alignment of the pair of sequences of maximum score; and

M5 is the maximum sum of the lengths of all the blocks of matches having a length of at least 3, for any alignment of the pair of sequences of maximum score, wherein:

the score of an alignment is determined according to the equation (A x *m*) - (B x *mm*) - (C x (*og* + *eg*)) - (D x *eg*)), wherein:

for each of (i) to (iv):

(i) *m* = 6, *mm* = 6, *og* = 0 and *eg* = 6,

(ii) $m = 6$, $mm = 6$, $og = 5$ and $eg = 1$,

(iii) $m = 6$, $mm = 2$, $og = 5$ and $eg = 1$, and

(iv) $m = 6$, $mm = 6$, $og = 6$ and $eg = 0$,

A is the total number of matched pairs of bases in the alignment;

B is the total number of internal mismatched pairs in the alignment;

C is the total number of internal gaps in the alignment; and

D is the total number of internal indels in the alignment minus the total number of internal gaps in the alignment; and

wherein the maximum score is determined separately for each of (i), (ii), (iii) and (iv).

30. A composition comprising a set of 150 molecules for use as tags or tag complements wherein each molecule comprises an oligonucleotide having a sequence of at least sixteen nucleotide bases wherein for any pair of sequences of the set:

$M1 \leq 19$, $M2 \leq 17$, $M3 \leq 21$, $M4 \leq 18$, and $M5 \leq 20$, where:

M1 is the maximum number of matches for any alignment of the pair of sequences in which there are no internal indels;

M2 is the maximum length of a block of matches for any alignment of the pair of sequences;

M3 is the maximum number of matches for any alignment of the pair of sequences having a maximum score;

M4 is the maximum sum of the lengths of the longest two blocks of matches for any alignment of the pair of sequences of maximum score; and

M5 is the maximum sum of the lengths of all the blocks of matches having a length of at least 3, for any alignment of the pair of sequences of maximum score, wherein:

the score of a said alignment is determined according to the equation $3A - B - 3C - D$, wherein:

A is the total number of matched pairs of bases in the alignment;

B is the total number of internal mismatched pairs in the alignment;

C is the total number of internal gaps in the alignment; and

D is the total number of internal indels in the alignment minus the total number of internal gaps in the alignment.

31. The composition of claim 29 or 30, wherein each said sequence has up to fifty bases.

32. The composition of claim 31, wherein each said sequence is between sixteen and forty bases in length, or between sixteen and thirty-five bases in length, or between eighteen and thirty bases in length, or between twenty and twenty-eight bases in length, or between twenty-one and twenty-seven bases in length, or between twenty-two and twenty-six bases in length.

33. The composition of any of claims 29 to 32, wherein each said sequence is of the same length as every other said sequence.

34. The composition of claim 33, wherein each said sequence is twenty-four bases in length.

35. The composition of any of claims 29 to 34 wherein no said sequence contains more than four contiguous bases that are identical to each other.

36. The composition of any of claims 29 to 35 wherein the number of G's in each said sequence does not exceed L/4 where L is the number of bases in said sequence.

37. The composition of claim 36, wherein the number of G's in each said sequence does not vary from the average number of G's in all of the sequences of the set by more than one.

38. The composition of claim 37, wherein the number of G's in each said sequence is the same as every other sequence of the set.

39. The composition of claim 37, wherein each said sequence is twenty-four bases in length and each said sequence contains 6 G's.

40. The composition of any of claims 29 to 39 wherein, for each said sequence, there is at most six bases other than G between every pair of neighboring pairs of G's.

41. The composition of any of claims 29 to 40, wherein at the 5'-end of each said sequence at least one of the first, second, third, fourth, fifth, sixth and seventh bases of the sequence is a G.

42. The composition of any of claims 29 to 40, wherein at the 3'-end of each said sequence at least one of the first, second, third, fourth, fifth, sixth and seventh bases of the sequence is a G.

43. The composition of claim 41, wherein at the 3'-end of each said sequence at least one of the first, second, third, fourth, fifth, sixth and seventh bases of the sequence is a G.

44. The composition of any of claims 29 to 43, wherein under a defined set of conditions, the maximum degree of hybridization between a said oligonucleotide and any complement of a different oligonucleotide of the composition does not exceed about 30% of the degree of hybridization between said oligonucleotide and its complement, more preferably 20%, more preferably 15%, more preferably 10%, more preferably 6%.

45. The composition of claim 44, wherein said set of conditions results in a level of hybridization that is the same as the level of hybridization obtained when hybridization conditions include 0.2 M NaCl, 0.1 M Tris, 0.08% Triton X-100, pH 8.0 at 37°C, and the oligonucleotides are covalently linked to microparticles.

46. The composition of claim 45, wherein under said defined set of conditions, the degree of hybridization between each oligonucleotide and its complement varies by a factor of between 1 and up to 8, more preferably up to 7, more preferably up to 6, more preferably up to 5.

47. The composition of any of claims 29 to 46, comprising one hundred and sixty said molecules, or comprising one hundred and seventy said molecules, or comprising one hundred and eighty said molecules, or comprising one hundred and ninety said molecules, or comprising two hundred said molecules, or comprising two hundred and twenty said molecules, or comprising two hundred and forty said molecules, or comprising two hundred and sixty said molecules, or comprising two hundred and eighty said molecules, or comprising three hundred said molecules, or comprising four hundred said molecules, or comprising five hundred said molecules, or comprising six hundred said molecules, or comprising seven hundred said molecules, or comprising eight hundred said molecules, or comprising nine hundred said molecules, or comprising one thousand said molecules.

48. A composition of any of claims 29 to 47, wherein each said molecule is linked to a solid phase support so as to be distinguishable from a mixture of other said molecules by hybridization to its complement.

49. The composition of claim 48, wherein each molecule is linked to a defined location on a said solid phase support, the defined location for each said molecule being different than the defined location for different other said molecules.

50. The composition of claim 49, wherein each said solid phase support is a microparticle and each said molecule is covalently to a different microparticle than each other different said molecule.

51. A composition comprising one hundred and fifty minimally cross-hybridizing molecules for use as tags or tag complements wherein each molecule comprises an oligonucleotide comprising a sequence of nucleotide bases for which, under a defined set of conditions, the maximum degree of hybridization between a said oligonucleotide and any complement of a different oligonucleotide does not exceed about 20% of the degree of hybridization between said oligonucleotide and its complement.

52. The composition of claim 51, wherein each said sequence has between ten and fifty bases.

53. The composition of claim 52, wherein each said sequence is between sixteen and forty bases in length, or between sixteen and thirty-five bases in length, or between eighteen and thirty bases in length, or between twenty and twenty-eight bases in length, or between twenty-one and twenty-seven bases in length, or between twenty-two and twenty-six bases in length.

54. The composition of any of claims 51 to 53, wherein each said sequence is of the same length as every other said sequence.

55. The composition of claim 54, wherein each said sequence is twenty-four bases in length.

56. The composition of any of claims 51 to 55 wherein no said sequence contains more than four contiguous bases that are identical to each other.

57. The composition of any of claims 51 to 56 wherein the number of G's in each said sequence does not exceed L/4 where L is the number of bases in said sequence.

58. The composition of claim 57, wherein the number of G's in each said sequence does not vary from the average number of G's in all of the sequences of the set by more than one.

59. The composition of claim 58, wherein the number of G's in each said sequence is the same as every other sequence of the set.

60. The composition of claim 58, wherein each said sequence is twenty-four bases in length and each said sequence contains 6 G's.

61. The composition of any of claims 51 to 60 wherein, for each said sequence, there is at most six bases other than G between every pair of neighboring pairs of G's.

62. The composition of any of claims 51 to 61, wherein at the 5'-end of each said sequence at least one of the first, second, third, fourth, fifth, sixth and seventh bases of the sequence is a G.

63. The composition of any of claims 51 to 61, wherein at the 3'-end of each said sequence at least one of the first, second, third, fourth, fifth, sixth and seventh bases of the sequence is a G.

64. The composition of claim 62, wherein at the 3'-end of each said sequence at least one of the first, second, third, fourth, fifth, sixth and seventh bases of the sequence is a G.

65. The composition of any of claims 61 to 64, wherein under a said defined set of conditions, the maximum degree of hybridization between a said oligonucleotide and any complement of a different oligonucleotide of the composition does not exceed about 15%, more preferably 10%, more preferably 6%.

66. The composition of claim 65, wherein said set of conditions results in a level of hybridization that is the same as the level of hybridization obtained when hybridization conditions include 0.2 M NaCl, 0.1 M Tris, 0.08% Triton X-100, pH 8.0 at 37°C, and the oligonucleotides are covalently linked to microparticles.

67. The composition of claim 66, wherein under said defined set of conditions, the degree of hybridization between each oligonucleotide and its complement varies by a factor of between 1 and up to 8, more preferably up to 7, more preferably up to 6, more preferably up to 5.

68. The composition of any of claims 51 to 67, comprising one hundred and sixty said molecules, or comprising one hundred and seventy said molecules, or comprising one hundred and eighty said molecules, or comprising one hundred and ninety said molecules, or comprising two hundred said molecules, or comprising two hundred and twenty said molecules, or comprising two hundred and forty said molecules, or comprising two hundred and sixty said molecules, or comprising two hundred and eighty said molecules, or comprising three hundred said molecules, or comprising four hundred said molecules, or comprising five hundred said molecules, or comprising six hundred said molecules, or comprising seven hundred said molecules, or comprising eight hundred said molecules, or comprising nine hundred said molecules, or comprising one thousand said molecules.

69. A composition of any of claims 51 to 68, wherein each said molecule is linked to a solid phase support so as to be distinguishable from a mixture of other said molecules by hybridization to its complement.

70. The composition of claim 69, wherein each molecule is linked to a defined location on a said solid phase support, the defined location for each said molecule being different than the defined location for different other said molecules.

71. The composition of claim 70, wherein each said solid phase support is a microparticle and each said molecule is covalently to a different microparticle than each other different said molecule.

72. A composition according to any of claims 1 to 71, wherein any base is substituted by an analogue thereof.

73. A composition according to any of claims 1 to 72, wherein each said molecule comprises a tag complement.

74. A kit for sorting and identifying polynucleotides, the kit comprising one or more solid phase supports each having one or more spatially discrete regions, each such region having a uniform population of substantially identical tag complements covalently attached, and the tag complements each being selected from the set of oligonucleotides as defined in any of claims 1 to 73.

75. A kit according to claim 74, wherein there is a tag complement for each said oligonucleotide of a said composition.

76. A kit according to claim 74 or 75 wherein said one or more solid phase supports is a planar substrate and wherein said one or more spatially discrete regions is a plurality of spatially addressable regions.

77. A kit according to any of claims 74 to 76 wherein said one or more solid phase supports is a plurality of microparticles.

78. A kit according to claim 77 wherein said microparticles each have a diameter in the range of from 5 to 40 μm.

79. A kit according to claim 77 or 78, wherein each microparticle is spectrophotometrically unique from each other microparticle having a different oligonucleotide attached thereto.

80. A method of analyzing a biological sample comprising a biological sequence for the presence of a mutation or polymorphism at a locus of the nucleic acid, the method comprising:

(A) amplifying the nucleic acid molecule in the presence of a first primer having a 5'-sequence having the sequence of a tag complementary to the sequence of a tag complement belonging to a family of tag complements as defined in claim 73 to form an amplified molecule with a 5'-end with a sequence complementary to the sequence of the tag;

(B) extending the amplified molecule in the presence of a polymerase and a

second primer having 5'-end complementary the 3'-end of the amplified sequence, with the 3'-end of the second primer extending to immediately adjacent said locus, in the presence of a plurality of nucleoside triphosphate derivatives each of which is: (i) capable of incorporation during transciption by the polymerase onto the 3'-end of a growing nucleotide strand; (ii) causes termination of polymerization; and (iii) capable of differential detection, one from the other, wherein there is a said derivative complementary to each possible nucleotide present at said locus of the amplified sequence;

(C) specifically hybridizing the second primer to a tag complement having the tag complement sequence of (A); and

(D) detecting the nucleotide derivative incorporated into the second primer in (B) so as to identify the base located at the locus of the nucleic acid.

81. A method of analyzing a biological sample comprising a plurality of nucleic acid molecules for the presence of a mutation or polymorphism at a locus of each nucleic acid molecule, for each nucleic acid molecule, the method comprising:

(A) amplifying the nucleic acid molecule in the presence of a first primer having a 5'-sequence having the sequence of a tag complementary to the sequence of a tag complement belonging to a family of tag complements as defined in claim 73 to form an amplified molecule with a 5'-end with a sequence complementary to the sequence of the tag;

(B) extending the amplified molecule in the presence of a polymerase and a second primer having 5'-end complementary the 3'-end of the amplified sequence, the 3'-end of the second primer extending to immediately adjacent said locus, in the presence of a plurality of nucleoside triphosphate derivatives each of which is: (i) capable of incorporation during transciption by the polymerase onto the 3'-end of a growing nucleotide strand; (ii) causes termination of polymerization; and (iii) capable of differential detection, one from the other, wherein there is a said derivative complementary to each possible nucleotide present at said locus of the amplified molecule;

(C) specifically hybridizing the second primer to a tag complement having the tag complement sequence of (A); and

(D) detecting the nucleotide derivative incorporated into the second primer in (B) so as to identify the base located at the locus of the nucleic acid;

wherein each tag of (A) is unique for each nucleic acid molecule and steps (A) and (B) are carried out with said nucleic molecules in the presence of each other.

82. A method of analyzing a biological sample comprising a plurality of double stranded complementary nucleic acid molecules for the presence of a mutation or polymorphism at a locus of each nucleic acid molecule, for each nucleic acid molecule, the method comprising:

(A) amplifying the double stranded molecule in the presence of a pair of first primers, each primer having an identical 5'-sequence having the sequence of a tag complementary to the sequence of a tag complement belonging to a family of tag complements as defined in claim 73 to form amplified molecules with 5'-ends with a sequence complementary to the sequence of the tag;

(B) extending the amplified molecules in the presence of a polymerase and a pair of second primers each second primer having a 5'-end complementary a 3'-end of the amplified sequence, the 3'-end of each said second primer extending to immediately adjacent said locus, in the presence of a plurality of nucleoside triphosphate derivatives each of which is: (i) capable of incorporation during transciption by the polymerase onto the 3'-end of a growing nucleotide strand; (ii) causes termination of polymerization; and (iii) capable of differential detection, one from the other;

(C) specifically hybridizing each of the second primers to a tag complement having the tag complement sequence of (A); and

(D) detecting the nucleotide derivative incorporated into the second primers in (B) so as to identify the base located at said locus;

wherein the sequence of each tag of (A) is unique for each nucleic acid molecule and steps (A) and (B) are carried out with said nucleic molecules in the presence of each other.

83. A method of analyzing a biological sample comprising a plurality of nucleic acid molecules for the presence of a mutation or polymorphism at a locus of each nucleic acid molecule, for each nucleic acid molecule, the method comprising:

(a) hybridizing the molecule and a primer, the primer having a 5'-sequence having the sequence of a tag complementary to the sequence of a tag complement belonging to a family of tag complements as defined in claim 73 and a 3'-end extending to immediately adjacent the locus;

(b) enzymatically extending the 3'-end of the primer in the presence of a

plurality of nucleoside triphosphate derivatives each of which is: (i) capable of enzymatic incorporation onto the 3'-end of a growing nucleotide strand; (ii) causes termination of said extension; and (iii) capable of differential detection, one from the other, wherein there is a said derivative complementary to each possible nucleotide present at said locus;

(c) specifically hybridizing the extended primer formed in step (b) to a tag complement having the tag complement sequence of (a); and

(d) detecting the nucleotide derivative incorporated into the primer in step (b) so as to identify the base located at the locus of the nucleic acid molecule;

wherein each tag of (a) is unique for each nucleic acid molecule and steps (a) and (b) are carried out with said nucleic molecules in the presence of each other.

84. The method of claim 81 wherein each said derivative is a dideoxy nucleoside triphosphate.

85. The method of claim 83, wherein each respective complement is attached as a uniform population of substantially identical complements in a spacially discrete region on one or more said solid phase supports.

86. The method of claim 85, each said tag complement comprises a label, each such label being different for respective complements, and step (d) includes detecting the presence of the different labels for respective hybridization complexes of bound tags and tag complements.

87. The hybridized molecule and primer of step (A) of any of claims 83 to 86.

88. A method of determining the presence of a target suspected of being contained in a mixture, the method comprising the steps of:

(i) labelling the target with a first label;

(ii) providing a first detection moiety capable of specific binding to the target and including a first tag;

(iii) exposing a sample of the mixture to the detection moiety under conditions suitable to permit (or cause) said specific binding of the molecule and target;

(iv) providing a family of tag complements as defined in claim 73 wherein the family contains a first tag complement having a sequence

complementary to that of the first tag;

(v) exposing the sample to the family of tag complements under conditions suitable to permit (or cause) specific hybridization of the first tag and its tag complement;

(vi) determining whether a said first detection moiety hybridized to a first said tag complement is bound to a said labelled target in order to determine the presence or absence said target in the mixture.

89. The method of claim 88 wherein said first tag complement is linked to a solid support at a specific location of the support and step (vi) includes detecting the presence the first label at said specified location.

90. The method of claim 88 wherein said first tag complement comprises a second label and step (vi) includes detecting the presence of the first and second labels in a hybridized complex of the moiety and the first tag complement.

91. The method of claim 88 wherein said target is selected from the group consisting of organic molecules, antigens, proteins, polypeptides, antibodies and nucleic acids.

92. The method of claim 91, wherein said target is an antigen and said first molecule is an antibody specific for said antigen.

93. The method of claim 92, wherein the antigen is a polypeptide or protein and the labelling step includes conjugation of fluorescent molecules, digoxigenin, biotinylation and the like.

94. The method of claim 93, wherein said target is a nucleic acid and the labelling step includes incorporation of fluorescent molecules, radiolabelled nucleotide, digoxigenin, biotinylation and the like.